STATE OF NORTH CAROLINA DEPARTMENT OF ENVIRONMENT AND NATURAL RESOURCES

REPORT OF PROCEEDINGS TO THE NORTH CAROLINA ENVIRONMENTAL MANAGEMENT COMMISSION FOR PROPOSED REVISIONS TO THE 15A NCAC 02L .0113 (VARIANCE) AND .0202 (GROUNDWATER QUALITY STANDARDS) GROUNDWATER RULES

PUBLIC HEARING

May 23, 2012, 6:30 PM – Archdale Building, Ground Floor Hearing Room, 512 N. Salisbury Street, Raleigh, NC

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I. INTRODUCTION

This report is the official record of the proceedings related to the North Carolina Department of Environment and Natural Resources (DENR) Division of Water Quality's (DWQ) proposal to revise the Variance Procedures in 15A North Carolina Administrative Code (NCAC) 02L .0113 and the Groundwater Quality Standards in 15A NCAC 02L .0202. This report includes written comments received during the public comment period, relevant exhibits, and the final recommendation of the hearing officer as to the proposed revisions to the Groundwater Rules for consideration by the North Carolina Environmental Management Commission (EMC).

II. BACKGROUND

In March 2011, McGuireWoods, on behalf of Rhodia, Inc., (Rhodia) submitted a rulemaking petition to the DWQ Director requesting amendment of the groundwater quality standard for 1,1-dichloroethylene (1,1-DCE) in 15A NCAC 02L .0202(g)(59) from 7 ug/L to 350 ug/L. The petition was submitted in accordance with North Carolina General Statute 150B-20 and 15A NCAC 02I .501, which allows any person to petition the Director to adopt, amend or repeal an existing rule of the EMC.

The groundwater quality standards, as specified in 15A NCAC 02L .0202, are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of the state, which may be tolerated without creating a threat to human health or which would otherwise render the groundwater unsuitable for its intended best usage as a source of drinking water.

Current regulations establish groundwater quality standards as the *least* of the six criteria contained in 15A NCAC 02L .0202(d)(1) - (6):

- 1. Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (0.10 for inorganics; 0.20 for organics)]/[2liters/day (avg. water consumption)];
- 2. Concentration which corresponds to an incremental lifetime cancer risk of 1x 10⁻⁶;
- 3. Taste threshold limit value:
- 4. Odor threshold limit value:
- 5. Maximum contaminant level; or
- 6. National secondary drinking water standard.

Further, Paragraph 15A NCAC 02L .0202(e) requires that the following references, in order of preference, be used in establishing concentrations of substances which correspond to levels described in Paragraph (d) above:

- 1. U.S. EPA Integrated Risk Information System (IRIS).
- 2. Health Advisories (U.S. EPA Office of Water).
- 3. Other health risk assessment data published by U.S. EPA.

4. Other relevant, published health risk assessment data and scientifically valid peer-reviewed published toxicological data.

The current groundwater standard for 1,1-DCE is the maximum contaminant level (MCL) of 7 ug/L, the *least* of the six criteria available for 1,1-DCE. MCLs are federal drinking water standards established by the U.S. EPA Office of Water and are applicable to public water systems regulated under the Federal Safe Drinking Water Act (http://water.epa.gov/lawsregs/rulesregs/sdwa/index.cfm). North Carolina has adopted federal MCLs by reference as drinking water standards applicable to public water systems in Title 15A, Subchapter 18C, Section .1500 of the North Carolina Administrative Code (http://www.ncwater.org/pws/rules/SECTION_1500.pdf).

Rhodia requested amendment of the 1,1-DCE groundwater quality standard because the 1,1-DCE MCL of 7 ug/L is based on outdated U.S. EPA IRIS health effects data that has since been removed from the IRIS database and updated with new information. The DWQ and the U.S. EPA both acknowledge that updated health effects data available in U.S. EPA's IRIS database (http://www.EPA.gov/iris/subst/0039.htm) support calculation of a less stringent MCL. However, based on U.S. EPA's second Six Year Review of Drinking Water Standards, U.S. EPA does not plan to update the MCL because any potential revision is not likely to provide a meaningful opportunity for cost-savings or health risk reduction to public water systems and their customers (http://www.gpo.gov/fdsys/pkg/FR-2010-03-29/pdf/2010-6624.pdf).

1,1- DCE is an industrial chemical *not* found naturally in the environment. Companies use 1,1- DCE to make plastics, such as flexible films like food wrap, flame retardant coatings, adhesives, and packaging materials. The major source of 1,1-DCE in drinking water is discharge from industrial chemical factories. Long-term exposure to 1,1-DCE in drinking water may cause liver, kidney and lung damage.

At its May 2011 meeting, the Groundwater Committee (GWC) of the EMC heard presentations on the rulemaking petition from Rhodia and DWQ staff. DWQ staff recommended that the rulemaking petition be denied and that Rhodia address deficiencies in their request for a variance to the 1,1-DCE groundwater standard. Rather than amending the 02L rules to deal with this specific standard issue, the variance rules in 15A NCAC 02L .0113 may be used to allow a less restrictive 1,1-DCE standard while requiring site-specific requirements necessary to protect public water systems that are required to meet the federal MCLs. After discussion, the GWC passed a motion recommending that the full EMC proceed with rulemaking as proposed by the petitioner to amend the 1,1-DCE standard from 7 ug/L to 350 ug/L. The Committee acknowledged that, per legal counsel, new rule language is needed to allow deviation from 15A NCAC 02L .0202(d), which requires that the groundwater standard be established at the lowest of the six criteria. The lowest of the six criteria for 1,1-DCE is the MCL of 7 ug/L. In its rulemaking petition, Rhodia provided a legal opinion that 15A NCAC 02L .0202(d) and (e), in tandem, are sufficiently broad to establish the groundwater standard at 350 ug/L under 15A NCAC 02L .0202(d)(1), based on the current toxicity data published in the U.S. EPA's Integrated Risk Information System (IRIS) database, rather than establishing the standard at the federal MCL, which was calculated prior to the updated toxicity data being published.

The EMC approved Rhodia's petition at its July 14, 2011 meeting and directed the DWQ to initiate rulemaking for three rule options:

- 1) A change in 02L .0202 (g)(59) to amend the 1,1-DCE standard from 7 ug/L to 350 ug/L;
- 2) A change in 02L .0202 (d) and (f) to allow the EMC to establish a standard less stringent than the maximum contaminant level (MCL) or national secondary drinking water standard when:
 - a. the MCL or secondary drinking water standard is not based on the most recent U.S. EPA health effects data listed in paragraph (e),
 - b. such a standard would not endanger public health and safety, and,
 - c. compliance with a standard based on the MCL would produce serious hardship without equal or greater public benefit.
- 3) A change in 02L .0113 to:
 - a. update the Division of Water Quality mailing address in .0113 (b),
 - b. allow the EMC to issue a state-wide variance to the 02L rules in .0113 (d), and,
 - c. clarify the existing variance requirements in .0113(i).

The EMC also approved taking the Fiscal Note out for public comment. Information in regard to the EMC's proceedings, including Rhodia's rulemaking petition, is available at http://portal.ncdenr.org/web/emc/july-14-2011 (Agenda Item 11-27).

III. SUMMARY OF PROPOSALS

The rule text for the three proposed options is located in Attachments A, B and C, respectively. The proposed changes to the rules have been highlighted.

Option 1: Amend 02L .0202 (g) (59)

Rhodia, a global specialty chemical manufacturer that formerly operated as Rhone-Poulenc in Gastonia, North Carolina, submitted a rulemaking petition to amend the 1,1-DCE groundwater standard in 02L .0202(g)(59) from 7 ug/L to 350 ug/L based on the availability of more recent U.S. EPA health effects data. A change in this standard may result in lower compliance costs for facilities that have a release of 1,1-DCE to groundwater. The purpose of changing the 1,1-DCE groundwater standard from 7 ug/L to 350 ug/L is to incorporate the most up-to-date U.S. EPA IRIS health effects data in the calculation of a health-based groundwater standard. During the EMC's deliberations, legal counsel advised that this option is not legally viable unless the modifications in Option 2 are also adopted. It is Rhodia's position that the current rules will allow a change in the standard without legal challenge.

Option 2: Amend 02L .0202 (d) and (f)

The purpose of the proposed changes to 02L .0202(d) and (f) is three-fold:

- 1) To ensure that the most recent U.S. EPA health effects data are used in establishing groundwater quality standards;
- 2) To ensure that the standard is protective of public health and safety; and,
- 3) To ensure that the standard is not overly burdensome to regulated parties.

If the lowest concentration of the six regulatory criteria for establishing a standard in 15A NCAC .0202(d) is the MCL or national secondary drinking water standard and it is not based on the most recent U.S. EPA health effects data in .0202(e), then this proposed rule option will allow the MCL or national secondary drinking water standard to be eliminated from consideration as the groundwater standard. At this time, 1,1-DCE is the only standard for which a change has been requested; however, this proposed rule change might lead to additional groundwater quality standard changes in the future.

The EMC's legal counsel believes that Option 2 will allow the EMC to set the 1,1-DCE standard above the MCL without legal challenge. It is Rhodia's position that the current rules will allow a change in the standard without legal challenge.

Option 3: Amend 02L .0113 (b) through (i)

Proposed changes under Option 3 include an update to the DWQ mailing address, clarification of the existing variance requirements, and would allow the EMC to consider a state-wide variance request for a less restrictive groundwater standard when the existing standard is based on outdated health effects data, such as the case with the existing 1,1-DCE standard. A state-wide variance to a groundwater standard would not change the fundamental way standards are currently established in 15A NCAC 02L .0202(d); however, it would ultimately result in the establishment of a second standard in addition to the one established in 02L .0202(g). The party requesting a state-wide variance would incur all of the cost of gathering the necessary data requirements.

IV. FISCAL ANALYSIS PROCESS

DWQ staff conducted outreach activities to potentially affected parties, including members of the regulated community, environmental groups and state agencies, to determine the impact of the proposed rule changes on their operations. The feedback received from these outreach activities was used to prepare a fiscal analysis, as required by the Administrative Procedure Act (N.C.G.S 150B-21-4). The fiscal analysis was approved by the DENR Budget Office on January 9, 2012, by the Office of State Management and Budget (OSBM) on March 28, 2012 and by the EMC at its March 8, 2012 meeting. It is included as Attachment P and is available on the OSBM website at http://www.osbm.state.nc.us/files/pdf files/DENR03282012.pdf.

The existing 15A NCAC 02L .0113 and .0202 groundwater rules serve as a baseline for the fiscal analysis. The analysis assumes that the adoption of any one of the three options proposed, or a combination of the options, would ultimately result in a higher 1,1-DCE standard and would result in the same fiscal impacts.

Summary of Costs and Benefits Associated with Proposed Rule Changes

Rhodia was the only private industry identified as being immediately impacted by the proposed rule based on outreach response. Rhodia, operating as Rhone-Poulenc, had a release of 1,1-DCE from an above ground storage tank in 1991 and began operating a pump-and-treat-groundwater remediation system in 1996 to remediate contaminated groundwater. Rhodia would see reduced groundwater cleanup and monitoring costs if the 1,1-DCE standard is changed from 7 ug/L to

350 ug/L. Other private industries with 1,1-DCE groundwater contamination above the current groundwater standard of 7 ug/L could also see reduced cleanup costs even though none were identified through outreach efforts.

DENR's Division of Waste Management (DWM) reported that it would realize decreased cost due to reduced regulatory oversight of Rhodia's cleanup activities.

The North Carolina Department of Transportation (NCDOT) reported that it would realize cost savings due to reduced 1,1-DCE reporting and mapping requirements for eighteen Asphalt Testing Lab sites that had releases of chlorinated solvents where 1,1-DCE is now below 350 ppb. NCDOT does not own or operate these labs; however, NCDOT used their laboratory services and are now assisting with remediation efforts at these lab sites.

No new costs were identified with the proposed rules as a result of outreach activities; however, the cost of decommissioning groundwater remediation equipment, including closure of groundwater monitoring wells, would be realized sooner, rather than later, in the cleanup process as a result of a higher 1,1-DCE standard.

No impacts to local or federal government or small businesses were reported. It is possible that water supply companies and local government would incur costs if they choose to use a contaminated water source after a remedial action plan is already approved.

No health-based benefits are expected as a result of changing the groundwater standard for 1,1-DCE from 7 ug/L to 350 ug/L because the proposed standard of 350 ug/L is based on the most recent U.S. EPA IRIS health effects data available at http://www.EPA.gov/iris/subst/0039.htm. According to the U.S. EPA IRIS database, the chemical is less toxic than previously thought and is no longer considered a carcinogen by the oral (drinking water) route.

If Rhodia is the only company immediately affected by this rule change and no additional costs are placed on drinking water suppliers, the costs of this proposed rule change will be approximately \$5,800 in FY2012-13 and \$27,000 in FY2027-28 (adjusted for inflation). Rhodia may save money through the immediate closure of 11 wells and 15 fewer years of operation and maintenance costs. The net present value of this cost savings could be as high as \$945,000 throughout the next 30 years. State benefits, in the form of less reporting and mapping for NCDOT and less oversight by DWM, have an estimated net present value of \$30,000 over 30 years.

The risk analysis section of the fiscal analysis examines additional costs and benefits that may be incurred by additional companies or the need for more drinking water remediation as a result of the rule change. The benefit amount for private companies with releases of 1,1-DCE to groundwater hinges on whether or not 1,1-DCE is the only groundwater contaminant that will be responsible for requiring environmental cleanup. A second possible risk is that 1,1-DCE pollution will affect a source of drinking water. This may create additional costs for public or private water systems.

V. PUBLIC HEARING PROCESS

The EMC approved DWQ's request to proceed to public hearing with the proposed revisions to 15A NCAC 02L .0113 & .0202 and the fiscal note at its July 14, 2011 meeting (Agenda Item 11-27). EMC member Dickson Phillips was appointed to serve as the hearing officer. The hearing officer designation memo from Chairman Stephen T. Smith is provided in Attachment D. A public hearing regarding the proposed revisions and fiscal note was held on May 23, 2012, 6:30 PM at the Archdale Building, Ground Floor Hearing Room, 512 N. Salisbury Street, Raleigh.

The announcement of the hearing was published in the May 1, 2012 edition of the *North Carolina Register* (http://www.oah.state.nc.us/rules/register/Volume26Issue21May12012.pdf). The public hearing announcement in Attachment E was mailed either electronically or in hard copy form to all individuals on the DWQ Rulemaking mailing list and those on the DWQ rules Listserv. A news release of the hearing was sent out via the DENR and DWQ Public Information Officers on May 15th, 2012. Additionally, the announcement and background information was made available in electronic form on the following websites:

- DENR's searchable *Proposed Rules* website (http://portal.ncdenr.org/web/guest/rules),
- DWQ's searchable *Proposed Rules* website (http://portal.ncdenr.org/web/wq/rules),
- EMC's website under DENR's Proposed Rules (http://portal.ncdenr.org/web/emc/), and
- DWQ's Groundwater Standards Information website (http://portal.ncdenr.org/web/wq/ps/csu/gwstandards).

The public hearing date, time and location was posted on the DENR Public Event Calendar (http://portal.ncdenr.org/web/guest/event-calendar).

The sixty-day comment period for the proposed groundwater rules revisions began on May 1, 2012 and ended on July 2, 2012.

VI. ORAL AND WRITTEN COMMENTS RECEIVED

The proposed rule changes were presented at the public hearing held in Raleigh on May 23, 2012. The hearing officer's remarks and DWQ staff presentation is presented in Attachment F and Attachment G, respectively. Seventeen people attended the public hearing in Raleigh, including DENR staff. No oral or written comments were received at the hearing. A list of those attending the public hearings is provided in Attachment H. A complete digital audio recording of the hearing is available for review from DWQ Planning Section.

Written comments were accepted, in accordance with the North Carolina Administrative Procedures Act, until the close of the public comment period on July 2, 2012. Seven written statements were received by the end of the comment period on July 2, 2012. A list of all persons who submitted written comments is provided in Attachment I and copies of all the written comments are provided in Attachment J.

VII. SUMMARY OF ORAL AND WRITTEN COMMENTS

The following is a summary of the major issues and comments that were brought forth in regard to the proposed changes to the groundwater quality standards along with a brief staff response.

Option 1: Comments and Response

A change in the 1,1-dichloroethylene standard from 7 ug/L to 350 ug/L is proposed under Option 1 as requested in a rulemaking petition approved by the EMC in order to incorporate the most recent U.S. EPA health effects information as published in the Integrated Risk Management System (IRIS) at http://www.epa.gov/IRIS/.

Three of seven comments received stated that the EMC should amend the groundwater standard for 1,1-dichloroethylene from 7 ug/L to 350 ug/l, as requested by Rhodia, Inc. in its petition for rule-making, based on the availability of updated health effects data for 1,1-DCE published in the U.S. EPA Integrated Risk Information System (IRIS) database. Regulated parties will see a cost savings in assessment and cleanup of 1,1-DCE groundwater plumes when 1,1-DCE is the groundwater contaminant that drives assessment and cleanup costs at the site. Only one private company, Rhodia, Incorporated, was identified in the fiscal analysis as impacted by a change in the 1,1-DCE from 7 ug/L to 350 ug/L; however, during public outreach one party was identified as potentially saving approximately \$75,000 by avoiding installation of an additional triple well nest to determine the horizontal and vertical extent of a 1,1-DCE groundwater plume if the 1,1-DCE standard is changed to 350 ug/L. The fiscal analysis addresses the possibility of reduced assessment cost to companies other than Rhodia as the contaminant plume based on a standard of 350 ug/L will be less extensive than a contaminant plume based on a standard of 7 ug/L; therefore, no modifications to the fiscal note based on public comments are proposed.

Several public comments were received that objected to the proposed change in the 1,1-DCE standard from 7 ug/L to 350 ug/L primarily because it could adversely impact private drinking water wells, as well as public water supply wells that must meet the federal maximum contaminant level (MCL) based on the federal Safe Drinking Water Act and North Carolina Public Water Supply standards (15A NCAC 18C .1518), and it would not be protective of public health. Specific comments and staff responses are provided below:

- 1. Comment: Option 1 would be in conflict with the current process adopted by the EMC for setting groundwater standards, as outlined in 15A NCAC 02L .0202(d).
 - Response: Changing the 1,1-DCE standard alone as proposed in Option 1 without changing the groundwater rules as proposed in Option 2 would represent a departure from the current rules in 02L .0202(d). Option 2 was proposed to allow the EMC to depart from 02L .0202(d) under specific circumstances set out in the 02L .0202(f) proposed language in Attachment B. The goal of the proposed 02L .0202(f) language is to ensure that groundwater standards are based on the most recent U.S. EPA health effects data, are protective of public health, and, do not result in unnecessary costs for the regulated community.
- 2. Comment: Permitted discharges of 1,1-DCE above the MCL will impair a source (or

future source) of drinking water, unfairly shifting costs from the polluter to innocent public and private water systems and users. Until EPA revises the MCL for 1,1-DCE, the EMC should incorporate the actual federal regulatory MCL value to ensure NC's groundwater standards are sufficiently protective of health and the environment and will allow all water supplies to meet federal Safe Drinking Water Act standards and North Carolina Public Water Supply standards (15A NCAC 18C .1518).

Response: As discussed in the Fiscal Analysis in Attachment P, it is anticipated that if the groundwater standard for 1,1-DCE were raised from 7 ug/L to 350 ug/L, the total number of water systems affected would be small. However, there are specific circumstances in which the 1,1-DCE standard change may affect groundwater sources that are used for drinking water and create costs for public drinking water treatment. This could happen if groundwater remediated to the new standard is used as a source of drinking water in the future or if a responsible party for the pollution cannot be identified. Each of these instances is predicted to be rare in the current environment. Since responsible parties are usually identified, any local government or private water supply system would be protected from having to bear the cost of additional water treatment from 1,1-DCE pollution. A rough estimate of the costs associated with 1,1-DCE contamination to source water is included in the above referenced fiscal analysis.

3. Comment: The EMC should correct the flaws in the current published MCL for 1,1-DCE to reflect a safe level of 35 ug/L, using a safety factor of 10 to account for 1,1-DCE's potential carcinogenicity. Option 1 is not protective of human health.

Response: In the results of the *Second Six-year Review of the National Primary Drinking Water Standards* published in the March 29, 2010 Federal Register (http://www.gpo.gov/fdsys/pkg/FR-2010-03-29/pdf/2010-6624.pdf - beginning on page 15535), and in the *October 2009 U.S. EPA Six-Year Review 2 Health Effects Assessment: Summary Report*

(http://water.epa.gov/lawsregs/rulesregs/regulatingcontaminants/sixyearreview/second_re_view/upload/Six-Year-Review-2-Health-Effects-Assessment-Summary-Report.pdf), U.S. EPA acknowledged that the federal maximum contaminant level (MCL) of 0.007 mg/L (7 ug/L) was calculated using an outdated 1987 U.S. EPA IRIS reference dose of 0.01 mg/kg-day, along with a 10-fold risk management factor to account for the potential that 1,1-DCE is a possible carcinogen. Since that time, U.S. EPA has reassessed the health effects data for 1,1-DCE and has published a revised reference dose of 0.05 mg/kg-day in its IRIS database (2002). U.S. EPA's reassessment also determined that the animal and human data are not sufficient to conclude that 1,1-DCE is a human carcinogen by the oral or inhalation route of exposure (http://www.epa.gov/iris/subst/0039.htm) and that the risk management factor of 10, applied to the current MCL, is no longer needed and a possible MCL of 0.35 mg/L (350 ug/L) could be calculated.

The EMC's proposed standard for 1,1-DCE of 350 ug/L in Option 1 is the non-cancer systemic threshold concentration calculated in accordance with 15A NCAC 02L .0202(d)(1) using the current reference dose of 0.005 mg/kg/day published in EPA's IRIS database (8/13/2002). A cancer potency factor has not been established for 1,1-DCE;

therefore, a concentration which corresponds to an incremental lifetime cancer risk of one-in-a-million cannot be calculated in accordance with 15A NCAC 02L .0202(d)(2).

Based on U.S. EPA's reassessment of 1,1-DCE's toxicity and discussions in its *Second Six-year Review of the Primary Drinking Water Standards*, the proposed 1,1-DCE standard of 350 ug/L is protective of public health.

4. Comment: Option 1 is preferable to Options 2 and 3 in that it does not apply to contaminants other than 1,1-DCE or create state-wide exceptions from groundwater standards.

Response: Changing the 1,1-DCE standard alone as proposed in Option 1 without changing the groundwater rules as proposed in Option 2 could represent a departure from the current rules adopted by the EMC for setting groundwater quality standards in 15A NCAC 02L .0202(d) and result in a legal challenge to the proposed standard. It should be noted that Rhodia provided a legal opinion in its rulemaking petition that 15A NCAC 02L .0202(d) and (e), in tandem, are sufficiently broad to establish the groundwater standard at 350 ug/L under 02L .0202(d)(1), based on the current toxicity data published in the U.S. EPA's Integrated Risk Information System (IRIS) database, rather than establishing the standard at the federal MCL, which was calculated prior to the updated toxicity data being published. After consultation with its legal counsel, the EMC determine that 02L .0202(d) and (e) were not sufficiently broad to allow a change to the 1,1-DCE standard from7 ug/L to 350 ug/L as requested. If the EMC adopts Option 1, then Option 2 would need to be adopted as well to avoid the proposed 1,1-DCE standard of 350 ug/L being legally challenged.

It should be noted that adoption of Option 2 would still require rulemaking to update any other groundwater quality standards where an updated MCL would be higher than the current MCL.

5. Comment: Option 1 may provoke rulemaking petitions to change standards for other chemicals.

Response: Rulemaking petitions submitted in accordance with 15A NCAC 2I .0501 are an appropriate avenue to challenge 15A NCAC 02L groundwater standards.

To date, the 1,1-DCE standard is the only 15A NCAC 02L .0202 (g) groundwater standard for which a rulemaking petition has been submitted requesting a change in the standard. The request is based on the availability of more recent health effects data published in U.S. EPA's IRIS database that was not used to establish the current MCL, the level at which the current 1,1-DCE 2L groundwater standard is set. A review of U.S. EPA's IRIS database identified two other chemicals, tetrachloroethylene and methylene chloride, with updated health effects information that may result in a request to change the 02L groundwater standards to concentrations above their published federal MCLs.

6. Comment: Any issue with 1,1-DCE can be dealt with using the current variance process.

Response: In November 2010, Rhodia requested a variance to the 1,1-DCE groundwater standard. The Division of Waste Management (DWM), the regulatory authority over the site, reviewed the request and determined it to be incomplete based on the requirements in 15A NCAC 02L .0113. Attachment K is a copy of the January 21, 2011 DWM letter to Rhodia outlining the deficiencies in the variance request. Rhodia withdrew the 1,1-DCE standard variance request, stating that the variance approach was not an appropriate mechanism for seeking relief from a standard that was not based on current health effects information. Attachment L is a copy of Rhodia's March 13, 2011 letter to DWM withdrawing the variance request. The subsequent rulemaking petition held merit and was considered by the EMC.

Previous to the petition and in regard to the 1,1-DCE issue (May 2005), the EMC Groundwater Committee (GWC) directed the DWQ to establish a Groundwater Stakeholder Group (GWSG) to discuss ways to amend the groundwater rules to ensure the use of the most recent toxicity information when developing groundwater standards. The GWSG consisted of representatives from various stakeholder groups, such as, Department of Environment and Natural Resources (DENR) regulatory agencies, city and county governments, major industries, environmental groups, agricultural interests, and public health. After much discussion, the GWSG could not reach consensus on how to amend the groundwater regulations and ultimately, the DWQ recommended to the GWC that the issue be dealt with by the variance process in 15A NCAC 02L .0113. At its July 12, 2006 meeting, the GWC unanimously accepted and concurred with the DWQ recommendation that in individual site-specific cases, a variance granted under 02L .0113 could be approved that would allow less restrictive Groundwater Quality Standards while providing the site specific requirements necessary to protect public water supplies.

7. Comment: The Commission retains authority to set more stringent standards where necessary to protect North Carolina groundwaters, and is not constrained by the criteria set forth in 02L .0202(d). See 15A N.C. Administrative Code 02L .0103(b) (expressing the intent to protect all [North Carolina] groundwaters "to a level of quality at least as high as that required under the [groundwater] standards established in Rule .0202.") (Emphasis added). Accordingly, there is no legal basis for the EMC to adopt the less stringent groundwater standard as proposed.

Response: The basis for establishing groundwater standards at health-based limits is provided in 15A NCAC 02L .0202(a) which states "...They are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of state, which may be tolerated without creating a threat to human health or which would render the groundwater unsuitable for its intended best usage."

Option 2: Comments and Staff Response

Proposed changes to 15A NCAC 02L .0202 (d) and (f) under Option 2 would allow the EMC to establish a standard less stringent than the maximum contaminant level (MCL) or national secondary drinking water standard, after public notice and opportunity for hearing, when:

- a. MCL is not based on the most recent U.S. EPA health effects data,
- b. Such a standard would not endanger public health and safety, and,

c. Compliance with a standard based on the MCL would produce serious hardship without equal or greater public benefit.

No comments were received that fully supported this option. Rather, the comments recommended rejecting the proposed language in Option 2 or alternative language was proposed.

- 1. Comment: A rule adjustment such as Option 2 will allow the EMC to correct the standard referenced in Option 1. However, it may not be the best solution to the fundamental problem of overly restrictive standard selection rules using arbitrary criteria beyond the control of our regulatory agency. The following revision to 02L .0202(d) was proposed:
 - (d) Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of using evaluation of the reliability, relative costs and benefits of:
 - (1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)[/ [2 liters/day (avg. water consumption)]:
 - (2) Concentration which corresponds to an incremental lifetime cancer risk of 1 x 10⁻⁶;
 - (3) Taste threshold limit value;
 - (4) Odor threshold limit value;
 - (5) Maximum contaminant level; or
 - (6) National secondary drinking water standard.

Response: Use of the terms "reliability, relative costs and benefits of" in the proposed rule language would be problematic for several reasons. The definition for those terms and how they would be used to determine the appropriate, health-based, cost-effective standard would have to be included in the rule. Without clarification of the terms and their implementation, the Rules Review Committee (RRC) would not likely approve the proposed language based on General Statutes 150B-19.1(a)(3) and 150B-21.9 which require rules to be written in a clear and unambiguous manner.

Although the groundwater standards are established at health-based levels, there are provisions in 15A NCAC 02L .0106 that consider "relative costs and benefits" associated with restoration of groundwater to the level of the standards when remediation is required.

2. Comment: Data sources should not be restricted to only those sources that are from EPA, as is stated in the proposed (f) language. They should include references other than EPA as provided in .0202 (e)(4) of the current 02L rules.

Response: For establishing a groundwater standard less stringent than the existing MCL, the use of high quality, science-based and peer-reviewed human health assessments prepared and maintained by the U.S. EPA is preferred because U.S. EPA is also the agency that establishes MCLs.

3. Comment: Commenter disagrees with the requirement that "compliance with a standard

based on the maximum contaminant level or national secondary drinking water standard would produce serious hardship without equal or greater public benefit." They further state that "if the new standard is based on the most up-to-date science and is protective of public health, safety and the environment, there should not have to be a "hardship" test. The hardship test should be maintained as an individual variance requirement and not be made a requirement for a state-wide standard change based on more recent data and studies. We recommend that in Option 2, if adopted, (f)(l) be amended to reference all of the sources of data listed in (e), and we recommend that (f)(3) be deleted."

Response: The proposed language in 02L .0202 (f) sets out the criteria the EMC will use to decide when it is appropriate to establish a groundwater standard higher than an MCL. MCLs are enforceable drinking water standards for public water systems regulated under the federal Safe Drinking Water Act. If the EMC establishes a groundwater standard above an MCL, the potential exists to negatively impact source water used by public water systems that must meet MCLs. In order to protect groundwater at the highest possible level, the hardship test is needed to further aid the EMC in its decision to allow a groundwater standard to be set above an MCL.

4. Comment: Option 2 is overbroad in that it is not limited to 1,1-DCE, but applies to all 147 of the contaminants listed in current Subsection (g). Option 2 opens the door for future weakening rule revisions to any and all of the other 146 contaminants by applying the overly flexible standard suggested by this option.

Response: The purpose of Option 2 is to ensure that the 02L groundwater standards are based on the most up-to-date health effects data published by U.S EPA. Only those 02L standards set at an MCL or national secondary drinking water standard that are not based on the most recent U.S. EPA health effects data would be affected.

Currently there are 147 groundwater standards, fifty-nine (59) of which also have an MCL. Twenty (20) groundwater standards are equal to the MCL. Thirty-seven (37) groundwater standards are set lower than the MCL. Two (acrylamide and epichlorohydrin) of the 59 MCLs are set at treatment technique limits while the 02L groundwater standard for these two are established at a health-based concentration.

There are currently 58 interim maximum allowable concentrations (IMACs), eleven (11) of which have MCLs. Five (5) IMACs are set at the MCL and the remaining six (6) IMACs are set lower than the MCL.

There are ten (10) 02L groundwater standards established at the secondary drinking water standard. Secondary drinking water standards are based on organoleptic effects such as taste, odor and staining. None of the current IMACs are set at the secondary drinking water standard.

U.S. EPA IRIS health effects information published since the end of the last groundwater triennial review (January 1, 2010) was examined to determine if new data would affect a 15A NCAC 02L groundwater standard or IMAC. It appears that the most recent health

effects data published in the IRIS data for methylene chloride and tetrachloroethylene could result in a groundwater standard less stringent than the MCL if Option 2 is adopted. Any changes to those two standards would have to proceed through the rulemaking process.

The methylene chloride MCL and 02L groundwater standard is 5 ug/L (based on the cancer endpoint). In accordance with 15A NCAC 02L .0202(d)(2), updated cancer health effects data would result in a one-in-a-million risk concentration of 20 ug/L.

The tetrachloroethylene MCL is 5 ug/L and 02L groundwater standard is 0.7 ug/L (based on the cancer endpoint). In accordance with 15A NCAC 02L .0202(d)(2), updated cancer health effects data would result in a one-in-a-million risk concentration of 20 ug/L.

The updated health effects data for tetrachloroethylene and methylene chloride were not available for the U.S. EPA's Second Six-year Review of Drinking Water Standards. The U.S. EPA has set the maximum contaminant level goal (MCLG), the level at which no adverse health effects are expected, to zero for these chemicals because they are classified as carcinogens. However, the current MCL for these chemicals is established at their laboratory practical quantitation limit of 5 ug/L. If updating the MCLs based on the new IRIS data presents a meaningful opportunity for cost savings while maintaining, or improving, the level of public health protection, then it is possible that U.S. EPA will update the MCLs for these chemicals. The next six-year review is not scheduled until 2015 and not likely to be complete before 2016

(http://water.epa.gov/lawsregs/rulesregs/regulatingcontaminants/sixyearreview/index.cfm).

5. Comment: This particular revision (Option 2) cannot be implemented without placing human health at risk. The current rule (15A NCAC 02L .0202) provides in Subsection (d) six criteria for the EMC to compare in establishing groundwater quality standards, the most stringent of which is to be adopted. Since the MCL represents the highest level at which a contaminant might exist in the groundwater without resulting in short-or long-term health risks, there is no way to allow a weaker, riskier standard to usurp the MCL and not endanger the public health and safety.

Response: Proposed Option 2 would allow the EMC to adopt a groundwater standard less stringent that an MCL *only* when the MCL is not based on the most recent U.S. EPA health effects data. A groundwater standard protective of noncancer and cancer effects would be established in accordance with 15A NCAC 02L .0202 d(1) and d(2) criteria even though it may be higher than an MCL that has not been adjusted to reflect the most up-to-date health effects data. In the case of 1,1-DCE, U.S. EPA acknowledges in its second six-year review of primary drinking water standards that a safe level could be established as high as 350 ug/L, using the most recent health effects data published in its IRIS database.

6. Comment: It makes little sense to target both the MCL and the national secondary drinking water standards, as Option 2 does, given that the focus of the current rulemaking efforts is 1,1-DCE, a contaminant without a national secondary drinking water standard.

Response: The national secondary drinking water standards are established by the U.S. EPA as guidelines to assist public water systems in managing their drinking water for aesthetic considerations, such as taste, color and odor. Updated U.S. EPA sources of taste and odor thresholds may be published in the future that provide a more appropriate threshold than the national secondary drinking water standard.

- 7. Comment: Option 2 should be rejected and any rule change should be limited to 1,1-DCE. While the EMC may be comfortable with the result in the present situation involving Rhodia's discharges of 1,1-DCE, there is no way of knowing what the result would be with other groundwater standard changes that might be made pursuant to Option 2 in the future, nor what the cost might be for water suppliers and users to comply with the Safe Drinking Water Act standards. For reference:
 - Safe Drinking Water Act
 (http://water.epa.gov/lawsregs/rulesregs/sdwa/index.cfm)
 - NC Drinking Water Standards (15A NCAC 18C .1518)
 http://ncrules.state.nc.us/ncac/title%2015a%20-
 https://ncrules.state.nc.us/ncac/title%2015a%20-
 https://ncrules.state.nc.us/ncac/title%2015a%20-
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Response: Any future increase in a groundwater standard above an MCL as a result of changes to 15A NCAC 02L .0202(d) and (f) will be supported by the use of the most recent health effects data; therefore, increased adverse health effects are not expected. Potential costs to water suppliers in regard to 1,1-DCE were addressed in the risk analysis section of the fiscal analysis for this proposed rule. For future changes to a groundwater standard above an MCL, the potential costs to a water supplier will be evaluated and considered during the rulemaking process. See also Response to Comment 2 under Option 1.

8. Comment: The only company known to be affected by the 1,1-DCE standard is Rhodia. Unlike Rhodia, which according to DWQ has direct releases of 1,1-DCE, most companies that discharge 1,1-DCE release it together with other more toxic chlorinated solvents above their respective groundwater standards, and in much higher concentrations than 1,1-DCE. The presence of these other regulated chemicals would likely trigger more complex and costly environmental cleanup efforts, thereby eliminating most, if not all, of the benefits of amending the 1,1-DCE groundwater standard for most companies because it is not the pollutant driving their cleanups.

Response: Agreed.

9. Comment: Rhodia itself admits that it is not requesting a change in the 1,1-DCE standard or any variance "on the grounds that compliance with the 1,1-DCE standard [is] technically infeasible or would cause undue economic hardship"; rather, the company is requesting the rule revisions to reduce its compliance costs, including remediation costs

and groundwater investigation and monitoring costs. But reducing compliance costs does not fall within the stated goals of North Carolina groundwater policy (15A NCAC 02L .0103(a), (b)) and there are no health-based benefits of the proposed rule changes. The rule revisions are therefore unjustified.

Response: 15A NCAC 02L rules do not list reduction of cost as a goal. However, these proposed rule revisions were based upon a Rulemaking Petition submitted by Rhodia. Under the state's Administrative Procedures Act, NC General Statute 150B-19.1 (Requirements for agencies in the rule-making process), any potential changes to existing rules are subject to "be designed to achieve the regulatory objective in a cost efficient and timely manner". NC GS150B-19.1 also requires the agency to "seek to reduce the burden upon those persons or entities who must comply with the rule." The proposed rules are designed to comply with the General Statutes and also protect public health.

Option 3: Comments and Staff Response

Changes to 15A NCAC 02L .0113 under Option 3 are proposed as follows:

- (b) Update the DWQ mailing address to *Division Water Quality*, 1617 Mail Service Center, Raleigh, N.C. 27699-1617,
- (c) Add "for site-specific variances" language to distinguish between proposed language in paragraph (d) that would allow a state-wide variance.
- (d) Add language to allow the EMC to approve a state-wide variance to a groundwater standard established in Section .0202 after consideration of information required in the application pertaining to public health and safety, potentially affected sites and increased cost for treatment for wells or water supply sources due to the proposed variance.
 - (h) (i) Clarify language. No substantive change.
- (i) Remove current paragraph (i) that states: "A variance shall not operate as a defense to an action at law based upon a public or private nuisance theory or any other cause of action."

Three comments were received in support of Option 3 if it is modified as requested. One comment was received in opposition to Option 3. The majority of comments received were in regard to the proposed language in 15A NCAC 02B .0113(d) that would allow the EMC to approve a state-wide variance to groundwater standards. These comments identified a number of potential problems with the approval and implementation of a state-wide variance to groundwater standards; therefore, this portion of Option 3 is not considered a viable option for addressing groundwater standards that are not based on the most recent health effects data, and is not recommended for adoption by the EMC. Specific comments and DWQ staff responses are provided below.

1. Comment: Option 3 is a cumbersome and unnecessary response to poorly worded current regulations that may impede reasonable and truly protective standards.

Response: There is significant information required related to the request for a state-wide variance. Information on all known potentially affected sites and a list of increased costs for treatment of groundwater drinking water sources, while time consuming and costly to produce, provides clarity for all regulatory agencies involved with protection of groundwater regarding the location and potential impacts that might be expected. The

required information is important and necessary in order for the EMC to make an informed decision on the impact of a requested state-wide variance to a groundwater standard.

2. Comment: The use of the term "state-wide" in 02L .0113 (d) implies that the variance would apply everywhere within the state. However, Paragraph (2) requires a list of all known potentially affected sites, implying that the variance would only be applied to those specific sites.

Response: The intent of the proposed language is to allow a state-wide variance from a standard that would apply to all sites throughout the state. If a site is not listed in the application, it would not preclude the variance from being applied to that site.

3. Comment: It should be clarified how the variance would apply to sites discovered after the variance is adopted or known by any other than the applicant.

Response: The variance would apply to sites discovered after a state-wide variance to a groundwater standard was approved by the EMC. An approved state-wide variance would be public noticed in the NC Register, posted on the DWQ web site, and notices would be sent out via email to the DWQ rulemaking and groundwater stakeholder list servers.

4. Comment: Paragraph (3) requires a "list of increased costs for treatment ...due to the proposed variance." Normally, a variance is requested to allow a lower cost to the applicant, but may also result in additional expense for those who use affected wells for water supply. Both factors should be considered. Rather than using a confusing "statewide variance", it is much more practical to establish appropriate state standards to begin with, and provide for specific corrections when an incorrect standard is encountered.

Response: It is implicit that a request for a state-wide variance would be made to allow for lower cost to the applicant and others with potentially affected sites; however, the 15A NCAC 02L .0113 (d) requirements, as written, may be overly burdensome for the applicant.

5. Comment: Option 3 in (d) requires an "application." That word appears to preclude the Commission from granting a variance "on its own initiative" [02L .0113(a)]. Option 3 should state that, in addition to an application from another party, the Commission may initiate a state-wide variance without an "application" pursuant to 02L .0113(a).

Response: This language was not modified from the current rule language and has not been considered to preclude the Commission from moving forward with variances under its own initiative per 02L .0113(a).

6. Comment: Paragraph (d)(2) may be impossible for an applicant or set of applicants (or even the Division of Water Quality) to produce because it may be impossible to produce a list of all known "potentially" affected sites for common substances. Also, (d)(3) is as equally difficult to produce in an application. For (d)(3), if it is necessary to retain this, an

estimated range of costs of treatment for different water sources might be more appropriate than trying to develop cost estimates for "all known potentially affected sites."

Response: The information requirements in 02L .0113 (d)(2) may be difficult, and costly, for an applicant to produce; Conversely, not identifying state-wide sites could be more costly should wells or water supply sources not identified become contaminated. An estimated range of costs of treatment for water supply sources identified in (d)(2) would be considered an acceptable response to (d)(3).

7. Comment: If up-to-date science and data evaluated by the Commission shows no endangerment to public health and safety and to the environment, then that should be sufficient for a state-wide variance. The Commission should make (d)(2) and/or (d)(3) optional at the discretion of the Commission. The Commission should do a preliminary evaluation of the proposed state-wide variance and determine if the Commission wishes the applicant to produce the information asked for in (d)(2) and (d)(3). Also the rule should state that the Commission has the discretion to request such information and to determine how much information it wants the applicant to produce for each. The Commission should be able to exercise discretion regarding what amount (if any) of the information in (d)(2) and (d)(3) the Commission feels it needs to make the determinations necessary for a state-wide variance.

Response: There may be merit to providing the EMC discretion to determine how much information it wants an applicant to produce when applying for a state-wide variance; however, if a state-wide variance applicant requests a groundwater standard that contravenes an MCL, then the information requested in 02L .0113 (d)(3) is needed to address impacts to water supply systems. However, approval of this type of language by the Rules Review Commission could be problematic as well under North Carolina General Statute (NCGS) 150B-19.1(a)(3) and NCGS 150B-21.9, both of which requires rules to be written in a clear and unambiguous manner.

8. Comment: The current variance process is sufficient to resolve the issue with 1,1-DCE and all proposed rules should be rejected (See Response to Comment # 6 under Option 1). If the EMC determines a rule revision is appropriate, the commenter urges it to select Option 3 with the following modifications to limit its scope and impact to the greatest degree possible. For the most part, in the event the EMC decides, against our recommendation, and selects Option 2, then these modifications could be made to that option as well.

Modification 1: First and foremost, because of the unique situation regarding the 1,1-DCE groundwater standard, any rule change should be expressly limited to 1,1-DCE. This will avoid unintended consequences by preventing any revised rule from allowing changes to groundwater standards for other regulated chemicals that do not share 1,1-DCE's circumstances.

Response: The EMC's counsel has stated that Option 1 is not a legally viable option

by itself.

Two groundwater standards that share 1,1-DCE's unique circumstance have been identified as a result of staff review of the U.S. EPA IRIS health effects information published since the end of the last groundwater triennial review (January 1, 2010). The most recent health effects data published in the IRIS data for methylene chloride and tetrachloroethylene could result in a groundwater standard less stringent than its MCL if Option 2 or Option 3 is adopted.

Modification 2: As the rule revisions are currently drafted, neither Option 2 nor Option 3 imposes the same preference for the EPA's IRIS database as does the standard for adopting groundwater standards and setting concentrations in the current 15A NCAC 02L .0202(e). As Rhodia acknowledges in its petition, "IRIS is the most preferred reference source for the establishment of groundwater standards. If adopted, the proposed rule should be revised to clarify that (1) the hierarchy, or order of preference, of EPA health references listed in section 2L .0202(e) applies as well in the proposed new section 2L; .0113(d) (or the new 2L .0202(f) if the EMC selects Option 2); and (2) the value from the EPA health reference that is most protective of human health is the preferred source of information."

Response: There is merit to revising the language in Option 2 (f)(1) and Option 3 (d)(1) to require that the hierarchy of health references be the same as required in 02L .0202(e).

Modification 3: When considering whether a variance would endanger public health and safety, we urge the EMC to require the consideration of North Carolina's most vulnerable populations and the impacts that a variance (or loosened groundwater standard) might have on these individuals. This approach is consistent with the approach taken under the Safe Drinking Water Act (SDWA), which requires the MCLG (maximum contaminant level goal) to be set at "the level of a contaminant in drinking water below which there is no known or expected health risk," with a margin of safety. The MCLGs must "take into account the risks of exposure for certain sensitive populations, such as infants, the elderly, and persons with comprised immune systems." Consistent with the policy behind NC's groundwater standards to protect drinking water for all of the state's residents, this precautionary approach should be incorporated in any rule change. (Safe Drinking Water Act regulations are located at: http://water.epa.gov/lawsregs/rulesregs/sdwa/index.cfm)

Response: When establishing groundwater standards, it is important to protect North Carolina's most sensitive populations. This protection is provided when a systemic threshold concentration as required in 15A NCAC .0202(d)(1) is calculated using a reference dose, which includes a margin of safety (usually a factor of 10) to protect sensitive subgroups (http://www.epa.gov/iris/rfd.htm). Supporting information required in 15A NCAC 02B.0113 (d)(1) should provide for adequate protection of sensitive populations.

Modification 4: If revised, the rule should also take into account the concept of "relative source contribution," that is, the idea that people can be exposed to additional amounts of a contaminant from sources other than their drinking water, such as food or air. Just as EPA accounts for these other contributions when calculating MCLGs, the EMC should revise the proposed rule so that relative source contribution is considered when determining whether a variance is appropriate.

Response: Relative source contribution is the fraction of total intake of the contaminant that is typically associated with water (as opposed to food, air, and other specific sources). The noncancer threshold concentration for a groundwater standard in 02L .0202(d)(1) is calculated using a default relative source contribution of 0.2 (twenty percent) for organic contaminants and 0.1 (ten percent) for inorganic contaminants, which are conservative estimates. Supporting information required in 15A NCAC 02L.0113 (d)(1) should include the application of an appropriate relative source contribution.

Modification 5: If Option 3 is selected, a state-wide variance should apply only to the particular applicant and only to those sites identified by the applicant in a variance application. Site-specific information would be central to EMC's evaluation of whether a state-wide variance would endanger public health and safety. Therefore, it is not appropriate to apply any state-wide variance to companies or sites not included in the application and its analysis.

Response: The intent of the proposed language in Option 3- 02L.0113 (d) is to allow a variance to a groundwater standard that would be applicable to all affected sites throughout the state as long as the requirements of the proposed language have been met. The commenter makes a good point in that site-specific information for all affected sites is critical when evaluating an application to ensure the state-wide variance will not endanger the public health and safety. Under this proposed option, the applicant will not be able to provide the needed information for future sites not yet identified at the time a state-wide variance is approved.

9. Comment: Similar to Option 2, the impacts of Option 3 would reach well beyond the groundwater standard for 1,1-DCE. The proposed state-wide variance provision would allow a chemical company to obtain a blanket exception to any groundwater quality standard, not just 1,1-DCE.

Response: The intent of the proposed variance provision is not to allow a responsible party to obtain a blanket exception to all groundwater standards. Rather, the purpose of allowing the EMC to approve a state-wide variance to a groundwater standard established in accordance with 02L .0202(d) is to make an exception to a groundwater standard only when it is not based on the most recent health effects data.

10. Comment: If Option 3 is selected, no variance should be permitted to operate as a defense to an action at law. Section 2L .01 13(i), as currently codified, provides that "a variance shall not operate as a defense to an action at law based upon a public or private nuisance

theory or any other cause of action." The proposed rule changes would remove this provision and create uncertainty for affected parties regarding available relief under the law for injuries sustained from discharges made under a variance. We do not support the removal of this language.

Response: The Hearing Officer and Division Staff concur that removal of this language is not in the public's best interest and support maintaining 02L .0113 as currently written.

General Comments and Staff Response:

Comment: The introduction of 1,1-DCE into the environment can lead to a more expansive, complicated cleanup necessitated by the breakdown products of 1,1-DCE, including the formation of vinyl chloride, a much more toxic chemical.

Response: Agreed.

Comment: 1,1-DCE evaporates or volatilizes from water into the air. Citizens who use groundwater as a source of drinking water can be exposed to 1,1-DCE not only from drinking the groundwater but also from inhalation of volatile contaminates when cooking, laundering or bathing. 1,1-DCE can be absorbed by the skin; if home water supplies are contaminated, people may absorb the chemical through their skin when bathing or washing dishes.

Response: Agreed. The current 02L rules do not address dermal or inhalation routes of exposure.

VIII. SUMMARY OF ALTERNATE PROPOSALS

In addition to collecting comments on the three proposed rule options, the EMC also collected comments on other proposals that allow flexibility in implementation of 15A NCAC 02L .0202(d).

Three of the proposals received recommended flexibility in the use of the six criteria used to establish in 02L .0202(d). Rather than setting standards "as the least of" the six criteria, these three proposals offered language that would allow the consideration of costs and benefits and best available scientific guidance in determining which of the criteria to use to establish the standard. The language for these alternate proposals is presented in Attachment M.

Response: This proposed language is too broad in scope and does not provide transparency and clarity in the establishment of groundwater standards. Approval of this type of language by the Rules Review Commission could be problematic as well under General Statutes 150B-19.1(a)(3) and 150B-21.9 which require rules to be written in a clear and unambiguous manner.

A fourth proposal suggested two approaches that the EMC could use to allow flexibility in establishing a 1,1-DCE groundwater standard. The first approach involves the EMC's status as a quasi-judicial entity and ability to take judicial notice provided under General Statute 143B-282.1. The EMC, as a quasi-judicial entity, could take "judicial notice" of the fact that the MCL currently published by U.S. EPA is technically incorrect and use the same formula as U.S. EPA

in calculating a corrected MCL of 35 ug/L. The full comments for this approach can be found in Attachment K as part of the comments received from the North Carolina Conservation Network.

Response: It is not clear if this approach will actually work because judicial notice does not appear to be available until a decision enters a quasi-judicial phase. In order to enter into the quasi-judicial phase, Rhodia may have to apply for a permit based on the 35 ug/L standard, have it denied and then appeal the decision to the Division Director.

The second approach would add the following language as a new Subsection (f) in order to revise the current 1,1-DCE standard to the technically correct MCL:

In cases where the U.S. EPA has delayed revising the maximum contaminant level despite having updated reference dose and other applicable data, and because of the administrative costs of a federal rulemaking proceeding, the Commission may adjust the Class GA and Class GSA standards in Subsections (h) and (i), respectively, to reflect the technically correct maximum contaminant level. Upon U.S. EPA revisions, the EMC will ensure that the adjusted maximum contaminant level remains consistent with the U.S. EPA's revised standard.

Response: The proposed language, while eloquently stating the current situation and preferred outcome, is too broad to implement. In addition, this proposed approach is basically embodied in Option 2.

Comment: One commenter requested that the EMC consider revising the criteria listed in 15A NCAC 02L .0202(d)(1) to reflect updated science on rates of human water consumption.

Response: The value of 2 liters per day for drinking water is currently used by the Office of Water in setting drinking water standards. In addition, 2 liters is close to the 90th percentile for drinking water ingestion according to the U.S. EPA 2011 Exposure Factors Handbook (http://www.epa.gov/ncea/efh/pdfs/efh-complete.pdf) and is comparable to the 8 glasses of water per day historically recommended by health authorities. The water ingestion rate appropriately includes water ingested directly as a beverage and water added to food.

NCDENR Alternate Proposal:

After evaluation of public comments, staff of the Divisions of Water Quality, Waste Management and Water Resources (hereinafter, DENR) propose a combination of Options 1 and 2, as originally noticed, with additional language to require the use of the federal MCL as the groundwater standard where public water systems and private drinking wells may be impacted. This combination addresses Rhodia's Rulemaking Petition request, results in a change to the 1,1-DCE groundwater standard from 7 ug/L to 350 ug/L and adds rule language to allow a groundwater standard to be established at a concentration less stringent than the MCL as long as private wells and public water systems are not impacted above the MCL. The change would also allow other groundwater standards to be established above their respective MCLs where updated health effects data is available. For example, tetrachloroethylene and methylene chloride could be revised through future rulemaking from 5 ug/L to 20 ug/L.

DENR received comments on the potential impacts to public water supplies and private wells associated with the proposed rules. After consideration of those comments, DENR acknowledges issues for private and public water supplies if they are contaminated at levels above the MCL. Primarily, additional treatment costs may fall upon impacted private and public entities instead of being addressed by the responsible party. Subsequently, DENR recommends including additional language in 02L .0202 (b) that provides the Division director an exception from the standards established in 02L .0202 for the explicit protection of public water systems and private drinking wells impacted in exceedance of the MCL. When listed as an exception to the statewide standard, with specific conditions as to its appropriate applicability, DENR believes that the rule provides a suitable mechanism of considering the use of the water for public consumption and addresses a potential economic issue for any affected water supply in the future. It should be noted that preliminary research used to develop the fiscal analysis for this rulemaking indicated that the total number of water systems potentially affected by the change to 1,1-DCE would be small or none.

The additional language proposed in .0202(b)(4) is as follows:

Where the groundwater standard for a substance is greater than the Maximum Contaminant Level (MCL), the Director shall apply the MCL as the groundwater standard at any private drinking water well or public water system that may be impacted.

Attachment N contains this DENR suggested rule language in combination with Options 1 and 2 as discussed above.

Implementation of subparagraph (b)(4) would be achieved through the corrective action process in 15A NCAC 02L .0106. Rule .0106 includes requirements to conduct site assessments which include receptor surveys (.0106(g)(3)) that would identify any impacted drinking water well. Similar requirements for receptor surveys are included in other groundwater remediation programs administered by DWM.

IX. RECOMMENDATIONS

Hearing Officer's Recommendation:

Following a careful and comprehensive review of all of the submitted written comments, supporting data, and attachments to this record, the hearing officer recommends that the North Carolina Environmental Management Commission adopt a variation of the Option 3 rule language in 02L .0113. This recommendation entails modifying the variance rules to allow for a site-specific variance approach specifically for the 02L .0200 Groundwater standards rules. This would change the originally proposed 02L .0113 to allow the EMC to grant a "site-specific" variance in cases where the MCL is not established using the most recent US EPA health effects information. This is accomplished by adding language publicly noticed in Option 2. In addition, it includes a requirement that the variance not result in an exceedance of a MCL in an impacted drinking water well or public water system.

Attachment O contains the hearing officer's recommended rule language.

This proposal keeps in place the current way groundwater quality standards are established in .0202 (d), which is a predictable and consistent process that adequately protects public health and groundwater for use as a drinking water source. Other benefits to this proposal include:

- Protection of private drinking wells and public water systems so that they do not bear the cost of litigation and/or treatment if source groundwater is contaminated above an MCL;
- Identification of private drinking wells and public water systems that could be impacted so that they can be protected;
- Requires no rulemaking; and,
- Addresses public concerns brought forward during public comment period.

The hearing officer acknowledges that this proposal does not satisfy Rhodia's rulemaking petition.

The hearing officer also recommends that the Commission adopt the following proposals:

- Amendment to .0113 (b) to update the Division's mailing address.
- Amendment to .0113 (h) to clarify that a variance applicant may appeal the Commission's final decision.

The hearing officer recommends that the Commission *not* adopt the following proposals:

- Amendment to .0113 that would allow the Commission to grant a state-wide variance to groundwater standards.
- Amendment to .0113(i) that would delete the current rule language.
- Option 1 amendment to .0202 that would change the 1,1-DCE groundwater standard from 7 ug/L to 350 ug/L.
- Option 2 amendments to .0202 that would allow the Commission to establish a groundwater standard that is less stringent than the MCL.

Notwithstanding the foregoing, should the Environmental Management Commission not adopt the Hearing Officer's recommendation of a site specific variance option, the Hearing Officer recommends that the Commission adopt the NCDENR alternate proposal outlined on pages 23-24 of this report and as provided in Attachment N of this report.

X. ATTACHMENTS

15A NCAC 02L .0202 is proposed for amendment as follows: (Option 1)

15A NCAC 02L .0202 GROUNDWATER QUALITY STANDARDS

- (a) The groundwater quality standards for the protection of the groundwaters of the state are those specified in this Rule. They are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of the state, which may be tolerated without creating a threat to human health or which would otherwise render the groundwater unsuitable for its intended best usage.
- (b) The groundwater quality standards for contaminants specified in Paragraphs (g) and (h) of this Rule are as listed, except that:
 - (1) Where the standard for a substance is less than the practical quantitation limit, the detection of that substance at or above the practical quantitation limit constitutes a violation of the standard.
 - (2) Where two or more substances exist in combination, the Director shall consider the effects of chemical interactions as determined by the Division of Public Health and may establish maximum concentrations at values less than those established in accordance with Paragraphs (c), (g), or (h) of this Rule. In the absence of information to the contrary, in accordance with Paragraph (d) of this Rule, the carcinogenic risks associated with carcinogens present shall be considered additive and the toxic effects associated with non-carcinogens present shall also be considered additive.
 - (3) Where naturally occurring substances exceed the established standard, the standard shall be the naturally occurring concentration as determined by the Director.
- (c) Except for tracers used in concentrations which have been determined by the Division of Public Health to be protective of human health, and the use of which has been permitted by the Division, substances which are not naturally occurring and for which no standard is specified shall not be permitted in concentrations at or above the practical quantitation limit in Class GA or Class GSA groundwaters. Any person may petition the Director to establish an interim maximum allowable concentration for a substance for which a standard has not been established under this Rule. The petitioner shall submit relevant toxicological and epidemiological data, study results, and calculations necessary to establish a standard in accordance with Paragraph (d) of this Rule. Within three months after the establishment of an interim maximum allowable concentration for a substance by the Director, the Director shall initiate action to consider adoption of a standard for that substance.
- (d) Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of:
 - (1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)];
 - (2) Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6;
 - (3) Taste threshold limit value;
 - (4) Odor threshold limit value;
 - (5) Maximum contaminant level; or
 - (6) National secondary drinking water standard.
- (e) The following references, in order of preference, shall be used in establishing concentrations of substances which correspond to levels described in Paragraph (d) of this Rule.
 - (1) Integrated Risk Information System (U.S. EPA).
 - (2) Health Advisories (U.S. EPA Office of Drinking Water).
 - (3) Other health risk assessment data published by U.S. EPA.
 - (4) Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.
- (f) Groundwater quality standards specified in Paragraphs (g) and (h) of this Rule and interim maximum allowable concentrations established pursuant to Paragraph (c) of this Rule shall be reviewed on a triennial basis. Appropriate modifications to established standards shall be made in accordance with the procedure prescribed in Paragraph (d) of this Rule where modifications are considered appropriate based on data published subsequent to the previous review.
- (g) Class GA Standards. Where not otherwise indicated, the standard refers to the total concentration in micrograms per liter of any constituent in a dissolved, colloidal or particulate form which is mobile in groundwater. This does not apply to sediment or other particulate matter which is preserved in a groundwater sample as a result of well construction or sampling procedures. The Class GA standards are:
 - (1) Acenaphthene: 80;
 - (2) Acenaphthylene: 200;

1	(3)	Acetone: 6 mg/L;
2	(4)	Acrylamide: 0.008;
3	(5)	Anthracene: 2 mg/L;
4	(6)	Arsenic: 10;
5	(7)	Atrazine and chlorotriazine metabolites: 3;
6	(8)	Barium: 700;
7	(9)	Benzene: 1;
8	(10)	Benzo(a)anthracene (benz(a)anthracene): 0.05;
9	(11)	Benzo(b)fluoranthene: 0.05;
10	(12)	Benzo(k)fluoranthene: 0.5;
11	(13)	Benzoic acid: 30 mg/L;
12	(14)	Benzo(g,h,i,)perylene: 200;
13	(15)	Benzo(a)pyrene: 0.005;
14	(16)	Bis(chloroethyl)ether: 0.03;
15	(17)	Bis(2-ethylhexyl) phthalate (di(2-ethylhexyl) phthalate): 3;
16	(18)	Boron: 700;
17	(19)	Bromodichloromethane: 0.6;
18	(20)	Bromoform (tribromomethane): 4;
19	(21)	n-Butylbenzene: 70;
20	(22)	sec-Butylbenzene: 70;
21	(23)	tert-Butylbenzene: 70;
22	(24)	Butylbenzyl phthalate: 1 mg/L;
23	(25)	Cadmium: 2;
24	(26)	Caprolactam: 4 mg/L;
25	(27)	Carbofuran: 40;
26	(28)	Carbon disulfide: 700;
27	(29)	Carbon tetrachloride: 0.3;
28	(30)	Chlordane: 0.1;
29	(31)	Chloride: 250 mg/L;
30	(32)	Chlorobenzene: 50;
31	(33)	Chloroethane: 3,000;
32	(34)	Chloroform (trichloromethane): 70;
33	(35)	Chloromethane (methyl chloride): 3;
34	(36)	2-Chlorophenol: 0.4;
35	(37)	2-Chlorotoluene (o-chlorotoluene): 100;
36	(38)	Chromium: 10;
37	(39)	Chrysene: 5;
38	(40)	Coliform organisms (total): 1 per 100 milliliters;
39	(41)	Color: 15 color units;
40	(42)	Copper: 1 mg/L;
41	(43)	Cyanide (free cyanide): 70;
42	(44)	2, 4-D (2,4-dichlorophenoxy acetic acid): 70;
43	(45)	DDD: 0.1;
44	(46)	DDT: 0.1;
45	(47)	Dibenz(a,h)anthracene: 0.005;
46	(48)	Dibromochloromethane: 0.4;
47	(49)	1,2-Dibromo-3-chloropropane: 0.04;
48	(50)	Dibutyl (or di-n-butyl) phthalate: 700;
49	(51)	1,2-Dichlorobenzene (orthodichlorobenzene): 20;
50	(52)	1,3-Dichlorobenzene (metadichlorobenzene): 200;
51	(53)	1,4-Dichlorobenzene (paradichlorobenzene): 6;
52	(54)	Dichlorodifluoromethane (Freon-12; Halon): 1 mg/L;
53	(55)	1,1-Dichloroethane: 6;
54	(56)	1,2-Dichloroethane (ethylene dichloride): 0.4;
55	(57)	1,2-Dichloroethene (cis): 70;

```
1
               (58)
                        1,2-Dichloroethene (trans): 100;
 2
               (59)
                       1,1-Dichloroethylene (vinylidene chloride): 7350;
 3
               (60)
                        1,2-Dichloropropane: 0.6;
 4
               (61)
                        1,3-Dichloropropene (cis and trans isomers): 0.4;
 5
               (62)
                       Dieldrin: 0.002;
 6
                       Diethylphthalate: 6 mg/L;
               (63)
 7
               (64)
                        2,4-Dimethylphenol (m-xylenol): 100;
 8
               (65)
                       Di-n-octyl phthalate: 100;
 9
               (66)
                        1,4-Dioxane (p-dioxane): 3;
10
                       Dioxin (2,3,7,8-TCDD): 0.0002 ng/L;
               (67)
                        1,1- Diphenyl (1,1,-biphenyl): 400;
11
               (68)
12
               (69)
                       Dissolved solids (total): 500 mg/L;
13
               (70)
                        Disulfoton: 0.3;
                       Diundecyl phthalate (Santicizer 711): 100;
14
               (71)
15
               (72)
                        Endosulfan: 40;
16
               (73)
                       Endrin, total: (includes endrin, endrin aldehyde and endrin ketone): 2;
17
               (74)
                        Epichlorohydrin: 4;
18
               (75)
                       Ethyl acetate: 3 mg/L;
19
               (76)
                       Ethylbenzene: 600;
20
               (77)
                       Ethylene dibromide (1,2-dibromoethane): 0.02;
21
                       Ethylene glycol: 10 mg/L;
               (78)
22
               (79)
                       Fluoranthene: 300;
23
               (80)
                       Fluorene: 300;
24
                       Fluoride: 2 mg/L;
               (81)
25
               (82)
                       Foaming agents: 500;
26
               (83)
                        Formaldehyde: 600;
27
               (84)
                        Gross alpha (adjusted) particle activity (excluding radium-226 and uranium): 15 pCi/L;
28
               (85)
                        Heptachlor: 0.008;
29
               (86)
                       Heptachlor epoxide: 0.004;
30
               (87)
                        Heptane: 400;
31
               (88)
                        Hexachlorobenzene (perchlorobenzene): 0.02;
32
               (89)
                        Hexachlorobutadiene: 0.4;
33
               (90)
                        Hexachlorocyclohexane isomers (technical grade): 0.02;
                        n-Hexane: 400;
34
               (91)
35
               (92)
                        Indeno(1,2,3\text{-cd})pyrene: 0.05;
36
               (93)
                        Iron: 300;
37
               (94)
                        Isophorone: 40;
38
               (95)
                       Isopropylbenzene: 70;
39
               (96)
                        Isopropyl ether: 70;
40
               (97)
                       Lead: 15;
41
               (98)
                        Lindane (gamma hexachlorocyclohexane): 0.03;
42
               (99)
                        Manganese: 50;
43
                        Mercury: 1;
               (100)
                        Methanol: 4 mg/L;
44
               (101)
45
               (102)
                        Methoxychlor: 40;
                        Methylene chloride (dichloromethane): 5;
46
               (103)
                        Methyl ethyl ketone (2-butanone): 4 mg/L;
47
               (104)
48
               (105)
                        2-Methylnaphthalene: 30;
                        3-Methylphenol (m-cresol): 400;
49
               (106)
50
               (107)
                        4-Methylphenol (p-cresol): 40;
51
                        Methyl tert-butyl ether (MTBE): 20;
               (108)
52
               (109)
                        Naphthalene: 6;
53
               (110)
                        Nickel: 100;
54
                        Nitrate: (as N) 10 mg/L;
               (111)
55
               (112)
                        Nitrite: (as N) 1 mg/L;
```

```
1
               (113)
                        N-nitrosodimethylamine: 0.0007;
 2
               (114)
                        Oxamvl: 200:
 3
                        Pentachlorophenol: 0.3;
               (115)
                        Petroleum aliphatic carbon fraction class (C5 - C8): 400;
 4
               (116)
 5
               (117)
                        Petroleum aliphatic carbon fraction class (C9 - C18): 700;
 6
                        Petroleum aliphatic carbon fraction class (C19 - C36): 10 mg/L;
               (118)
 7
               (119)
                        Petroleum aromatics carbon fraction class (C9 - C22): 200;
 8
               (120)
                        pH: 6.5 - 8.5;
 9
               (121)
                        Phenanthrene: 200:
10
                        Phenol: 30;
               (122)
                        Phorate: 1;
11
               (123)
                        n-Propylbenzene: 70;
12
               (124)
13
               (125)
                        Pyrene: 200;
                        Selenium: 20;
14
               (126)
                        Silver: 20;
15
               (127)
16
               (128)
                        Simazine: 4:
                        Styrene: 70;
17
               (129)
                        Sulfate: 250 mg/L;
18
               (130)
19
               (131)
                        1,1,2,2-Tetrachloroethane: 0.2;
                        Tetrachloroethylene (perchloroethylene; PCE): 0.7;
20
               (132)
21
                        2,3,4,6-Tetrachlorophenol: 200;
               (133)
22
                        Toluene: 600;
               (134)
                        Toxaphene: 0.03;
23
               (135)
24
                        2, 4, 5,-TP (Silvex): 50;
               (136)
                        1,2,4-Trichlorobenzene: 70;
25
               (137)
26
               (138)
                        1,1,1-Trichloroethane: 200;
27
               (139)
                        Trichloroethylene (TCE): 3;
28
                        Trichlorofluoromethane: 2 mg/L;
               (140)
29
               (141)
                        1,2,3-Trichloropropane: 0.005;
30
               (142)
                        1,2,4-Trimethylbenzene: 400;
31
               (143)
                        1,3,5-Trimethylbenzene: 400;
                        1,1,2-Trichloro-1,2,2-trifluoroethane (CFC-113): 200 mg/L;
32
               (144)
33
                        Vinyl chloride: 0.03;
               (145)
34
               (146)
                        Xylenes (o-, m-, and p-): 500; and
35
               (147)
                        Zinc: 1 mg/L.
36
      (h) Class GSA Standards. The standards for this class are the same as those for Class GA except as follows:
37
               (1)
                        chloride: allowable increase not to exceed 100 percent of the natural quality concentration; and
38
               (2)
                        total dissolved solids: 1000 mg/l.
39
      (i) Class GC Waters.
40
                        The concentrations of substances which, at the time of classification, exceed the standards applicable
               (1)
41
                        to Class GA or GSA groundwaters shall not be caused to increase, nor shall the concentrations of other
42
                        substances be caused to exceed the GA or GSA standards as a result of further disposal of
43
                        contaminants to or beneath the surface of the land within the boundary of the area classified GC.
44
                        The concentrations of substances which, at the time of classification, exceed the standards applicable
               (2)
                        to GA or GSA groundwaters shall not be caused to migrate as a result of activities within the boundary
45
                        of the GC classification, so as to violate the groundwater or surface water quality standards in
46
47
                        adjoining waters of a different class.
                        Concentrations of specific substances, which exceed the established standard at the time of
48
               (3)
49
                        classification, are listed in Section .0300 of this Subchapter.
50
                        Authority G.S. 143-214.1; 143B-282(a)(2);
51
      History Note:
52
                        Eff. June 10, 1979;
53
                        Amended Eff. November 1. 1994: October 1. 1993: September 1. 1992: August 1. 1989:
54
                        Temporary Amendment Eff. June 30, 2002:
55
                        Amended Eff. August 1, 2002;
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1	Temporary Amendment Expired February 9, 2003;
2	Amended Eff. <u>July 2012;</u> January 1, 2010; April 1, 2005.
3	

15A NCAC 02L .0202 is proposed for amendment as follows: (Option 2)

1 2 3

15A NCAC 02L .0202 GROUNDWATER QUALITY STANDARDS

(a) The groundwater quality standards for the protection of the groundwaters of the state are those specified in this Rule. They are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of the state, which may be tolerated without creating a threat to human health or which would otherwise render the groundwater unsuitable for its intended best usage.

8 9 (b) The groundwater quality standards for contaminants specified in Paragraphs (g) and (h) of this Rule are as listed, except that:

10 11 12

Where the standard for a substance is less than the practical quantitation limit, the detection of that (1) substance at or above the practical quantitation limit constitutes a violation of the standard.

13 14 15 (2) Where two or more substances exist in combination, the Director shall consider the effects of chemical interactions as determined by the Division of Public Health and may establish maximum concentrations at values less than those established in accordance with Paragraphs (c), (g), or (h) of this Rule. In the absence of information to the contrary, in accordance with Paragraph (d) of this Rule, the carcinogenic risks associated with carcinogens present shall be considered additive and the toxic effects associated with non-carcinogens present shall also be considered additive.

16 17 18

Where naturally occurring substances exceed the established standard, the standard shall be the (3) naturally occurring concentration as determined by the Director.

19 20

21

22

23

24

(c) Except for tracers used in concentrations which have been determined by the Division of Public Health to be protective of human health, and the use of which has been permitted by the Division, substances which are not naturally occurring and for which no standard is specified shall not be permitted in concentrations at or above the practical quantitation limit in Class GA or Class GSA groundwaters. Any person may petition the Director to establish an interim maximum allowable concentration for a substance for which a standard has not been established under this Rule. The petitioner shall submit relevant toxicological and epidemiological data, study results, and calculations necessary to establish a standard in accordance with Paragraph (d) of this Rule. Within three months after the establishment of an interim maximum allowable concentration for a substance by the Director, the Director shall initiate action to consider adoption of a standard for that substance.

(d) Except as provided in Paragraph (f), groundwater Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of:

31 32

30

Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult (1) body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)]/[2 liters/day (avg. water consumption)];

33

Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6; (2)

34 35

(3) Taste threshold limit value;

36 37

(4) Odor threshold limit value; (5) Maximum contaminant level; or

38 39 (6)National secondary drinking water standard.

40 41 (e) The following references, in order of preference, shall be used in establishing concentrations of substances which correspond to levels described in Paragraph (d) of this Rule.

42

(1) Integrated Risk Information System (U.S. EPA).

43 44

Health Advisories (U.S. EPA Office of Drinking Water). (2) Other health risk assessment data published by U.S. EPA. (3)

45

(4) Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.

46 47

(f) The Commission may establish groundwater standards less stringent than existing maximum contaminant levels or national secondary drinking water standards if it finds, after public notice and opportunity for hearing, that

48 49

(1) more recent data published in any of the EPA health references listed in paragraph (e) results in a standard which is protective of public health, taste threshold, or odor threshold,

50 51

(2) such a standard will not endanger the public health and safety, including health and environmental effects from exposure to groundwater contaminants, and

52 53 (3) compliance with a standard based on the maximum contaminant level or national secondary drinking water standard would produce serious hardship without equal or greater public benefit.

54 55

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1
      (fyg) Groundwater quality standards specified in Paragraphs (g)(h) and (h)(i) of this Rule and interim maximum
 2
      allowable concentrations established pursuant to Paragraph (c) of this Rule shall be reviewed on a triennial basis.
 3
      Appropriate modifications to established standards shall be made in accordance with the procedure prescribed in
 4
      Paragraph (d) of this Rule where modifications are considered appropriate based on data published subsequent to the
 5
      previous review.
 6
      (g)(h) Class GA Standards. Where not otherwise indicated, the standard refers to the total concentration in micrograms
 7
      per liter of any constituent in a dissolved, colloidal or particulate form which is mobile in groundwater. This does not
 8
      apply to sediment or other particulate matter which is preserved in a groundwater sample as a result of well construction
      or sampling procedures. The Class GA standards are:
 9
10
                        Acenaphthene: 80;
               (1)
                        Acenaphthylene: 200;
11
               (2)
                        Acetone: 6 mg/L;
12
               (3)
13
               (4)
                        Acrylamide: 0.008;
                        Anthracene: 2 mg/L;
14
               (5)
15
               (6)
                        Arsenic: 10;
16
               (7)
                        Atrazine and chlorotriazine metabolites: 3;
                        Barium: 700;
17
               (8)
18
               (9)
                        Benzene: 1;
19
               (10)
                        Benzo(a)anthracene (benz(a)anthracene): 0.05;
                        Benzo(b)fluoranthene: 0.05;
20
               (11)
21
                        Benzo(k)fluoranthene: 0.5;
               (12)
22
                        Benzoic acid: 30 mg/L;
               (13)
23
               (14)
                        Benzo(g,h,i,)perylene: 200;
24
                        Benzo(a)pyrene: 0.005;
               (15)
                        Bis(chloroethyl)ether: 0.03;
25
               (16)
                        Bis(2-ethylhexyl) phthalate (di(2-ethylhexyl) phthalate): 3;
26
               (17)
27
               (18)
                        Boron: 700;
28
                        Bromodichloromethane: 0.6;
               (19)
29
               (20)
                        Bromoform (tribromomethane): 4;
30
               (21)
                        n-Butylbenzene: 70;
                        sec-Butylbenzene: 70;
31
               (22)
32
                        tert-Butylbenzene: 70;
               (23)
33
                        Butylbenzyl phthalate: 1 mg/L;
               (24)
34
               (25)
                        Cadmium: 2;
35
                        Caprolactam: 4 mg/L;
               (26)
                        Carbofuran: 40;
36
               (27)
37
               (28)
                        Carbon disulfide: 700;
38
               (29)
                        Carbon tetrachloride: 0.3;
39
               (30)
                        Chlordane: 0.1;
40
               (31)
                        Chloride: 250 mg/L;
41
               (32)
                        Chlorobenzene: 50;
42
               (33)
                        Chloroethane: 3,000;
43
               (34)
                        Chloroform (trichloromethane): 70;
                        Chloromethane (methyl chloride): 3;
44
               (35)
45
               (36)
                        2-Chlorophenol: 0.4;
                        2-Chlorotoluene (o-chlorotoluene): 100;
46
               (37)
                        Chromium: 10;
47
               (38)
48
               (39)
                        Chrysene: 5;
                        Coliform organisms (total): 1 per 100 milliliters;
49
               (40)
50
               (41)
                        Color: 15 color units;
51
               (42)
                        Copper: 1 mg/L;
52
               (43)
                        Cyanide (free cyanide): 70;
53
               (44)
                        2, 4-D (2,4-dichlorophenoxy acetic acid): 70;
```

54

55

(45)

(46)

DDD: 0.1; DDT: 0.1;

```
1
               (47)
                        Dibenz(a,h)anthracene: 0.005;
 2
               (48)
                        Dibromochloromethane: 0.4:
 3
               (49)
                        1,2-Dibromo-3-chloropropane: 0.04;
 4
               (50)
                        Dibutyl (or di-n-butyl) phthalate: 700;
 5
               (51)
                        1,2-Dichlorobenzene (orthodichlorobenzene): 20;
 6
                        1,3-Dichlorobenzene (metadichlorobenzene): 200;
               (52)
 7
               (53)
                        1,4-Dichlorobenzene (paradichlorobenzene): 6;
 8
               (54)
                        Dichlorodifluoromethane (Freon-12; Halon): 1 mg/L;
 9
               (55)
                        1.1-Dichloroethane: 6:
10
                        1,2-Dichloroethane (ethylene dichloride): 0.4;
               (56)
                        1,2-Dichloroethene (cis): 70;
11
               (57)
12
               (58)
                        1,2-Dichloroethene (trans): 100;
13
               (59)
                        1,1-Dichloroethylene (vinylidene chloride): 7;
14
               (60)
                        1,2-Dichloropropane: 0.6;
15
               (61)
                        1,3-Dichloropropene (cis and trans isomers): 0.4;
16
               (62)
                        Dieldrin: 0.002;
17
               (63)
                        Diethylphthalate: 6 mg/L;
18
               (64)
                        2,4-Dimethylphenol (m-xylenol): 100;
19
               (65)
                        Di-n-octyl phthalate: 100;
20
               (66)
                        1,4-Dioxane (p-dioxane): 3;
21
                        Dioxin (2,3,7,8-TCDD): 0.0002 ng/L;
               (67)
22
                        1,1- Diphenyl (1,1,-biphenyl): 400;
               (68)
23
               (69)
                        Dissolved solids (total): 500 mg/L;
24
               (70)
                        Disulfoton: 0.3;
25
               (71)
                        Diundecyl phthalate (Santicizer 711): 100;
26
               (72)
                        Endosulfan: 40;
27
               (73)
                        Endrin, total: (includes endrin, endrin aldehyde and endrin ketone): 2;
28
               (74)
                        Epichlorohydrin: 4;
29
               (75)
                        Ethyl acetate: 3 mg/L;
30
               (76)
                        Ethylbenzene: 600;
31
               (77)
                        Ethylene dibromide (1,2-dibromoethane): 0.02;
32
               (78)
                        Ethylene glycol: 10 mg/L;
33
               (79)
                        Fluoranthene: 300;
34
               (80)
                        Fluorene: 300;
35
                        Fluoride: 2 mg/L;
               (81)
36
               (82)
                        Foaming agents: 500;
37
               (83)
                        Formaldehyde: 600;
38
               (84)
                        Gross alpha (adjusted) particle activity (excluding radium-226 and uranium): 15 pCi/L;
39
               (85)
                        Heptachlor: 0.008;
40
               (86)
                        Heptachlor epoxide: 0.004;
41
               (87)
                        Heptane: 400;
                        Hexachlorobenzene (perchlorobenzene): 0.02;
42
               (88)
43
               (89)
                        Hexachlorobutadiene: 0.4;
                        Hexachlorocyclohexane isomers (technical grade): 0.02;
44
               (90)
45
               (91)
                        n-Hexane: 400;
               (92)
                        Indeno(1,2,3\text{-cd})pyrene: 0.05;
46
                        Iron: 300;
47
               (93)
48
               (94)
                        Isophorone: 40;
                        Isopropylbenzene: 70;
49
               (95)
                        Isopropyl ether: 70;
50
               (96)
51
               (97)
                        Lead: 15;
52
               (98)
                        Lindane (gamma hexachlorocyclohexane): 0.03;
53
               (99)
                        Manganese: 50;
54
               (100)
                        Mercury: 1;
                        Methanol: 4 mg/L;
55
               (101)
```

```
1
               (102)
                        Methoxychlor: 40;
 2
               (103)
                        Methylene chloride (dichloromethane): 5;
 3
                        Methyl ethyl ketone (2-butanone): 4 mg/L;
               (104)
 4
               (105)
                        2-Methylnaphthalene: 30;
 5
               (106)
                        3-Methylphenol (m-cresol): 400;
 6
                        4-Methylphenol (p-cresol): 40;
               (107)
 7
               (108)
                        Methyl tert-butyl ether (MTBE): 20;
 8
               (109)
                        Naphthalene: 6;
 9
               (110)
                        Nickel: 100;
10
                        Nitrate: (as N) 10 mg/L;
               (111)
                        Nitrite: (as N) 1 mg/L;
11
               (112)
                        N-nitrosodimethylamine: 0.0007;
12
               (113)
13
               (114)
                        Oxamyl: 200;
14
                        Pentachlorophenol: 0.3;
               (115)
15
               (116)
                        Petroleum aliphatic carbon fraction class (C5 - C8): 400;
16
               (117)
                        Petroleum aliphatic carbon fraction class (C9 - C18): 700;
17
               (118)
                        Petroleum aliphatic carbon fraction class (C19 - C36): 10 mg/L;
18
               (119)
                        Petroleum aromatics carbon fraction class (C9 - C22): 200;
19
               (120)
                        pH: 6.5 - 8.5;
20
               (121)
                        Phenanthrene: 200;
21
                        Phenol: 30;
               (122)
22
                        Phorate: 1;
               (123)
23
               (124)
                        n-Propylbenzene: 70;
24
                        Pyrene: 200;
               (125)
25
                        Selenium: 20;
               (126)
26
               (127)
                        Silver: 20;
27
               (128)
                        Simazine: 4;
28
                        Styrene: 70;
               (129)
29
               (130)
                        Sulfate: 250 mg/L;
30
               (131)
                        1,1,2,2-Tetrachloroethane: 0.2;
31
               (132)
                        Tetrachloroethylene (perchloroethylene; PCE): 0.7;
32
                        2,3,4,6-Tetrachlorophenol: 200;
               (133)
                        Toluene: 600;
33
               (134)
34
               (135)
                        Toxaphene: 0.03;
35
                        2, 4, 5,-TP (Silvex): 50;
               (136)
36
                        1,2,4-Trichlorobenzene: 70;
               (137)
37
               (138)
                        1,1,1-Trichloroethane: 200;
38
               (139)
                        Trichloroethylene (TCE): 3;
39
               (140)
                        Trichlorofluoromethane: 2 mg/L;
40
               (141)
                        1,2,3-Trichloropropane: 0.005;
41
               (142)
                        1,2,4-Trimethylbenzene: 400;
42
               (143)
                        1,3,5-Trimethylbenzene: 400;
43
                        1,1,2-Trichloro-1,2,2-trifluoroethane (CFC-113): 200 mg/L;
               (144)
44
               (145)
                        Vinyl chloride: 0.03;
45
               (146)
                        Xylenes (o-, m-, and p-): 500; and
                        Zinc: 1 mg/L.
46
               (147)
47
      (h)(i) Class GSA Standards. The standards for this class are the same as those for Class GA except as follows:
48
               (1)
                        chloride: allowable increase not to exceed 100 percent of the natural quality concentration; and
49
               (2)
                        total dissolved solids: 1000 mg/l.
50
      (i)(i) Class GC Waters.
51
                        The concentrations of substances which, at the time of classification, exceed the standards applicable
               (1)
52
                        to Class GA or GSA groundwaters shall not be caused to increase, nor shall the concentrations of other
53
                        substances be caused to exceed the GA or GSA standards as a result of further disposal of
54
                        contaminants to or beneath the surface of the land within the boundary of the area classified GC.
```

1	(2)	The concentrations of substances which, at the time of classification, exceed the standards applicable
2		to GA or GSA groundwaters shall not be caused to migrate as a result of activities within the boundary
3		of the GC classification, so as to violate the groundwater or surface water quality standards in
4		adjoining waters of a different class.
5	(3)	Concentrations of specific substances, which exceed the established standard at the time of
6		classification, are listed in Section .0300 of this Subchapter.
7		
8	History Note:	Authority G.S. 143-214.1; 143B-282(a)(2);
9		Eff. June 10, 1979;
10		Amended Eff. November 1, 1994; October 1, 1993; September 1, 1992; August 1, 1989;
11		Temporary Amendment Eff. June 30, 2002;
12		Amended Eff. August 1, 2002;
13		Temporary Amendment Expired February 9, 2003;
14		Amended Eff. November 1, 2012; January 1, 2010; April 1, 2005.
15		

Option 3: Proposed Rule Text: 15A NCAC 02L .0113

1 15A NCAC 02L .0113 is proposed for amendment as follows: (Option 3) 2 3 15A NCAC 02L .0113 VARIANCE 4 (a) The Commission, on its own initiative or pursuant to a request under G.S. 143-215.3(e), may grant variances to the rules 5 of this Subchapter. 6 (b) Requests for variances are filed by letter from the applicant to the Environmental Management Commission. The 7 application shall be mailed to the chairman of the Commission in care of the Director, Division of Environmental Management, Post Office Box 29535, Raleigh, N.C. 27626 0535. Water Quality, 1617 Mail Service Center, Raleigh, N.C. 8 9 27699-1617. 10 (c) For site-specific variances, the The application shall contain the following information: 11 Applications filed by counties or municipalities must include a resolution of the County Board of (1) Commissioners or the governing board of the municipality requesting the variance. 12 13 (2) A description of the past, existing or proposed activities or operations that have or would result in a 14 discharge of contaminants to the groundwaters. Description of the proposed area for which a variance is requested. A detailed location map, showing the 15 (3) orientation of the facility, potential for groundwater contaminant migration, as well as the area covered by 16 the variance request, with reference to at least two geographic references (numbered roads, named 17 streams/rivers, etc.) must be included. 18 19 (4) Supporting information to establish that the variance will not endanger the public health and safety, 20 including health and environmental effects from exposure to groundwater contaminants. (Location of wells 21 and other water supply sources including details of well construction within 1/2 mile of site must be shown 22 on a map). 23 (5) Supporting information to establish that requirements of this Rule cannot be achieved by providing the best 24 available technology economically reasonable. This information must identify specific technology 25 considered, and the costs of implementing the technology and the impact of the costs on the applicant. Supporting information to establish that compliance would produce serious financial hardship on the 26 (6) 27 applicant. 28 (7) Supporting information that compliance would produce serious financial hardship without equal or greater 29 public benefit. 30 (8) A copy of any Special Order that was issued in connection with contaminants in the proposed area and supporting information that applicant has complied with the Special Order. 31 32 A list of the names and addresses of any property owners within the proposed area of the variance as well as any property owners adjacent to the site covered by the variance. 33 34 (d) For state-wide variances to groundwater standards established in Section .0202, the application shall contain the 35 following information: (1) Supporting information to establish that the variance will not endanger the public health and safety, including 36 37 health and environmental effects from exposure to groundwater at the proposed constituent levels. This should 38 include information obtained from the following references. 39 (a) Integrated risk Information System (U.S. EPA). (b) Health Advisories (U.S. EPA Office of Drinking Waters). 40 (c) Other health risk assessment data published by U.S. EPA. 41 (d) Other relevant, published health and ecological risk assessment data, and scientifically valid peer-42 reviewed published toxicological data. 43 (2) A list of all known potentially affected sites, to include permitted sites and incident sites. For each site listed, a 44 map for each site with the location of wells and other water supply sources within $\frac{1}{2}$ mile of the affected site must be 45 46 provided. (3) A list of increased costs for treatment for any of the wells or water supply sources listed in Paragraph (2) above 47 48 due to the proposed variance to Section .0202. 49 (d)(e) Upon receipt of the application, the Director will review it for completeness and request additional information if 50 necessary. When the application is complete, the Director shall give public notice of the application and schedule the matter 51 for a public hearing in accordance with G.S. 143-215.4(b) and the procedures set out in Paragraph $\frac{(e)(f)}{(f)}$ of this Rule. 52 (e)(f) Notice of Public Hearing:

Notice of public hearing on any variance application shall be circulated in the geographical areas of the

proposed variance by the Director at least 30 days prior to the date of the hearing:

53 54 (1)

Option 3: Proposed Rule Text: 15A NCAC 02L .0113

1	(A)	by publishing the notice one time in a newspaper having general circulation in said county;	
2	(B)	by mailing to the North Carolina Department of Environment, Health, and Natural Resources,	
3		Division of Environmental Health and appropriate local health agency;	
4	(C)	by mailing to any other federal, state or local agency upon request;	
5	(D)	by mailing to the local governmental unit or units having jurisdiction over the geographic area	
6		covered by the variance;	
7	(E)	by mailing to any property owner within the proposed area of the variance, as well as any property	
8		owners adjacent to the site covered by the variance; and	
9	(F)	by mailing to any person or group upon request.	
10	(2) The 6	contents of public notice of any hearing shall include at least the following:	
11	(A)	name, address, and phone number of agency holding the public hearing;	
12	(B)	name and address of each applicant whose application will be considered at the meeting;	
13	(C)	brief summary of the variance request;	
14	(D)	geographic description of a proposed area for which a variance is requested;	
15	(E)	brief description of activities or operations which have or will result in the discharge of	
16	` '	contaminants to the groundwaters described in the variance application;	
17	(F)	a brief reference to the public notice issued for each variance application;	
18	(G)	information regarding the time and location for the hearing;	
19	(H)	the purpose of the hearing;	
20	(I)	address and phone number of premises at which interested persons may obtain further	
21	` '	information, request a copy of each application, and inspect and copy forms and related	
22		documents; and	
23	(J)	a brief description of the nature of the hearing including the rules and procedures to be followed.	
24	(* /	The notice shall also state that additional information is on file with the Director and may be	
25		inspected at any time during normal working hours. Copies of the information on file will be made	
26		available upon request and payment of cost or reproduction.	
27	(f)(g) All comments received within 30 days following the date of the public hearing shall be made part of the application file		
28	and shall be considered by the Commission prior to taking final action on the application.		
29	(g)(h) In determining whether to grant a variance, the Commission shall consider whether the applicant has complied with any		
30	Special Order, or Special Order by Consent issued under G.S. 143-215.2.		
31		n's final decision is unacceptable, the applicant may file The applicant may appeal the Commission's	
32		a petition for a contested case in accordance with Chapter 150B of the General Statutes. If the petition	
33		ays, the Commission's decision on the variance shall be final and binding.	
34		ot operate as a defense to an action at law based upon a public or private nuisance theory or any other	
35	cause of action.		
36			
37	History Note: Auth	ority G.S. 143-215.3(a)(1); 143-215.3(a)(3); 143-215.3(a)(4); 143-215.3(e); 143-215.4;	
38	Eff. A	August 1, 1989;	
39		nded Eff. <mark>November <u>1, 2012;</u> October 1, 1993.</mark>	
40			

ATTACHMENT D **Hearing Officer Appointment**

ENVIRONMENTAL MANAGEMENT COMMISSION

A39

Stephen T. Smith Chairman Charles Peterson Vice Chairman

NORTH CAROLINA DEPARTMENT OF ENVIRONMENT AND NATURAL RESOURCES

Beverly Eaves Perdue, Governor Dee Freeman, Secretary

Christopher J. Ayers Donnie Brewer Marvin S. Cavanaugh, Sr. Thomas F. Cecich Marion Deerhake Tom Ellis William L. Hall, Jr. Steve P. Keen Ernest W. Larkin

Kevin Martin Jeffrey V. Morse Darryl D. Moss David B. Peden Dickson Phillips III Amy E. Pickle Clyde E. Smith, Jr. Steve W. Tedder

April 12, 2012

MEMORANDUM

TO:

DICKSON PHILLIPS

ENVIRONMENTAL MANAGEMENT COMMISSION

FROM:

STEPHEN T. SMITH

SUBJECT: HEARING OFFICER APPOINTMENT

I hereby appoint you to serve as the hearing officer for the public hearing to be held for the Proposed Changes to Groundwater Rules. Proposed changes to 15A NCAC 2L .0202 and .0113 regulations will be presented to the interested public by staff of the Planning Section of the Division of Water Quality. The public hearing is scheduled for May 23rd at 6:30 p.m. in the Archdale Building, Ground Floor Hearing Room, 512 N. Salisbury Street, Raleigh. Sandra Moore (919-807-6417) will provide staff support for you. Please present your findings and recommendations to the Environmental Management Commission.

Thank you for your assistance and service.

CC:

Chuck Wakild

Lois Thomas Sandra Moore

Hearing Record File



PROPOSED CHANGES TO GROUNDWATER RULES

The Environmental Management Commission (EMC) will hold a public hearing to collect comments on proposed revisions to rules regulating groundwater quality standards and variance procedures codified in Title 15A, Subchapter 02L, Sections .0202 and .0113 of the North Carolina Administrative Code (http://reports.oah.state.nc.us/ncac.asp) and the fiscal note associated with the revisions. A hearing is scheduled as follows:

May 23, 6:30 PM, in Raleigh – Archdale Building, Ground Floor Hearing Room, 512 N. Salisbury Street, Raleigh. Speaker registration begins at 6:00 PM.

The purpose of revising the rules is to ensure that groundwater standards are established using the most recent U.S. EPA health effects information. This revision was initiated by a rulemaking petition submitted by a representative for Rhodia, Inc., a global specialty chemical manufacturer that formerly operated as Rhone-Poulenc in Gastonia, North Carolina. A change in the 1,1-dichloroethylene standard from 7 ug/L to 350 ug/L is proposed in order to incorporate the most recent U.S EPA health effects information as published in the Integrated Risk Management System at http://www.epa.gov/IRIS/ (Option 1). A change in the criteria used to establish a standard is proposed in order to allow the EMC to establish a standard less stringent than the federal maximum contaminant level (MCL) when the MCL is not established using the most recent U.S. EPA IRIS health effects information (Option 2). A change in the variance procedure is proposed to allow the EMC to consider a request for a statewide variance from the groundwater rules and to make editorial corrections (Option 3).

In addition, the EMC seeks other proposals that allow flexibility in implementation of 15A NCAC 02L .0202(d) while maintaining or achieving appropriate water quality and public health standards, recognizing that any such proposal, if acted upon, might constitute a substantial change from the proposed rule amendments described in detail in this public notice, and might require an additional rule-making procedure.

Further explanations and details on the proposed rules, including the fiscal analysis, may be obtained by visiting the Division of Water Quality/Planning Section Groundwater Standards website at http://portal.ncdenr.org/web/wg/groundwaterrulesrevisions or writing or calling the contact person listed below.

SUMMARY OF PROPOSED RULES

There are three rule change options proposed:

Option 1. A change in 02L .0202 (g)(59) to amend the 1,1-dichloroethylene standard from 7 ug/L to 350 ug/L.

Option 2. A change in 02L .0202 (d) and (f) to allow the EMC to establish a standard less stringent than the maximum contaminant level (MCL) when:

- a. the MCL is not based on the most recent U.S. EPA health effects data as published in U.S. EPA's Integrated Risk Information System (http://www.U.S. EPA.gov/IRIS/);
- b. such a standard would not endanger public health and safety; and
- c. compliance with a standard based on the MCL would produce serious hardship without equal benefit.

Option 3. A change in 02L .0113 to:

- a. update the Division of Water Quality mailing address in .0113 (b),
- b. allow the EMC to issue a state-wide variance to the 02L rules in .0113 (d), and
- c. clarify the existing variance requirements in .0113(i).

HOW TO SUBMIT COMMENTS

The EMC is interested in all comments pertaining to these proposed rule changes and fiscal note. It is very important that all interested and potentially affected persons or parties make their views known to the EMC whether in favor of or opposed to any or all of the proposed amendments.

You may attend the public hearing and make relevant verbal comments and/or submit written comments, data or other relevant information on the proposed amendments, the fiscal note or standard development methodology. The Hearing Officer may limit the length of time that you may speak at the public hearing, if necessary, so that all those who wish to speak will have that opportunity. Written copies are requested for any oral comments presented at the public hearings. Comments may be presented at the public hearings or submitted in writing to the Planning Section of the NCDENR-Division of Water Quality by July 2, 2012. Such correspondence should be brought to the attention of:

Sandra Moore DENR/DWQ Planning Section 1617 Mail Service Center Raleigh, N.C. 27699-1617 Phone: (919) 807-6417 Fax: (919) 807-6497

Sandra.moore@ncdenr.gov

In the case of inclement weather on the day of the scheduled public hearing, please contact the above telephone number for a recorded message regarding any changes to the location, day or time of the hearing.

ATTACHMENT F

Hearing Officers Speech

Proposed Changes to Groundwater Rules 15A NCAC 02L .0113 & .0202

Title 15A of the North Carolina Administrative Code, Subchapter 2L, Sections .0113 & .0202

Good evening. It is now 6:30 p.m. and this public hearing is officially called to order. My name is <u>Dickson Phillips</u> and I am a member of the Environmental Management Commission. I have been designated the hearing officer for the *Proposed Changes to Groundwater Rules* rulemaking effort.

This hearing is being held under the authority of North Carolina General Statutes, Chapter 150B-21.2. In accordance with the General Statutes, a public notice of this hearing was published in the May 1, 2012 edition of the North Carolina Register. Additionally, notices were sent to persons thought to be interested in tonight's hearing and those who have requested to be placed on the water quality rulemaking mailing list and e-mail notification list. Notice to the public was also provided through the Division of Water Quality's website and a press release was issued by the Division of Water Quality on May 15th.

The purpose of tonight's Hearing is to obtain public comment on <u>three proposed options</u> to amend the regulations governing groundwater quality standards and variance procedures. The EMC is also interested in other proposals that allow flexibility in establishing standards while maintaining or achieving appropriate water quality and public health standards.

Copies of the public notice and the proposed rule text are available at the registration table. A presentation will be given shortly to go over the proposed changes. A written record of this hearing will be prepared which will include all the relevant comments, questions and discussions. For this reason, the hearing is being tape-recorded. Written comments received by July 2, 2012 will also be included in the record.

Based on public comments and input by the Division of Water Quality staff, I will make a recommendation to the Environmental Management Commission. In making the final decision, the Environmental Management Commission will consider the written record, the recommendations of Division staff, the recommendations of the hearing officer, and any concerns of other commission members.

The recommendation for the proposed rules may be to adopt them as proposed or to adopt a modified version of the proposals. The EMC may not recommend a rule that differs substantially from the text of the proposed rule unless the EMC publishes the text with modifications and then holds another public hearing.

At this time, I will recognize the government officials that are here tonight:

Let's also recognize members of the staff of the Division of Water Quality present:

Now, Sandra Moore will present a brief overview of the proposed rules, which are the subject of this hearing. After Sandra's presentation, comments from the audience will be allowed.

The Environmental Management Commission wants to hear your comments on the proposed rules and on other proposals that allow flexibility in establishing standards in 15A NCAC 02L .0202 while maintaining or achieving appropriate water quality and public health standards.

Page 1 of 2

All interested and potentially affected persons or parties are encouraged to make their opinion known to the Commission, whether in favor of or opposed to any or all provisions of the proposed rules. Remember, your comments are important and will enable the Commission to act in the best interest of the public.

We will now accept comments on the proposed rules from the audience. If you have written copies of your comments, I would appreciate receiving a copy of them. I may question speakers if the need arises. When your name is called, please come up to the microphone and state your name and any business or group affiliation.

All comments should be limited to matters that are relevant to the proposed rulemaking on the groundwater quality standards. After all registered speakers have had an opportunity to comment, I will allow comments from additional speakers if time permits. DWQ staff will be available after the hearing to answer any questions that you may have.

I will now call on the first speaker. (Call speakers in the order that they registered).

If there is time: Are there any additional comments?

If there are no more comments: I declare the hearing closed.

The hearing record will remain open until close of business on July 2, 2012. That means that anytime between tonight and close of business on July 2, 2012, anyone can submit further comments on the proposed rules in writing. Written comments received by US Mail or by e-mail during this time period will be made a part of the public record.

Written comments should be addressed to Sandra Moore, NCDENR-Division of Water Quality, 1617 Mail Service Center, Raleigh, NC 27699-1617, phone (919) 807-6417, fax (919) 807-6497, email Sandra.Moore@ncdenr.gov

After the comment period has ended, I will present my recommendations to the Environmental Management Commission at one of the regularly scheduled meetings. As I noted earlier, the Commission may not make substantial changes in the final rules without re-notice and rehearing. If the Commission adopts the proposed rules, then the expected effective date for the rules would be November 1, 2012.

It is the desire of the commission to always act in the best possible interest of the public. Public participation is a very important part of the rulemaking process. We would like to thank everyone for being here tonight and offering your comments.

Staff will be around for a few minutes to answer any additional questions you might have.

ATTACHMENT G

DWQ STAFF PUBLIC HEARING PRESENTATION

REQUEST FOR COMMENTS

Proposed Changes to NC Groundwater Rules 15A NCAC 2L .0202 (Groundwater Standards) and .0113 (Variance)

Public Hearing May 23, 2012

Sandra Moore, Standards Co-coordinator Division of Water Quality Planning Section

Division of Water Quality



OVERVIEW OF PRESENTATION

- Groundwater Standards
- Proposed Rule Revisions & Purpose
- · Costs & Benefits
- · Rulemaking Schedule

upmort und changes

Groundwater Standards

Per 15A NCAC 2L .0202 (d) = the lower of:

- 1. Non-cancer threshold
- 2. 1/million cancer risk
- 3. Taste threshold
- 4. Odor threshold
- 5. Maximum Contaminant Level (MCL)
- 6. Secondary Drinking Water Standard

Division of Islater Quality

To protect and onlance
North Curolina's restrem...

Data Sources Per 15A NCAC 2L .0202(e)

- 1. EPA Integrated Risk Information System IRIS.
- 2. EPA Health Advisories.
- 3. Published EPA Health risk assessment data.
- 4. Other published scientifically valid health effects data.

Division of 1thter Quality

To protest and unbones North Curolina's renters.

Basis for 1,1-DCE Groundwater Standard

- 1) 350 ppb = Non-cancer Threshold
- 2) No cancer potency factor
- 3) No taste threshold
- 4) No odor threshold
- 5) 7 ppb = MCL
- 6) No secondary Drinking Water Std

Division of Hister Quality

To protect and enhance
North Corollar's vertices...

PROPOSED REVISIONS & PURPOSE

Option 1: 2L .0202(g)(59)

- 1,1-DCE (7 ppb to 350 ppb)

Option 2: 2L .0202(d) & (f)

 Allows a 2L standard less stringent than the MCL when certain requirements are met

Option 3: 2L .0113

- Statewide variance to GW Standards

Divisian of Hluter Quality

To protect and enhance.

OPTION 1 1,1-DCE (**7 ppb to 350 ppb**)

- Rhodia Rulemaking Petition
- Current 2L = 7 ppb federal MCL
- EPA IRIS data supports 350 ppb



OPTION 2: REVISE 2L .0202

- Allow a standard less stringent than MCL (or a secondary MCL) when:
 - MCL not based on recent information
 - Standard would not endanger public
 - Hardship without benefit



OPTION 3: VARIANCE

- State-wide variance to GW Standard
- · Updated language

Oivision of White Quality
To protest and anhance South Condinate tourism.

EMC SEEKING OTHER PROPOSALS TO GW RULES

- · Allow flexibility
- Maintain water quality
- · Protect public health

Orisina of Water Quality

*To protect and askins:
Ratio Curoline's restorm.

10

FISCAL ANALYSIS BREAKDOWN OF IMPACTS

G.S. 150B 19.1 (e)

- Private Yes (Rhodia)
- State Yes (NCDENR & NCDOT)
- Local No
- Federal No
- Small business No

SUMMARY OF COSTS & BENEFITS OVER 30 YEARS

No costs identified

• Rhodia savings: ≈ \$945,000

• State savings: ≈ \$30,000

 No increased adverse health effects are expected

Division of Huter Quality

To produce and evaluance to the Carolina's custom.

ESTIMATED RULEMAKING SCHEDULE (2012)

- July 2: Public comment period ends
- September 13: Environmental Management Commission
- October 18: Rules Review Commission
- November 1: Effective Date

Direction of White Quality

Thy product unit advances

Aberli Circulater a waleres.

SUBMIT COMMENTS BY JULY 2, 2012

- Sandra.moore@ncdenr.gov
- Sandra Moore
 NCDENR-Division of Water Quality, 1617 Mail Service Center
 Raleigh, NC 27699-1617

Orision of Stater Quality
Thi potent and redunce Rooth Challen's unders.

ATTACHMENT H

Public Hearing Summary

Proposed Changes to Groundwater Rules 15A NCAC 02L .0202 (Groundwater Quality Standards) & .0113 (Variance) May 23, 2012

The hearing was called to order at 6:30 p.m. by Mr. Dickson Phillips, the hearing officer and an Environmental Management Commission member. No oral or written comments were received. The hearing was adjourned at 7:00 p.m. Below is a list of attendees.

Hearing Officer

Mr. Dickson Phillips, EMC member

NCDENR/ Division of Water Quality

Sandra Moore

Betty Wilcox

Steve Kroeger

Keith Larick

Susan Massengale

Debra Watts

Connie Brower

Nikki Schimizzi

Elizabeth Kountis

Gary Kreiser

NCDENR/Division of Waste Management

Linda Culpepper

Environmental Management Commission

Marion Deerhake

Members of the Public

Benne Hutson, McGuireWoods, Charlotte, NC Steve Stadelman, Novozymes, Franklinton, NC Mark Fogel, Attorney, Raleigh, NC C. C. Wheeler, Progress Energy, Raleigh, NC

Groups or Individuals That Submitted Comments on the Proposed 2L Groundwater Rules and Fiscal Note

Name of Group or Individual	Stakeholders Represented
Southern Environmental Law Center and others	Group of organizations that advocate for thousands of
(SELC et al.):	North Carolinians who rely on groundwater as their
Peter Raabe, NC Conservation Director	source of drinking water, who place a high value on the
AMERICAN RIVERS	quality of NC's groundwater, and who will be adversely
Sam Perkins, Director of Technical Programs	affected by the proposed rules.
CATAWBA RIVERKEEPER FOUNDATION	
Katie Hicks, Assistant Director	
CLEAN WATER FOR NORTH CAROLINA	
Heather Jacobs Deck, Pamlico-Tar Riverkeeper	
PAMLICO-TAR RIVER FOUNDATION	
Julia F. Youngman, Senior Attorney	
Kelly F. Moser, Staff Attorney	
SOUTHERN ENVIRONMENTAL LAW CENTER	
David Emmerling, EdD, Executive Director	
Erin Riggs, Associate Executive Director	
WATERKEEPERS CAROLINA	
Julie Mayfield, Executive Director	
Hartwell Carson, French Broad Riverkeeper	
WESTERN NORTH CAROLINA ALLIANCE	
Mark E. Fogel, Attorney at Law	Retained by unnamed client who may be affected by the
5 West Hargett Street	proposed rules.
Suite 510	
Raleigh, NC 27601	
North Carolina Conservation Network (NCCN)	NCCN works with a statewide network of over one
19 East Martin St. Suite 300	hundred environmental, community, and environmental
Raleigh, NC 27601	justice organizations focused on protecting NC's
Grady McCallie, Shannon Arata	environment and public health. Members of these
M. C	organizations may be affected by rule outcome
Manufacturers and Chemical Industry Council of North	MCIC is a non-profit corporation that seeks to preserve,
Carolina (MCIC)	protect and promote the interests of manufacturers in
620 N. West Street, Suite 101	North Carolina. Many MCIC member companies, and
Raleigh, NC 27603	other North Carolina manufacturers, will be directly
A. Preston Howard, Jr., President	affected by the actions that the Environmental
Progress Energy Service Company, LLC (PEC)	Management Commission takes on the subject rules. PEC is a regional energy company serving the Southeast
PO Box 14042	region of the country.
St. Petersburg, FL 33733	region of the country.
Michael Olive	
Director, Environmental Services	
North Carolina Farm Bureau Federation (NCFB)	NCFB represents the interests of farm and rural people
NC Farm Bureau Federation	in NC.
5301 Glenwood Avenue	III 110.
Raleigh, NC 27612	
Anne Coan, Director of Environmental Affairs	
NC League of Municipalities (NCLM)	NCLM is a membership organization of over 550 NC
215 North Dawson Street	municipalities and affiliate organizations, many of
Raleigh, NC 27603	which conduct permitted operations affected by the
PO Box 3069	state's groundwater quality standards in 15A NCAC
Erin Wynia	02L .0202(d).

Comments to the Fiscal Note Received Prior to the Comment Period

North Carolina Department of Transportation (NCDOT)	NCDOT has been active since the early 1990's in the
Roadside Environmental Unit	assessment and remediation of solvent releases related to
1557 Mail Service Center	asphalt testing activities across the State, despite the fact
Raleigh, NC 27699-1567	that NCDOT never owned, operated, or controlled any
Thomas C. Niver, Environmental Operations Engineer	of the asphalt testing lab sites.

Mark E. Fogel
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Suite 510 Raleigh, North Carolina 27601 (919) 833-8852 MAILING ADDRESS: P. O. BOX 1189 RALEIGH, NC 27602

May 29, 2012

Sandra Moore, DENR/DWQ Planning Section 1617 Mail Service Center Raleigh, NC 27699-1617

RE: Proposed Rule Change to 2L.0202 (g)(59)

Dear Ms. Moore:

Enclosed please find my written comments pursuant to the Rule Making held May 23, 2012, with respect to the above referenced Rule.

Please make the enclosed comment part of the Rule Making Record for consideration by the Honorable Dixon Philips.

I enjoyed speaking with you at the Hearing.

Sincerely,

Mark E. Fogel, Attorney

Enclosure

MEF/nmm

NORTH CAROLINA DEPARTMENT OF ENVIRONMENT AND NATURAL RESOURCES

IN THE MATTER OF:) .
)
PROPOSED RULE MAKING) COMMENTS TO RULE MAKING RECORD
CHANGE TO 2L-0202(G)(59))
)

My name is Mark E. Fogel. I am an Environmental Attorney taking the opportunity to submit comments in relation to the Benefits of the proposed change to the above referenced Rule. I wish these comments be made part of the Rule Making Record for consideration by the Honorable Dixon Philips, the Rules Review Commission and the Environmental Management Commission.

I have a Client into the REGISTERED Environmental Consultant Program engaged in the Remedial Investigation Phase. At this point the remaining element is the determination of the horizontal and vertical extent of the DCE plume. As of the latest Quarterly Monitoring Report the DCE exceeded the 7 ppb Ground Water Standard in bedrock wells at al level of 68.7 ppb. With the proposed change to 350 ppb, I estimate a savings of at least \$75,000.00 to avoid an additional triple well nest. In addition, there is no question in my mind that there could be very significant savings to my Client in the carrying out of the Remedial Action under the REC Program. Finally, I foresee no measurable negative impact on the environment if the REC Program is carried out under the proposed 2L change for DCE.

Sincerely Submitted,

Mark E. Fogel, Attorney

5 West Hargett Street, Suite 510

Raleigh, NC 27601

919-833-8852

mefogel@mindspring.com

American Rivers • Catawba Riverkeeper Foundation • Clean Water for North Carolina • Pamlico-Tar River Foundation • Southern Environmental Law Center • Waterkeepers Carolina • Western North Carolina Alliance

July 2, 2012

VIA U.S. MAIL AND E-MAIL

Sandra Moore
DWQ Planning Section
N.C. Department of Environment & Natural Resources
1617 Mail Service Center
Raleigh, NC 27699-1617
sandra.moore@ncdenr.gov

Re: Proposed Changes to Groundwater Rules

Dear Ms. Moore:

We appreciate the opportunity to comment on the proposed changes to North Carolina's groundwater rules, specifically the standard for 1,1-dichloroethylene, the manner in which standards are revised, and the manner in which variances are granted (15A N.C. Admin. Code 02L .0202 and .0113). Collectively, our organizations advocate for thousands of North Carolinians who rely on groundwater as their source of drinking water, who place a high value on the quality of North Carolina's groundwater, and who will be adversely affected by the proposed rules.

We are concerned about the proposed rules and the likely unintended consequences for North Carolina's water quality and public health. Although the stated intent of the proposed rules is to change the groundwater standard for 1,1-dichloroethylene, the proposal would relax the process for setting groundwater standards for other chemicals and could threaten the health of North Carolina's groundwaters. We urge the Environmental Management Commission ("EMC") either to reject the proposed rules altogether or, at most, to adopt Option 3 (state-wide variance) with the modifications described below.

Background

According to the United States Environmental Protection Agency ("EPA"), 1,1-dichloroethylene ("1,1-DCE") is an organic liquid that "is used in making adhesives, synthetic

fibers, refrigerants, food packaging and coating resins such as the saran types." Humans rapidly absorb 1,1-DCE through inhalation and oral exposure. Health effects from prolonged exposure include liver and kidney damage. People exposed to high concentrations of 1,1-DCE can experience the depression of their central nervous systems, inebriation, convulsions, spasms, and unconsciousness. 1,1-DCE also impacts the human respiratory system causing inflammation of the mucous membranes, and is considered to be a possible cancer-causing substance.

Under the Federal Safe Drinking Water Act ("SDWA"), EPA must set maximum contaminant level goals ("MCLGs"); promulgate national primary drinking water regulations; and set maximum contaminant levels ("MCLs") for contaminants (1) that may have an adverse health effect on humans, (2) that are known to occur or are likely to occur in public water systems, and (3) the regulation of which "presents a meaningful opportunity for health risk reduction for persons served by public water systems." 42 U.S.C. § 300g-1(b). The SDWA also requires EPA to review each national primary drinking water regulation at least once every six years and to revise them, if appropriate. The purpose of this "Six-Year Review" is to identify those national primary drinking water regulations "for which current health effects assessments, changes in technology, and/or other factors provide a health or technical basis to support a regulatory revision that will maintain or strengthen public health protection."

EPA set the MCLG for 1,1-DCE at 0.007 mg/L or 7 ppb, and set an enforceable MCL for it at the same level, effective as of 1989. 40 C.F.R. § 141.50 (setting MCLG); 40 C.F.R. § 141.61 (setting MCL). EPA set these levels of protection "based on the best available science

¹ U.S. EPA, *Basic Information About 1,1-dichloroethylene in Drinking Water, available at* http://water.epa.gov/drink/contaminants/basicinformation/1-1-dichloroethylene.cfm. (last visited July 1, 2012).

² Bob Benson, Environmental Protection Agency, *Concise International Chemical Assessment Document* 51, 1,1-DICHLOROETHENE (VINYLIDENE CHLORIDE), World Health Organization et al. (2003), available at www.who.int/entity/ipcs/publications/cicad/en/cicad51.pdf (last visited July 2, 2012).

³ Id.

⁴ U.S. EPA, *Vinylidene Chloride (1,1-Dichloroethylene)* (hereinafter "Vinylidene Chloride"), *available at* http://www.epa.gov/ttn/atw/hlthef/di-ethyl.html (last visited July 2, 2012).

⁶ 1,1-Dichloroethylene, § II.A.1., available at http://www.epa.gov/IRIS/subst/0039.htm (last visited July 2, 2012).

⁷ U.S. EPA, Six-Year Review of Drinking Water Standards, available at http://water.epa.gov/lawsregs/rulesregs/regulatingcontaminants/sixyearreview/index.cfm (last visited July 1, 2012).

to prevent potential health problems." As of its last six-year review, EPA determined that the 0.007 mg/L standard was still appropriate and protective of human health.⁹

States such as North Carolina that have "primary enforcement responsibility for public water systems" must adopt "drinking water regulations that are no less stringent than the national primary drinking water regulations." 42 U.S.C. § 300g-2(a); 40 C.F.R. § 142.10(a). North Carolina has complied, setting its drinking water standard for 1,1-DCE at 7 μ g/L (which is the equivalent of 0.007 mg/L). 15A N.C. Admin. Code 18C .1518.

North Carolina also decided by regulation that its groundwater standards must be established as the lowest concentration (that is, the most stringent) of six criteria, one of which is the federal MCL. 15A N.C. Admin Code 2L .0202(d)(1)-(6). North Carolina's groundwater standards are intended to "maintain and preserve the quality of groundwaters, prevent and abate pollution and contamination of waters of the state, protect public health, and permit management of the groundwaters for their best usage by the citizens of North Carolina." 15A N.C. Admin. Code 02L .0103(a). The EMC sets North Carolina's groundwater standards pursuant to its authority under chapter 143, article 21, of the North Carolina General Statutes, which provides, among other things, that:

- (a) It is hereby declared to be the public policy of this State to provide for the conservation of its water and air resources [and] . . . to achieve and to maintain for the citizens of the State a total environment of superior quality. Recognizing that the water and air resources of the State belong to the people, the General Assembly affirms the State's ultimate responsibility for the preservation and development of these resources in the best interest of all its citizens
- (b) It is the public policy of the State to maintain, protect, and enhance water quality within North Carolina
- (c) . . . Standards of water and air purity shall be designed to protect human health, to prevent injury to plant and animal life, to prevent damage to public and private property, . . . and to secure for the people of North Carolina, now and in the future, the beneficial uses of these great natural resources.

N.C. Gen. Stat. § 143-211. The rules governing groundwater should be amended only in ways that further these objectives and ensure the highest water quality for all North Carolinians.

⁸ U.S. EPA, *Basic Information About 1,1-dichloroethylene in Drinking Water, available at* http://water.epa.gov/drink/contaminants/basicinformation/1-1-dichloroethylene.cfm (last visited July 1, 2012); *see also* 42 U.S.C. § 300g-1(b)(ii)(II) (requiring the use of the "best available public health information").

⁹ See National Primary Drinking Water Regulations; Announcement of the Results of EPA's Review of Existing Drinking Water Standards and Request for Public Comment and/or Information on Related Issues, 75 Fed. Reg. 15,500, 15,535-36 (Mar. 29, 2010) (hereinafter, "Results of Six-Year Review"), available at http://edocket.access.gpo.gov/2010/pdf/2010-6624.pdf (last visited July 1, 2012).

The EMC has determined that the best usage of North Carolina's groundwaters "is as a source of drinking water," and that North Carolina's groundwater standards are in place to ensure the state's groundwaters remain suitable as future drinking water sources. 15A N.C. Admin. Code 02L .0103(a). More than 50 percent of North Carolinians rely on groundwater as a source of drinking water.¹⁰

The proposed rule revisions that are the subject of these comments contain three "options." Under Option 1, the EMC would drastically weaken the groundwater standard for 1, 1-DCE 50-fold, from 7 μ g/L to 350 μ g/L. Option 2 would also permit the EMC to revise an MCL-based groundwater standard downward despite the fact that the MCL is set by EPA by regulation and has not been changed by EPA. Finally, Option 3 of the proposed rules would provide a mechanism by which the EMC could issue a state-wide variance to a groundwater standard. The proposed rules jeopardize the health of North Carolina's residents and drinking water supply. The proposal would unfairly shift costs from the polluter to innocent end users and make uncertain whether a variance would shield polluters from legal action by injured end users. The rule revisions are also unnecessary because any issue with the 1,1-DCE groundwater standard could be resolved through the current variance process. Accordingly, the EMC should reject the requested rule revisions in whole; in the alternative, the EMC should adopt Option 3 with modifications, as set forth below.

Comments

I. The Proposed Rule Revisions Unfairly Shift the Cost of Cleanup and Compliance with the National Drinking Water Standard for 1,1-DCE to the Ultimate User of the Water.

One risk of the proposed rule revision is that permitted discharges of 1,1-DCE above the MCL will impair a source (or future source) of drinking water, unfairly shifting costs from the polluter to innocent public and private water systems and users. Private water users would be saddled with the costs of treating their water supplies to remove the health risks created by the polluter. And, if discharges of 1,1-DCE permitted under the proposed rules result in detections in public water systems of 1,1-DCE above the MCL required under the SDWA (0.007 mg/L), the public water system would be responsible for taking actions necessary to meet the MCL. See 15A N.C. Admin. Code 18C .1518. According to the North Carolina Department of Environment and Natural Resources Public Water Supply Section, there are over 2,000 public water systems that could be unfairly burdened by the rule revisions. 12

The proposed rule revisions open the door for the EMC to revise other groundwater standards in addition to the 1,1-DCE standard. Each potential future standard change would

¹⁰ N.C. Dept. of Water Quality ("DWQ"), *Fiscal Note for Proposed Rules 15A NCAC 02L-Groundwater* (hereinafter, "Fiscal Note"), at 5.

¹¹ *Id.* at 20-21.

¹² Fiscal Note at 21.

likewise burden private and public water suppliers and users with the cost of treating their water to meet the federal MCLs under the SDWA and to make it safe for human consumption. Even assuming Rhodia's 1,1-DCE discharges do not reach groundwater used by any public water system, the same would not necessarily be true of other groundwater standards changed in the future under the proposed revised rules. The proposed rules do not contain any terms that would prevent a change in a groundwater standard if the new standard otherwise met the new proposed criteria, but would affect human water supplies and would force public and private entities to bear the cost of remediation.

We therefore urge the EMC to reject the rules outright on principles of equity.

II. The Proposed Rule Revisions Are Unjustified; Any Issue With the Current 1,1-DCE Groundwater Standard Can Be Dealt with by Using 2L .0113's Current Variance Process.

The EMC has considered requested changes to the 1,1-DCE groundwater standard before. The EMC Groundwater Committee concluded in 2010 that the groundwater rules do not need revision because any "1,1-DCE issue could be dealt with using the variance process." We agree, and urge the EMC to reject again the proposal to change the standard for 1,1-DCE. There is no reason for the EMC to depart from its prior decision, and it would be arbitrary for it to do so.

The only company known to be affected by the 1,1-DCE standard is Rhodia. ¹⁵ Unlike Rhodia, which according to DWQ has direct releases of 1,1-DCE, most companies that discharge 1,1-DCE release it together with other more toxic chlorinated solvents above their respective groundwater standards, and in much higher concentrations than 1,1-DCE. The presence of these other regulated chemicals would likely trigger more complex and costly environmental cleanup efforts, thereby eliminating most, if not all, of the benefits of amending the 1,1-DCE groundwater standard for most companies because it is not the pollutant driving their cleanups. ¹⁶

Rhodia itself admits that it is not requesting a change in the 1,1-DCE standard or any variance "on the grounds that compliance with the 1,1-DCE standard [is] technically infeasible or would cause undue economic hardship"; rather, the company is requesting the rule revisions to reduce its compliance costs, including remediation costs and groundwater investigation and monitoring costs.¹⁷ But reducing compliance costs does not fall within the stated goals of North

¹⁶ Fiscal Note at 18-19; *see also* Results of Six –Year Review, 75 Fed. Reg. 15,500, 15,536 ("compliance technologies for [1,1-DCE reduce concentrations of] other co-occurring contaminants"), *available at* http://edocket.access.gpo.gov/2010/pdf/2010-6624.pdf (last visited July 1, 2012).

¹³ *Id.*, Appx. A, at 92.

¹⁴ *Id.*, Appx. A, at 92.

¹⁵ *Id.*, at 12-13.

¹⁷ Fiscal Note, Appx. A, at 6-12.

Carolina's groundwater policy (15A N.C. Admin. Code 02L. 0103(a), (b)) and there are no health-based benefits of the proposed rule changes. ¹⁸ The rule revisions are therefore unjustified.

As the EMC has already determined, Rhodia appears to be the only company affected by the 1,1-DCE standard, and a site-specific variance under the current rules "could be approved that would allow less restrictive Groundwater Quality Standards [for Rhodia] while providing the site-specific requirements necessary to protect Public Water Supplies." In other words, we understand that the agency has determined that a variance could be given to Rhodia to allow it to exceed groundwater standards while still ensuring that public water supplies are protected elsewhere.

For the reasons stated above, the current variance process is sufficient to resolve any issues with the 1,1-DCE standard, and the proposed rule revisions should be rejected in full. If, however, the EMC does not reject the proposed rules altogether, it should adopt Option 3, with the modifications discussed below. Selecting Option 3 would not revise North Carolina's protective groundwater standards for all permitted parties; rather the onus would rightly be on the party requesting the variance to present the necessary data to justify the variance.

III. The Proposed Change in the Groundwater Standard for 1,1-DCE Under Option 1 Is Unauthorized.

Even if the EMC finds that a change in the 1,1-DCE standard may be appropriate, it cannot summarily adopt the 350 μ g/L 1,1-DCE standard as proposed. Doing so would be an unauthorized departure from the current process adopted by the EMC for setting groundwater standards, as outlined in 2L .0202. This process requires that the groundwater standard be set at the lowest of six criteria, in this case the MCL; thus, for a change below the federal MCL to be legal, the EMC would need to establish, and follow, a reasoned and protective regulatory process by which to deviate from 2L .0202(d). Although proposed Option 2 sets forth a potential process by which to deviate from 2L .0202(d), a rule change of this nature is, as explained more fully above, unnecessary, unjustified, and unfair to end users of the water. Moreover, while the EMC may be comfortable with the result in the present situation involving Rhodia's discharges of 1,1-DCE, there is no way of knowing what the results would be with other groundwater standard changes that might be made pursuant to Option 2 in the future, nor what the cost might be for water suppliers and users to comply with the SDWA. It, therefore, should be rejected.

In its petition, Rhodia argues that a change in the 1,1-DCE standard is legally required under the current rules because the toxicity data in EPA's Integrated Risk Information System ("IRIS") database, which is identified by the EMC as the preferred reference for toxicity values in establishing groundwater standards in 2L .0202(e), no longer "corresponds" to the MCL because EPA increased the reference dose (RfD) for 1,1-DCE in the IRIS database without a

¹⁹ *Id.*, Appx. A, at 96.

¹⁸ Fiscal Note at 16.

²⁰ See id., Appx. A, at 3.

corresponding increase to the MCL. Rhodia is wrong, and the EMC has already rejected this argument. Section 2L .0202(d) is clear: groundwater standards should be established as the least of the six identified criteria, including the MCL set under the SWDA. The least of the identified criteria for 1,1-DCE is the MCL of 7 μ g/L; therefore, 7 μ g/L is the value that should remain the groundwater standard for 1,1-DCE.

As the EMC has already concluded, the references set forth in 2L .0202(e), including IRIS, are not to be used to override criteria such as the MCL or the national secondary drinking water standard, which are developed by EPA according to regulation. Instead, the references set out in 2L .0202(e) are for use in calculating the criteria dependent on data, such as the systemic threshold concentration or the incremental lifetime cancer risk. Even so, the Commission retains authority to set more stringent standards where necessary to protect North Carolina groundwaters, and is not constrained by the criteria set forth in 2L .0202(d). See 15A N.C. Admin. Code 2L .0103(b) (expressing the EMC's intent "to protect all [North Carolina] groundwaters "to a level of quality at least as high as that required under the [groundwater] standards established in Rule .0202.") (emphasis added). Accordingly, there is no legal basis for the EMC to adopt the less stringent groundwater standard as proposed.

Finally, EPA reviewed but did not revise the standards related to 1,1-DCE in its most recent Six-Year Review. EPA stated that updated health effects data *may* support increasing the 1,1-DCE MCL to 0.350 mg/L (the equivalent of 350 μ g/L); however, it decided not to update the MCL because the revision "would not likely result in cost-savings or health risk reduction to public water systems and their customers." We understand that the proposed – but rejected – change to 350 μ g/L was based on two pieces of information. First, in 2002, EPA updated the reference dose for 1,1-DCE from 0.010 mg/kg/day to 0.050 mg/kg/day. This change potentially could have justified a five-fold increase in the MCL to 35 μ g/L, had EPA elected to do so, but it did not. Second, EPA made an indefinite and unverifiable statement that a "risk management factor of ten *may no longer be needed.*" The elimination of that risk factor could have increased the MCL from 35 μ g/L to 350 μ g/L, but EPA chose not to do so, and, in any event, the elimination was not justified by such an indefinite statement. Accordingly, if EMC considers any change at this time (which we strongly oppose), it should only be to 35 μ g/L

²¹ *Id.*, Appx. A, at 3-4.

²² *Id.*, Appx. A, at 92-93.

²³ *Id.*, Appx. A, at 92-93.

²⁴ Id.

²⁵ See Results of Six-Year Review, 75 Fed. Reg. 15500, 15535-36 (Mar. 29, 2010), available at http://edocket.access.gpo.gov/2010/pdf/2010-6624.pdf (last visited July 1, 2012).

²⁶ *Id.* at 15,535; Fiscal Note at 26-27; *see also* "National Primary Drinking Water Regulations; Announcement of Completion of EPA's Review of Existing Drinking Water Standards," 68 Fed. Reg. 42.908, 42.918-19 (July 18, 2003).

²⁷ See Results of Six-Year Review, 75 Fed. Reg. 15,500, 15535 (emphasis added), available at http://edocket.access.gpo.gov/2010/pdf/2010-6624.pdf (last visited July 1, 2012).

(based on the updated reference dose) and should not include the unjustified elimination of the risk factor.

In sum, EPA elected not to make either of these changes. Thus, unless and until EPA revises the MCL for 1,1-DCE, the EMC should incorporate the actual federal regulatory MCL value to ensure North Carolina's groundwater standards are sufficiently protective of human health and the environment and will allow all water supplies to meet SDWA standards. An increase to $350~\mu g/L$ would be absolutely unjustified at this time.

IV. If Option 3 (or Option 2) Is Selected, It Should Be Revised so that It Is as Protective of Human Health as Possible.

As stated above, we urge the EMC to reject the proposed revisions in full. If, however, the EMC determines a rule revision is appropriate, we urge it to select Option 3 with the following modifications to limit its scope and impact to the greatest degree possible. For the most part, in the event the EMC decides, against our recommendation, to select Option 2, these modifications could be made to that option as well.

First and foremost, because of the unique situation regarding the 1,1-DCE groundwater standard, any rule change should be expressly limited to 1,1-DCE. This will avoid unintended consequences by preventing any revised rule from allowing changes to groundwater standards for other regulated chemicals that do not share 1,1-DCE's circumstances.

Next, as the rule revisions are currently drafted, neither Option 2 nor Option 3 imposes the same preference for the EPA's IRIS database as does the standard for adopting groundwater standards and setting concentrations in the current 2L .0202(e). EPA's IRIS database provides high-quality, science-based human health assessments and, at least where more stringent sources do not exist, should remain the preferred source for decision-making in the groundwater context. EPA's process for developing IRIS assessments includes a call for scientific information from the public through a Federal Register announcement and IRIS internet site, a review of the current scientific literature, internal agency review, consultation with other federal agencies and White House offices, external expert peer review by entities such as the National Academy of Sciences, and public comment. 28 As Rhodia acknowledges in its petition, "IRIS is the most preferred reference source for the establishment of groundwater quality standards."²⁹ If adopted, the proposed rule should be revised to clarify that (1) the hierarchy, or order of preference, of EPA health references listed in section 2L .0202(e) applies as well in the proposed new section 2L .0113(d) (or the new 2L .0202(f) if the EMC selects Option 2); and (2) the value from the EPA health reference that is most protective of human health is the preferred source of information.

²⁹ Fiscal Note, Appx. A, at 4.

²⁸ U.S. EPA. Integrated Risk Information System, *available at* http://www.epa.gov/iris/help_ques.htm#whatiris (last visited July 2, 2012).

When considering whether a variance would endanger public health and safety, we also urge the EMC to require the consideration of North Carolina's most vulnerable populations and the impacts that a variance (or loosened groundwater standard) might have on these individuals. This approach is consistent with the approach taken under the SDWA, which requires the MCLG to be set at "the level of a contaminant in drinking water below which there is no known or expected health risk," with a margin of safety. The MCLGs must "take into account the risks of exposure for certain sensitive populations, such as infants, the elderly, and persons with compromised immune systems." Consistent with the policy behind North Carolina's groundwater standards to protect drinking water for all of the state's residents, this precautionary approach should be incorporated in any rule change.

If revised, the rule should also take into account the concept of "relative source contribution," that is, the idea that people can be exposed to additional amounts of a contaminant from sources other than their drinking water, such as food or air. ³² Just as EPA accounts for these other contributions when calculating MCLGs, ³³ the EMC should revise the proposed rule so that relative source contribution is considered when determining whether a variance is appropriate.

Finally, if Option 3 is selected, a state-wide variance should apply only to the particular applicant and only to those sites identified by the applicant in a variance application. Site-specific information would be central to EMC's evaluation of whether a state-wide variance would endanger public health and safety. Therefore, it is not appropriate to apply any state-wide variance to companies or sites not included in the application and its analysis.

V. If Option 3 is Selected, No Variance Should Be Permitted to Operate as a Defense to an Action at Law.

Section 2L .0113(i), as currently codified, provides that "a variance shall not operate as a defense to an action at law based upon a public or private nuisance theory or any other cause of action." The proposed rule changes would remove this provision and create uncertainty for affected parties regarding available relief under the law for injuries sustained from discharges made under a variance. We do not support the removal of this language. As explained previously, a risk of the proposed rule revisions is that permitted discharges of 1,1-DCE above the MCL will impair a source of drinking water, unfairly shifting the treatment and compliance

³⁰ U.S. EPA, *Drinking Water Standards and Health Effects*, *available at* http://water.epa.gov/lawsregs/rulesregs/sdwa/index.cfm (last visited July 2, 2012).

³¹ *Id.*

³² U.S. EPA, *What is the relative source contribution (RSC) with regard to development of drinking water standards?*, http://safewater.supportportal.com/link/portal/23002/23015/Article/20103/What-is-the-relative-source-contribution-RSC-with-regard-to-development-of-drinking-water-standards (last visited July 2, 2012).

 $^{^{33}}$ /d.

³⁴ See Proposed Changes to Groundwater Rules, Option 3, 2L .0113(d).

costs from the variance holder to public and private water systems and users. These innocent parties cannot be left without recourse if they are burdened with taking corrective action due to contaminants discharged under a variance. Therefore, we urge the EMC to reject this gratuitous change.

Conclusion

We appreciate DWQ's and the EMC's ongoing efforts to protect our state's valuable groundwater resources. Thank you for the opportunity to comment on the proposed revisions to North Carolina's groundwater rules and for your consideration of our comments.

Sincerely,

Peter Raabe, NC Conservation Director AMERICAN RIVERS

Sam Perkins, Director of Technical Programs CATAWBA RIVERKEEPER FOUNDATION

Katie Hicks, Assistant Director CLEAN WATER FOR NORTH CAROLINA

Heather Jacobs Deck, Pamlico-Tar Riverkeeper PAMLICO-TAR RIVER FOUNDATION

Julia F. Youngman, Senior Attorney Kelly F. Moser, Staff Attorney SOUTHERN ENVIRONMENTAL LAW CENTER

David Emmerling, EdD, Executive Director Erin Riggs, Associate Executive Director WATERKEEPERS CAROLINA

Julie Mayfield, Executive Director Hartwell Carson, French Broad Riverkeeper WESTERN NORTH CAROLINA ALLIANCE





STATE OF NORTH CAROLINA DEPARTMENT OF TRANSPORTATION

BEVERLY EAVES PERDUE
GOVERNOR

EUGENE A. CONTI, JR. SECRETARY

April 24, 2012

VIA US MAIL & EMAIL

Ms. Sandra Moore
Planning Section, Classifications & Standards Unit
NCDENR Division of Water Quality
1617 Mail Service Center
Raleigh, NC 27604
sandra.moore@ncdenr.gov

Subject:

Environmental Management Commission – March 8, 2012

March 2012 Agenda Action Item II, 12-07Request for Approval of the Fiscal Analysis of Proposed Changes to Groundwater Rules 15A NCAC 2L.0202 (1,1-DCE groundwater standard) and .0113 (variance procedures)

Dear Ms. Moore,

As a follow-up to requested information the North Carolina Department of Transportation ("NCDOT") provided to your office on September 12, 2011, regarding the above-noted proposed rule change, I have reviewed the information that was presented to the Environmental Management Commission on March 8, 2012. (For your convenience, a copy of the EMC's March 2012 Agenda is enclosed.) The purpose of this letter is to bring to your attention the fact that some of the information presented to the EMC is not accurate or consistent with the NCDOT Asphalt Testing Lab Program ("ATL Program"). As you are aware, the ATL Program encompasses a very complex set of issues. The ATL program consists of conducting comprehensive environmental site assessments along with the development and implementation of corrective action plans at formal asphalt testing labs. As such, it is important that correct and accurate information is reflected in the public record.

It is important to clarify the information – specifically on page A8 of "Attachment A: Fiscal Impacts of Proposed 15A NCAC 02L Groundwater Rules, as it relates to the ATL Program. (For your convenience, a copy of page A8 is enclosed.) In 1989, the North Carolina General Assembly recognized potential environmental and human health hazards at scores of

Re: Environmental Management Commission - March 8, 2012

privately-owned ATL sites throughout North Carolina. Asphalt paving companies, private engineering/testing firms, and federal, state, and local government agencies (including NCDOT) used the ATL's for years for quality control testing of asphalt products. It is believed that the release of chlorinated aliphatic compounds used in tests by one or more of ATL users may have resulted in soil and groundwater contamination at some of the ATL sites. The General Assembly directed NCDOT to participate in the investigation and cleanup of the ATL's by furnishing personnel, equipment, or other materials and resources as necessary. NCDOT's participation in the ATL site investigations and clean-up was not an indication or admission of liability by NCDOT. The intent of the Legislature, as outlined in the 1995 legislative minutes, was for NCDOT and NCDENR to work shoulder-to-shoulder in the assessment and development of cleanup plans at ATL sites.

Finally, it must be noted that NCDOT use of chlorinated solvents for asphalt quality control testing was terminated in North Carolina in 1996. Ignition ovens are now used to test asphalt samples. NCDOT is currently very active, and has been since the early 1990's, in the assessment and remediation of solvent releases related to asphalt testing activities across the State – despite the fact that NCDOT never owned, operated, or controlled any of the ATLs. The parties potentially responsible for soil and groundwater contamination at some former ATLs include the laboratory personnel, asphalt paving companies, private engineering/testing firms, and federal, state, and local government agencies. NCDOT is certainly not the only party potentially responsible for the contamination.

If you have any questions or need additional information, I may be reached at (919) 835-8481.

Sincerely,

Thomas C. Niver, P.G., CHMM Environmental Operations Engineer

Momas C. This

Enclosures

cc: North Carolina Environmental Management Commission Terry Gibson, PE, State Highway Administrator Ricky Greene, PE, Director of Asset Management Don Lee, CPESC, State Roadside Environmental Engineer Ken Pace, P.E., State Environmental Operations Engineer



2 July 2012

Sandra Moore Water Quality Planning Section Division of Water Quality 1617 Mail Service Center Raleigh, NC 27699-1617

Subject: Comments on Proposed Groundwater Rules 15A NCAC 02L .0202 and .0113

Dear Ms. Moore:

I am writing today to submit comments on behalf of the Manufacturers and Chemical Industry Council of North Carolina (MCIC). MCIC is a non-profit corporation that seeks to preserve, protect and promote the interests of manufacturers in North Carolina. Many MCIC member companies, and other North Carolina manufacturers, will be directly affected by the actions that the Environmental Management Commission takes on the subject rules.

Without regard to any other actions that the EMC may take in response to this rule-making action, the EMC should amend the groundwater standard for 1,1-dichloroethylene (DCE) from 7 ug/l to 350 ug/l, as requested by Rhodia, Inc. In its petition for rule-making, Rhodia has clearly established that the existing standard for DCE is outdated.

As to the other actions that the EMC should take, I offer the following comments. In the past several years, the EMC has found itself frustrated by its own groundwater rules. First, the EMC was frustrated in establishing the correct, scientifically defensible standard for fluoride. That frustration led to the formation of a stakeholders work group to look into the manner in which the EMC establishes its groundwater standards. I had the pleasure of serving on this stakeholders work group. Unfortunately, the deliberations of the work group were dominated by DWQ staff who insisted upon maintaining the same old approach to setting groundwater standards; an approach that dictates that the most stringent of all possible options be selected, even when current scientific information suggests that such actions are inappropriate and unsupportable.

In September 2005, as the stakeholders work group was completing its work, I submitted a recommendation (again, on behalf of MCIC) on how to amend the rule to avoid similar rule-making frustrations in the future. Unfortunately, the EMC chose not to amend its rules, and not surprisingly, found itself similarly frustrated over how to handle Rhodia's petition. Had the EMC amended its rules as recommended by me and several other members of the stakeholders work group back in 2005, it would have been far simpler to manage the Rhodia petition (and all future petitions, as there surely will be others).

Therefore, I submit the following recommended revisions to the EMC groundwater rule:

- (d) <u>In establishing groundwater</u> Groundwater quality standards for substances in Class GA and Class GSA groundwaters <u>the Commission shall consider each of the following values:</u> are established as the least of:
 - 1. Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)];
 - 2. Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6;
 - 3. Taste threshold limit value;
 - 4. Odor threshold limit value:
 - 5. Maximum contaminant level; or
 - 6. National secondary drinking water standard.
- (e) The following references, in order of preference, shall be considered used in establishing concentrations of substances which correspond to levels described in Paragraph (d) of this Rule:
 - 1. Integrated Risk Information System (U.S. EPA).
 - 2. Health Advisories (U.S. EPA Office of Drinking Water).
 - 3. Other health risk assessment data published by U.S. EPA.
 - 4. Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.

This approach will provide the EMC with sufficient regulatory flexibility to manage the types of issues that arose with Flouride and DCE, and will allow the EMC to make the correct, most scientifically supportable decision, every time.

Thank you for the opportunity to submit these comments. If you have any questions or need any additional information, please contact me (phone 919-740-8834 / email preston.howard@mcicnc.org).

Sincerely,

Original Signed by A. Preston Howard, Jr.

A. Preston Howard, Jr., P.E. President

July 2, 2012

Sandra Moore Water Quality Planning Section 1617 Mail Service Center Raleigh, NC 27699-1617

Delivered via e-mail: Sandra.Moore@ncdenr.gov

Dear Ms. Moore:

The North Carolina Farm Bureau Federation (NCFB) is this state's largest general farm organization representing the interests of farm and rural people in North Carolina. This letter is to comment on the proposed changes to the North Carolina Groundwater Standards as published in the NC Register on May 1, 2012, pages 1768 – 1776.

The proposal offers three options for comment but also solicits other proposals that allow flexibility in implementation of 15A NCAC 2L .0202(d) while maintaining or achieving appropriate water quality and public health standards.

In these comments, we will not address Option 1 due to our not being familiar with the particulars of this substance. Of the three options, we favor Option 3 if it is amended as we propose. Because we do not know which option might be selected, we are also providing the following comments about Option 2.

If Option 2 is adopted we recommend Option 2 be changed. We disagree with restricting the data sources to only those sources that are from EPA, as is stated in (f)(3). This precludes using the references described in (e)(4) which could be more up-to-date and relevant than the listed EPA sources. Also, we disagree with (f)(3). It requires that "compliance with a standard based on the maximum contaminant level or national secondary drinking water standard would produce serious hardship without equal or greater public benefit." If the new standard is based on the most up-to-date science and is protective of public health, safety and the environment, there should not have to be a "hardship" test. The hardship test should be maintained as an individual variance requirement and not be made a requirement for a statewide standard change based on more recent data and studies. We recommend that in Option 2, if adopted, (f)(1) be amended to reference all of the sources of data listed in (e), and we recommend that (f)(3) be deleted.

We favor Option 3 with some changes. Option 3 in (d) requires an "application." That word appears to preclude the Commission from granting a variance "on its own initiative" [2L .0113(a)]. Option 3 should state that, in addition to an application from another party, the Commission may initiate a statewide variance without an "application" pursuant to 2L .0113(a). Also, (d)(2) and (d)(3) may be impossible for an applicant or set of applicants (or even the Division of Water Quality) to produce because it may be impossible to produce a list of all known "potentially" affected sites for common substances.

Sandra Moore Division of Water Quality Page 2

This then leads in Option 3 to (d)(3) being as equally difficult to produce in an application. For (d)(3), if it is necessary to retain this, an estimated range of costs of treatment for different water sources might be more appropriate than trying to develop cost estimates for "all known potentially affected sites."

Regarding Option 3, if up-to-date science and data evaluated by the Commission shows no endangerment to public health and safety and to the environment, that should be sufficient for a state-wide variance. The Commission should make (d)(2) and/or (d)(3) optional at the discretion of the Commission. The Commission should do a preliminary evaluation of the proposed statewide variance and determine if the Commission wishes the applicant to produce the information asked for in (d)(2) and (d)(3). Also the rule should state that the Commission has the discretion to request such information and to determine how much information it wants the applicant to produce for each. The Commission should be able to exercise discretion regarding what amount (if any) of the information in (d)(2) and (d)(3) the Commission feels it needs to make the determinations necessary for a statewide variance.

Finally, we assume that DWQ would want to update its mailing address under all three options [2L .0113(a)], not just under Option 3.

Thank you for the opportunity to submit these comments. If you have questions, please do not hesitate to contact me. (phone 919-788-1005)

Sincerely,

Anne Coan Director of Environmental Affairs NC Farm Bureau Federation 5301 Glenwood Avenue Raleigh, NC 27612



T: 919.857.4699 F: 919.833.8819

19 East Martin St. Suite 300 Raleigh, NC 27601

www.ncconservationnetwork.org

July 2, 2012

Sandra Moore
Planning Section, Classifications & Standards Unit
DENR Division of Water Quality
1617 Mail Service Center
Raleigh, NC 27699-1617
Sandra.moore@ncdenr.gov

RE: Groundwater Rules Proposed Revisions (15A NCAC .02L .0202 and .0113)

Dear Ms. Moore:

The North Carolina Conservation Network appreciates the opportunity to comment on proposed changes to North Carolina's groundwater rules, codified in Title 15A, Subchapter 02L, Sections .0202 and .0113 of the North Carolina Administrative Code (NCAC).

Our organization works with a statewide network of over one hundred environmental, community, and environmental justice organizations focused on protecting North Carolina's environment and public health. To the extent that the proposed rulemaking reaches beyond the narrow issue that prompted it, thousands of our activists and members of these organizations may be affected by its outcome.

The Environmental Management Commission's (EMC) proposal, a response to Rhodia Incorporated's petition for rulemaking, requests comment on three alternative provisions: Option 1, which modifies 15A NCAC 02L .0202(g)(59) by changing the groundwater quality standard for 1,1-DCE from 7 μ g/L to 350 μ g/L; Option 2, which modifies the same rule by granting the Commission permission to establish groundwater quality standards less stringent than those currently required; and Option 3, which establishes a new, statewide variance provision in 15A NCAC 02L .0113. Of these three options, Option 1 is preferable since it does not explicitly apply to contaminants other than 1,1-DCE or create statewide exceptions. However, without further limiting language, this option does have the potential to impact other groundwater quality standards.

As an alternative to the three options in the proposal, we suggest two more limited – and, we hope, faster – approaches to solving Rhodia's problem while retaining North Carolina's strong framework for protecting groundwater quality.

Background

Although the rulemaking options proposed by the EMC could have far-reaching consequences, shaping standards and future clean-up targets for scores of contaminants, the spur for the proposal was a specific, narrow issue involving a single toxic: 1,1- dichloroethylene. For that reason, it is worth considering the health impacts of this specific chemical before turning to the larger implications of the changes sought by Rhodia and the proposals offered by the EMC.

1,1-dichloroethylene (also known as 1,1-DCE or vinylidene chloride) is a volatile organic compound¹ not known to occur naturally.² It is used in the manufacture of adhesives, synthetic fibers, refrigerants, food packaging, and coating resins such as the saran types.³ When used in manufacturing, 1,1-DCE is a colorless liquid with a mild, sweet, chloroform-like odor.⁴

1,1-DCE enters the environment through releases into the water or onto the land. Specifically, 1,1-DCE is released into the environment via emissions or in wastewater during manufacturing processes, such as the breakdown of polyvinylidene chloride (PVDC) products and from the biotic or abiotic breakdown of 1,1,1-trichloroethane, tetrachloroethene, 1,1,2-trichloroethene, and 1,1-dichloroethane. The chemical is often found at hazardous waste sites.

Upon release into the water or on land, 1,1-DCE evaporates, or volatizes, rapidly into the air. ⁹ In fact, volatilization is the major transport process from water, soil, and sediment because of 1,1-DCE's high vapor pressure and low water solubility. ¹⁰ The chemical's atmospheric half-life, or the time it takes for half of a quantity of 1,1-DCE vapor to degrade into other chemicals, is estimated at 16 hours. ¹¹

The exact rate that 1,1-DCE volatizes is dependent on its specific environment. One wastewater biodegradation study found that 45% to 78% of 1,1-DCE was lost in 7 days, with a sizeable fraction of the loss attributable to volatilization. ¹² Further, ninety-seven percent of 1,1-DCE was removed in a wastewater treatment plant, but again, the exact fraction lost via volatilization is unknown. ¹³

¹ New Jersey Dept. of Health & Senior Services, Div. of Environmental. & Occupational Health Services, *Volatile Organic Compounds in Drinking Water*.

² Bob Benson, Environmental Protection Agency, <u>Concise International Chemical Assessment Document</u> <u>51, 1, 1-DICHLOROETHENE (VINYLIDENE CHLORIDE)</u>, World Health Organization et al. (2003) [hereinafter Benson].

³ Environmental Protection Agency, <u>Basic Information about 1, 1-Dichloroethylene in Drinking Water</u> (last visited June 25, 2012) [hereinafter <u>Basic Information</u>].

⁴ Id.

⁵ Environmental Protection Agency, <u>Technical Fact Sheet on: 1, 1-DICHLOROETHYLENE</u> (last visited June 25, 2012) [hereinafter *Technical Fact Sheet*].

⁶ Id

⁷ Benson, *supra* note 2.

⁸ *Id*.

⁹ *Id*.

¹⁰ *Id*.

¹¹ Benson, *supra* note 2.

¹² Technical Fact Sheet, supra note 5.

¹³ *Id*.

1,1-DCE in groundwater or landfills, where it cannot volatize, undergoes reductive dechlorination and becomes vinyl chloride, ¹⁴ a gas that is slightly water soluble and quite flammable. ¹⁵ Vinyl chloride is oncogenic, or tumor-causing. ¹⁶ Remediation techniques for vinyl chloride are different than those suggested for 1,1-DCE, and include microbial oxidation, ¹⁷ use of oxygen release compounds, ¹⁸ and *in situ* chemical oxidation. ¹⁹ Therefore, the introduction of 1-1,DCE into the environment can lead to a more expansive, complicated cleanup necessitated by the formation of vinyl chloride.

When non-volatized 1,1-DCE remains in the groundwater, 50% of the 1,1-DCE disappears in 5 to 6 months. ²⁰ In landfills, 1,1-DCE degrades in 1 to 3 weeks. ²¹

The principal sources of environmental exposure to 1,1-DCE for humans are contaminated ambient air and drinking water, ²² with the latter likely resulting from discharges from industrial chemical factories. ²³

Most exposures occur when people breathe 1,1-DCE vapors, such as those emanating from products containing 1,1-DCE. ²⁴ For example, 1,1-DCE vapor inhalation can occur when a contaminated home water supply is used for cooking, laundering, or bathing. ²⁵ Further exposure occurs by drinking contaminated water. ²⁶ Additionally, people who work or play around contaminated soils might be exposed if they touch their mouths or eat with dirty hands, two very likely possibilities for children. ²⁷ Finally, 1,1-DCE also can be absorbed by the skin; if home water supplies are contaminated, people may absorb the chemical through their skin when bathing or washing dishes. ²⁸

The human body rapidly absorbs 1,1-DCE following either inhalation or oral exposure. ²⁹ Although the body distributes 1,1-DCE to all tissues, it especially accumulates in the liver and kidneys. ³⁰ People exposed to sufficiently high concentrations of 1,1-DCE experience central nervous system depression

¹⁴ *Id*.

¹⁵ Environmental Protection Agency, *Vinyl Chloride* (last visited July 2, 2012).

¹⁶ Terminology Services, *Terms & Acronyms – <u>Search Results</u>*, Environmental Protection Agency (last visited June 25, 2012).

¹⁷ P.M. Bradley & F.H. Chapelle, <u>Microbial Oxidation of Vinyl Chloride Under Iron-Reducing Conditions</u>, U.S. Geological Survey, (last visited July 2, 2012).

¹⁸ Remediation of Vinyl Chloride, ORC Technical Bulletin # 2.2.2.3, (last visited July 2, 2012).

¹⁹ Richard Wong & Mark Bonsavage, <u>Pilot Scale Remediation of Dissolved Vinyl Chloride Plume Using In Situ Chemical Oxidation</u> (May 9, 2011).

²⁰ Technical Fact Sheet, supra note 5.

²¹ *Id*.

²² Benson, *supra* note 2.

²³ Id.

²⁴ Wisconsin Department of Health Services, <u>1,1-DICHLOROETHYLENE</u> (last visited June 19, 2012) [hereinafter Wisconsin].

²⁵ *Id*.

²⁶ *Id*.

²⁷ Id.

²⁸ Id.

²⁹ Benson, *supra* note 2.

³⁰ *Id*.

and symptoms of inebriation, convulsions, spasms, and unconsciousness.³¹ Respiratory effects include inflammation of the mucous membranes.³² Some individuals who consume water containing 1,1-DCE well in excess of the maximum contaminant level (MCL)³³ over a period of years could experience liver problems.³⁴ 1,1-DCE has been detected at low levels in a number of drinking water supplies across the United States.³⁵

Animal testing confirms that 1,1-DEC can harm the liver, kidneys, and the Clara cells of the lung. ³⁶ After low-dose, long-term exposure, the liver is the major target organ in rats following oral or inhalation exposure, but the kidney is the major target organ in mice following inhalation exposure. ³⁷

Epidemiological evidence is limited.³⁸ Still, 1,1-DCE is considered to be a possible cancer-causing substance.³⁹ Following high doses of 1,1-DCE, laboratory animals developed kidney and adrenal gland tumors.⁴⁰

As of EPA's 2012 Edition of the Drinking Water Standards and Health Advisories, ⁴¹ 1,1-DCE was listed with an "S" class cancer descriptor, meaning that it is "suggestive of carcinogenic potential." The uncertainty factor for 1,1-DCE remains 10.0; 1,1-DCE previously was categorized as a "C" class contaminant with a 10.0 uncertainty factor. ⁴³

The Proposed Options Generally

This rulemaking stems from a breakdown in a single cleanup under North Carolina's existing groundwater remediation rules. Rhodia, Incorporated (Rhodia) is a specialty chemical company that previously operated a chemical manufacturing facility in Gastonia. In the course of operations, the facility introduced 1,1-DCE into groundwater at the site. Rhodia has not owned the plant since 1997, but has operated a 'groundwater remediation system' there for over a decade. In 2009, during the triennial review of groundwater standards, Rhodia asked the Division of Water Quality to relax the

³¹ Environmental Protection Agency, <u>Vinylidene Chloride (1, 1-Dichloroethylene)</u>(last visited June 18, 2012) [hereinafter *Vinylidene Chloride*].

³² *Id*.

³³ As established by the Environmental Protection Agency and explained below

³⁴ Benson, *supra* note 2.

³⁵ Vinylidene Chloride, supra note 31.

³⁶ Benson, *supra* note 2.

³⁷ Id.

³⁸ Benson, *supra* note 2.

³⁹ Wisconsin, *supra* note 24.

⁴⁰ Id

⁴¹ Environmental Protection Agency, <u>2012 Edition of the Drinking Water Standards and Health Advisories</u> (April 2012) [hereinafter *2012 Standards*].

⁴² Jon B. Marshack, <u>Water Boards Training Academy Course WQ120</u>, Water Quality Goals (February 2012).

⁴³ Id.

⁴⁴ Letter from Benne C. Hutson, Attorney, McGuireWoods LLP, to Sandra Moore, NC Division of Water Quality (May 28, 2009) [hereinafter <u>Rhodia Letter 1</u>].

standard for 1,1-DCE. The agency, while sympathetic to the company's concerns, felt legally constrained to retain the current state standard. Subsequently, Rhodia sought a variance from the standards, but determined a variance – which would have required site by site analysis – was not really a remedy for a mismatch in the cleanup standard. Rhodia now seeks an amendment to the 1,1-DCE groundwater quality standard that would allow well owners to avoid remediation obligations for 47 of the 66 wells in North Carolina that are known to exceed the current standard.s.

Rhodia's argument hinges on a shortcoming of EPA's published maximum contaminant level (MCL) 45 for 1,1-DCE. In setting groundwater standards, state rules require the Division of Water Quality to choose the most stringent of several different methods or numbers, of which the MCL is one. 46 In the case of 1,1-DEC, the MCL is (by far) the most stringent. 47 However, when EPA sets an MCL, it relies on formulas built around other numbers. In the case of 1,1-DCE, EPA freely admits that the currently-published MCL of 7 μ g/L has been flawed since 2002, when EPA, in a rulemaking, changed the official numbers that fed into the agency's MCL formula but did not make the necessary correction to the MCL itself. However, the agency has not found it worthwhile to fix the MCL, a process that would require federal rulemaking. As a result, North Carolina's rules appear to trap the state (and Rhodia) with a groundwater standard that is inconsistent with EPA's own health data and that requires an unnecessarily strict cleanup.

At Rhodia's urging, the EMC has suggested three alternative rule revisions, each of which involves abandoning EPA's MCL and falling back to one of the other five, more lenient criteria mentioned in state rules:

• Option 1 modifies 15A NCAC 02L .0202(g)(59) by changing the groundwater quality standard for 1,1-DCE from 7 μ g/L to 350 μ g/L;

⁴⁵ A maximum contaminant level, or MCL, is the "highest level of a contaminant that EPA allows in drinking water. MCLs ensure that drinking water does not pose either a short-term or long-term health risk. EPA sets MCLs at levels that are economically and technologically feasible. Some states set MCLs which are more strict than EPA's." Environmental Protection Agency, <u>Drinking Water Glossary</u> [hereinafter *Drinking Water Glossary*].

⁴⁶ 15A NCAC 02L .0202(d). The six criteria are: the systemic threshold concentration; the concentration which corresponds to an incremental lifetime cancer risk of 1x10-6; the taste threshold limit value; the odor threshold limit value; the maximum contaminant level; or the national secondary drinking water standard.

 $^{^{47}}$ Of the six criteria listed in 15A NCAC 02L .0202 and described in the previous footnote, only the maximum contaminant level and the systemic threshold concentration exist for 1,1-DCE. Presentation by Sandra Moore, North Carolina Division of Water Quality, to the North Carolina Environmental Management Commission (July 14, 2011), [hereinafter Moore]. The technically correct maximum contaminant level for 1,1-DCE is 35 $\mu g/L$; the systemic threshold concentration is 350 $\mu g/L$.

 $^{^{48}}$ This is an extraordinarily rare situation. First, it is unusual for EPA to have made the technical mistake in 2002 of changing inputs to its formula without correcting the output, the MCL. Second, the volatility of this particular chemical, noted above, means that drinking water systems will almost never face a violation of the incorrect 7 μ g/L standard, so for EPA's purposes, the incorrect standard does not hurt any public utilities. It would be a mistake to conclude that North Carolina cannot safely rely on EPA's MCLs for the vast majority of contaminants.

- Option 2 modifies 15A NCAC 02L .0202 by adding a new Subsection (f), which allows the Commission to establish less stringent groundwater quality standards than would be established by using the six criteria in Subsection (d) and prioritized references in Subsection (e); and
- Option 3 modifies the 15A NCAC 02L .0113 variance rule by expanding the current site-specific variance provision and adding a state-wide variance provision for groundwater standards established in Section .0202.

We do not think abandoning the MCL is either necessary or wise. Instead, the EMC can and should take an approach that effectively corrects the flaws in the current published MCL to reflect a genuinely safe level of 35 μ g/L. Below, we explain both why this is scientifically correct and how it is legally permissible. Here, we observe that allowing Rhodia to leave groundwater concentrations of 1,1-DCE of up to 350 μ g/L – which all three options would permit – does not protect human health. On a broader level, we also note that options 2 and 3 unnecessarily open the door to lax discharge and remediation requirements for other toxics in the future.

The currently published MCL, the technically correct MCL, and Rhodia's request

Since the value of 35 μ g/L has been largely invisible in the record so far, before we address the three proposed options, we will explain why we believe the EMC should recognize it as the 'technically correct' MCL, and use it as a basis for regulating Rhodia.

As noted briefly above, EPA sets maximum contaminant levels (MCL) as enforceable drinking water quality standards for public water systems under the federal Safe Drinking Water Act. ⁴⁹ MCLs balance public health goals against fiscal and technical constrains on the ability of water systems to detect and remove contaminants. For purposes of the current rulemaking, it is vital to understand how the MCL is usually calculated. ⁵⁰

In calculating an MCL, EPA starts with the oral reference dose (RfD), the agency's best estimate of a safe daily dose of the chemical over a lifetime. ⁵¹

Second, EPA determines the drinking water equivalent level (DWEL):52

⁴⁹ 42 U.S.C. §§ 300g-1(b); Environmental Protection Agency, <u>Safe Drinking Water Act Amendments of 1996</u> (August 1996).

⁵⁰ Environmental Protection Agency, <u>1,1-DICHLOROETHYLENE Health Advisory</u> (1987)(explaining how to derive the MCL).

⁵¹ An RfD is "[a]n estimate (with uncertainty spanning perhaps an order of magnitude) of a daily oral exposure to the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a lifetime." Environmental Protection Agency, <u>IRIS Glossary/Acronyms & Abbreviations</u>, (last visited June 20, 2012) [hereinafter IRIS Glossary].

⁵² A DWEL is "a drinking water lifetime exposure level, assuming 100% exposure from that medium, at which adverse, noncarcinogenic health effects would not be expected to occur." *2012 Standards*, *supra* note 41.

[RfD mg/kg/day x 70 kg (the assumed body weight of an adult)] / [2 L/day (assumed daily water consumption of an adult)] = DWEL

From those two numbers, EPA derives the MCL:

[DWEL mg/L x 0.20 (assumed relative source contribution from water for organics)] / [10 (the additional uncertainty factor for class "S" carcinogens)] = MCL

Keeping the above general calculations in mind, one can calculate both the currently published and technically correct MCLs for 1,1-DCE. The difference between the two values results from differences in the RfDs. The 1987 RfD of 0.010 mg/kg/day yields an MCL of 7 μ g/L. The current and updated RfD of 0.050 mg/kg/day yields an MCL of 35 μ g/L. The full calculations are as follows:

Using the 1987 RfD for 1,1-DCE:

RfD: 0.010 mg/kg/day

DWEL: $[0.010 \text{ mg/kg/day x } 70 \text{ kg}] / [2 \text{ L/day}] = 0.350 \text{ mg/L or } 350 \text{ } \mu\text{g/L}$

MCL: $[0.350 \text{ mg/L} \times 0.20] / [10] = 0.007 \text{ mg/L} \text{ or } 7 \text{ µg/L}$

The revised, current RfD:

RfD: 0.050 mg/kg/day

DWEL: [0.050 mg/kg/day x 70 kg] / [2 L/day] = 1.75 mg/L or 1750 µg/L

MCL: $[1.75 \text{ mg/L} \times 0.20] / [10] = 0.035 \text{ mg/L} \text{ or } 35 \text{ } \mu\text{g/L}$

As the calculations demonstrate, the correct value of the MCL for 1,1-DCE, using EPA's corrected inputs and standard method, is 35 μ g/L. That EPA forgot or neglected to change the MCL when it changed the RfD in 2002 does not eliminate the benefit of protections that a correctly calculated MCL can provide. Rhodia asks the EMC to set aside the MCL altogether, but as a matter of science, that is not protective of public health. The remaining question – which we tackle below, after comments on the three options proposed by the EMC – is to explore how, legally, the EMC can move forward with a solution built around the health-protective standard of 35 μ g/L.

Option 1

The proposed rule change specific to Option 1 is problematic because it conflicts with the applicable NCAC provisions found in 15A NCAC 02L .0202(d) and (e). Per these provisions, data from sources listed in (e) must be used to determine the most stringent of the 6 criteria listed in (d), which in turn is used to

⁵³ National Primary Drinking Water Regulations; Announcement of Completion of EPA's Review of Existing Drinking Water Standards, 68 Fed. Reg. 42,908, 42,918 (July 18, 2003) (noting that the RfD is 0.046 mg/kg/day; this number then is rounded to 0.050 mg/kg/day) [hereinafter 2003 Federal Register].

establish groundwater quality standards. Of the six criteria, only two have established values for 1,1-DCE: the MCL and the systemic threshold concentration. ⁵⁴ The systemic threshold concentration formula yields a value of 350 μ g/L; Rhodia proposes that the EMC scrap the MCL and instead rely on this number. ⁵⁵ In doing so, the company urges the EMC to set the state groundwater standard at a level *ten* times higher than the technically correct MCL.

Option 1 is certainly preferable to Options 2 and 3 in that it does not apply to contaminants other than 1,1-DCE or create statewide exceptions from groundwater protections. That said, adoption of Option 1 may provoke rulemaking petitions to change standards for other chemicals by fiat (rather than as required by 15A NCAC 02L .0202). In any event, as noted above, Option 1 is not protective of human health.

Option 2

Option 2, which applies to all 147 listed contaminants rather than to just 1,1-DCE, allows the EMC to establish groundwater quality standards less stringent than the MCL or national secondary drinking water standards, an allowance that renders the six standards in subsection (d) moot. .

We recommend the EMC reject this option for three reasons.

- This option is overbroad. That is, it is not limited to 1,1-DCE, but applies to all 147 of the contaminants listed in current Subsection (g). Option 2 opens the door for future weakening rule revisions to any and all of the other 146 contaminants by applying the overly-flexible standard suggested by this option.
- This particular revision cannot be implemented without placing human health at risk. The
 current rule provides in Subsection (d) six criteria for the EMC to compare in establishing
 groundwater quality standards, the most stringent of which is to be adopted. Since the MCL
 represents the highest level at which a contaminant might exist in the groundwater without
 resulting in short- or long-term health risks,⁵⁶ there is no way to allow a weaker, riskier standard
 to usurp the MCL and not endanger the public health and safety.
- It makes little sense to target *both* the MCL and the national secondary drinking water standards, as Option 2 does, given that the focus of the current rulemaking efforts is 1,1-DCE, a contaminant without a national secondary drinking water standard.

Option 3

Similar to Option 2, the impacts of Option 3 would reach well beyond the groundwater standard for 1,1-DCE. The proposed statewide variance provision would allow a chemical company to obtain a

⁵⁴ Moore, *supra* note 47.

⁵⁵ Letter from Benne C. Hutson, Attorney, McGuireWoods LLP, to Coleen Sullins, Director, NC Division of Water Quality (March 14, 2011) [hereinafter <u>Rhodia Letter 2</u>].

⁵⁶ Drinking Water Glossary, supra note 45.

blanket exception to any groundwater quality standard, not just 1,1-DCE. Since there is no indication that similar technical errors exist for any of the other 146 listed contaminants in 15A NCAC 02L .0202, it seems unwise for the EMC to create such an invitation to circumvent the protections currently built into the rule.

Rhodia's arguments for 350 µg/L

Beyond commenting on the options, we want to address two rationales offered by Rhodia for raising the groundwater standard for 1,1-DCE to $350 \mu g/L$.

Reliance on EPA's MCL. Over the past several years, Rhodia has claimed that it is illegal⁵⁷ under North Carolina regulations to use the current MCL in establishing groundwater quality standards for 1,1-DCE. This argument is based on the idea that an agency "must follow its own regulations...Failure to follow such regulations and guidelines is generally an arbitrary and capricious action." However, Rhodia's categorization is based on an inaccurate reading of the applicable rules and a misleading statement of the law.

First, and as explained above, both the currently published and technically correct MCLs are the "least of" the six criteria listed in Subsection (d). Whether one uses 7 μ g/L or 35 μ g/L, both numbers derive from IRIS data (the first being the currently published MCL and the latter being the MCL that results from using the currently published RfD). By using the MCL and therefore utilizing IRIS data, the EMC also is complying with Subsection (e), which requires that priority be given to IRIS reference information. Therefore, EMC's use of the MCL has been both functionally appropriate and protective of human health and groundwater quality. As noted above, adopting the systemic threshold concentration would not only be in conflict with state rules; it would not fully protect human health.

Additionally, the provision in Subsection (e) that requires that the listed references be used in determining the six criteria in Subsection (d) merely point to the sources of information that EMC must consult. The provision does not specify the result the EMC must reach, or that the information provided by the Subsection (e) sources exactly mirror the values in Subsection (d). The restrictive reading suggested by Rhodia⁵⁹ would conflict with the broad authority granted to the EMC.⁶⁰ The current groundwater quality standard complies with this rule, but it would also continue to comply using our suggested alternatives.

Cancer risk concentration. Rhodia has also argued that 1,1-DCE's cancer risk categorization somehow disallows cancer criteria from being used to establish such a standard. Specifically, Rhodia claims that "U.S. EPA removed the oral cancer slope factor for 1,1-DCE and concluded that the compound was only

⁵⁷ "[U]nder North Carolina regulations, the current MCL legally cannot be used to establish the 2L groundwater quality standard for 1,1-DCE." Rhodia Letter 2, *supra* note 55.

⁵⁸ Glendale Neighborhood Association v. Greensboro Housing Authority, 901 F. Supp. 996, 1003 (M.D.N.C. 1995).

⁵⁹ Rhodia Letter 1, *supra* note 44.

⁶⁰ See, e.g., Gen. Stat. § 143B-282(d).

a possible human carcinogen. As a result, there is no cancer risk concentration that can be used to establish a groundwater quality standard." 61

First, and possibly most importantly, EPA has *not* removed the oral cancer slope factor for 1,1-DCE. The oral cancer slope remains listed in IRIS and numerous other recent EPA documents.⁶²

Further, Rhodia wrongly equates the specific oral cancer slope factor with *any* "cancer risk concentration." An oral cancer slope factor is:

An upper bound, approximating a 95% confidence limit, on the increased cancer risk from a lifetime oral exposure to an agent. This estimate, usually expressed in units of proportion (of a population) affected per mg/kg-day, is generally reserved for use in the low-dose region of the dose-response relationship, that is, for exposures corresponding to risks less than 1 in 100.63

Somewhat confusingly, the only logical reason to mention the oral cancer slope would be to eliminate 15A NCAC 02L .0202(d)(2) – incremental lifetime cancer risk – from consideration in establishing the 1,1-DCE groundwater quality standard, something that DWQ already has done.⁶⁴

Similarly, Rhodia exaggerates the scope of a 2003 EPA 1,1-DCE exposure analysis by suggesting that "EPA concluded that the risks previously cited for 1,1-DCE had been *greatly overstated*."⁶⁵ This exposure analysis was based on a 16-state cross section did not include North Carolina and included states where 1,1-DCE was "unlikely to occur at concentrations above 0.007 mg/L."⁶⁶ It goes without saying that 1,1-DCE does occur in North Carolina at concentrations above .007 mg/L, as is evidenced by Rhodia's rulemaking petition and the 66 wells with 1,1-DCE concentrations above this threshold.⁶⁷ Therefore, it is neither accurate nor reasonable to say that EPA has concluded that 1,1-DCE risks have been "greatly overstated" for North Carolina.

At most, EPA has acknowledged that "[a]Ithough there are new health effects data that *might* support calculation of a less stringent standard for 1,1-dichloroethylene, EPA does not believe a revision...is appropriate at this time." EPA reached this same conclusion in its most recent 2010 review as well. ⁶⁹ As explained below, our suggested alternative comports with this conclusion.

⁶¹ Rhodia Letter 2, *supra* note 55.

⁶² Environmental Protection Agency, <u>1,1-Dichloroethylene (CASRN 75-35-4)</u>, Integrated Risk Information System, (last visited June 21, 2012); *Vinylidene Chloride*, *supra* note 31 (citing IRIS).

⁶³ IRIS Glossary, supra note 51.

⁶⁴ Moore, *supra* note 47.

⁶⁵ Rhodia Letter 2, *supra* note 55 (emphasis added).

⁶⁶ 2003 Federal Register, *supra* note 53, at 42,919-42,921; Environmental Protection Agency, <u>Occurrence Estimation Methodology and Occurrence Findings Report for the Six-Year Review of Existing National Primary Drinking Water Regulations</u> (2003).

⁶⁷ Rhodia Letter 2, *supra* note 55.

⁶⁸ 2003 Federal Register, *supra* note 53, at 42,921.

⁶⁹ National Primary Drinking Water Regulations; Announcement of the Results of EPA's Review of Existing Drinking Water Standards and Request for Public Comment and/or Information on Related Issues, 75 Fed. Reg. 15,500, 15,519, 15,536 (March 29, 2010) [hereinafter 2010 Federal Register].

Ultimately, these arguments offered by Rhodia are a distraction from the real issues: that there is no scientific basis on which to modify the groundwater quality standard to 350 μ g/L; that doing so will result in a standard far higher than the technically correct MCL of 35 μ g/L; and that adopting the higher standard would put human health and the environment at risk.

Remediation Feasibility and Costs

Remediating low 1,1-DCE concentrations is entirely feasible.

Granular activated carbon. EPA has identified granular activated carbon (GAC) adsorption and packed tower aeration as the best available treatment technologies for removing 1, 1-dichloroethylene from drinking water. Granular activated carbon (GAC) used in conjunction with packed tower aeration has been proven to remove 1,1-DCE to below 7 μg/L. More broadly, GAC adsorption is a widely accepted treatment technology for removing VOCs from groundwater. The acceptance of GAC adsorption is, in part, based on its long history of effectively removing organic contaminants from water and wastewater. GAC adsorption is an extremely versatile technology, and for many water treatment applications, it is the least expensive treatment option. Granular activated carbon is relatively inexpensive and readily available. For example, it can be purchased either online or from aquarium supply shops. Of relevance here, carbon adsorption is particularly effective in treating low concentration waste streams, allowing those responsible for cleanup to meet stringent treatment levels. One of the major attributes of activated carbon treatment is its ability to remove a wide variety of toxic organic compounds to non-detectable levels (99.99%). Cleanup costs using this method typically depend on the amount of carbon consumed.

Additional GAC adsorption cost savings might be realized if the MCL were increased to 35 μ g/L from 7 μ g/L, ⁸⁰ a fact that supports raising the currently published MCL to the technically correct MCL. It is reasonable to assume that this cost-saving change would gain the regulated community's support.

⁷⁰ Commonwealth of Pennsylvania, Department of Environmental Protection, <u>Citizen's Guide to Volatile</u> <u>Synthetic Organic Chemicals in Drinking Water</u> (last visited June 18, 2012).

⁷¹ Basic Information, supra note 3.

⁷² Mark H. Stenzel & William J. Merz, <u>Use of Carbon Adsorption Processes in Groundwater Treatment</u>, Calgon Carbon Corporation, (last visited June 18, 2012).

⁷⁴ Austin R. Shepherd, <u>Granular Activated Carbon for Water & Wastewater Treatment</u>, Carbtrol Corporation (1992), [hereinafter Shepherd].

⁷⁵ Josh Kearns, <u>Five-Gallon Bucket Filter for Rooftop Harvested Rainwater</u>, Aqueous Solutions (2008).

⁷⁷ Shepherd, *supra* note 74.

⁷⁸ Id.

⁷⁹ Id

⁸⁰ See Environmental Protection Agency, <u>Occurrence Analysis for Potential Source Waters for the Six-Year Review of National Primary Drinking Water Regulations</u> (2009).

Packed tower aeration. A second preferred method of remediating VOC-contaminated groundwater is packed tower aeration, also known as air stripping. ⁸¹ Air stripping is a simple, reliable, and effective water treatment technology for removing dissolved VOCs. ⁸² VOC removals in excess of 99.99% are possible with this treatment method. ⁸³ Additionally, air stripping reduces organic chemical concentrations from as high as 50 parts per million to below detection limits. ⁸⁴ The detection limit for 1,1-DCE is 0.50 μg/L. ⁸⁵Packed air stripping costs, while proportional to air flow, are less expensive than similar water treatment methods. ⁸⁶

Based on the above discussion, it appears that although the regulated community will incur *some* 1,1-DCE remediation costs at an MCL of 35 μ g/L, those costs are low relative to other remediation methods, ⁸⁷ and even lower compared to those likely absorbed by public water supply systems. ⁸⁸ Alternatively, EPA found that revising the existing MCL for 1,1-DCE would not provide a meaningful opportunity for cost savings. ⁸⁹ Therefore, even if remediation costs were to fluctuate as a result of modifying the MCL, the cost impacts would be negligible.

Despite the presumably low remediation costs associated with 1,1-DCE, Rhodia still suggests increasing the 1,1-DCE's groundwater quality standard to 350 μ g/L. Per a letter to the director of DWQ from Rhodia's attorney, Rhodia and others have 66 wells with 1,1-DCE concentrations of greater than 7 μ g/L, and 19 wells with concentrations greater than 350 μ g/L. ⁹⁰ The result of the petitioned-for increase would be to allow the abandonment of 47 groundwater wells with 1,1-DCE concentrations somewhere between 7 μ g/L and 350 μ g/L. ⁹¹ It is reasonable to assume that at least *some* of these 47 wells contain 1,1-DCE concentrations of greater than 35 μ g/L, and above the safe threshold for drinking water.

Rhodia also argues that increasing the 1,1-DCE groundwater standard to 350 μ g/L would "achieve more clean up of the environment quicker than the current standard of 7 μ g/L will ever provide." It goes without saying that looser regulatory requirements will be easier for industry to meet. However, it is erroneous to imply that meeting a standard that is ten times greater than the technically correct standard will somehow provide greater remediation. There is a clear difference between simply meeting

⁸¹ Motorola, *Air Stripping Water Treatment Technology*, (last visited June 22, 2012).

⁸² Id

⁸³ Jaeger Products, Inc., <u>Air Stripping of VOCs from Water</u>, (last visited July 2, 2012) [hereinafter Jaeger Products].

⁸⁴ Wireless Information System for Emergency Responders, <u>1,1-Dichloroethylene</u> (last visited June 22, 2012).

⁸⁵ Environmental Protection Agency, <u>U.S. EPA's Methods and Minimum Detection Limits</u>, (last visited June 22, 2012).

⁸⁶ Jaeger Products, *supra* note 83.

⁸⁷ See, e.g., Environmental Protection Agency, <u>Cost Analysis for Selected Groundwater Cleanup Projects:</u>
<u>Pump and Treat Systems and Permeable Reactive Barriers</u> (2001); Fredric Hoffman, <u>Ground Water</u>
<u>Remediation Using "Smart Pump and Treat"</u>, 31 Ground Water 1 (1993).

⁸⁸ Fiscal Impacts of Proposed 15A NCAC 02L Groundwater Rules (2012).

⁸⁹ 2010 Federal Register, *supra* note 69.

⁹⁰ Rhodia Letter 2, *supra* note 55.

⁹¹ *Id*.

⁹² *Id*.

a more lenient standard and actually achieving the greater and more meaningful cleanup that would be achieved by remediating to a 35 μ g/L standard.

The Preferable Alternatives

Rather than adopting any of the three proposed options, we suggest adopting either of the two alternatives described below.

Judicial notice. First, the EMC might take judicial notice of the fact that the MCL currently published by EPA is technically incorrect, and that the correct MCL is $35 \,\mu\text{g/L}$. The EMC's status as a quasi-judicial entity and ability to take judicial notice are confirmed both by General Statute 143B-282.1 and by EMC and court precedent. The advantage of this option is that it would allow the EMC to revise the current 1,1-DCE standard quickly and without engaging in the formalities of the rulemaking process. This is an option that we or others could have suggested at the time of Rhodia's previous variance request, but, not being a party to that quasi-judicial proceeding, any comment would have been disallowed as an ex parte communication. However, the EMC might now make up for lost time by adopting this option.

Should this approach interest the EMC, two points merit further discussion: how to deploy it, and whether it opens the door to future abuses. On the first, 'judicial notice' does not seem available until a decision enters a quasi-judicial phase. We think, if Rhodia applies for a permit based on the 35 μ g/L standard and it is denied, and Rhodia appeals that decision to the Division Director, the matter at that stage would become quasi-judicial, and the Director could take notice of and correct the technical error in EPA's standard.

On the second question, we think the terms of judicial notice limit the potential abuse. Judicial notice can only be taken of indisputable facts; in this case, of the fact putting the updated RfD into the EPA's formula for a MCL yields a value of 35 μ g/L, not 7 μ g/L. It cannot be used to replace one standard with another on the basis that the replacement is more protective, more reasonable, or more scientifically valid. For example, one cannot take judicial notice that the correct standard should be 350 μ g/L; that is a policy choice, not a correction of a mathematical error. But where, as here, a failure to correctly complete the formula in EPA's 2002 rulemaking has resulted in a standard of 7 μ g/L instead of 35 μ g/L, invocation of judicial notice may be appropriate.

Narrow rule change. A second alternative is for the EMC to revise the current rule by using the technically correct MCL, as calculated and described in previous sections. Rhodia has argued (without reference to specific rule provisions) that the "regulations require that groundwater quality standards

⁹³ "Acting in its quasi-judicial authority, the Commission may take judicial notice of changes in the law or regulations and matters of common knowledge relevant to the issue in a contested case." <u>United States</u> <u>Fish & Wildlife Service v. North Carolina Division of Air Quality</u>, 08 EHR 1067 (Environmental Management Commission May 27, 2011) (citing N.C.G.S. § 8C-1, Rule 201(b) (1992); *State v. Thompson*,

⁵⁰⁸ S.E.2d 277 (1998); *In Re Spivey*, 480 S.E.2d 693 (1997)).

94 Rhodia Letter 2, *supra* note 55. Although Rhodia submitted a variance request application in 2010, the application did not make it to stage at which the EMC accepts public comments.

must be based on current, valid scientific data,"⁹⁵ and adopting this alternative will accomplish this desired requirement.

In order to accomplish this, EMC also should revise 15A NCAC 02L .0202 by adding the following language as a new Subsection (f):

In cases where the U.S. EPA has delayed revising the maximum contaminant level despite having updated reference dose and other applicable data, and because of the administrative costs of a federal rulemaking proceeding, the Commission may adjust the Class GA and Class GSA standards in Subsections (h) and (i), respectively, to reflect the technically correct maximum contaminant level. Upon U.S. EPA revisions, the EMC will ensure that the adjusted maximum contaminant level remains consistent with the U.S. EPA's revised standard.

This language would allow the EMC to make such common sense revisions when warranted, and until EPA revises the MCL to accord with its own data.

Conclusion

We appreciate this opportunity to comment on the proposed rulemaking and thank you for your time and consideration. We appreciate both the EMC's and DWQ's attention to our state's groundwater quality standards and work toward protecting this vital resource. Ultimately, we hope that the final rule will continue to rigorously protect North Carolina's groundwater by sensibly incorporating EPA's technically correct and up-to-date data into the groundwater rules.

Sincerely,

Grady McCallie Shannon Arata

⁹⁵ <u>Presentation by Benne C. Hutson</u>, Attorney for Rhodia, Inc., McGuireWoods LLP, to the North Carolina Environmental Management Commission (July 14, 2011).



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To: Sandra Moore, Classifications & Standards Unit

From: Erin Wynia, Legislative & Regulatory Issues Manager
Re: Groundwater Quality Standards Proposed Changes

Date: July 2, 2012

Dear Ms. Moore,

The NC League of Municipalities is a membership organization of over 550 N.C. municipalities and affiliate organizations, many of which conduct permitted operations affected by the state's groundwater quality standards in 15A NCAC 02L .0202(d). These operations include disposal of treated wastewater to the subsurface; land application of residuals, biosolids and other waste treatment solids; use of treated wastewaters via spray irrigation; use of high capacity infiltration systems such as rotary distributors; operation of landfills; and use of underground storage tanks.

League members uphold a central tenet of environmental stewardship, as stated in the member-adopted Core Municipal Principles: "Local governments are partners with state and federal agencies in protecting the environment and quality of life for our citizens." Cities and towns understand their responsibility in ensuring the highest possible water quality in their communities. Therefore, they allocate extensive staff time and public financial resources to activities that protect water quality in an efficient manner based on sound science.

On behalf of the League's members, I appreciate the opportunity to provide feedback on the request by the N.C. Environmental Management Commission (EMC) to receive proposals that allow flexibility in the implementation of the groundwater quality standards, and we look forward to working with you and other stakeholders on possible revisions to the method of setting groundwater quality standards.

Respectfully submitted,

Erin L. Wynia

Erin L. Wynia

Legislative & Regulatory Issues Manager

ewynia@nclm.org

(919) 715-4126

NCLM commentsGroundwater Quality Standards Proposed ChangesJuly 2, 2012

Background

North Carolina's groundwater quality standards program exists to ensure the state's groundwaters are preserved for their best usage. This policy goal for the program is enshrined in the N.C. Administrative Code, where each of the three classifications of groundwaters in 15A NCAC 02L .0201 defines "best usage" of groundwater as human consumptive uses, such as drinking water or other domestic uses:

- <u>Class GA groundwaters</u>: Existing or potential source of drinking water supply for humans
- <u>Class GSA groundwaters</u>: Existing or potential source of water supply for potable mineral water and conversion to freshwater
- <u>Class GC groundwaters</u>: A source of water supply for purposes other than drinking, including other domestic uses by humans

Given that the aim of the program is to protect groundwaters for human drinking water consumption or domestic uses, the level at which standards are set should match that level of protection. But in many cases, current rules dictate groundwater standards that are far more protective than those needed for human drinking water or domestic uses. The EMC is hamstrung by the current inflexibility of 15A NCAC 02L .0202(d), which requires the EMC to set groundwater quality standards at the least of six values, regardless of whether the resulting standard exceeds the level of protection established by science as being protective of human drinking water consumption or domestic uses. The lowered standards result in unnecessary increased operational costs for the variety of permitted municipal operations affected by the standards.

Recommendations

For the reasons stated above, the League urges the EMC to revise 15A NCAC 02L .0202(d) to allow flexibility in setting groundwater quality standards. These recommendations should allow EMC to better align the standards with the program's goal of protecting groundwaters for human drinking water consumption and other domestic uses.

<u>Recommendation #1</u>: Rewrite the rule to read, "Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established according to the best available scientific guidance and include considerations such as..."

Recommendation #2: Consider revising the criteria listed in 15A NCAC 02L .020(d)(1) to reflect updated science on rates of human water consumption.



June 29, 2012

Sent Via E-mail (Sandra.Moore@ncdenr.gov)

Ms. Sandra Moore North Carolina Division of Water Quality – Planning Section 1617 Mail Service Center Raleigh, North Carolina 27699-1617

Re: Proposed North Carolina Groundwater Rule Revision

Dear Ms. Moore:

Progress Energy, Inc. is a regional energy company serving the Southeast region of the country. Progress Energy, Inc. is organized as a holding company that owns Carolina Power & Light Company, doing business as Progress Energy Carolinas, Inc. (PEC or the Company), a regulated electric utility operating in North and South Carolina that serves approximately 1.5 million homes, businesses and industries. On behalf of Progress Energy Carolinas, I offer the following comments on the proposed revisions to groundwater quality regulations as described the NC State Register Volume 26, Issue 21, published on May 1, 2012.

Notwithstanding our general objection to development and use of a statewide groundwater standard variance, the Company respectfully submits specific comments on each of the three rule change options proposed:

Option 1. A change in 02L .0202 (g) (59) to amend the 1,1-dichloroethylene standard from 7 ug/L to 350 ug/L;

PEC agrees with the proposed change in the standard as specified. The current standard is clearly incompatible with the intent of the NC regulations due to EPA's unwillingness to correct an MCL they know to be flawed.

Option 2. A change in 02L .0202 (d) and (f) to allow the EMC to establish a standard less stringent than the maximum contaminant level (MCL) when: (a) the MCL is not based on the most recent U.S. EPA health effects data as published in U.S. EPA's Integrated Risk Information System: (b). such a standard would not endanger public health and safety; and, (c) compliance with a standard based on the MCL would produce serious hardship without equal benefit.

PEC agrees that a rule adjustment such as Option 2 will allow the EMC to correct the erroneous standard referenced in Option 1. It may not be the best solution to the fundamental problem of overly restrictive standard selection rules using arbitrary criteria beyond the control of our regulatory agency. The Company suggests that if Option 2 is implemented by the Commission, it be considered a short term measure to be effectively replaced by the change proposed in our comments to Option 3.

Ms. Sandra Moore June 29, 2012 Page 2

Option 3: A change in 02L .0113 to allow the EMC to issue a state-wide variance to the 02L rules in .0113 (d).

PEC considers option 3 to be a cumbersome and unnecessary response to poorly worded current regulations that may impede reasonable and truly protective standards. The use of the term "state-wide" in .01113 (d) implies that the variance would apply everywhere within the state. However, Paragraph (2) requires a list of all known potentially affected sites, implying that the variance would only be applied to those specific sites. It should be clarified how the variance would apply to sites discovered after the variance is adopted, or known by any other than the applicant. Additionally, Paragraph (3) requires a "list of 'increased costs for treatment...due to the proposed variance." Normally, a variance is requested to allow a lower cost to the applicant, but may also result in additional expense for those who use affected wells for water supply. Both factors should be considered. Rather than using a confusing "state-wide variance", it is much more practical to establish appropriate state standards to begin with, and provide for specific corrections when an incorrect standard is encountered. An alternative proposal is discussed below.

In addition to the options outlined above, the EMC solicited other proposals that allow flexibility in implementation of 15A NCAC 2L .0202(d) while maintaining or achieving appropriate water quality and public health standards, recognizing that any such proposal, if acted upon, might constitute a substantial change from the proposed rule amendments described in detail in this public notice, and might require an additional rule-making procedure.

PEC proposes a more comprehensive and realistic solution for the dilemma that is encountered by DENR and the EMC when overly restrictive wording such as that in 02 L .0202 (d) must be implemented, requiring any standard be the least of several widely varying criteria that may be poorly or incorrectly defined, as in the case of 1,1-dichloroethylene. The rule directing the method to identify and select a value for a standard should not tie the hands of the EMC and DWQ in such a way that even when they know it is not correct or inappropriate, they must impose it regardless. Specifically, .0202(d) should be revised to read:

- (d) Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of using evaluation of the reliability, relative costs and benefits of:
 - (1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)];
 - (2) Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6;
 - (3) Taste threshold limit value;
 - (4) Odor threshold limit value;
 - (5) Maximum contaminant level; or
 - (6) National secondary drinking water standard.

Ms. Sandra Moore June 29, 2012 Page 3

Progress Energy's environmental policy promotes compliance with laws and regulations and it would be our preference to achieve compliance with standards rather than to rely upon variancees. For the sake of brevity, the details of the obvious flaws of the current version of .0202 (d) will not be elaborated here except for the simple specific example of requiring equal weight to both the subjective aesthetically-based guidance of the national secondary drinking water standards and the systemic threshold concentration or cancer risk when choosing a standard.

PEC appreciates the opportunity to provide comments on this important regulatory initiative. Please do not hesitate to contact Cam Wheeler at (919) 546-4655) if you have any questions.

Sincerely,

Michael Olive

Director, Environmental Services

ATTACHMENT K



North Carolina Department of Environment and Natural Resources

Division of Waste Management Dexter R. Matthews

Director

Dee Freeman Secretary

Beverly Eaves Perdue Governor

CERTIFIED MAIL
Return Receipt Requested

January 21, 2011

Michael J. Shatynksi Rhodia, Inc. 8 Cedar Brook Drive Cranbury, New Jersey 08512

Re: 1,1-DichloroetheneVariance Application

Rhodia (Former Rhone-Poulenc Facility)

207 Telegraph Road Gastonia, Gaston County, North Carolina IHSB # NONCD0002395

Dear Mr. Shatynksi:

The Division of Waste Management (Division) of the North Carolina Department of Environment and Natural Resources (NCDENR) is in receipt of your November 1, 2010 "Application for a Variance to the 2L Groundwater Quality Standard (GWQS) for 1,1-Dichloroethylene (1,1-DCE)." The Environmental Management Commission forwarded your application to the Division as this Division is overseeing the remedial action at the above referenced site (Site). I understand that you are requesting a variance to a 15A NCAC 2L standard (GWQS) for purposes of securing an alternate cleanup standard under the Division's Inactive Hazardous Sites Program. Division staff have reviewed your application and reported the following:

1. While a petroleum release from the on-site underground storage tank system and a catastrophic release of 1,1-dichloroethene from an aboveground storage tank in 1991 are known to have contributed to the Site's groundwater contamination plume, the application does not contain descriptions of other past activities and operations that have resulted in discharges of contaminants to the Site's environmental media. Other areas of soil and groundwater contamination previously documented in file records indicate other discharges of hazardous substances have occurred at the Site, which were not specified and adequately described in the variance application.



- 2. The lateral extent of the known 1,1-dichloroethene groundwater contamination plume is not fully depicted in Figure 6 of Exhibit L and Figure 7 of Exhibit M of the application. Note the absence in the figures of known 1,1-dichloroethene groundwater contamination at monitor well MW-34B (located immediately south and adjacent to monitor well MW-34A), which was reported in the Site's 2009 *Groundwater Monitoring Report*. Further, the November 1997 *Summary of Sampling Results* reported 1,1-dichloroethene concentrations that exceed the GWQS in groundwater samples from two temporary monitor wells within the vicinity of the SBR Polymer Plant. The two wells were located approximately 120 feet south and 180 feet southwest of monitor well MW-1A, and this data is also not illustrated on the aforementioned figures. These examples demonstrate the assessment to determine the extent of 1,1-dichloroethene contamination remains incomplete.
- 3. File records also indicate that the primary objective of the current corrective action at the Site was hydraulic control of the groundwater contamination plume using a groundwater recovery and treatment system (which began in 1996). The Revised February 1998 Performance Monitoring Plan reported that hydraulic containment failure would be indicated if contaminants are detected at groundwater monitor wells where they were not detected during previous sampling events prior to system startup (e.g. MW-38, MW-39, MW-35), or increases of contaminant concentrations are observed at groundwater monitor wells from the levels detected prior to system startup (e.g. MW-31A, MW-32, MW-33). The 2009 Groundwater Monitoring Report indicates that concentrations of 1,1-dichloroethene began to be detected in groundwater samples from monitor wells MW-38 in March 2007 and MW-39 in October 2006, and the reported concentrations of 1,1-dichloroethene from both monitor wells exceeded the GWQS for two consecutive sampling events in 2009. Since 2006, the concentration of 1,1dichloroethene in groundwater from monitor well MW-33 has consistently been reported at two orders of magnitude higher when compared to the initial 1,1-dichloroethene concentration in 1994. The 2009 Groundwater Monitoring Report clearly demonstrates that the current corrective action has failed to maintain hydraulic control, and the 1,1-dichloroethene contamination plume has continued to migrate beyond the established containment zone boundary. The figures included in the application (Figures 6 and 7) do not acknowledge the failure of the current corrective action strategy and do not show the potential for further groundwater plume migration.
- 4. The applicant's proposed variance does not consider potential exposure to contaminants in groundwater through structural vapor intrusion. The present groundwater contamination plume is known to extend into the surrounding residential area. The concentration of 1,1-dichoroethene reported in the 2009 *Groundwater Monitoring Report* exceeds the current Inactive Hazardous Sites Program Residential Screening Level (RSL) for an acceptable groundwater concentration. The exceedance of the RSL indicates a potential for vapors from groundwater contamination to enter residential structures. Additional assessment is needed to determine if there is in fact a vapor exposure concern. Rhodia has not performed this assessment.
- 5. A map that shows the location of wells and other water supply sources including details of well construction within ½ mile of the Site is required. A map that complies with this application requirement was not observed within the variance application.

Michael J. Shatynksi January 21, 2011 Page 3 of 3

- 6. The variance application reports that Rhodia, Inc. (Rhodia), on its own initiative, has routinely evaluated existing and emerging remedial technologies to assess their potential applicability at the Site. The application does not include these evaluations. Our file records indicate that the Division of Water Quality (DWQ) directed Rhodia to prepare an updated analysis of remedial alternatives each time a groundwater monitoring report was submitted since at least March 2003. Staff reviewed these reports and, in particular, the more recent Summary of Remedial Technologies listed in the 2009 Groundwater Monitoring Report. The data within the summary is insufficient to establish that achieving compliance with the current 1,1-dichloroethene GWQS with best available technology would be economically unreasonable. No supporting information of specific technologies including (without limitation) remedial pilot tests, economic cost estimates, and scientific literature references were observed. A review of the referenced monitoring report and variance application found that these documents do not contain supporting information that identifies the impact of costs on the applicant or that establishes compliance with the 15A NCAC 2L rule would produce serious financial hardship on the applicant. As a result, the application fails to demonstrate that compliance with the current GWQS would produce a serious financial hardship without equal or greater public benefit.
- 7. As a result of an incomplete assessment of the Site, a Notice of Public Hearing pursuant to 15A NCAC 2L .0113(e) that requires notification to any property owner within the proposed area of the variance, as well as any property owners adjacent to the Site covered by the variance, could be highly problematic. "Site", as defined under the Inactive Hazardous Sites Response Act, includes any area where a hazardous substance has been deposited, stored, disposed of, or placed, or otherwise come to be located. In other words, the "Site" includes all areas where discharges of hazardous substances have occurred and all areas of environmental media to which contamination (including groundwater) has migrated. The list of names and addresses included in Exhibit O of the application as to where the Notice would be sent cannot be considered complete until the location of groundwater contamination is known.

Based on the information provided by Rhodia for the variance, the Division cannot proceed with Rhodia's request for a variance to the 1,1-dichloroethene GWQS. The assessment of the nature and extent of the groundwater contamination problem is incomplete. Rhodia will need to conduct the needed sampling and provide the data outlined in this letter. Once that information is submitted, the Division can conduct a subsequent review of a rule variance request for your Site upon receipt of a complete application. If you have questions regarding this matter, please contact George Adams or Bruce Parris at (704) 663-1699.

Sincerely,

Dexter R. Matthews, Director Division of Waste Management

cc: Paul E. Linskey
Benne C. Hutson
Tegwyn Williams

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SUPERFUND SECTION

PAGE 02/03

McGuireWoods LLP 201 North Tryon Street P.O. Box 31247 (28231) Charlotte, NC 28202 Phone: 704.343.2000 Fax: 704.343.2300

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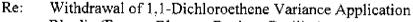
Benne C. Hutson

Direct: 704,343,2060

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March 14, 2011

Dexter R. Matthews
Director
Division of Waste Management
North Carolina Department of Environment
and Natural Resources
1646 Mail Service Center
Raleigh, North Carolina 27699-1646



Rhodia (Former Rhone – Poulenc Facility)

207 Telegraph Road

Gastonia, Gaston County, North Carolina

IHSB # NONCD 0002395-Former APS No. 8919



We represent Rhodia Inc. with regard to this matter. I am writing in response to your January 21, 2011 letter regarding the Division staff's review of Rhodia's variance request and the Division's conclusion that it cannot proceed with that request at this time.

Rhodia's variance request was not focused on the premise that it was technically or economically infeasible to comply with the current standard. Rather, the variance request was focused on the fact that under applicable North Carolina laws and regulations, the current groundwater quality standard is legally wrong. What Rhodia was attempting to do was to have the correct standard set – at least for Rhodia's site – and then plan, perform and evaluate its remediation activities based on that standard.

The Division staff's comments do not even mention, let alone address Rhodia's contention and ample support for the conclusion (a conclusion that has been shared for years by the State's toxicologist and DWQ staff responsible for the most recent triennial reviews) that the current standard is wrong. Rather, the sole and exclusive focus is on alleged site-specific lack of information to determine whether or not Rhodia can meet that standard. For the record, Rhodia continues to disagree with many of the site-specific points raised in your letter; however, we will address these issues with the Inactive Hazardous Sites Branch at a later time.

Because of the site specific nature of the variance regulation, Rhodia has never viewed the variance process as the appropriate approach to obtain a change to a standard that should apply globally to all North Carolina sites and would never have even pursued it but for the EMC's



Dexter R. Matthews March 14, 2011 Page 2

9197334811

Groundwater Committee's statement "that the 1,1-DCE issue could be dealt with using the variance process," an inapposite but convenient rationalization for their failure to act to change the standard. Based on this, Rhodia dutifully filed the application to comply with the EMC's suggestion as one step in satisfying its legal obligation to exhaust its administrative remedies.

From your letter, you have clearly confirmed Rhodia's presupposition that the variance approach is not an appropriate mechanism for seeking relief from a legally incorrect 2L standard for 1,1-DCE. Rather, that change will have to be obtained either through other mechanisms, potentially including the judicial system.

In light of your written confirmation of Rhodia's position in this matter, by this letter Rhodia formally withdraws its November 1, 2010 Application for a Variance to the 2L Groundwater Quality Standard for 1,1-Dichloroethylene.

By copy of this letter to the Recording Clerk for the Environmental Management Commission, Rhodia provides notice of the formal withdrawal of its November 1, 2010 Application for a Variance to the 2L Groundwater Quality Standard for 1,1-Dichloroethylene.

Sincerely,

BCH

Recording Clerk, Environmental Management Commission cc: Rhodia Management Team

<u>ATTACHMENT M: ALTERNATE PROPOSALS TO 2L .0202 RECEIVED DURING PUBLIC COMMENT PERIOD</u>

Proposed Rule Change 15A NCAC 02L .0202(d)	North Carolina Conservation Network	Manufacturers and Chemical Industry Council of North Carolina	Progress Energy Corporation	North Carolina League of Municipalities
(d) Except as provided in Paragraph (f),	d) Except as provided in Paragraph (f),	(d) In establishing groundwater	(d) Groundwater quality standards for	(d) Groundwater quality standards for
groundwater Groundwater quality	groundwater Groundwater quality	Groundwater quality standards for	substances in Class GA and Class GSA	substances in Class GA and Class GSA
standards for substances in Class GA and	standards for substances in Class GA and	substances in Class GA and Class GSA	groundwaters are established as the least	groundwaters are established as the least
Class GSA groundwaters are established	Class GSA groundwaters are established	groundwaters the Commission shall	of using evaluation of the reliability,	of according to the best available scientific
as the least of:	as the least of:	consider each of the following values: are	relative costs and benefits of:	guidance and include considerations such
(1) Systemic threshold concentration	(1) Systemic threshold concentration	established as the least of:	(1) Systemic threshold concentration	<u>as</u> :
calculated as follows: [Reference Dose	calculated as follows: [Reference Dose	Systemic threshold concentration	calculated as follows: [Reference Dose	(1) Systemic threshold concentration
(mg/kg/day) x 70 kg (adult body	(mg/kg/day) x 70 kg (adult body	calculated as follows: [Reference Dose	(mg/kg/day) x 70 kg (adult body	calculated as follows: [Reference Dose
weight) x Relative Source Contribution	weight) x Relative Source Contribution	(mg/kg/day) x 70 kg (adult body	weight) x Relative Source Contribution	(mg/kg/day) x 70 kg
(.10 for inorganics; .20 for organics)] /	(.10 for inorganics; .20 for organics)] /	weight) x Relative Source Contribution	(.10 for inorganics; .20 for organics)[/	(adult body weight) x Relative Source
[2 liters/day (avg. water consumption)];	[2 liters/day (avg. water consumption)];	(.10 for inorganics; .20 for organics)] /	[2 liters/day (avg.water consumption)]:	Contribution (.10 for inorganics; .20 for
(2) Concentration which corresponds to	(2) Concentration which corresponds to	[2 liters/day (avg. water consumption)];	(2) Concentration which corresponds to	organics)[/ [2 liters/day
an incremental lifetime cancer risk of	an incremental lifetime cancer risk of	2. Concentration which corresponds to	an incremental lifetime cancer risk of 1	(avg. water consumption)]:
1x10-6;	1x10-6;	an incremental lifetime cancer risk of	x 10 ⁻⁶ ;	(2) Concentration which corresponds to
(3) Taste threshold limit value;	(3) Taste threshold limit value;	1xlO-6;	(3) Taste threshold limit value'	an incremental lifetime cancer risk of 1
(4) Odor threshold limit value;	(4) Odor threshold limit value;	3. Taste threshold limit value;	(4) Odor threshold limit value;	x 10-6;
(5) Maximum contaminant level; or	(5) Maximum contaminant level; or	4. Odor threshold limit value;	(5) Maximum contaminant level; or	(3) Taste threshold limit value'
(6) National secondary drinking water	(6) National secondary drinking water	5. Maximum contaminant level; or	(6) National secondary drinking water	(4) Odor threshold limit value;
standard.	standard.	6. National secondary drinking water	standard.	(5) Maximum contaminant level; or
		standard.		(6) National secondary drinking water
				standard.
(e) The following references, in order of	(e) The following references, in order of	(e) The following references, in order of	No Change Recommended	No Change Recommended
preference, shall be used in establishing	preference, shall be used in establishing	preference, shall be considered used in		
concentrations of substances which	concentrations of substances which	establishing concentrations		
correspond to levels described in	correspond to levels described in	of substances which correspond to levels		
Paragraph (d) of this Rule.	Paragraph (d) of this Rule.	described in Paragraph (d) of this Rule:		
(1) Integrated Risk Information System	(1) Integrated Risk Information System	1. Integrated Risk Information System		
(U.S. EPA).	(U.S. EPA).	(U.S. EPA).		
(2) Health Advisories (U.S. EPA Office	(2) Health Advisories (U.S. EPA Office	2. Health Advisories (U.S. EPA Office		
of Drinking Water).	of Drinking Water).	of Drinking Water).		
(3) Other health risk assessment data	(3) Other health risk assessment data	3. Other health risk assessment data		
published by U.S. EPA.	published by U.S. EPA.	published by U.S. EPA.		
(4) Other relevant, published health risk	(4) Other relevant, published health risk	4. Other relevant, published health risk		
assessment data, and scientifically valid	assessment data, and scientifically valid	assessment data, and scientifically valid		
peer-reviewed published toxicological	peer-reviewed published toxicological	peer-reviewed published toxicological		
data.	data.	data.		

Proposed Rule Change 15A NCAC 02L .0202(d)	North Carolina Conservation Network	Manufacturers and Chemical Industry Council of North Carolina	Progress Energy Corporation	North Carolina League of Municipalities
(f) The Commission may establish	(f) In cases where the U.S. EPA has	No Change Recommended	No Change Recommended	No Change Recommended
groundwater standards less stringent than	delayed revising the maximum	_	-	_
existing maximum contaminant levels or	contaminant level despite having updated			
national secondary drinking water	reference dose and other applicable data,			
standards if it finds, after public notice and	and because of the administrative costs of			
opportunity for hearing, that	a federal rulemaking proceeding, the			
(1) more recent data published in any of	Commission may adjust the Class GA and			
the EPA health references listed in	Class GSA standards in Subsections (h)			
paragraph (e) results in a standard which	and (i), respectively, to reflect the			
is protective of public health, taste	technically correct maximum contaminant			
threshold, or odor threshold,	level. Upon U.S. EPA revisions, the EMC			
(2) such a standard will not endanger the	will ensure that the adjusted maximum			
public health and safety, including	contaminant level remains consistent with			
health and environmental effects from	the U.S. EPA's revised standard.			
exposure to groundwater contaminants,				
<u>and</u>				
(3) compliance with a standard based on				
the maximum contaminant level or				
national secondary drinking water				
standard would produce serious hardship				
without equal or greater public benefit.				

15A NCAC 02L .0202 is proposed for amendment as follows: (Option 1 and Option 2 combined with additional language)

15A NCAC 02L .0202 GROUNDWATER QUALITY STANDARDS

- (a) The groundwater quality standards for the protection of the groundwaters of the state are those specified in this Rule. They are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of the state, which may be tolerated without creating a threat to human health or which would otherwise render the groundwater unsuitable for its intended best usage.
- (b) The groundwater quality standards for contaminants specified in Paragraphs (g) (h) and (h) (i) of this Rule are as listed, except that:
 - (1) Where the standard for a substance is less than the practical quantitation limit, the detection of that substance at or above the practical quantitation limit constitutes a violation of the standard.
 - Where two or more substances exist in combination, the Director shall consider the effects of chemical interactions as determined by the Division of Public Health and may establish maximum concentrations at values less than those established in accordance with Paragraphs (c), (g), (h), or (h) (i) of this Rule. In the absence of information to the contrary, in accordance with Paragraph (d) of this Rule, the carcinogenic risks associated with carcinogens present shall be considered additive and the toxic effects associated with non-carcinogens present shall also be considered additive.
 - (3) Where naturally occurring substances exceed the established standard, the standard shall be the naturally occurring concentration as determined by the Director.
 - (4) Where the groundwater standard for a substance is greater than the Maximum Contaminant Level (MCL), the Director shall apply the MCL as the groundwater standard at any private drinking water well or public water system well that may be impacted.
- (c) Except for tracers used in concentrations which have been determined by the Division of Public Health to be protective of human health, and the use of which has been permitted by the Division, substances which are not naturally occurring and for which no standard is specified shall not be permitted in concentrations at or above the practical quantitation limit in Class GA or Class GSA groundwaters. Any person may petition the Director to establish an interim maximum allowable concentration for a substance for which a standard has not been established under this Rule. The petitioner shall submit relevant toxicological and epidemiological data, study results, and calculations necessary to establish a standard in accordance with Paragraph (d) of this Rule. Within three months after the establishment of an interim maximum allowable concentration for a substance by the Director, the Director shall initiate action to consider adoption of a standard for that substance.
- (d) Except as provided in Paragraph (f), groundwater Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of:
 - (1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)];
 - (2) Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6;
 - (3) Taste threshold limit value;
 - (4) Odor threshold limit value;
 - (5) Maximum contaminant level; or
 - (6) National secondary drinking water standard.
- (e) The following references, in order of preference, shall be used in establishing concentrations of substances which correspond to levels described in Paragraph (d) of this Rule.
 - (1) Integrated Risk Information System (U.S. EPA).
 - (2) Health Advisories (U.S. EPA Office of Drinking Water).
 - (3) Other health risk assessment data published by U.S. EPA.
 - (4) Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.
- (f) The Commission may establish groundwater standards less stringent than existing maximum contaminant levels or national secondary drinking water standards if it finds, after public notice and opportunity for hearing, that
 - (1) more recent data published in any of the EPA health references listed in paragraph (e)(1),(2), and (3) results in a standard which is protective of public health, taste threshold, or odor threshold,
 - (2) such a standard will not endanger the public health and safety, including health and environmental effects from exposure to groundwater contaminants, and

```
(3) compliance with a standard based on the maximum contaminant level or national secondary drinking water
 1
 2
               standard would produce serious hardship without equal or greater public benefit.
 3
      (f)(g) Groundwater quality standards specified in Paragraphs (g)(h) and (h)(i) of this Rule and interim maximum
 4
      allowable concentrations established pursuant to Paragraph (c) of this Rule shall be reviewed on a triennial basis.
 5
      Appropriate modifications to established standards shall be made in accordance with the procedure prescribed in
 6
      Paragraph (d) of this Rule where modifications are considered appropriate based on data published subsequent to the
 7
      previous review.
 8
      (g)(h) Class GA Standards. Where not otherwise indicated, the standard refers to the total concentration in micrograms
 9
      per liter of any constituent in a dissolved, colloidal or particulate form which is mobile in groundwater. This does not
10
      apply to sediment or other particulate matter which is preserved in a groundwater sample as a result of well construction
11
      or sampling procedures. The Class GA standards are:
12
               (1)
                        Acenaphthene: 80;
13
               (2)
                        Acenaphthylene: 200;
14
               (3)
                        Acetone: 6 mg/L;
15
               (4)
                        Acrylamide: 0.008;
16
               (5)
                        Anthracene: 2 mg/L;
17
               (6)
                        Arsenic: 10;
                        Atrazine and chlorotriazine metabolites: 3;
18
               (7)
19
                        Barium: 700;
               (8)
20
               (9)
                        Benzene: 1;
21
                        Benzo(a)anthracene (benz(a)anthracene): 0.05;
               (10)
22
                        Benzo(b)fluoranthene: 0.05;
               (11)
23
               (12)
                        Benzo(k)fluoranthene: 0.5;
24
               (13)
                        Benzoic acid: 30 mg/L;
25
                        Benzo(g,h,i,)perylene: 200;
               (14)
26
               (15)
                        Benzo(a)pyrene: 0.005;
27
               (16)
                        Bis(chloroethyl)ether: 0.03;
28
                        Bis(2-ethylhexyl) phthalate (di(2-ethylhexyl) phthalate): 3;
               (17)
29
               (18)
                        Boron: 700;
30
               (19)
                        Bromodichloromethane: 0.6;
31
               (20)
                        Bromoform (tribromomethane): 4:
                        n-Butylbenzene: 70;
32
               (21)
33
                        sec-Butylbenzene: 70;
               (22)
                        tert-Butylbenzene: 70;
34
               (23)
35
               (24)
                        Butylbenzyl phthalate: 1 mg/L;
36
               (25)
                        Cadmium: 2;
                        Caprolactam: 4 mg/L;
37
               (26)
38
               (27)
                        Carbofuran: 40;
39
                        Carbon disulfide: 700;
               (28)
40
               (29)
                        Carbon tetrachloride: 0.3;
41
               (30)
                        Chlordane: 0.1;
                        Chloride: 250 mg/L;
42
               (31)
43
                        Chlorobenzene: 50;
               (32)
44
                        Chloroethane: 3.000:
               (33)
45
               (34)
                        Chloroform (trichloromethane): 70;
46
               (35)
                        Chloromethane (methyl chloride): 3;
47
                        2-Chlorophenol: 0.4;
               (36)
48
               (37)
                        2-Chlorotoluene (o-chlorotoluene): 100;
                        Chromium: 10;
49
               (38)
                        Chrysene: 5;
50
               (39)
51
               (40)
                        Coliform organisms (total): 1 per 100 milliliters;
52
               (41)
                        Color: 15 color units;
53
               (42)
                        Copper: 1 mg/L:
54
                        Cyanide (free cyanide): 70;
               (43)
55
                        2, 4-D (2,4-dichlorophenoxy acetic acid): 70;
               (44)
```

```
1
               (45)
                        DDD: 0.1;
 2
               (46)
                        DDT: 0.1:
 3
               (47)
                        Dibenz(a,h)anthracene: 0.005;
 4
               (48)
                        Dibromochloromethane: 0.4;
 5
               (49)
                        1,2-Dibromo-3-chloropropane: 0.04;
 6
               (50)
                        Dibutyl (or di-n-butyl) phthalate: 700;
 7
               (51)
                        1,2-Dichlorobenzene (orthodichlorobenzene): 20;
 8
               (52)
                        1,3-Dichlorobenzene (metadichlorobenzene): 200;
 9
               (53)
                        1,4-Dichlorobenzene (paradichlorobenzene): 6;
10
               (54)
                        Dichlorodifluoromethane (Freon-12; Halon): 1 mg/L;
11
               (55)
                        1,1-Dichloroethane: 6;
12
               (56)
                        1,2-Dichloroethane (ethylene dichloride): 0.4;
13
               (57)
                        1,2-Dichloroethene (cis): 70;
14
               (58)
                        1,2-Dichloroethene (trans): 100;
15
               (59)
                        1,1-Dichloroethylene (vinylidene chloride): 7350;
16
               (60)
                        1,2-Dichloropropane: 0.6;
                        1,3-Dichloropropene (cis and trans isomers): 0.4;
17
               (61)
18
               (62)
                        Dieldrin: 0.002;
19
               (63)
                        Diethylphthalate: 6 mg/L;
20
               (64)
                        2,4-Dimethylphenol (m-xylenol): 100;
21
               (65)
                        Di-n-octyl phthalate: 100;
22
                        1,4-Dioxane (p-dioxane): 3;
               (66)
23
               (67)
                        Dioxin (2,3,7,8-TCDD): 0.0002 ng/L;
24
               (68)
                        1,1- Diphenyl (1,1,-biphenyl): 400;
25
               (69)
                        Dissolved solids (total): 500 mg/L;
26
               (70)
                        Disulfoton: 0.3;
27
               (71)
                        Diundecyl phthalate (Santicizer 711): 100;
28
               (72)
                        Endosulfan: 40;
29
               (73)
                        Endrin, total: (includes endrin, endrin aldehyde and endrin ketone): 2;
30
               (74)
                        Epichlorohydrin: 4;
31
               (75)
                        Ethyl acetate: 3 mg/L;
32
               (76)
                        Ethylbenzene: 600;
33
                        Ethylene dibromide (1,2-dibromoethane): 0.02;
               (77)
34
               (78)
                        Ethylene glycol: 10 mg/L;
35
               (79)
                        Fluoranthene: 300;
36
               (80)
                        Fluorene: 300;
37
               (81)
                        Fluoride: 2 mg/L;
38
               (82)
                        Foaming agents: 500;
39
               (83)
                        Formaldehyde: 600;
40
               (84)
                        Gross alpha (adjusted) particle activity (excluding radium-226 and uranium): 15 pCi/L;
41
               (85)
                        Heptachlor: 0.008;
42
               (86)
                        Heptachlor epoxide: 0.004;
43
               (87)
                        Heptane: 400;
               (88)
                        Hexachlorobenzene (perchlorobenzene): 0.02;
44
45
               (89)
                        Hexachlorobutadiene: 0.4;
46
               (90)
                        Hexachlorocyclohexane isomers (technical grade): 0.02;
47
               (91)
                        n-Hexane: 400;
48
               (92)
                        Indeno(1,2,3\text{-cd})pyrene: 0.05;
49
               (93)
                        Iron: 300;
50
               (94)
                        Isophorone: 40;
51
               (95)
                        Isopropylbenzene: 70;
52
               (96)
                        Isopropyl ether: 70;
53
               (97)
                        Lead: 15:
54
               (98)
                        Lindane (gamma hexachlorocyclohexane): 0.03;
55
               (99)
                        Manganese: 50;
```

```
1
               (100)
                        Mercury: 1;
 2
               (101)
                        Methanol: 4 mg/L;
 3
               (102)
                        Methoxychlor: 40;
                        Methylene chloride (dichloromethane): 5;
 4
               (103)
 5
               (104)
                        Methyl ethyl ketone (2-butanone): 4 mg/L;
 6
               (105)
                        2-Methylnaphthalene: 30;
 7
               (106)
                        3-Methylphenol (m-cresol): 400;
 8
               (107)
                        4-Methylphenol (p-cresol): 40;
 9
               (108)
                        Methyl tert-butyl ether (MTBE): 20;
10
               (109)
                        Naphthalene: 6;
                        Nickel: 100;
11
               (110)
12
               (111)
                        Nitrate: (as N) 10 mg/L;
13
                        Nitrite: (as N) 1 mg/L;
               (112)
                        N-nitrosodimethylamine: 0.0007;
14
               (113)
15
               (114)
                        Oxamyl: 200;
16
               (115)
                        Pentachlorophenol: 0.3;
                        Petroleum aliphatic carbon fraction class (C5 - C8): 400;
17
               (116)
18
               (117)
                        Petroleum aliphatic carbon fraction class (C9 - C18): 700;
19
               (118)
                        Petroleum aliphatic carbon fraction class (C19 - C36): 10 mg/L;
20
               (119)
                        Petroleum aromatics carbon fraction class (C9 - C22): 200;
21
                        pH: 6.5 - 8.5;
               (120)
22
                        Phenanthrene: 200;
               (121)
               (122)
23
                        Phenol: 30;
24
               (123)
                        Phorate: 1:
25
                        n-Propylbenzene: 70;
               (124)
26
               (125)
                        Pyrene: 200;
27
               (126)
                        Selenium: 20;
28
                        Silver: 20;
               (127)
29
               (128)
                        Simazine: 4;
30
               (129)
                        Styrene: 70;
31
               (130)
                        Sulfate: 250 mg/L;
32
               (131)
                        1,1,2,2-Tetrachloroethane: 0.2;
33
                        Tetrachloroethylene (perchloroethylene; PCE): 0.7;
               (132)
                        2,3,4,6-Tetrachlorophenol: 200;
34
               (133)
35
               (134)
                        Toluene: 600;
36
               (135)
                        Toxaphene: 0.03;
37
               (136)
                        2, 4, 5,-TP (Silvex): 50;
38
               (137)
                        1,2,4-Trichlorobenzene: 70;
39
               (138)
                        1,1,1-Trichloroethane: 200;
40
               (139)
                        Trichloroethylene (TCE): 3;
41
               (140)
                        Trichlorofluoromethane: 2 mg/L;
42
               (141)
                        1,2,3-Trichloropropane: 0.005;
43
                        1,2,4-Trimethylbenzene: 400;
               (142)
44
                        1,3,5-Trimethylbenzene: 400;
               (143)
45
               (144)
                        1,1,2-Trichloro-1,2,2-trifluoroethane (CFC-113): 200 mg/L;
46
               (145)
                        Vinyl chloride: 0.03;
47
               (146)
                        Xylenes (o-, m-, and p-): 500; and
48
               (147)
                        Zinc: 1 mg/L.
49
      (h)(i) Class GSA Standards. The standards for this class are the same as those for Class GA except as follows:
50
               (1)
                        chloride: allowable increase not to exceed 100 percent of the natural quality concentration; and
51
               (2)
                        total dissolved solids: 1000 mg/l.
52
      (i)(j) Class GC Waters.
53
               (1)
                        The concentrations of substances which, at the time of classification, exceed the standards applicable
54
```

to Class GA or GSA groundwaters shall not be caused to increase, nor shall the concentrations of other

1 2 3 4 5 6	(2)	substances be caused to exceed the GA or GSA standards as a result of further disposal of contaminants to or beneath the surface of the land within the boundary of the area classified GC. The concentrations of substances which, at the time of classification, exceed the standards applicable to GA or GSA groundwaters shall not be caused to migrate as a result of activities within the boundary of the GC classification, so as to violate the groundwater or surface water quality standards in adjoining waters of a different class.
/	(3)	Concentrations of specific substances, which exceed the established standard at the time of
8		classification, are listed in Section .0300 of this Subchapter.
9		
10	History Note:	Authority G.S. 143-214.1; 143B-282(a)(2);
11		Eff. June 10, 1979;
12		Amended Eff. November 1, 1994; October 1, 1993; September 1, 1992; August 1, 1989;
13		Temporary Amendment Eff. June 30, 2002;
14		Amended Eff. August 1, 2002;
15		Temporary Amendment Expired February 9, 2003;
16		Amended Eff. November 1, 2012; January 1, 2010; April 1, 2005.
17		

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15A NCAC 02L .0113 is proposed for amendment as follows: (Option 3 with additional language noticed as Option 2)

15A NCAC 02L .0113 VARIANCE

- (a) The Commission, on its own initiative or pursuant to a request under G.S. 143-215.3(e), may grant variances to the rules of this Subchapter.
- (b) Requests for variances are shall be filed by letter from the applicant to the Environmental Management Commission. The application shall be mailed to the chairman of the Commission in care of the Director, Division of Environmental Management, Post Office Box 29535, Raleigh, N.C. 27626 0535 Division of Water Quality, 1617 Mail Service Center, Raleigh, N.C. 27699-1617.
- (c) The application for variances to the rules of Section .0100 shall contain the following information:
 - (1) Applications filed by counties or municipalities must include a resolution of the County Board of Commissioners or the governing board of the municipality requesting the variance.
 - (2) A description of the past, existing or proposed activities or operations that have or would result in a discharge of contaminants to the groundwaters.
 - (3) Description of the proposed area for which a variance is requested. A detailed location map, showing the orientation of the facility, potential for groundwater contaminant migration, as well as the area covered by the variance request, with reference to at least two geographic references (numbered roads, named streams/rivers, etc.) must be included.
 - (4) Supporting information to establish that the variance will not endanger the public health and safety, including health and environmental effects from exposure to groundwater contaminants. (Location of wells and other water supply sources including details of well construction within 1/2 mile of site must be shown on a map).
 - (5) Supporting information to establish that requirements of this Rule cannot be achieved by providing the best available technology economically reasonable. This information must identify specific technology considered, and the costs of implementing the technology and the impact of the costs on the applicant.
 - (6) Supporting information to establish that compliance would produce serious financial hardship on the applicant.
 - (7) Supporting information that compliance would produce serious financial hardship without equal or greater public benefit.
 - (8) A copy of any Special Order that was issued in connection with contaminants in the proposed area and supporting information that applicant has complied with the Special Order.
 - (9) A list of the names and addresses of any property owners within the proposed area of the variance as well as any property owners adjacent to the site covered by the variance.
- (d) Site-specific Variances to Groundwater Standards in Section .0200:
 - (1) The Commission may grant a site-specific variance to a groundwater standard established in Rule .0202 that is set at the maximum contaminant level or the national secondary drinking water standard in paragraph (d) of Rule .0202 if it finds, after public notice and opportunity for hearing, that
 - (A) The existing maximum contaminant level or national secondary drinking water standard was established using outdated health effects information and more recent data published in any of the U.S. EPA health references listed in Rule .0202 (e)(1), (2) and (3) results in a standard which is protective of public health, taste threshold, or odor threshold;
 - (B) The variance will not endanger the public health and safety, including health and environmental effects from exposure to groundwater contaminants;
 - (C) The variance will not cause an exceedance of a maximum contaminant level in an impacted private drinking water well or public water system; and,
 - (D) Compliance with a standard based on the maximum contaminant level or national secondary drinking water standard would produce serious hardship without equal or greater public benefit.
 - (2) The site-specific variance application shall contain the following information:
 - (A) A description of the past, existing or proposed activities or operations that have or would result in a discharge of contaminants to the groundwaters;
 - (B) A description of the proposed area for which a variance is requested. A detailed location map, showing the orientation of the facility, potential for groundwater contaminant migration, as well as the area covered by the variance request, and the location of private drinking wells and public water system wells within ½ mile of the affected site, with at least two geographic references (numbered roads, named streams/rivers, etc.);

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- (C) A list of the names and addresses of any property owner within the proposed area of the variance as well as any property owners adjacent to the site covered by the variance; and,
- (D) Supporting information to establish that the variance will comply with the criteria in subparagraph (d)(1) of this Rule.

(d)(e) Upon receipt of the application, the Director will review it for completeness and request additional information if necessary. When the application is complete, the Director shall give public notice of the application and schedule the matter for a public hearing in accordance with G.S. 143-215.4(b) and the procedures set out in Paragraph (e) of this Rule.

(e)(f) Notice of Public Hearing:

- Notice of public hearing on any variance application shall be circulated in the geographical areas of the proposed variance by the Director at least 30 days prior to the date of the hearing:
 - (A) by publishing the notice one time in a newspaper having general circulation in said county;
 - (B) by mailing to the North Carolina Department of Environment, Health, and Natural Resources,
 Division of Environmental Health and appropriate local health agency;
 - (C) by mailing to any other federal, state or local agency upon request;
 - (D) by mailing to the local governmental unit or units having jurisdiction over the geographic area covered by the variance;
 - (E) by mailing to any property owner within the proposed area of the variance, as well as any property owners adjacent to the site covered by the variance; and
 - (F) by mailing to any person or group upon request.
- (2) The contents of public notice of any hearing shall include at least the following:
 - (A) name, address, and phone number of agency holding the public hearing;
 - (B) name and address of each applicant whose application will be considered at the meeting;
 - (C) brief summary of the variance request;
 - (D) geographic description of a proposed area for which a variance is requested;
 - (E) brief description of activities or operations which have or will result in the discharge of contaminants to the groundwaters described in the variance application;
 - (F) a brief reference to the public notice issued for each variance application;
 - (G) information regarding the time and location for the hearing;
 (H) the purpose of the hearing;
 - (I) address and phone number of premises at which interested persons may obtain further information, request a copy of each application, and inspect and copy forms and related documents; and
 - (J) a brief description of the nature of the hearing including the rules and procedures to be followed. The notice shall also state that additional information is on file with the Director and may be inspected at any time during normal working hours. Copies of the information on file will be made available upon request and payment of cost or reproduction.

(f)(g) All comments received within 30 days following the date of the public hearing shall be made part of the application file and shall be considered by the Commission prior to taking final action on the application.

(g)(h) In determining whether to grant a variance, the Commission shall consider whether the applicant has complied with any Special Order, or Special Order by Consent issued under G.S. 143-215.2.

(h)(i) If the Commission's final decision is unacceptable, the applicant may file The applicant may appeal the Commission's final decision by filing a petition for a contested case in accordance with Chapter 150B of the General Statutes. If the petition is not filed within 60 days, the Commission's decision on the variance shall be final and binding.

(i)(j) A variance shall not operate as a defense to an action at law based upon a public or private nuisance theory or any other cause of action.

 $\textit{History Note:} \qquad \textit{Authority G.S. 143-215.3(a)(1); 143-215.3(a)(3); 143-215.3(a)(4); 143-215.3(e); 143-215.4;}$

Eff. August 1, 1989;

Amended Eff. November 1, 2012; October 1, 1993.

ATTACHMENT P

Fiscal Impacts of Proposed 15A NCAC 02L Groundwater Rules

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- II. Introduction and Purpose of Rule Changes
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 - b. 15A NCAC 02L .0113 Variance
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- VI. Appendices
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 - B. Petition Summary and Background Information
 - C. 15A NCAC 02L .0202 (g)(59) Proposed Text
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Fiscal Impacts of Proposed Rules

Rule Citation: 15A NCAC 02L .0202 – Groundwater Quality Standards

15A NCAC 02L .0113 - Variance

DENR Division/

Commission: Division of Water Quality (DWQ)/ Environmental Management

Commission (EMC)

Agency Contact: Sandra Moore, Planning Section, Classifications & Standards Unit

DENR Division of Water Quality

1617 Mail Service Center Raleigh, NC 27699-1617

(919) 807-6417

sandra.moore@ncdenr.gov

Impact Summary: State government: Yes

Local government: No Private industry: Yes Substantial impact: No Federal government: No Small business: No

Authority: G.S. 143-214.1; 143B-282(a)(2)

G.S. 143-215.3(a)(1); 143-215.3(a)(3); 143-215.3(a)(4); 143-

215.3(e); 143-215.4

Necessity: The proposed rule amendments incorporate the most recent U.S.

EPA health effects data into the 1,1-Dichloroethylene (1,1-DCE) groundwater quality standard and clarify existing groundwater rule requirements. This will make the cost of regulatory compliance lower without sacrificing public health and safety. The North Carolina Environmental Management Commission (EMC) approved these proposed amendments on July 14, 2011.

I. Summary

There are three rule change options proposed:

- 1) A change in 02L.0202 (g)(59) to amend the 1,1-DCE standard from 7 ug/L to 350 ug/L;
- 2) A change in 02L .0202 (d) and (f) to allow the EMC to establish a standard less stringent that the maximum contaminant level (MCL) when:
 - a. the MCL is not based on the most recent U.S. EPA health effects data as published in U.S. EPA's Integrated Risk Information System (http://www.U.S. EPA.gov/IRIS/),
 - b. such a standard would not endanger public health and safety, and,
 - c. compliance with a standard based on the MCL would produce serious hardship without equal benefit.

- 3) A change in .0113 to:
 - a. update the Division of Water Quality mailing address, in .0113 (b),
 - b. allow the EMC to issue a state-wide variance to the 02L rules in .0113 (d), and,
 - c. clarify the existing variance requirements in .0113(i).

Following public notification in the North Carolina Register, a public hearing and a 60-day public comment period, the EMC will decide which of the above options, or combination of options, to adopt.

Option 1:

Rhodia, Inc., a global specialty chemical manufacturer that formerly operated as Rhone-Poulenc in Gastonia, North Carolina, submitted a rulemaking petition to amend the 1,1-DCE groundwater standard in 02L .0202(g)(59) from 7 ug/L to 350 ug/L based on the availability of more recent U.S. EPA health effects data. A change in this standard may result in lower compliance costs for facilities that have a release of 1,1-DCE to groundwater. However, potential compliance costs may increase for public water supply systems that use 1,1-DCE-contaminated groundwater as a source of drinking water. Parties responsible for 1,1-DCE groundwater contamination may not realize any cost savings for this change because 1,1-DCE seldom is the only pollutant that motivates cleanup activities and is often found with other chlorinated solvents. In addition, if contaminated water is currently, or could in the future be, impacting a public water supply groundwater source regulated by the NC Drinking Water Act, the company would still have to treat the water to the 7 ug/L drinking water standard. Also, there is an unresolved question regarding whether Option 1 is a legally viable solution, as some believe that the EMC might not have the authority to change the standard alone.

Option 2:

The DWQ and EMC seek to amend 02L .0202(d) and (f) on the advice of the EMC's legal counsel that rule language is needed to allow deviation from 2L .0202(d), which requires that the groundwater standard be established at the lowest of the six criteria, one of which is the federal maximum contaminant level (MCL). DWQ staff believe that this option would have the same impact as Option 1 because it will allow the 1,1-DCE standard to be set above the MCL but without legal challenge.

Option 3:

Proposed changes to 02L .0113 include the addition of a statewide variance option that would allow the EMC to consider a request for a less restrictive groundwater standard when the existing standard is based on outdated health effects data, such as the case with the existing 1,1-DCE standard. DWQ staff anticipate that the EMC will adopt Options 1 and 2, and not Option 3; however, if the EMC adopts Option 3 and not Options 1 and 2, then Rhodia, Inc. will most likely request a statewide variance to the 1,1-DCE groundwater standard because this contaminant is solely responsible for cleanup requirements and costs at the Rhodia site. 1,1-DCE has not been identified as the sole contaminant driving cleanup requirements and cost at any other sites in the state. DWQ staff assumes that the benefits of adopting Option 3 would essentially be the same as adopting Options 1 and 2. The inclusion of a statewide variance may reduce the

number of future variances submitted to DENR because a statewide variance would apply to sites across the state. Staff time spent reviewing and processing a single statewide variance would likely be less than staff time spent reviewing multiple variances for the same request. The party requesting a statewide variance will incur the cost of gathering the necessary data requirements.

Other proposed changes to the variance procedures in 02L .0113 include an update to the DWQ mailing address and clarification of the existing variance requirements that are not expected to result in any additional costs or benefits.

The approximate effective date of the proposed rules is November 1, 2012.

Based on outreach response from potentially impacted parties and information provided by state regulatory agencies, Rhodia is the only company immediately affected by the proposed rule changes. If Rhodia is the only company immediately affected by this rule change, and no additional costs are placed on drinking water suppliers, the costs of this proposed rule change will be approximately \$5,800 in FY2012-13 and \$27,000 in FY27-28 (adjusted for an assumed 2% annual inflation). Benefits, in the form of opportunity cost-savings for NCDENR and less monitoring for NCDOT in the next 30 years will have an approximate net present value of \$30,000 (using 7% discount rate). Rhodia may experience a cost savings of up to \$945,000 in the next 30 years (in net present value terms). The total 30-year net present value of the proposed rule change would be approximately \$960,000. Net present value is presented over a period of 30 years since this is the estimated time it would take Rhodia to complete cleanup at the site under existing rules using pump-and-treat remediation. The risk analysis section examines additional costs and benefits that may be incurred by additional companies and water supply systems or the need for more water remediation as a result of the rule change. Table 1 is a partial representation of total costs and benefits associated with the proposed rule changes. The full table is presented in Appendix K.

Table 1:
Partial Representation of Total Costs and Benefits Associated with Proposed Rule Changes to
15A NCAC 02L 0202 Groundwater Quality Standards with Two Percent Inflation

Fiscal Year	2011-12	2012-13	2013-14	2014-15	2015-16
Year Number	0	1	2	3	4
Costs					
Private Company Well Closure Costs	\$0	\$5,834	\$0	\$0	\$0
Total Costs	\$0	\$5,834	\$0	\$0	\$0
Benefits					
State Benefits					
NCDOT Reduced Reporting	\$0	\$3,672	\$3,537	\$3,396	\$3,247
NCDENR Opportunity Cost Savings	\$0	\$0	\$0	\$0	\$0
Private Company Benefits					
Monitoring Cost Savings	\$0	\$5,969	\$6,088	\$6,210	\$6,334
Well Closure Cost Savings	\$0	\$0	\$0	\$0	\$0
Operations and Maintenance Cost					
Savings	\$0	\$0	\$0	\$0	\$0
Total Benefits	\$0	\$9,641	\$9,626	\$9,606	\$9,582
Net Impact (benefits-costs)	\$0	\$3,807	\$9,626	\$9,606	\$9,582
Total Impact (benefits+costs)	\$0	\$15,475	\$9,626	\$9,606	\$9,582
30-year Net Present Value (@7%					
discount rate)	\$960,152				

^{*} The proposed rule would expedite the closure of the 11 wells; while the company would incur closure costs in Year 1, it would also experience an equal cost-savings (no accounting for inflation) in Year 15 from not needing to close these 11 wells at that point.

II. Introduction and Purpose of Rule Changes

Groundwater Classifications and Standards in 15A NCAC 02L .0200 are intended to "maintain and preserve the quality of the groundwaters, prevent and abate pollution and contamination of the waters of the state, protect public health, and permit management of the groundwaters for their best usage by the citizens of NC." It is the policy of the North Carolina EMC that the best usage of groundwaters of the state is as a source of drinking water. More than 50 percent of North Carolinians rely on groundwater as a source of drinking water.

By regulation, groundwater standards are established as the lowest concentration of the following six criteria contained in 15A NCAC 02L .0202(d)(1) - (6):

(1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) × 70 kg (adult body weight) × Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)];

- (2) Concentration which corresponds to an incremental lifetime cancer risk of 1×10^{-6} ;
- (3) Taste threshold limit value;
- (4) Odor threshold limit value;
- (5) Maximum contaminant level; or
- (6) National secondary drinking water standard.

The maximum contaminant level (MCL) of 7 ug/L for 1,1-DCE is the lowest concentration of the six criteria in 02L .0202(d) and was used to establish the groundwater standard. MCLs are federal drinking water standards established by the U.S. EPA Office of Water and are applicable to public water supply systems regulated under the Federal Safe Drinking Water Act.

In March 2011, McGuireWoods, on behalf of Rhodia Inc., submitted a rulemaking petition to the Division of Water Quality Director requesting amendment of the groundwater quality standard for 1,1-DCE contained in 15A NCAC 02L .0202(g)(59) from 7 ug/L to 350 ug/L. The Petition was submitted in accordance with N.C.G.S. 150B-20 and 15A NCAC 02I .501, which allows any person to petition the Director to adopt, amend or repeal an existing rule of the EMC. A copy of the Petition is included Appendix A. A summary of the Petition and background information is included in Appendix B.

1,1-Dichloroethene (DCE) is an industrial chemical not found naturally in the environment. Companies use 1,1-DCE to make plastics, such as flexible films like food wrap, flame retardant coatings, adhesives, and packaging materials. Long term or chronic exposure to 1,1-DCE by drinking 1,1-DCE-contaminated groundwater may cause liver toxicity. 1,1-DCE shows equivocal evidence of carcinogenicity by the oral route of exposure; therefore, it is not known if exposure to 1,1-DCE increases the risk of cancer in humans (http://www.U.S. EPA.gov/iris/subst/0039.htm http://water.U.S. EPA.gov/drink/contaminants/basicinformation/1-1-dichloroethylene.cfm#one).

The major source of 1,1-DCE in drinking water is discharge from industrial chemical factories.

The U.S. EPA, the federal agency that establishes MCLs, acknowledges that updated health effects data support increasing the 1,1-DCE MCL to 350 ug/L. However, U.S. EPA decided not to update the MCL for 1,1-DCE citing that any potential revision is not likely to provide a meaningful opportunity for cost-savings or health risk reduction to public water systems and their customers http://edocket.access.gpo.gov/2010/pdf/2010-6624.pdf.

Rhodia's Petition was presented at the May 2011 EMC Groundwater Committee meeting and the July 2011 EMC meeting. Information is available on the EMC Web site at http://portal.ncdenr.org/web/emc/agenda/2011/home. On July 14, 2011, the EMC approved Rhodia's petition and initiated rulemaking to amend the 1,1-DCE groundwater standard as requested. At the July meeting, the EMC granted approval to the DWQ to

initiate rulemaking to adopt proposed rule language in one or more of the three options discussed in this fiscal note:

Option 1: 02L .0202 (g) (59), Option 2: 02L .0202 (d) and (f), Option 3: 02L .0113 (b) through (i).

Option 1:

The purpose of changing the 1,1-DCE groundwater standard from 7 ug/L to 350 ug/L is to incorporate the most up-to-date health effects data. The proposed change to 02L .0202(g)(59), would have the same impact as the one anticipated for the proposed changes in .0202(d) and (f), assuming that the latter change would only lead to the relaxation of the 1,1-DCE standard to 350ug/L. To this end, only the impact from .0202(d) and (f) is discussed in this fiscal note.

Option 2:

The purpose of the proposed changes to 02L .0202(d) and (f) is three-fold: 1) to ensure that the most recent U.S. EPA health effects data are used in establishing groundwater quality standards; 2) to ensure that the standard is protective of public health and safety; and, 3) to ensure that the standard is not overly burdensome to regulated parties. If the lowest concentration of the six regulatory criteria for establishing a standard in .0202(d) is the Maximum Contaminant Level (MCL) and the MCL is not based on the most recent U.S. EPA health effects data in .0202(e), then the proposed rule will allow the MCL to be eliminated for consideration as the groundwater standard. At this time, 1,1-DCE is the only standard that is being changed, but this proposed rule change may lead to additional groundwater quality standard changes in the future.

Option 3:

The purpose of the proposed amendments to 02L .0113 is to update the DWQ mailing address, clarify the existing variance requirements and to allow the EMC to issue a statewide variance to the 02L rules when requested. The allowance of a statewide variance presents an alternate option to Options 1 and 2 that would not change the fundamental way standards are currently established in 2L .0202(d).

The three proposed amendments are located in Appendices C, D and E, respectively. The proposed changes to the rules have been highlighted in yellow. In addition, Appendix F includes a summary of the proposed amendments and the potential economic impact.

Support letters for Rhodia's Rulemaking Petition were received from Radiator Specialty Company, Indian Trail, NC and Duncklee & Dunham Environmental Consulting & Engineering, Cary, NC. Copies are located in Appendix G and H, respectfully.

III. Costs and Benefits by Rule

Each proposed rule revision is listed below with a description of the rule, the proposed changes, and the estimated economic impact expected for various public and private entities. The existing rules serve as the baseline from which economic impacts are evaluated.

The DWQ has collected information from a number of potentially affected parties including members of the regulated community, such as power utility companies, chemical manufacturers, dry-cleaning associations, local governments, state government, treated wood industries, the poultry and pork federations, furniture manufacturers and state regulatory agencies. A list of contacted parties is located in Appendix I. Parties identified during the outreach activities that are potentially affected by the proposed rules are discussed below.

a. 15A NCAC 02L .0202 - Groundwater Quality Standards (Option 1 and 2)

15A NCAC 02L .0202 sets out the criteria used to establish groundwater standards and provides a list of established groundwater standards. There are two proposed options to revise this language: an increase in the 1,1-DCE standard in .0202(g)(59) from 7 ug/L to 350 ug/L and a revision to .0202(d) and (f) to allow a groundwater standard to be established above an MCL, if that MCL was established using outdated U.S. EPA IRIS health effects data. Either of the option would lead to the same impact, at least in the foreseeable future.

Costs and Benefits Associated With Propose Changes to Rule 15A NCAC 02L .0202

These costs and benefits were estimated using the assumption that the change in the 2L groundwater standard would not alter the number of drinking water sources contaminated with 1,1-DCE. The Division of Water Quality believes that this is the most probable scenario. In the risk analysis section, we consider what would happen if more drinking water sources are contaminated as a result of the rule change.

i. Federal Government Impact

No increased or decreased expenditures were identified as a result of the proposed rule changes.

ii. State Impact

DWQ contacted state government agencies potentially affected by this proposal including the Department of Transportation, Department of Agriculture, Division of Waste Management, Division of Air Quality, Division of Water Quality, and Division of Environmental Health.

NCDENR reported that it would realize decreased cost due to reduced regulatory oversight. NCDOT reported that it would realize decreased expenditures due to reduced reporting (text discussions and mapping) requirements and the other agencies reported no anticipated direct impact.

The North Carolina Department of Transportation (NCDOT)

The NCDOT has identified and attempted to quantify the economic impacts associated with the proposed 15A NCAC 02L rule changes. The program within the NCDOT that will be principally affected by this change is the Asphalt Testing Program. The NCDOT Asphalt Testing Program performs on-site testing of asphalt for department construction activities using ASTM Method D2172-88. This method requires the use of a solvent, such as trichloroethylene, 1,1,1-tetrachloroethane, or carbon tetrachloride. Solvents stored, spilled, or disposed of on-site near operating labs resulted in releases of chlorinated solvents to the environment. 1,1-DCE is a breakdown product of chlorinated solvents and has been detected in the groundwater at Asphalt Testing Sites.

Twenty-three Asphalt Testing Program sites may potentially be impacted by a change in the groundwater standard for 1,1-DCE. Groundwater at five of the 23 sites exceeds the proposed 1,1-DCE standard of 350 ug/L. NCDOT does not anticipate a significant reduction in compliance costs because other chlorinated solvents are present in the groundwater and these would have to be cleaned up regardless of the change in the standard for 1,1-DCE. However, the reporting (text discussions and mapping of 1,1-DCE) may be reduced by a limited extent at 18 sites where the 1,1-DCE concentration is below 350 ug/L. NCDOT estimates an annual savings of approximately \$200 per site. DOT further estimates that one facility will cease testing each year. Savings in the first year would be \$3,600 and decrease by \$200 in each following year.

The NCDOT determined that no additional work efforts or cost savings would be realized as a result of the proposed revisions to .0202(d) & (f) and 02L .0113.

Division of Water Quality (DWQ)

The DWQ Aquifer Protection Section (APS) is authorized under 15A NCAC 02L and 15A NCAC 2T to issue permits that allow the discharge of waste onto land or into the subsurface under conditions outlined in the permit (non-discharge permits). If permitted facilities experience a change as a result of the rule amendment, this could potentially affect the Division's workload. Staff examined the Basinwide Information Management System (BIMS) database to estimate the number of potentially affected sites and to determine if there are any current cleanup activities on permitted sites related to the contaminant 1,1-DCE. There are no reported cleanup activities underway as a result of permitted activities. No Notices of Violation were reported for exceedances of the current standard outside the compliance boundary. Compliance boundaries at a typical DWQ permitted waste site are illustrated in Appendix J. In addition, there are 171 DWQ permitted facilities monitoring groundwater for volatile organic compounds that could include 1,1-DCE, however, there were no reports of 1,1-DCE exceeding the current standard. This information suggests that the change in standards would have no direct impacts on the division.

Division of Waste Management (DWM)

The Division of Waste Management has four sections that manage and regulate specific types of waste: The Hazardous Waste, Superfund, Solid Waste and Underground Storage Tank Sections. While 1,1-DCE is one of several constituents found in groundwater at sites regulated by DWM cleanup programs, according to DWM staff and two

independent consultants, it is seldom the only driver for the assessment and/or cleanup of contaminated groundwater. Only Rhodia, Inc., was identified by the DWM as being primarily impacted by the proposed 1,1-DCE standard change.

The Superfund Section's Inactive Hazardous Waste Sites Branch is the agency with regulatory oversight of Rhodia, Inc. Increasing the 1,1-DCE groundwater standard will most likely reduce the time it takes Rhodia to come into compliance with the groundwater standard, and reduce staff time and resources needed for oversight of the facility's cleanup responsibilities.

In Figure 1 of its Petition, Rhodia estimates that it will take 30 years to remediate 1,1-DCE to 7 ug/L using pump-and-treat technology. Rhodia also estimates that it will take 15 years to remediate 1,1-DCE to 350 ug/L using an alternate cleanup technology, in-situ chemical oxidation. For purposes of this analysis, it is assumed that there will be a fifteen-year reduction in the time it will take for Rhodia to cleanup 1,1-DCE groundwater contamination if the standard is changed from 7 ug/L to 350 ug/L.

The annual cost-savings of staff time is \$774, assuming 22 hours of staff time associated with report review and correspondence and an annual site visit for a mid-range engineer position with a total hourly compensation of \$35.18.

The estimated mileage cost-saving of a yearly site visit is \$60, assuming a maximum distance of 120 mile from the Mooresville Regional Office to the Rhodia site and a mileage rate of \$0.50 per mile for a state-owned Ford Explorer, 4X4 at the state Motor Fleet mileage rate.

http://www.ncmotorfleet.com/documents/NewRateSheetMay2010.pdf

The total cost-savings is estimated to be \$834 per year.

For Superfund sites, the 02L standard is the Applicable or Relevant and Appropriate Requirement (ARAR) for groundwater cleanup. If the proposed rule language is adopted the ARAR standard would become 350 ug/L. However, if the cleanup affects groundwater that is also a regulated drinking water source, the drinking water standard (7 ug/L) would be the ARAR. Rules and regulations, including drinking water standards, applicable to public water systems for the State of North Carolina are found in Title 15A, Subchapter 18C of the North Carolina Administrative Codes (see rule 15A NCAC 18C .1518). The party responsible for the pollution would have to clean up the groundwater to the drinking water standard. This means that no additional water treatment costs would be placed on water supply companies or local governments.

It is possible that water supply companies and local government would incur costs if they choose to use a contaminated water source after a remedial action plan is already approved. This seems highly unlikely though because these groups seek the cleanest possible source waters in an effort to contain water treatment costs.

iii. Local Government Impact

DWQ staff contacted local governments through various associations such as the NC League of Municipalities, NC Councils of Government, NC Association of County Commissioners, and state programs that regulate local government activities such as environmental cleanup and operation of publically owned wastewater treatment plants, public water supply systems and solid waste landfills. DWQ received eleven comments on the potential economic impacts of the proposed rules either directly from or on behalf of local governments. No direct costs or benefits were identified as a result of the propose rule revisions.

The proposed change to groundwater standards does not affect drinking water standards. The drinking water standard for 1,1-DCE would remain at 7 ug/L. This difference in groundwater and drinking water standards potentially may lead to future costs for publically owned and operated public water supply systems if groundwater used as a source water is contaminated above the MCL of 7 ug/L and treatment is required. DWQ staff anticipates this to be an unlikely outcome. The Division of Water Resources has identified current and future needs and resources for drinking water, including groundwater, throughout the state so most current/future drinking water sources are known (see link to plans –

http://www.ncwater.org/Water_Supply_Planning/NC_Water_Supply_Plan/). Further, there have been very few MCL violations reported for 1,1-DCE. Both state and federal drinking water program data support that 1,1-DCE is not a likely problem even if the groundwater standard is raised to 350 ug/L and the MCL remains at 7 ug/L. The Risk Analysis section contains a discussion of this potential cost.

iv. Private Industry Impact

Companies that pollute groundwater in excess of the 02L .0202 Groundwater Quality Standards may be required to take corrective action in accordance with 15A NCAC 02L .0106. A 50 fold change in the 15A NCAC 02L .0202(g)(59) groundwater standard for 1,1-DCE from 7 ug/L to 350 ug/L could reduce compliance cost at sites with known groundwater contamination above the current standard of 7 ug/L and at sites where future 1,1-DCE groundwater contamination might occur or be discovered. Private companies performing groundwater remediation may experience a reduction of compliance costs in the following ways:

- As a result of a higher standard, the groundwater plume will be smaller and the length of time to cleanup will be shorter.
- A smaller plume and higher cleanup level may allow the use a more economical cleanup technology.
- A lower number of groundwater wells may be needed to determine the boundaries of the contamination.
- Monitoring wells that meet the proposed standard may be closed and no longer monitored.

The type of cleanup technology employed to reduce contaminant levels to the groundwater standard is site-specific and will depend on a number of factors, including, but not limited to, the number and types of contaminants, contaminant properties, extent

of contamination, hydrogeologic properties (soil and rock type) and cleanup goals. These factors, including the type of remediation employed at a site, will affect the time and cost to cleanup groundwater to the standard. 1,2,3,4

One private company, Rhodia, Inc., was identified as impacted by the proposed 2L rules.

In its Rulemaking Petition, Rhodia states that it will save money if the new standard is adopted. A release of 1,1-dichloroethylene from an above ground storage tank in 1991 is the source of the site's 1,1-DCE groundwater contaminant plume. Division of Waste Management staff verified that this pollutant is the primary factor affecting assessment and cleanup costs at the Rhodia site (Appendix N).

Rhodia began operating a pump-and-treat groundwater remediation system at the site in September 1996. The primary objective of the groundwater extraction system is to hydraulically contain and control the movement of the groundwater contaminant plume to prevent further migration according to Rhodia's 2010 Annual Groundwater and Surface Water Sampling Results and 2010 Annual Groundwater Extraction System Performance Report (Appendix O). The secondary objective is to reduce the concentration and mass of dissolved volatile organic contaminants, primarily 1,1-DCE, in the groundwater.

In 1996, 1,1-DCE groundwater concentrations were greater than 100,000 ug/L in wells near the source (132,000 ug/L in MW-16A and 161,000 ug/L in MW 17-B). In 2010, the concentrations were orders of magnitude lower in the same general area (830 ug/L in MW-16A and 3,800 ug/L in monitoring well 17-B), indicating that the pump-and-treat system has been effective in significantly reducing the dissolved 1,1-DCE concentration in groundwater. However, the report also indicates that contaminant removal rates are leveling out.

While pump and treat systems are often effective in controlling the migration and reducing the size of the plume, the effectiveness is limited by contaminant tailing and rebound problems associated with this technology. Contaminant "tailing" is the progressively slow decline (leveling out) in groundwater contaminant concentration in the extracted water with pumping duration. Tailing results in longer remediation times since larger and large volumes of water have to be extracted to remove the smaller and smaller concentrations of a contaminant. Contaminant "rebounding" refers to the increase in contaminant concentration in groundwater after a period of time once treatment stops.

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¹ Cost Analyses for Selected Groundwater Cleanup Projects: Pump & Treat Systems and Permeable Reactive Barriers. USEPA OSWER EPA 542-R-00-013 February 2001 http://cluin.org/download/remed/542R00013.pdf

² Groundwater Cleanup: Overview of Operating Experience at 28 Sites. USEPA OSWER EPA 542-R-99-006 September 1999 http://www.epa.gov/tio/download/remed/ovopex.pdf

³ A Citizens Guide to Pump & Treat: http://cluin.org/download/citizens/pump_and_treat.pdf

⁴ A Citizens guide to Chemical Oxidation: http://cluin.org/download/citizens/oxidation.pdf

Tailing and rebound are associated with different physical and chemical processes, such as dissolution, diffusion and desorption that take place in the groundwater aquifer. Thus, prediction of cleanup duration cannot be determined by examination of concentration versus time data alone

(http://www.epa.gov/superfund/health/conmedia/gwdocs/pdfs/ptmethods.pdf).

For purposes of this analysis, the assumption is made that it will take Rhodia thirty years to cleanup 1,1-DCE groundwater contamination to 7 ug/L and fifteen years to cleanup to the proposed standard of 350 ug/L using pump-and-treat technology. This assumption is based on remediation time estimates provided by Rhodia in Figure 1 of its Petition.

Rhodia asserts in its Petition that eleven monitoring wells can be closed immediately and monitoring costs saved if the 1,1-DCE standard is amended to 350 ug/L. Cost-savings due to reduced monitoring for these eleven wells would likely be realized over the next fifteen years, the estimated time it would take to reduced the levels in these eleven wells to below the current groundwater standard of 7 ug/L.

There are 44 monitoring wells and 5 extraction wells that are currently being monitored at the site according to Rhodia's 2010 Groundwater Monitoring Report (Appendix O). This analysis assumes that Rhodia would close eleven wells once the proposed rule becomes effective. These are the wells where the 1,1-DCE groundwater concentration is already less than 350 ug/L. Furthermore, it is assumed that the remaining 38 wells would stay open until the entire site is cleaned up, given the uncertainty related to the speed with which Rhodia would be able to close all the wells. This assumption might lead to the overestimation of the savings portrayed in the analysis. Cost-savings for the remaining 38 wells would be realized for fifteen years beginning in year 16 and ending in year 30 of the analysis, based on the difference between a 30-year estimated remediation time if the standard remains at 7 ug/L compared to a 15-year estimated remediation time if the standard is amended 350 ug/L. Potential cost savings for Rhodia due to reduced monitoring are illustrated in Table 2 below.

Table 2: Potential Cost Savings to Rhodia Due to Reduced Monitoring				
Number of wells that can be Estimated monitoring cost				
closed saving per year				
11 \$5,852				
38	\$11,970			

Monitoring costs include the cost to sample the well (labor costs) and analyze the groundwater sample (analytical costs). In Figure 1 of its Petition, Rhodia estimated the analytical cost per sample at \$111 (\$15,000 total analytical cost/135 samples = \$111). The labor cost for well monitoring is estimated to be \$155 per well, taken from the DWM UST Program's 2010 reasonable rate document at

<u>http://portal.ncdenr.org/web/wm/ust/rrd</u>. The total monitoring cost per sample per well is calculated as follows: \$111 + \$155 = \$266 per well.

Based on Rhodia's monitoring requirements in the 2010 Groundwater Monitoring Report, 31 of the wells are monitored once a year and 18 are monitored twice a year. The number of monitoring events for the 11 wells that could be closed immediately was assumed to be two per year. Seven of the remaining 38 wells were assumed to be monitored twice a year and 31 are assumed to be monitored once a year.

The monitoring costs for the 11 wells that could be closed immediately under the proposed rule are calculated as follows: 11 wells \times \$266 per well \times 2 monitoring events per year = \$5,852 per year (not adjusted for inflation).

The 38 monitoring wells that could be closed in 15 years if the standard is amended to 350 ug/L is calculated as follows:

7 wells \times \$266 per well \times 2 monitoring events per year = \$3,724

31 wells \times \$266 \times 1 monitoring event per year = \$8,246.

Total monitoring cost for 38 wells that could be closed in 15 years = \$3,724 + \$8,246 = \$11,970 per year (not adjusted for inflation).

There are costs associated with closing monitoring wells in accordance with North Carolina regulations in Title 15A NCAC 02C .0113. The cost of a well closure is estimated to be \$520 per well as determined by averaging the estimated cost provided by DWM staff (\$584) and an independent consultant (\$455). In year 1 of the analysis, the one-time cost of properly closing the 11 wells, as required by NC regulations, is approximately \$5,720 (not adjusted for inflation). The estimated cost of closing the remaining 38 wells in year 16 of the analysis is \$19,760 (not adjusted for inflation). Note that, Rhodia will incur these well closure costs at some point in time, regardless of the standard. The difference is that if the standard is amended to 350 ug/L, then all of the wells would be closed 15 years earlier than expected and, as a result, costs would be incurred 15 years earlier. Essentially, while Rhodia would experience a cost from closing 11 wells in year 1 and 38 wells in year 16, if would incur an equal saving (not accounting for the time value of money, i.e. discounting, or for inflation) from not having to close anymore those 11 wells in year 15 and 38 wells in year 30.

In its petition, Rhodia estimated that the operation and maintenance costs of a pump-and-treat system would be \$4,800,000 over a 30-year period, which represents a yearly cost of \$160,000 (not adjusted for inflation). If the standard is amended to 350 ug/L and the site is closed in 15 years, rather than in 30 years if the standard remains at 7 ug/L, then Rhodia will benefit from a 15-year reduction in operation and maintenance costs.

Assuming 1,1-DCE groundwater concentrations will decrease to 350 ug/L in the next 15 years, the estimated cost-savings to Rhodia as a result of amending the groundwater standard from 7 ug/L to 350 ug/L due to reduced monitoring and operation and maintenance costs over a thirty-year period is estimated to be around \$930,000. The cost-savings estimate assumes that it would be cost prohibitive for Rhodia at this stage in

its clean-up to decommission its current pump-and-treat technology in favor a different technology, which might have been more cost effective if employed from the beginning. Table 3 is a partial table that represents the estimated cost savings to private industry (Rhodia). The full table is presented in Appendix L.

Table 3							
Private Industry Costs	Private Industry Costs and Benefits with the Proposed Rule Change						
W	ith Two Perd	ent Inflation	1				
Fiscal Year	2011-12	2012-13	2013-14	2014-15	2015-16		
Year Number	0	1	2	3	4		
Costs							
Well Closure Costs	\$0	\$5,834	\$0	\$0	\$0		
Total Costs	\$0	\$5,834	\$0	\$0	\$0		
Benefits							
Monitoring Cost Savings	\$0	\$5,969	\$6,088	\$6,210	\$6,334		
Well Closure Cost Savings	\$0	\$0	\$0	\$0	\$0		
Operation and Maintenance							
Costs	\$0	\$0	\$0	\$0	\$0		
Total Benefits	\$0	\$5,969	\$6,088	\$6,210	\$6,334		
Net Impact (benefits-costs)	Net Impact (benefits-costs) \$0 \$135 \$6,088 \$6,210 \$6,33						
Total Impact (benefits+costs)	\$11,803	\$6,088	\$6,210	\$6,334			
30-year Net Present Value (@7%	30-year Net Present Value (@7%						
discount rate)	\$930,522						

Rhodia submitted a list of nine facilities, including the Rhodia site, known to have groundwater contamination above the current 1,1-DCE groundwater standard of 7 ug/L, as well as the number of monitoring wells at each site with contamination above the proposed 1,1-DCE standard of 350 ug/L. According to Rhodia, if 1,1-DCE is the only constituent exceeding a groundwater quality standard and the standard is changed from 7 ug/L to 350 ug/L, monitoring of 47 groundwater wells could cease. DWQ staff contacted the agency that regulates these sites to determine if 1,1-DCE is present above the current and proposed groundwater standard of 7 ug/L and 350 ug/L, respectively, and if 1,1-DCE is the only contaminant being remediated.

Based on information provided by DWM in Appendix N, 1,1-DCE contamination at the Rhodia site was the result of a 1,1-DCE storage tank release and was the sole motivation for the remediation. The company probably will be able to reduce the number of monitoring wells and the number of years needed for remediation. While 1,1-DCE was present at many of the other eight sites, other chlorinated solvents, such as tetrachoroethylene, trichloroethylene, 1,1,1-trichloroethane, commonly found in groundwater along with 1,1-DCE, were also present above the groundwater standard and

are predicted to drive assessment and cleanup. It is unlikely that the assessment and cleanup costs for these sites will be reduced by a change in the 1,1-DCE standard.

For companies, like Rhodia, currently undertaking remediation activities, the cost to decommission an existing system and replace it with a different technology may likely be higher than any potential cost savings. For sites where groundwater contaminated with 1,1-DCE has not yet been discovered or remediation has not yet begun, the proposed standard may result in reduced assessment cost as the contaminant plume based on a standard of 350 ug/L will be less extensive than a contaminant plume based on a standard of 7 ug/L. In addition, a higher standard may give companies more flexibility in the type of remediation system used. Any future benefits resulting from changes in technology or remediation time resulting from this proposed rule change are contingent on the presence of other chemicals, selected technologies and other factors. DWQ does not attempt to estimate them in this analysis.

Public Benefits

The groundwater regulations in 15A NCAC 02L .0202(e) require the use of the following references, in order of preference, to be used in establishing groundwater standards:

- 1) U.S. EPA's Integrated Risk Information System (IRIS);
- 2) U.S. EPA Office of Drinking Water Health Advisories;
- 3) Other health risk assessment data published by U.S. EPA;
- 4) Other relevant, published health risk assessment data and scientifically valid peer-reviewed published toxicological data.

U.S. EPA's IRIS database provides high quality science-based human health assessments to support the Agency's regulatory activities. The IRIS database contains information for more than 550 chemical substances containing information on human health effects that may result from exposure to various substances in the environment.

No health-based benefits are expected as a result of changing the groundwater standard for 1,1-DCE from 7 ug/L to 350 ug/L because the proposed standard of 350 ug/L is based on the most recent U.S. EPA IRIS health effects data available at http://www.epa.gov/iris/subst/0039.htm. The current groundwater standard of 7 ug/L is based on the federal MCL, which was calculated prior to the updated toxicity data being published. According to the U.S. EPA IRIS database, the chemical is less toxic than previously thought and is no longer considered a carcinogen by the oral route.

The revised language in .0202(d) and (f) would allow the EMC to eliminate the use of the federal MCL as a criterion for establishing a standard when the MCL is not based on the most recent EPA IRIS health effects data. Therefore, any future increase in a groundwater standard as a result of changes to .0202(d) and (f) will be supported by the use of the most recent health effects data and increased adverse health effects are not expected.

15A 02L .0113-Variance (Option 3)

The variance rules in 15A 02L .0113 allow an applicant to request a variance to the 02L Groundwater Rules. Variance requests are submitted to the EMC for approval. Proposed revisions to the variance rules update the DWQ mailing address, allow the EMC to issue a statewide variance to the 02L rules and clarify the existing variance requirements. DWQ staff assumes that the benefits of this proposed option would essentially be the same as adopting Options 1 and 2.

The inclusion of a statewide variance may reduce the number of future variances submitted to DENR because a statewide variance would apply to sites across the state. Staff time spent reviewing and processing a single statewide variance would likely be less than staff time spent reviewing multiple variances for the same request. Although, given that in recent years there has been less than one variance request per year, the annualized savings might be minimal. The party requesting a statewide variance, however, will incur the cost of gathering the necessary data requirements. It is unclear what the net effect of this particular proposal would be on the costs the private sector would incur from going through the variance process, which could take as long as 2 year.

Summary of Costs and Benefits

If Rhodia is the only company immediately affected by this rule change and no additional costs are placed on drinking water suppliers, the costs of this proposed rule change will be approximately \$5,800 in year 1 and \$27,000 in year 16 (adjusted for inflation). State benefits, in the form of less monitoring for NCDOT and less oversight by DWM, have an estimated net present value of \$30,000 over 30 years. Rhodia may save money through the immediate closure of 11 wells and 15 fewer years of operation and maintenance costs. The net present value of this cost savings could be as high as \$945,000 throughout the next 30 years, in net present value terms. The total 30-year net present value of impact from the proposed rule is estimated at \$960,000. The risk analysis section examines additional costs and benefits that may be incurred by additional companies and wells or the need for more drinking water remediation as a result of the rule change. Below is a partial representation of total costs and benefits associated with the proposed rule changes. The full table is presented in Appendix K .

Table 4:						
Partial Representation of Total Costs and Benefits Associated with Proposed Rule Changes to 15A NCAC 02L .0202 Groundwater Quality Standards with Two Percent Inflation						
Fiscal Year	2011-12	2012-13	2013-14	2014-15	2015-16	
Year Number						
Costs						
Private Company Well Closure						
Costs	\$0	\$5,834	\$0	\$0	\$0	
Total Costs	\$0	\$5,834	\$0	\$0	\$0	
Benefits						
State Benefits						
DOT Reduced Monitoring	\$0	\$3,672	\$3,537	\$3,396	\$3,247	
DWM Opportunity Cost Savings	\$0	\$0	\$0	\$0	\$0	
Private Company Benefits						
Monitoring Cost Savings	\$0	\$5,969	\$6,088	\$6,210	\$6,334	
Well Closure Cost Savings	\$0	\$0	\$0	\$0	\$0	
Operations and Maintenance Cost						
Savings	\$0	\$0	\$0	\$0	\$0	
Total Benefits	\$0	\$9,641	\$9,626	\$9,606	\$9,582	
Net Impact (benefits-costs)	\$0	\$3,807	\$9,626	\$9,606	\$9,582	
Total Impact (benefits+costs)	\$0	\$15,475	\$9,626	\$9,606	\$9,582	
30-year Net Present Value(@7%						
discount rate)	\$960,152					

IV. Risk Analysis

The proposed change to the groundwater standard for 1,1-DCE from 7 ug/L to 350 ug/L is responsible for the majority of benefits and costs. The benefit amount for private companies with releases of 1,1-DCE to groundwater hinges on whether or not 1,1-DCE is the only groundwater contaminant that will be responsible for requiring environmental cleanup which includes site characterization, installation of a treatment system, operation and maintenance of the treatment system and monitoring. A second possible risk is that 1,1-DCE pollution will affect a source of drinking water. This may create additional costs for public or private water systems.

While 1,1-DCE can be found in groundwater as a result of its direct release, as in Rhodia's case, it is commonly found as a breakdown product and in conjunction with other chlorinated solvents such as 1,1,1-trichloroethane, trichloroethylene and tetrachloroethylene (perchloroethylene). As noted in the previous section, none of the other companies cited by Rhodia has remediation projects that were motivated solely by 1,1-DCE contamination. Other more toxic breakdown products, such as vinyl chloride,

are usually present as well. The chlorinated solvents and breakdown products listed are generally found in much higher concentrations and have more stringent groundwater standards than 1,1-DCE as illustrated below.

Table 5:				
Groundwater Standa	rds for Chlorinated Solvents			
Contominant	2L .0202(g) Groundwater Standard			
Contaminant in ug/L				
1,1-Dichloroethylene	7 (350 proposed)			
Tetrachloroethylene (or	0.7			
perchloroethylene)	0.7			
1,1,1-Trichloroethane	200			
Trichloroethylene	3			
Vinyl Chloride	0.03			

The presence of more toxic chlorinated solvents above their respective groundwater standard, and in much higher concentrations than 1,1-DCE, would likely trigger more complex and costly environmental cleanup efforts. If this is the case, then little or no benefits will be realized as a result of amending the groundwater standard for 1,1-DCE to 350 ug/L because it is not necessarily the pollutant of greatest concern.

The first analysis made the assumption that one company, Rhodia, would benefit from the proposed rule change and that 1,1-DCE is the chemical motivating the cleanup effort. In addition to Rhodia, there could be current or future unidentified companies that have 1,1-DCE pollution that would benefit from the proposed rule change. For sites where groundwater contaminated with 1,1-DCE has not yet been discovered or remediation has not yet begun, the proposed standard may result in reduced assessment cost, as the contaminant plume based on a standard of 350 ug/L will be less extensive than a contaminant plume based on a standard of 7 ug/L.

In addition, a higher standard may give companies more flexibility in the type of remediation system used. Any future benefits resulting from changes in technology or remediation time resulting from this proposed rule change are contingent on the presence of other chemicals, selected technologies and other factors. DWQ does not attempt to estimate them in this analysis.

While this rule change would surely impact Rhodia, it is unclear how many more companies might be affected. If the proposed rule would impact other companies aside from Rhodia, the costs and benefits estimated in this analysis could increase significantly. This analysis assumed that under the proposed rule change it would take Rhodia 15 years to clean-up the site as opposed to 30 years under the current rules. The benefits to Rhodia estimated could change if this assumption does not hold true.

Table 6. Sensitivity Analysis of Years it Would Take Rhodia to Clean-up the Site Under the					
Proposed Rule					
	Number of Years				
	5 10 15 20				
30 year NPV of Private Impact	\$1,941,170	\$1,375,676	\$930,522	\$580,100	

Impacts on Sources of Drinking Water

There are some very specific circumstances in which the standard change may affect groundwater sources that are used for drinking water and create costs for public drinking water treatment. This could happen if groundwater remediated to the new standard is used as a source of drinking water in the future or if a responsible party for the pollution cannot be identified. Each of these instances would be rare in the current environment. Usually an existing or new water company would avoid using a contaminated source of water or would only use one if they believed treatment would be cost effective (benefits greater than costs). DENR knows of no local government that had to bear the cost of additional water treatment from 1,1-DCE pollution because responsible parties are usually identified. We present this analysis to better describe potentially impacted parties and to give a rough estimate of the costs associated with 1,1-DCE contamination to a source water.

Public water systems are defined as those that provide piped drinking water to at least 15 connections or 25 or more people sixty or more days per year. They are further characterized as Community Water Systems, Non-Transient Non-community Water Systems as follows:

A "Community Water System" (CWS) means a public water system which serves at least 15 service connections used by year-round residents or regularly serves at least 25 year-round residents.

A "Non-Transient Non-Community Water System" (NTNCWS) means a public water system that regularly serves at least 25 of the same nonresident persons per day for more than six months per year. Examples of such systems are those serving the same individuals (industrial workers, school children, church members) by means of a separate system.

A "Transient Non-Community Water System" (TNCWS) means a non-community public water system that does not serve 25 of the same nonresident persons per day for more than six months per year. Examples of such systems are those, RV park, diner or convenience store where the permanent nonresident staff number less than 25, but the number of people served exceeds 25.

Any of these systems could be adversely affected if 1,1-DCE is detected in their source water above 7 ug/L; however, the MCL and surveillance monitoring requirements only apply to Community and Nontransient Non-community systems. According to the DENR

Public Water Supply Section (PWS Section), as of September 29, 2011, there are 2,081 Community and 406 Non-transient Non-community active public water systems in North Carolina where groundwater is source water. The systems are further classified below as state, local, federal or private, along with the population served.

Table 7. Classification and Number of Public Water Systems					
Ownership Type	Community	Nontransient Non-	<u>Total</u>	Population Served	
		<u>community</u>			
Federal	8	8	16	158,484	
Local	549	141	690	6,676,495	
State	3	14	17	945	
Private	1,520	243	1,763	877,798	
<u>Total</u>	<u>2,081</u>	<u>406</u>	<u>2,487</u>	<u>7,713,722</u>	

Violations of the 1,1-DCE drinking water standard are not common. The PWS Section anticipates that if the groundwater protection standard for 1,1-DCE were raised from 7 to 350 ug/L, the total number of additional system affected would be small, perhaps one facility every ten years, as would the corresponding increase in workload for staff. Additional activities and cost associated with compliance, monitoring, document review, approvals, inspections and technical assistance were determined to be *de minimis* relative to the overall workload that currently exists.

According to the PWS Section, only three active systems have been in violation of the state and federal drinking water standard for 1,1-DCE since 2001. These systems are identified in Table below.

Table 8. Public Water Supply Systems Found in Violation of the 1,1-DCE MCL				
PWS System	System Type	County	Treatment Type	Year of Last Violation
Harbor House	Private Community	Mecklenburg	Carbon Filter	2009
American Truetzschler	Private NTNC*	Mecklenburg	Carbon Filter	2005
Middlesex Water System	Local Community	Nash	In process of installing treatment system	2011

^{*}Non-Transient Non-Community Water System

According to the PWS Section, the best available treatment technology for 1,1-DCE is a granular activated carbon filter system. Cost information was requested from the three facilities that have implemented or investigated this technology (Harbor House,

American Truetzschler, Middlesex), however, no response was received after numerous requests via email and phone.

The PWS Section referenced a 1989 Calgon Carbon Corporation publication (http://www.calgoncarbon.com/documents/UseofGroundwater.pdf) that estimates a total capital expenditure of approximately \$125,000 for a complete 300 gpm (gallons per minute) treatment system (which is an average size system). The same source quotes an operation and maintenance expense of about \$42,000 per year. In the table below, information from this publication was used, after adjusting for inflation, to estimate the cost of compliance for a typical water supply system regulated under the NC Drinking Water rules and found to be in violation of the 1,1-DCE maximum contaminant level (MCL). Since this estimate was done some time ago, it likely overstates the actual cost because pollution control technology tends to fall in price over time. Compliance costs for a period of five years are illustrated below, beginning ten years out in the future which is the estimated period of time that the first violation is expected to occur. The full table is presented in Appendix M. Potential costs to come into compliance include the following:

- Installation of a granular activated carbon treatment system;
- Annual operation and maintenance (O&M) of the system;
- Quarterly monitoring.

Table 9: Estimated Compliance Cost for Public Water Supply Systems with a 1,1-DCE Violation							
Estimated Compila		djusted for Inf		vicii a 1,1 DCL v	iolation		
Fiscal Year	2021-22	2022-23	2023-24	2024-25	2025-26		
Year Number	10	10 11 12 13 14					
Costs							
Capital Expenditure	\$251,418	\$0	\$0	\$0	\$0		
Annual Operation &	Annual Operation &						
Maintenance	\$84,476	\$86,166	\$87,889	\$89,647	\$91,440		
Annual Monitoring	Monitoring \$731 \$746 \$761 \$776 \$792						
Total Costs	\$336,625	\$86,912	\$88,650	\$90,423	\$92,232		

¹Capitol expenditure is a one-time cost estimated using the inflation-adjusted estimated cost of installing a typical (accepted standard size) granular activated carbon treatment system (\$125,000) from the 1989 Calgon publication "Use of Carbon Absorption Processes in Groundwater Treatment" (http://www.calgoncarbon.com/documents/UseofGroundwater.pdf).

²The annual operation and maintenance costs were taken from the 1989 Calgon publication (\$42,000) and adjusted for inflation.

³The NC Public Water Supply Section estimates the cost of monitoring to be \$150 per sample. A minimum of one sample per quarter (\$600/year) will be required.

^{*}Costs were initially adjusted for inflation from 1989 to present using the IHS Global Insight 10-year GDP deflator forecast, and then a 2% annual inflation factor was applied to compute future values.

Assumptions:

- If the 1,1-DCE groundwater standard is 350 ug/L and the maximum contaminant level for drinking water is 7 ug/L, then every ten years one Community or Nontransient Non-community public water supply system that uses groundwater as source water will have a 1,1-DCE MCL violation and will be required to take corrective action. This assumption is based on NC and USEPA 1,1-DCE MCL violation occurrence data. The first violation will occur in 2021, ten years after the groundwater standard is changed to 350 ug/L.
- A public water supply using groundwater as source water that is in violation of the 1,1-DCE MCL will be able to meet the MCL by installing the standard size carbon filter system described in the 1989 Calgon publication.
- The carbon system will be effective in reducing a 1,1-DCE groundwater concentration of 350 ug/L to 7 ug/L or less.
- The activated carbon will be replaced no more than once a year.
- The system will be monitored quarterly to determine compliance and to ensure the carbon system is working properly.
- Annual operation and maintenance costs will begin the year the carbon filter system is installed.

Based on the information provided by the PWS Section, approximately 71 percent of the potentially impacted water systems are privately owned. Another 28 percent of the systems are owned and operated by local government. The state and federal governments each own and operate less than one percent of all facilities. Below is a breakdown of the estimated total yearly costs that would attributed to private companies, local, state, and federal governments based on ownership share. The costs to Federal and State systems are considered to be negligible.

Table 10. Breakdown of Total Yearly Costs to Public Water Supply Systems (Adjusted for Inflation)						
System Type	Ownership Percentage ¹ 2021-22 2022-23 2023-24 2024-25 2025-26					2025-26
Private	71%	\$236,113	\$60,966	\$62,185	\$63,429	\$64,697
Local	28%	28% \$93,115 \$24,043 \$24,524 \$25,014 \$25,514				
Federal	0.50% \$1,663 \$429 \$438 \$447 \$456					
State	0.50%	\$1,663	\$429	\$438	\$447	\$456

¹ Ownership percentage was determined as follows: The number of systems per ownership type was divided by the total number of systems. For example, for private systems the ownership percentage was determined by dividing the number of private systems (1,763) by the total number of systems (2,487) = 0.71 or 71%.

To determine yearly cost distribution the total annual costs were multiplied by the system type ownership percentage. For example, the 2021-22 cost distribution for private systems was determined by multiplying 0.71 (71%) by the total annual cost (\$370,696) = \$263,194.

Other Issues

There are limitations to the type of information that can be obtained to develop fiscal and economic impacts. The following are important factors to consider in estimating costs and benefits:

- Incident response databases at state agencies may not contain enough information to be useful in this analysis about the status of sites, types of substances that need to be cleaned up, and cleanup technology used. Readily available data may not show detailed information on which substances appear at what sites. Most databases do not tell us if a site is cleaning groundwater with pump-and-treat or some other technology. General information about the type of release is shown in most databases. There is little consistency between state regulatory agencies with respect to the types of information collected.
- The actual duration of a groundwater cleanup varies based on many factors. The concentration of substances, vertical and lateral extent of contamination, solubility of substances, the ability of the substance to naturally degrade or attenuate, the type of cleanup technology employed and the potential threat to groundwater and health all play a role in determining the time needed to cleanup a site. The best information available is from the Underground Storage Tank Section and shows that most pump-and-treat groundwater cleanups will take approximately 10 to 15 years, although many of these sites may never meet the 15A NCAC 02L .0202 groundwater quality standards. Because the duration of cleanup varies, the overall cost/benefit for cleanup will vary from site to site.
- Raising a standard could result in a decrease in the number of years that a pumpand-treat cleanup operation is in place where a cleanup currently is underway.
 Therefore, the change could affect the overall cost of cleanup. There is no
 standard baseline data for the cost of cleaning up specific substances. The
 assessment of contamination and the duration of cleanup are the most significant
 factor in determining costs.

V. <u>Alternative Policies</u>

The proposed rulemaking to change the 1,1-DCE groundwater standard from 7 ug/L to 350 ug/L is the result of a rulemaking petition submitted by Rhodia, Incorporated. The health effects data in the U.S. EPA IRIS database has been updated and a revised health-based groundwater standard of 350 ug/L is considered a viable option to the current standard. The proposed standard of 350 ug/L will incorporate the most recent health effects data as published in the U.S. EPA IRIS database.

One alternative considered by DWQ was to leave the 1,1-DCE groundwater standard at 7 ug/L because the federal maximum contaminant level is 7 ug/L and is a regulatory criterion used to establish groundwater standards in 15A NCAC 02L .0202(d). However, the federal maximum contaminant level is not based on the updated health effects data in U.S. EPA's IRIS database; therefore, this alternative was not considered a viable option. Additional rule language is proposed in 2L .0202(d) and (f) to ensure that the Environmental Management Commission can establish a groundwater standard using the most recent U.S. EPA IRIS health effects data.

Another alternative, recommended by the NC Division of Public Health, State Toxicologist, Ken Rudo, was to change the 1,1-DCE standard to 35 ug/L based on the updated health effects data in the U.S. EPA IRIS database and a safety factor of 10 to account for its potential carcinogenicity. While the IRIS database lists 1,1-DCE as a Class C, potential human carcinogen, U.S. EPA has determined that data are inadequate for an assessment of human carcinogenic potential for the oral route, which includes drinking water. Therefore, this alternative was not considered a viable option.

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March 14, 2011

Ms. Coleen Sullins Director NC Division of Water Quality 1617 Mail Service Center Raleigh, NC 27699-1617

Re:

Rhodia Inc. - Petition for Rule-making, 15A N.C.A.C. 2L .0202(g)(59), Amendment to Groundwater Quality Standard for 1,1-Dichloroethylene

Dear Ms. Sullins:

On behalf of Rhodia Inc. (Rhodia), we are, by this letter, filing this Petition for Rule-Making (the "Petition") pursuant to and in accordance with N.C.G.S. § 150B-20 and 15A N.C.A.C. 2I .0501, which allow any person to petition the Director of the appropriate Division of the North Carolina Department of Environment and Natural Resources to adopt, amend or repeal an existing rule by submitting a rule-making petition. Specifically, Rhodia is seeking to amend the groundwater quality standard for 1,1-dichloroethylene ("1,1-DCE") as set forth at 15A N.C.A.C. 2L .0202(g)(59). Pursuant to 15A N.C.A.C. 2I.0501(a), we are addressing this Petition to you as the Director of the North Carolina Division of Water Quality ("DWQ") since DWQ (and specifically within DWQ, the Classification and Standards Unit of the Planning Section) is the Division responsible for developing and implementing groundwater quality standards as required by applicable law and regulations, including but not limited to 15A N.C.A.C. 2L .0202(d) and (e). As the petition and supporting documents exceed 10 pages in length, we are providing 20 copies of the whole petition and its attachments as required by 15A N.C.A.C. 2I.0501(c).

In the following sections, we provide the information that corresponds to the requirements for a rule-making petition as set forth in subparagraphs to 15A N.C.A.C. 2I For ease of reference, we have set forth in bold, italicized type the required information required by each subparagraph followed by the requisite information.

(1) Provide the text of the proposed rule(s) conforming to the Codifier of Rules' requirements for publication of proposed rules in the North Carolina Register.

The current groundwater quality standard for 1,1-dichloroethylene (in micrograms per liter) in 15A N.C.A.C. 2L .0202(g)(59) is:

(59) 1,1-Dichloroethylene (vinylidene chloride): 7

Rhodia requests that the groundwater quality standard for 1,1-dichloroethylene in 15A N.C.A.C. 2L .0202(g)(59) be amended to state:

- (59) 1,1-Dichloroethylene (vinylidene chloride): 7 350
- (2) Provide the statutory authority for the agency to promulgate the rule(s).

The statutory authority for the agency to promulgate the rule is N.C.G.S. § 143-214.1 and N.C.G.S. § 143B-282(a)(2).

(3) Provide a statement of the reasons for adoption of the proposed rule(s).

U.S EPA's Determination that the Risks Posed by 1,1-DCE Have Been Overstated

Historically, the groundwater quality standard for 1,1-DCE has been $7\mu g/L$, the same as the maximum contaminant level ("MCL") under the Federal Safe Drinking Water Act.

In the July 18, 2003 Federal Register, the United States Environmental Protection Agency ("U.S. EPA") published its review of existing drinking water standards for 15 different compounds, including 1,1-DCE. Based on this review, which included a revision to the file for 1,1-DCE in the Integrated Risk Information System ("IRIS"), U.S. EPA concluded that the risks previously cited for 1,1-DCE had been greatly overstated. Although the health effects data supported the calculation of a less stringent standard for 1,1-DCE, U.S. EPA decided not to change the MCL for totally non-scientific reasons—competing workload priorities, the administrative costs associated with a required rulemaking, and the burden on states and the regulated community to implement any regulatory change that resulted. A copy of the U.S. EPA notice from the July 18, 2003 Federal Register is attached as Exhibit A.

Since then, U.S. EPA has on two occasions – October 28, 2003 and June 22, 2005 – had the opportunity to review the IRIS standard for 1,1-DCE again. In each instance, U.S. EPA reaffirmed its previous conclusion that the health risks posed by 1,1-DCE had been overstated. (See Section VII. of U.S. EPA's current IRIS file for 1,1-DCE attached as Exhibit B.)

North Carolina Regulations Governing the Establishment of Groundwater Quality Standards

Two regulations govern the establishment of groundwater quality standards in North Carolina. First, under 15A N.C.A.C. 2L.0202(d), the lesser of six different standards is to be used to set a compound's groundwater limit. Second, 15A N.C.A.C. 2L.0202(e) requires the Environmental Management Commission ("EMC") and the North Carolina Department of Environment and Natural Resources ("NCDENR") to establish groundwater standards based upon four sources of toxicity data listed "in order of preference" in the regulation. (To avoid lengthy regulatory citations, these rules will be referred to as 202(d) and 202(e) throughout the remainder of this Petition.)

Subparagraphs 202(d) and 202(e) are expressly inter-related. They must be read in tandem because the sources of toxicity data for establishing standards listed in 202(e) must "correspond to levels described in Paragraph (d) of this Rule". (Emphasis added.) From this it follows that if a basis for a groundwater quality standard listed in 202(d) does not "correspond" to an accepted reference source listed in 202(e), then the standard in 202(d) cannot be used. (Copies of these two regulations are attached as Exhibit C.)

Why 202(d) and 202(e) Require that Groundwater Quality Standard for 1,1-DCE Must Be Changed from 7mg/L to 350mg/L.

202(e) states that the "following references, in order of preference, shall be used in establishing concentrations of substances which correspond to levels described in" 202(d):

- 1. Integrated Risk Information System (U.S. EPA).
- 2. Health Advisories (U.S. EPA Office of Drinking Water).
- 3. Other health risk assessment data published by U.S. EPA.
- 4. Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.

202(d) provides that groundwater quality standards are to be established as the least of the following for a compound:

- 1. Systemic threshold concentration;
- 2. Concentration corresponding to a 1 X 10⁻⁶ incremental lifetime cancer risk;

- 3. Taste threshold limit value;
- 4. Odor threshold limit value;
- 5. Maximum contaminant level; or
- 6. National secondary drinking water standard.

1,1-DCE does not have a taste threshold limit value, odor threshold limit value or national secondary drinking water standard. In its IRIS revisions, U.S. EPA removed the oral cancer slope factor for 1,1-DCE and concluded that the compound was only a possible human carcinogen. As a result, there is no cancer risk concentration that can be used to establish a groundwater quality standard.

Although the "least" of the six identified standards in 202(d) is the MCL of 7 µg/L, it legally cannot be used in this situation. The Federal Safe Drinking Water Act requires that determination of an MCL for a compound must be based in part on the oral reference dose (RfD) for that compound. In its IRIS re-evaluation of 1,1-DCE in 2003, U.S. EPA recalculated the RfD for 1,1-DCE, changing it from 0.009 mg/kg/day to 0.050 mg/kg/day. As cited above, pursuant to 202(e), IRIS is the most preferred reference source for the establishment of groundwater quality standards. However, even though it changed the RfD, U.S. EPA did not change the MCL for 1,1-DCE. Consequently, the MCL for 1,1-DCE no longer "corresponds" to the IRIS evaluation of 1,1-DCE. As a result, under North Carolina regulations, the current MCL legally cannot be used to establish the 2L groundwater quality standard for 1,1-DCE.

The only basis for establishing a groundwater quality standard for 1,1-DCE is the systemic threshold concentration for 1,1-DCE. According to 202(d)(1), the systemic threshold concentration is calculated as follows:

[Reference dose mg/kg/day x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganic; .20 for organics)] / [2 liters/day (avg. water consumption)]

Since this formula can use U.S. EPA's recalculated RfD for 1,1-DCE, the systemic threshold concentration "corresponds" with IRIS and thus complies with the requirements of 202(e). Performing the calculation using the recalculated RfD of 0.050 and a relative source contribution of .20 (as 1,1-DCE is an organic compound), the systemic threshold concentration for 1,1-DCE is 350 μ g/L, 50 times higher than the current 2L standard.

Based on the express language of 202(d) and (e), the proposed rule amendment must be approved.

Rhodia's Prior Requests for Changing the Standard for 1,1-DCE

2005 Triennial Review

During the two most recent triennial reviews of groundwater quality standards conducted by DWQ as required by 15A NCAC 2L .0202(f), Rhodia has submitted written public comments requesting that the EMC change the groundwater quality standard for 1,1-DCE. Rhodia first submitted public comments in a letter to David Hance of the Division of Water Quality dated September 28, 2004. (A copy of this letter is attached as Exhibit D.) Based on U.S. EPA's 2003 conclusions that the risks from 1,1-DCE had been overstated and an analysis of North Carolina's regulations governing the establishment of groundwater quality standards, Rhodia concluded that the 2L standard for 1,1-DCE should be changed from 7 $\mu g/L$ to at least 322 $\mu g/L$. Similar comments were also independently submitted by Dr. Shawn L. Sager, a principal scientist with ARCADIS G&M of North Carolina, Inc., in a letter dated September 13, 2004. (A copy of this letter is included as Exhibit E.) These letters were reviewed by Dr. Luanne K. Williams, a toxicologist with the Occupational Environmental Epidemiology Branch of the North Carolina Division of Public Health. In comments enclosed with a letter dated October 1, 2004, Dr. Williams, the State's toxicologist, concluded that based upon the information provided in Rhodia's public comments, "the recommended groundwater quality standard for 1,1-dichloroethylene is 350 µg/L and not 7 µg/L." (Emphasis added.) (A copy of Dr. Williams' letter is attached as Exhibit F.)

The hearing officers did not dispute the arguments presented by Rhodia and Dr. Sager in their public comments or the conclusion of Dr. Williams, the State's toxicologist. Nevertheless, the 2004 hearing officers' report concluded that they were legally prevented from changing the standard for 1,1-DCE. The hearing officers stated that in order for the 1,1-DCE standard to be changed, the groundwater quality regulations in 15 N.C.A.C. Subchapter 2L would have to be amended. The hearing officers, however, did not provide any specifics as to what amendments were required.

In response to the hearing officers' conclusions, Rhodia took its arguments directly to the EMC. In a January 18, 2005 letter to Dr. David H. Moreau, then chairman of the EMC, Rhodia explained why the EMC was required to change the standard for 1,1-DCE. (A copy of that letter is attached as Exhibit G.) However, at its February 2005 meeting, the EMC refused to do so.

2009 Triennial Review

In a letter dated May 28, 2009, Rhodia (this time joined by Radiator Specialty Company and Ashland Inc.) again requested that the groundwater quality standard for 1,1-DCE be changed from 7 μ g/L to 350 μ g/L as part of the 2009 triennial review. (A copy of this letter is attached as Exhibit H.) In addition to the reasons set forth in 2004, Rhodia noted that in a December 31, 2003 rulemaking, U.S. EPA had relied on the IRIS

revisions to change the recommended surface water quality standard for 1,1-DCE from 0.057 μ g/L to 330 μ g/L, an increase of nearly 6,000 times the previous standard.

The hearing officers' report, dated September 10, 2009, was nearly identical to the report of the hearing officers in 2004. Once again, the hearing officers did not disagree with Rhodia's contentions or conclusions. However, once again, despite that lack of disagreement, the hearing officers did not change the standard for 1,1-DCE. Rather, the hearing officers' report referenced the work of a Groundwater Stakeholder Work Group that had met between July 14 and December 1, 2005 which "discussed needed changes to the groundwater rules to update groundwater standards so that the most up-to-date toxicity information was being incorporated." However, apparently due to the fact that no such regulatory changes had been made, "the EMC Groundwater Committee decided not to propose any changes to the groundwater rules stating that the 1,1-DCE issue could be dealt with using the variance process." (A copy of the portion of the Hearing Officers' report relating to Rhodia's comments is attached as Exhibit I.) At its September 2009 meeting, the EMC did not change the standard for 1,1-DCE.

Rhodia's Variance Request

Pursuant to the Groundwater Committee's conclusion "that the 1,1-DCE issue could be dealt with through the variance process" and in fulfillment of its legal obligation to exhaust its administrative remedies, on November 1, 2010, Rhodia filed with the EMC an Application for a Variance to the 2L Groundwater Quality Standard for 1,1-Dichloroethylene (1,1-DCE).

N.C.G.S. §143-215.3(c) provides the statutory authority to grant variances, which authority is implemented by regulation at 15A NCAC 2L .0113. Pursuant to these authorities, an applicant for a variance must show (1) that the variance will not endanger human health or safety and (2) that compliance with the existing standard "cannot be achieved by application of the best available technology found to be economically reasonable . . . and would produce serious hardship without equal or greater benefits to the public." (N.C.G.S. §143-215.3(e)(2)) Rhodia pointed out that the first factor was satisfied for the reasons previously stated in this Petition, namely that based on U.S. EPA's revisions to its IRIS evaluation as applied to North Carolina regulations governing the establishment of groundwater quality standards, a standard of 350mg/L for 1,1-DCE in groundwater would be protective of public health and the environment, a conclusion with which the State's toxicologist and NCDENR staff responsible for the triennial review agreed.

As to the second factor, Rhodia contended that it was not applicable to or determinative of the variance request as Rhodia was not basing the request on the grounds that compliance with the 1,1-DCE standard was technically infeasible or would cause undue economic hardship but rather on the fact that the current standard for 1,1-DCE was wrong.

Although the establishment of groundwater quality standards based on North Carolina regulations is the responsibility of the Classification and Standards Unit of DWQ's Planning Section, the EMC forwarded the variance application to the Division of Waste Management ("DWM") as DWM has jurisdiction over Rhodia's cleanup activities at a site in Gastonia, North Carolina. DWM has advised Rhodia it could not proceed with the variance request for a number of site specific reasons, none of which are in any way relevant to Rhodia's central allegation that the standard is simply wrong as a matter of law and should not apply either at Rhodia's site or throughout the State of North Carolina. For clarity, Rhodia's Petition and rationale for this change in the groundwater standard is not based upon and has nothing whatsoever to do with any site specific issues or conditions at the Gastonia site that has been undergoing remediation for 13 years and running.

Over the past six plus years, Rhodia has submitted public comments as part of the triennial reviews and, at the direction of EMC, a variance application. None of these actions has resulted in the legally required change to the groundwater quality standard for 1,1-DCE, either statewide or as applied to Rhodia. Rhodia's only remaining administrative remedy is to submit this Petition for Rulemaking.

(4) Provide a statement of the effect on existing rules or orders.

Other than the rule to be amended, Rhodia is not aware of any effect on existing rules or orders.

(5) Provide copies of any documents and data supporting the proposed rule(s).

We have attached copies of documents supporting the proposed rule as Exhibits A through J.

(6) Provide a statement of the effect of the proposed rule(s) on existing practices in the area involved, including cost factors for persons affected by the proposed rule(s).

This rule would apply to sites with known groundwater contamination where the level of 1,1-DCE exceeds the current standard of $7 \mu g/L$ and sites where in the future groundwater contaminated by 1,1-DCE might occur or be discovered. The effects on these sites would be as follows:

¹ Since 1997, a groundwater remediation system has been operating to remediate groundwater contamination at a specialty chemical manufacturing facility located at 207 Telegraph Drive in Gastonia, North Carolina (the "Site") formerly owned and operated by Rhone-Poulenc Inc. Since January 1, 1998, Rhodia has been contractually obligated to perform this clean up. The primary contaminant of concern is 1,1-DCE.

The Area of 1,1-DCE Contamination Above State Standards Will Either Decrease or Cease to Exist Altogether.

The highest concentrations of contamination always exist at what is known as the source area, such as where a spill occurred or a tank leaked. As contamination migrates out from the source area in the groundwater, the concentration decreases as the contaminants are dispersed in the groundwater or as they start to break down due to the effect of natural conditions.

Under North Carolina law and regulations, the area where groundwater is considered contaminated is the area where the concentrations exceed the 2L standard. For a site with 1,1-DCE contamination, this currently is the area where the level of 1,1-DCE exceeds 7 μ g/L. Based on the principle described in the preceding paragraph, it follows that the point where the concentration is 7 μ g/L will be further from the source area than the point where the contamination is 350 μ g/L. As a result, if the standard is changed to 350 μ g/L, the area of contamination at all sites with 1,1-DCE in the groundwater will either be smaller or no longer exist.

Groundwater Investigation Costs Will Decrease

Under North Carolina law and regulations, a party responsible for groundwater contamination at a site must determine the vertical and horizontal extent of contamination, that is how far out the contamination extends and how deep it goes. As to 1,1-DCE, under the current standard, a party must continue to install wells until the levels are below 7 μ g/L. If the standard is changed to 350 μ g/L, the area of contamination will not be as large and it will take fewer wells and groundwater samples to determine the boundaries of the contamination. A smaller number of wells and groundwater samples will obviously decrease the cost of the overall investigation. These cost savings will apply to ongoing investigations which have not yet been completed and any investigations conducted in the future.

Groundwater Monitoring Costs Will Decrease

As part of groundwater remediation efforts, responsible parties periodically must sample permanent monitoring wells in the area of contamination to determine how the clean up is proceeding and when the 2L standards have been met and the clean up is done. Once a well is shown to meet the 2L standards consistently, monitoring of it is no longer required.

Parties remediating groundwater contaminated with 1,1-DCE currently have to sample any monitoring well where the concentration of 1,1-DCE exceeds 7 μ g/L. Whether a clean up is being done by an active remediation system (such as a pump-and-treat, air sparging or soil vapor extraction system) or by a process of monitored natural attenuation, levels of 1,1-DCE in groundwater will reach 350 μ g/L before reaching 7

 μ g/L. From this it follows that fewer monitoring wells will have to be sampled for a shorter period of time and can be taken out of service sooner if the groundwater quality standard for 1,1-DCE is 350 μ g/L instead of 7 μ g/L.

There are numerous sites in North Carolina where the concentration of 1,1-DCE exceeds the current groundwater quality standard of 7 $\mu g/L$. These include sites regulated under the Resource Conservation and Recovery Act, the Inactive Hazardous Sites Act, the Comprehensive Environmental Response, Compensation and Liability Act and the Dry-Cleaning Solvent Cleanup Act. The following table is a list of sites Rhodia's undersigned counsel is currently aware of where the level of 1,1-DCE in groundwater exceeds the current standard of 7 $\mu g/L$ and what the compliance status of each site would be if the standard was 350 $\mu g/L$.

Name and Address of Site	Number of Monitoring Wells $> 7 \mu g/L$	Number of Monitoring <u>Wells > 350 µg/L</u>
Rhodia Inc. 207 Telegraph Drive Gastonia, NC	21	11
Radiator Specialty Company 100 Radiator Road Indian Trail, NC	9	1
Suttle Avenue, LLC Intersection of Wilkinson Blvd and Suttle Ave. Charlotte, NC	1	0
Intersection of Pearson's Turnpike and Chicksaw Road Gastonia Township, NC	8	0
The Park Ministries, Inc. 2500 Independence Blvd. Charlotte, NC	1	0

Ms. Coleen Sullins March 14, 2011 Page 10	Appendix A	A13
Design Center Carolinas, LLC 118-120 West Worthington Ave. Charlotte, NC	1	0
Intersection of Highway 321 and Rankin Lake Road Gastonia, NC	3	0
Former Ashland Inc. Site 1415 South Bloodworth Street Raleigh, NC	11	1
Former Ashland Inc. Site 2802 Patterson Street Greensboro, NC		6
TOTAL NUMBER OF WELLS	66	19

Assuming that 1,1-DCE is the only constituent exceeding a groundwater quality standard in these wells, if the standard is changed from 7 μ g/L to 350 μ g/L, monitoring of 47 groundwater monitoring wells across all of these sites can likely cease and the wells can be abandoned.

More Remediation Technologies Can Potentially Be Used

There are two predominant geologic features in North Carolina that affect the remediation of contaminants like 1,1-DCE. First, the groundwater aquifer is unconfined meaning that water exists from shallow to deep depths without interruption by any confining layer. Second, the crystalline bedrock is fractured with any number of fissures and cracks running through it in sizes and directions that are not capable of precise determination. As a result, when a contaminant that is heavier than water like 1,1-DCE reaches the groundwater, it moves vertically through the water table eventually reaching the fractures in the crystalline bedrock. If a sufficient amount of a contaminant is released, what eventually collects in the bedrock is what is called a dense non-aqueous phase liquid or DNAPL which is a liquid that is not only denser than water but also does not dissolve readily in water. This geologic setting exists throughout the entirety of the Piedmont region.

There are a number of existing and emerging remedial technologies that theoretically can remediate 1,1-DCE in groundwater. Exhibit J summarizes the remedial technologies that have been considered for implementation at the Rhodia site.

Industry experience over approximately the last two decades has shown that attempts to remediate sites with relatively small DNAPL source areas (typically less than

Ms. Coleen Sullins March 14, 2011 Page 11

one acre) using the technologies described on Exhibit J, even under ideal hydrogeologic and site conditions, are typically only able to achieve about a one order of magnitude reduction in contaminant concentration (which is equal to a 90% reduction.)² For example, if the level of contamination is 1,000 μ g/L, these technologies will only be able to reduce the contamination to 100 μ g/L.

With the current standard of 7 μ g/L, if maximum level of contamination is 70 μ g/L or less, the use of these technologies is viable as the groundwater quality standard can likely be achieved and the remediation completed. On the other hand, if the

Kingston, J.L.T. 2008. "A Critical Evaluation of In-Situ Thermal Technologies." PhD Dissertation, Arizona State University, May. (Conducted a review of 182 sites where thermal technologies were used for source treatment and identified 14 sites with sufficient performance data to evaluate the impact of source treatment on groundwater concentration and mass discharge. For nine of these 14 sites, the thermal treatment resulted in 90% or less reduction in contaminants of concern concentrations in groundwater compared to pre-treatment conditions.)

Lebron, C., D. Major, and B. Kueper, 2008. "DNAPL Technology Evaluation Screening Tool (DNAPL TEST)," Version Beta1.0 2008-02-15. Developed by Geosyntec Consultants, NFESC, Queen's University, and University of Edinburgh. Developed for U.S. Department of Defense Environmental Security Technology Certification Program (ESTCP), Project ER-0424. (Reported groundwater chlorinated volatile organic compound (CVOC) concentration reductions for 72 DNAPL remediation projects. The median percent reduction in concentration achieved for all projects was 92%. Individual technologies were reported as having the following median concentration reductions due to treatment: bioremediation, 95% (n = 17); chemical oxidation, 91% (n = 40); and thermal, 96% (n = 13). Nine projects (13%) reported an increase in groundwater concentrations following DNAPL treatment. The study also reported DNAPL mass removal for 90 remediation projects. The median DNAPL mass removal achieved for all projects was 71%. Only thermal technologies using electrical resistive heating or steam were reported to have median DNAPL removal above 90%.)

McGuire, T.M., J.M. McDade, and C.J. Newell. 2006. "Performance of DNAPL Source Depletion Technologies at 59 Chlorinated Solvent-Impacted Sites," *Ground Water Monitoring and Remediation*, 26(1), 73-84. (Analyzed groundwater concentration data at 59 sites where DNAPL remediation had been performed using one of the four commonly applied technologies: enhanced bioremediation, in situ chemical oxidation, thermal treatment, and surfactant flushing. For these 59 sites, the median treatment volume was small, equivalent to 100 feet by 100 feet in area (<0.25 acres) and 10 feet thick (~3,800 cubic yards). The median distance between the treatment point (an injection well or thermal point) and the monitoring well where the data were collected was 7 feet. For these 59 sites, remediation performance was assessed by calculating the change (i.e., percent decrease or increase) in groundwater concentrations within the treatment zone from before treatment and after treatment was completed. Results indicated that the median percent reduction in parent CVOC concentrations due to treatment was 92%. The median reduction in parent CVOC concentrations by technology was: bioremediation, 95% (n = 26); chemical oxidation, 88% (n = 23); thermal, 97% (n = 6), and surfactant/cosolvent, 95% (n = 4).)

² GeoSyntec. 2004. Assessing the Feasibility of DNAPL Source Zone Remediation: Review of Case Studies.
Prepared by GeoSyntec Consultants. Prepared for Naval Facilities Engineering Services Center, Port Hueneme,
California. (Performed a technical survey of DNAPL technology performance at 175 sites. Most respondents
(vendors, users, and regulators) reported that remediation resulted in greater than 80% DNAPL mass removal. The
results from this study were obtained by asking site personnel to report the mass removal that was achieved by each
project, and was not an independent review of site data. The previous studies were based on mining monitoring data
from sites and performing an independent analysis of treatment technology performance. Despite the different
approaches, the GeoSyntec survey and the two data mining projects reached the similar conclusion that source zone
remediation resulted in about a one-order-of-magnitude or 90% reduction in initial concentration.)

maximum contaminant level is, for example, 3,500 μ g/L, the most these technologies can typically do is to reduce the level of contamination to 350 μ g/L, still far above the level needed to obtain a determination that no further clean up is required. In that situation, since the remediation will still continue for a number of years, the most cost effective and feasible option is often a system that will contain the area of contamination and gradually reduce the levels over time, like a pump-an-treat system.

However, if the groundwater quality standard is 350 μ g/L, then the use of other clean up technologies becomes viable at sites with much higher levels of contamination, as the likelihood of completing the clean up to the standard in a much shorter period of time greatly increases. This would achieve more clean up of the environment quicker than the current standard of 7 μ g/L will ever provide.

As a result, if the proposed rule amendment occurs, at all sites with 1,1-DCE contamination, a further review of alternate remedial technologies will be appropriate and may show that, because the plume is smaller and the standard is higher, some technologies have become more feasible and cost effective. Even if that review shows that other technologies are still not feasible or cost-effective, the current remediation system, whether it be pump-and-treat system or something else, will not have to be operated for as long a period of time to reach a 350 $\mu g/L$ standard as compared to a 7 $\mu g/L$ standard.

(7) Provide a statement explaining the computation of the costs factors.

At a site with 1,1-DCE groundwater contamination, two categories of costs will be incurred. The first includes all costs to assess and determine the vertical and horizontal extent of the contamination. This category will consist of costs to install and sample groundwater monitoring wells and evaluate the results of such sampling. The second category includes all costs to identify and evaluate viable technologies to remediate the contamination and then perform the clean up with the selected technology. This category will consist of consulting, engineering, construction, and operation and maintenance costs.

Figure 1 illustrates these cost factors for a hypothetical release of 1,500 gallons of 1,1-DCE in the saprolite portion of the groundwater aquifer of the Piedmont region of North Carolina with contamination extending to a depth of 60 feet below ground surface. The first box shows a comparison of the extent of contamination and the work necessary to define that extent for both a 7 μ g/L and a 350 μ g/L groundwater quality standard. The second box shows the estimated present value cost to assess and clean up to a standard of 7 μ g/L with the clean up using pump-and-treat technology and taking 30 years to complete. The third box shows the estimated present value cost to assess and clean up to a standard of 350 μ g/L with the clean up involving the injection of chemical oxidants into the contaminated groundwater and taking five years to complete.

(8) Provide a description, including the names and addresses, if known, of those most likely to be affected by the proposed rule(s).

See the sites identified in the table included in item 6 of this Petition.

(9) Provide the name(s) and address(es) of the petitioner(s).

Rhodia Inc.
To the Attention of Mike Shatynski
8 Cedar Brook Drive
Cranbury, NJ 08512

Benne C. Hutson McGuireWoods LLP P.O. Box 31247 Charlotte, NC 28231

We appreciate your consideration of this Petition. If you have any questions or need additional information, please call me.

Sincerely,

McGuireWoods LLP

Enclosures

cc: Lois Thomas, EMC Clerk

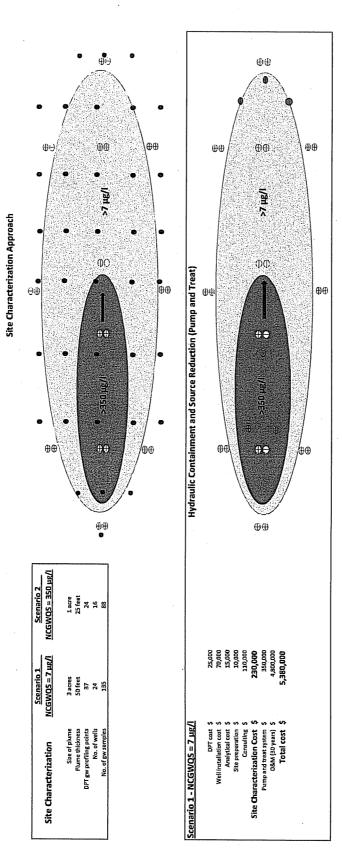
Rhodia – Gastonia Project Management Team

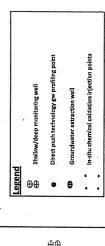
Amanda K. Short

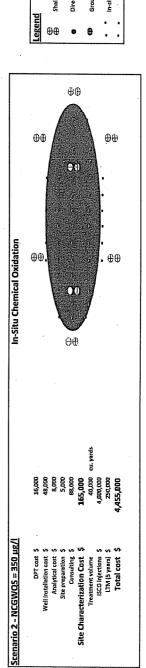
LIST OF EXHIBITS

- Figure 1: 1,1-Dichloroethylene Release Site
- Exhibit A: EPA Notice, National Primary Drinking Water Regulations:
 Announcement of Completion of EPA's Review of Existing Drinking
 Water Standards, Federal Register, Vol. 68, No. 138 (July 18, 2003)
- Exhibit B: US EPA Integrated Risk Information System, 1,1-Dichloroethylene (CASRN 75-35-4), last updated on March 16, 2010
- Exhibit C: Groundwater Quality Standards, 15A N.C.A.C. 2L .0202(d) and (e)
- Exhibit D: September 28, 2004 letter from Rhodia to David Hance, Division of Water Quality
- Exhibit E: September 13, 2004 letter from Dr. Shawn L. Sager, ARCADIS G&M of North Carolina, Inc. to David Hance
- Exhibit F: October 1, 2004 letter from Dr. Luanne K. Williams, Toxicologist, Occupational and Environmental Epidemiology Branch
- Exhibit G: January 18, 2005 letter from Rhodia to Dr. David H. Moreau, then chairman of the North Carolina Environmental Management Commission
- Exhibit H: May 28, 2009 letter from Rhodia, Radiator Specialty Company and Ashland Inc. to Sandra Moore, NCDENR, Division of Water Quality
- Exhibit I: Portion of Hearing Officer's Report of Proceedings to the North Carolina Environmental Management Commission for the Proposed Revisions to the 15A NCAC 02L .0202 Groundwater Quality Standards, September 10, 2009
- Exhibit J: Summary of Remedial Technologies, 207 Telegraph Drive, Gastonia, North Carolina

1,1-Dichloroethylene Release Site







ENVIRONMENTAL PROTECTION AGENCY

[FRL-7529-1]

RIN 2040-AD67

National Primary Drinking Water Regulations; Announcement of Completion of EPA's Review of Existing Drinking Water Standards

AGENCY: Environmental Protection Agency.

AСПОN: Notice.

SUMMARY: The Safe Drinking Water Act (SDWA) requires the United States Environmental Protection Agency (EPA) to conduct a periodic review of existing National Primary Drinking Water Regulations (NPDWRs). EPA has completed its review of 69 NPDWRs that were established prior to 1997, including 68 chemical NPDWRs and the Total Coliform Rule (TCR). The intended purpose of the review was to identify those NPDWRs for which current health risk assessments, changes in technology, and/or other factors, provide a health or technological basis to support a regulatory revision that will maintain or improve public health protection

EPA published its protocol for the review of NPDWRs and its preliminary revise/not revise decisions for the 69 NPDWRs in the April 17, 2002, edition of the Federal Register (67 FR 19030 (USEPA, 2002g)) in order to seek comment from the public. Today's action briefly describes the major comments, other new information, and EPA's current revise/not revise decisions for the 69 NPDWRs. ADDRESSES: The official public docket for this action is located at EPA West Building, Room B102, 1301 Constitution Avenue, NW., Washington, DC. FOR FURTHER INFORMATION CONTACT: Contact: Ken Rotert, (202) 564-5280,

e-mail: rotert.kenneth@epa.gov for inquiries regarding the TCR. For all other technical inquiries contact: Judy Lebowich, (202) 564-4884, e-mail: lebowich.judy@epa.gov, or Wynne Miller, (202) 564-4887, e-mail: miller.wynne@epa.gov. General information may also be obtained from the EPA Safe Drinking Water Hotline. Callers within the United States may reach the Hotline at (800) 426-4791. The Hotline is open Monday through Friday, excluding Federal holidays. from 9 a.m. to 5:30 p.m. Eastern Time. SUPPLEMENTARY INFORMATION:

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I. General Information

A. Does This Notice Apply to My Public Water System?

This action itself does not impose any requirements on anyone. Instead, it notifies interested parties of the availability of EPA's responses to comments received on EPA's Six-Year Review protocol and the Agency's current revise/not revise decisions for 69 NPDWRs.

B. How Can I Get Copies of Related Information?

1. Docket

EPA has established an official public docket for this action under Docket ID No. OW-2002-0012. The official public docket consists of the documents specifically referenced in this action, any public comments received, and other information related to this action. Although a part of the official docket, the public docket does not include Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. The official public docket is the collection of materials that is available for public viewing at the Water Docket in the EPA Docket Center, (EPA/DC) EPA West, Room B102, 1301 Constitution Ave., NW., Washington, DC. The EPA Docket Center Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566-1744, and the telephone number for the Water Docket is (202) 566-2426.

2. Electronic Access

You may access this Federal Register document electronically through the EPA Internet under the "Federal Register" listings at http:// www.epa.gov/fedrgstr/.

An electronic version of the public docket is available through EPA's electronic public docket and comment system, EPA Dockets. You may use EPA Dockets at http://www.epa.gov/edocket/ to view public comments, access the index listing of the contents of the official public docket, and access those documents in the public docket that are available electronically. Although not all docket materials may be available electronically, you may still access any of the publicly available docket. materials through the docket facility identified in section I.B.1. Once in the system, select "search," then key in the appropriate docket identification number

Abbreviations and Acronyms Used in This Action

BAT-best available technology CBI—confidential business information CCL—contaminant candidate list CFR—Code of Federal Regulations CMR—Chemical Monitoring Reform DACT—diaminochlorotriazine DEA-desethyl atrazine DEHA—di(2-ethylhexyl)adipate DEHP—di(2-ethylhexyl)phthalate DIA-desisopropyl atrazine EPA—United States Environmental Protection Agency

EPA/DC-EPA Docket Center FQPA—Food Quality Protection Act FR—Federal Register IOC—inorganic chemical IRED—interim reregistration eligibility decision IRIS—Integrated Risk Information System LCCA—Lead Contamination Control Act of 1988 LCR—Lead and Copper Rule MCL—maximum contaminant level MCLG-maximum contaminant level MDL—method detection limit mg/kg/day—milligram(s) per kilogram of body weight per day mg/L—milligram(s) per liter MYP—multi-year plan NAS—National Academy of Sciences NCOD-National Contaminant Occurrence Database NDWAC-National Drinking Water Advisory Council NPDWR-National Primary Drinking Water Regulation NRC-National Research Council NTP—National Toxicology Program NTNCWS-non-transient noncommunity water system Occurrence Methodology Document-Occurrence Estimation Methodology and Occurrence Findings Report for the Six-Year Review of Existing National Primary Drinking Water Regulations currence Summary Document-Occurrence Summary and Use Support Document for the Six-Year Review of Existing National Primary **Drinking Water Regulations** OPP—Office of Pesticide Programs OW-Office of Water PE—performance evaluation PHS—Public Health Service PQL—practical quantitation level Protocol Document—EPA Protocol for Review of Existing National Primary **Drinking Water Regulations**

RED—reregistration eligibility decision

ROS—regression on ordered statistics

RSC-relative source contribution

SAB—Science Advisory Board

SAP—Science Advisory Panel

SDWA—Safe Drinking Water Act

SOC—synthetic organic chemical TCR—Total Coliform Rule

TMDLs-total maximum daily loads

Treatment Technology Feasibility

Support Document for Chemical

Year Review of National Primary

Drinking Water Regulations

VOC—volatile organic chemical

VQP—water quality parameter

TT-treatment technique

-water supply

Treatment Feasibility Document-Water

Contaminants; In Support of EPA Six-

RID-reference dose

II. Background

A. What Is the Statutory Requirement for the Six-Year Review?

Under the Safe Drinking Water Act (SDWA), as amended in 1996, EPA must periodically review existing National Primary Drinking Water Regulations (NPDWRs) and, if appropriate, revise them. Section 1412(b)(9) of SDWA

The Administrator shall, not less often than every 6 years, review and revise, as appropriate, each national primary drinking water regulation promulgated under this title. Any revision of a national primary drinking water regulation shall be promulgated in accordance with this section, except that each revision shall maintain, or provide for greater, protection of the health of persons.

B. What Has the Agency Done To Address the Statutory Requirement?

The Agency developed a systematic process, or protocol, for the review of existing NPDWRs in accordance with the SDWA requirements and applied the protocol to the review of the NPDWRs for total coliforms and 68 inorganic and organic chemicals published prior to the SDWA 1996 Amendments (i.e., pre-1997 NPDWRs). In the April 17, 2002, Federal Register, EPA provided:

 A description of the review protocol;

 A detailed discussion of how the protocol was applied in assessing each of the 69 pre-1997 NPDWRs;

The preliminary results of each of the technical reviews, and the preliminary decision for each NPDWR; and

 A request for the public to comment on any aspect of the Agency's protocol and preliminary decisions.

Please refer to the April 17, 2002, Federal Register for the detailed discussion of EPA's revise/not revise decisions for each of the 69 NPDWRs. Today's action briefly summarizes the major public comments, other new information, and EPA's current revise/ not revise decisions for the 69 NPDWRs. Today's action only discusses in detail those decisions or rationales that were affected by public comments or other new information that has become available since April 2002.

In June 2002, EPA consulted with the Science Advisory Board (SAB) Drinking Water Committee and requested their review and comment on whether the protocol EPA developed based on the National Drinking Water Advisory Council (NDWAC) recommendations was consistently applied and appropriately documented. The SAB provided verbal feedback regarding the transparency and clarity of EPA's

decision criteria for making its revise/ not revise decisions under the current review. EPA has revised this protocol document to better explain how the decision criteria were applied and will also take the SAB comments into consideration when planning for the next review cycle.

III. EPA's Current Revise/Not Revise Decisions for the 69 Pre-1997 NPDWRs

EPA received comments from 44 commenters on its preliminary revise/ not revise decisions in the April 17, 2002, Federal Register. The Agency responded to these comments in the "Public Comment and Response Summary for the Six-Year Review of National Primary Drinking Water Regulations" (USEPA, 2003e), which is available in the Water Docket in the EPA Docket Center and at the EPA Dockets Web site http://www.epa.gov/ epadocket/. Other technical support documents 1 for the decisions discussed in today's action are also available in the Water Docket and at the EPA Dockets Web site http://www.epa.gov/ epadocket/ and the Safewater Web site http://www.epa.gov/safewoter/.

Based on the Agency's preliminary review, as well as the public comments received and other new information, EPA believes that it is appropriate to revise the Total Coliform Rule (TCR). The Agency also believes that it is not appropriate to revise the 68 chemical NPDWRs at this time. However, for the reasons discussed in sections IV.B.7, IV.B.13, and IV.B.14 of today's action, the Agency has modified the basis of its not revise findings for 1,1dichloroethylene, lead, and lindane, respectively. Table III-1 reflects the Agency's current revise/not revise decisions for the 69 NPDWRs. As indicated in Table III-1, EPA's decision not to revise an NPDWR at this time is

based on one of the following reasons: Health risk assessment is in process: As of December 31, 2002, the Agency is currently conducting, or has scheduled, a detailed review of current health effects information. Because the results of the assessment are not yet available, or were not available in time for consideration under the 1996–2002 review cycle, the Agency does not believe it is appropriate to revise the NPDWR at this time. In these cases, EPA will consider the results of the updated

¹ These include: (1) EPA's overall protocol for the review of NPDWRs (USEPA, 2003c); (2) health effects (USEPA, 2003f); (3) enalytical methods feasibility (USEPA, 2003a); (4) treatment technology (USEPA, 2003g); (5) consideration of other regulatory revisions (USEPA, 2003b); (6) occurrence and exposure (USEPA, 2003d; USEPA, 2002f); (7) and economic considerations (USEPA, 2002c).

health risk assessment during the 2002-2008 review cycle. If the results of the health risk assessment indicate a compelling reason to reconsider the maximum contaminant level goal (MCLG), EPA may decide to accelerate the review schedule for that contaminant's NPDWR.

- NPDWR remains appropriate after data/information review: The outcome of the review indicates that the current regulatory requirements remain appropriate, and therefore, no regulatory revisions are warranted. Any new information available to the Agency
- either supports the current regulatory requirements or does not justify a revision.
- New information, but no revision appropriate at this time because:
- —Low priority: In EPA's judgment, any resulting revisions to the NPDWR would not provide a meaningful opportunity for health risk reduction or result in meaningful cost-savings to public water systems and their customers. These revisions are a low priority activity for the Agency and, thus, are not appropriate for revision
- at this time because of one or more of the following considerations: competing workload priorities; the administrative costs associated with rulemaking; and the burden on States and the regulated community to implement any regulatory change that resulted.
- —Information gaps: Although results of the review support consideration of a possible revision, the available data are insufficient to support a definitive regulatory decision at this time.

BILLING CODE 6560-50-P

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IV: Summary of Major Comments and New Information and the Agency's Response

This section summarizes the major public comments, including the Agency's response, and other new information, and explains any modifications to EPA's preliminary revise/not revise decisions. For a more detailed summary of the comments and the Agency's response, please refer to the document: "Public Comment and Response Summary for the Six-Year Review of National Primary Drinking Water Regulation" (USEPA, 2003e).

A. What Did Commenters Say Regarding the Reasonableness and Appropriateness of EPA's Six-Year Review Approach?

1. Overall Approach and Decision

a. Adequacy of the Review. Commenters generally agreed that EPA had identified the appropriate key elements of the review. However, some commenters stated that the Agency could have done more in some areas (e.g., implementation) and a few commenters expressed the opinion that the Agency's review process contains weaknesses, or was not applied appropriately, because it did not identify any chemical NPDWRs for

vision at this time.

PA Response: The Agency believes mat its basic review protocol and decision rationale are reasonable and appropriate. Even though EPA's application of the protocol did not identify any chemical NPDWRs for revision at this time, that is not a reason to reject or modify the protocol. The review did result in the initiation of health risk assessments for three contaminants and efforts to address data gaps/research needs for several other contaminants. Health risk assessments are underway for approximately half of the chemical contaminants addressed in today's action. The Agency expects most of these assessments to be completed within the next few years. When completed, these assessments will support further analysis that may result in different revise/not revise decisions as part of the ongoing Six-Year Review

b. Criteria far Deciding that an MCLG/ MCL Revision is Appropriate and Definition of "Significant" and "Negligible." While some commenters agreed, others disagreed with the Agency's consideration of estimated changes in occurrence levels and available economic information

henever a health or technological basis to revise a standard. For example,

some commenters felt that EPA should revise the MCLG and, as appropriate, the maximum contaminant level (MCL), whenever a health basis exists, regardless of other considerations. A few commenters criticized the Agency for not defining what it considers "significant" and "negligible" gains in public health protection and/or costsavings in terms of regulatory revision.

EPĂ Response: Section 1412(b)(9) of SDWA, as amended in 1996, provides the Administrator with broad discretion to determine when a revision to an NPDWR is appropriate. As a part of this determination, the Agency believes it is reasonable to consider whether a potential revision is likely to provide a meaningful opportunity for health risk reduction. This criterion is consistent with the statutory provisions governing the regulatory determination process under section 1412(b)(1)(A) for contaminants not currently regulated. EPA also believes it is reasonable to consider the extent of potential costsavings for public water systems and their customers when determining whether revisions that potentially would result in a relaxed standard (i.e., where a health basis exists for a less stringent standard) or streamlined implementation are appropriate. These considerations allow the Administrator to better prioritize efforts that are most likely to result in a meaningful opportunity for health risk reduction or cost-savings to public water systems and their customers. Revisions that do not satisfy at least one of these criteria are a low priority activity for the Agency, and thus are not appropriate at this time because of one or more of the following considerations:

Competing workload priorities; The administrative costs associated with rulemaking; and

 The burden on States and the regulated community to implement any regulatory change that resulted.

EPA believes that the determination of whether the impact of a potential revision is "significant" or "negligible" is a matter of judgment that depends on a number of variables, not all of which are amenable to precise definition. However, EPA recognizes that the use of "negligible/significant" terminology may imply more precision and quantitation in the determination than is possible. The Agency also understands that the use of the term "negligible" may imply to some that the Agency is belittling small gains in health risk reduction. This is not the Agency's intent. Accordingly, in today's action, the Agency has explained its rationale more clearly in terms of the criteria noted in the previous paragraph.

c. Authority to Relax an Existing Standard and Deregulation of Low/Non-Occurring Contaminants. Some commenters argued that the Agency should never consider relaxing a standard because doing so, by definition, would lessen the level of public health protection. Other commenters encouraged the Agency to actively consider deregulating contaminants that have low occurrence or do not appear to be occurring in finished water or, at a minimum, to further reduce the frequency of monitoring for these contaminants.

EPA Response: EPA disagrees with those commenters who oppose relaxing a standard for any reason. The legislative history of the SDWA Amendments of 1996 makes clear that Congress envisioned the possibility that a relaxed standard might be appropriate under circumstances that would not result in a lessening of the level of public health protection. In its discussion of potential revisions to an existing drinking water standard, Senate Report Number 104-169 (available electronically at http://thomas.loc.gov/) states:

Amendments made by the bill require that any future standard issued for a contaminant already regulated must maintain or provide for greater protection of the health of persons. Generally, this will preclude the promulgation of a revised standard for a contaminant that is less stringent than the standard already in place. However, there are circumstances under which a standard may be relaxed. The maximum contaminant level goal for a contaminant is set at a level at which there is no adverse effect on the health of persons with an adequate margin of safety. New scientific information may cause the MCLG to be revised and in some cases these revisions may be to less stringent levels. This may lead to a revision of the maximum contaminant level since it need be no more stringent than the MCLG. New information may also allow for a smaller margin of safety because it narrows the range of uncertainty for estimates of health risks. Finally, some substances which have been regulated as carcinogens for ingestion in drinking water may be reclassified (as asbestos has been in the most recent revision) or assigned a threshold for the effect based on new scientific information. In each of these cases, EPA may issue a revised standard for a contaminant that is less stringent than the one it replaces.

(S. Rep. 104-169, 104th Cong., 1st Sess. (1995) at 38)

However, because section 1412(b)(9) of SDWA requires that any revision to an existing NPDWR maintain or improve the level of public health protection, EPA believes that a clear, technically-based demonstration

regarding the absence of potential risk is necessary to deregulate a contaminant. EPA does not believe it is appropriate to deregulate any currently regulated contaminant at this time because the Agency is not able to make a determination, pursuant to section 1412(b)(9) of SDWA, that there would be no lessening of public health protection if the contaminant were deregulated. The Agency disagrees that evaluation of finished water data is sufficient to consider deregulation of low or non-occurring contaminants. The apparent low or non-occurrence of these contaminants in finished water may be the result of effective treatment processes in place rather than the lack of occurrence in source water.

EPA believes that the existing waiver provisions in the SDWA regulations give States sufficient flexibility to reduce or potentially eliminate monitoring of a chemical contaminant, where appropriate. States that have primacy for the drinking water regulations are responsible for their waiver programs and can grant waivers if a particular pesticide or herbicide has not been previously used, manufactured, stored, transported, or disposed in the area, a system's source water is not susceptible to contamination from the chemical, or the State has determined the system is not ulnerable. The State can grant waivers

individual contaminants, a group of .itaminants, or issue an area-wide waiver (see 40 CFR 141.23 (b) and (c), and 141.24 (f) and (h)). In addition, States can adopt alternative monitoring strategies as long as the approach is as stringent as the Federal requirements (USEPA, 1997b).

2. Health Effects Technical Review

a. Contaminants Undergoing Health Risk Assessments. A few commenters raised issues with respect to the 36 chemical contaminants for which health risk assessments were underway when EPA published its preliminary revise/not revise decisions in the April 17, 2002, Federal Register. In particular, these commenters wanted to know the process that EPA plans to follow to review each NPDWR once the risk assessment is completed, including when that review would occur and when an accelerated review would be appropriate.

EPA Response: Between April and August 2002, the Agency completed health risk assessments for 2 of the 36 contaminants: 1,1-dichloroethylene and lindane. The results of those assessments and the impact on the ency's revise/not revise decisions are consed in sections IV.B.7 and

IV.B.14, respectively, of today's action. NPDWRs for the remaining contaminants for which health risk assessments are in process will be reviewed as a part of the 2002–2008 review cycle. However, if in the Agency's judgment, a compelling reason exists to revisit the "not revise" decision sooner, EPA may accelerate the review cycle for that NPDWR. In reviewing these regulations, EPA expects to apply an approach consistent with the protocol used for the current review. That is, the Agency will consider the same key elements and apply the same basic decision tree for making a revise/not revise decision. The key elements of the review include health effects technical review, technology review, other regulatory revisions review, and, if appropriate, occurrence/exposure analyses and consideration of available economic information (see 67 FR 19030 at 19038,

April 17, 2002 (USEPA, 2002g)).

b. Other Issues Related to the Health Effects Technical Review. One commenter stated that the Agency risk assessments underestimate risk because absorption of chemicals through the skin, lung, and nose is not "adequately" taken into account. Another commenter encouraged the Agency to evaluate the literature for potential reproductive and developmental effects for chemicals with zero MCLGs since risk management strategies, such as monitoring frequency or treatment requirements, may be affected by such information.

EPA Response: EPA disagrees that the Agency underestimates risk when deriving MCLGs. The Agency takes multiple routes of exposure into account by including a relative source contribution (RSC) in its calculation of an MCLG value. The RSC compares exposure from air, food, and drinking water and uses the data in allocating a portion of the total exposure to drinking water. When exposure data for the chemical are not available, EPA assumes that the RSC from drinking water is 20 percent of the total exposure. This allows 80 percent of the total exposure to come from sources other than drinking water, such as exposure from food, inhalation, or dermal contact.

EPA recognizes the possibility that some chemicals with zero MCLGs may also be of reproductive and/or developmental concern. EPA is investigating these endpoints and their potential impact on monitoring frequency or treatment requirements. However, the Agency does not believe the analysis can be completed during the current review cycle without significantly delaying the current revise/

not revise decisions. To the extent possible, EPA will consider the results of this analysis and any additional information during subsequent Six-Year Reviews.

Analytical Methods Feasibility Technical Review

Commenters generally supported the Agency's approach of using Performance Evaluation (PE) Water Supply (WS) data and the 10 times method detection limit (MDL) multiplier to evaluate possible changes in analytical feasibility for several of the contaminants under this Six-Year Review. A few commenters agreed that the WS data are a valuable source of information for evaluating interlaboratory performance and for developing practical quantitation levels (PQLs). However, the same commenters questioned whether the approach of using PE WS data will be possible for future reviews since the Agency's laboratory certification program that once collected this information has been externalized to private providers. These commenters questioned whether the externalized or privatized data would be sufficient for the determination and/or re-evaluation of PQLs. In addition, at least one commenter suggested that it may be appropriate (in the next Six-Year Review) to re-evaluate the policy of basing the PQL on only EPA Regional and State laboratory results, and recommended that the Agency include commercial and large utility laboratory results. According to the commenter, these laboratories (commercial and large utility) have demonstrated "significant innovation in method development and improved quantitation."

EPA Response: EPA agrees that the WS studies have been a valuable source of information for determining PQLs. At this time, the Agency has not determined whether the privatized data will be sufficient for the purposes mentioned by the commenter. In addition, the Agency has not yet determined how best to gather data to determine and/or reassess PQLs for future reviews. The Agency is in the process of evaluating acceptable options. The policy for determining the most appropriate methodology for calculating PQLs for drinking water contaminants is outside the scope of the Six-Year Review.

4. Review of Treatment Technologies and Related Issues

Commenters suggested that, while EPA's review of existing NPDWRs was generally consistent with the NDWAC recommendations to EPA (NDWAC, 2000), the Agency's review of treatment

technologies which support the regulations should be expanded. Specifically, commenters recommended that EPA review all treatment technique (TT) requirements and allow for changing or expanding these TT requirements where new information warrants such a change.

EPA Response: EPA continues to believe its approach to reviewing TT requirements is appropriate. The "EPA Protocol for the Review of Existing NPDWRs" (Protocol Document) discusses when it is appropriate for the Agency to consider revisions to TT-type regulations (see sections II.C and III.B of the Protocol documents) (USEPA, 2002d; USEPA, 2003c). The Agency discussed the review of the four chemical treatment technique NPDWRs (i.e., acrylamide, copper, epichlorohydrin, and lead) in both the draft and final "Water Treatment Technology Feasibility Support Document for Chemical Contaminants; In Support of EPA Six-Year Review of National Primary Drinking Water Regulations" (Treatment Feasibility Documents) (USEPA, 2002b; USEPA, 2003g). The Agency has no specific information that provides a basis for revisions to TT requirements at this time. However, EPA believes that research data in a number of treatmentrelated areas may be useful in future reviews of NPDWRs. The Agency is ommitted to working with stakeholders to identify and prioritize treatmentrelated research needs, and to work with EPA's research partners to address the highest priority needs.

5. Review of Implementation-Related Issues

While several commenters felt overall that EPA's Six-Year Review protocol was reasonable and appropriate, they encouraged EPA to consider implementation-related modifications (i.e., "other regulatory revisions") as a reason to revise a rule, even if there were no basis to revise the MCLG and/or MCL/TT requirements.

EPA Response: Implementation-related issues are the primary reason for the Agency's decision to revise the TCR at this time (67 FR 19030 at 19085, April 17, 2002 (USEPA, 2002g)), so it is clear that EPA considered implemented-related issues in its review. The Protocol Document (USEPA, 2002d; USEPA, 2003c) identifies the conditions under which the Agency will consider implementation-related revisions. EPA continues to believe these criteria are appropriate. During the current review, none of the identified potential

aplementation-related revisions raining to the chemical NPDWRs, in

EPA's judgment, met the stated criteria for reasons documented in EPA's final document, "Consideration of Other Regulatory Revisions for Chemical Contaminants in Support of the Six-Year Review of National Primary Drinking Water Regulations" (USEPA, 2003b).

6. Review of Occurrence and Exposure

a. Occurrence Database Concerns. A few commenters asked for information regarding next steps for the National Contaminant Occurrence Database (NCOD). Another commenter pointed out that States have been willing to assist EPA by providing occurrence data beyond what is required of them. However, the commenter raised concerns that he/she felt EPA needs to address to facilitate further data sharing.

Some commenters expressed concern about the completeness and representativeness of the 16-State data set used for the Six-Year Review. One commenter suggested that the Agency should have issued an Information Collection Request to obtain more complete data for the Six-Year Review analysis.

EPA Response: The Agency is updating the NCOD to provide sample data that have been quality checked and used in various EPA analyses. This update to NCOD includes unregulated occurrence data collected prior to 1999 as well as the latest Unregulated Contaminant Monitoring Rule data (64 FR 50556, September 17, 1999 (USEPA, 1999b)) reported by laboratories for public water systems required to report results. It also includes the data used for the Six-Year Review of regulated contaminants. EPA appreciates that some States are willing to share their full compliance monitoring records with the Agency, even though it is not required. The Agency and the States are continuing to work together to establish a protocol for data sharing, including safeguards to prevent misuse and misinterpretation of data.

The 16-State cross-section data set compiled for occurrence analyses for the Six-Year Review is the largest compliance monitoring data set for drinking water assembled by EPA to date. The design and construction of the 16-State cross-section data set was based on the fact that contaminant occurrence varies spatially (geographically) due to differing patterns of population, land use, chemical use, geology, hydrology, and climate. The detailed description of the "pollution-potential" and geographic diversity considerations, and the derived balanced cross-section of States (that was developed to be collectively indicative of national

occurrence) is included in the "Occurrence Estimation Methodology and Occurrence Findings Report for the Six-Year Review of Existing National Primary Drinking Water Regulations" (Occurrence Methodology Document) (USEPA, 2003d). EPA selected its 16-State cross-section to be as representative as possible of national contaminant occurrence. In EPA's judgment, these States provide a reasonable cross-section of agricultural and industrial pollution potential, as described in the Occurrence Methodology Document, and also provide geographic coverage of the United States. Therefore, EPA believes that the data assembled from these States is the most representative data currently available of national contaminant occurrence.

The Agency did receive occurrence data from States other than those in its 16-State cross-section. However, many State data sets contained incomplete records (e.g., no water type or population records specified) or had other quality problems. Therefore, they were not included in the analyzed data set.

b. Occurrence Analysis Methodology. One commenter noted that while the occurrence estimation methodology has several strengths, it also has a number of flaws. The commenter was concerned about the large proportion of nondetected observations in the occurrence data, and the difficulty of verifying the assumptions made by the Agency. The commenter agreed that EPA's occurrence analysis may represent a "decent" estimate given the limitations of the data. The commenter also noted that the occurrence estimation methodology is premised on "subjective decisions or qualitative observations * * rather than documented, statistically-based quantitative ones' and would like to have seen alternate approaches used to provide confirmation of the estimates. In addition, the commenter questioned why the Agency used "modeled data sets to test the model rather than a standard statistical strategy of basing the model on a portion of the data set and using the remainder to test the model."

One commenter stated that the Stage 2 analysis (Bayesian analysis) was poorly described and that this conflicts with the transparency requirements of the 1996 SDWA Amendments. In addition, the commenter asked EPA to clarify how the occurrence data from other survey efforts, which are summarized in "Occurrence Summary and Use Support Document for the Six-Year Review of Existing National Primary Drinking Water Regulations"

(Occurrence Summary Document) (USEPA, 2002f), were used to inform the modeling effort.

EPA Response: EPA's occurrence model development work was significantly revised to reflect peer review comments prior to the March 2002 Occurrence Methodology Document (USEPA, 2002e) and the April 17, 2002, Federal Register. The additional work involved the development of a detailed simulation study to evaluate the Bayesian model. EPA evaluated the performance of the Bayesian estimator and an alternative occurrence estimation approach, the Regression on Ordered Statistics (ROS) method, against synthetic data (i.e., data developed with known national contaminant occurrence distributions). This simulation study also enabled an explicit evaluation of the validity of the assumption of a log-normal distribution of the data.

The simulation study was conducted using varying conditions of a correctly and incorrectly specified model, and synthetic data sets developed with high and low amounts of non-detected data. The study findings indicated that the Bayesian estimator performed well at estimating the distributions of contaminant concentration means (especially in the upper tails), performed better than the alternate approach (i.e., the ROS method), and

curately estimated the uncertainty of a distributional estimates. The Agency believes that this analysis supports the validity of EPA's analytical approach. The Bayesian model was tested against the ROS approach because the ROS method is an accepted drinking water contaminant occurrence estimation approach and was used to estimate occurrence for the recent arsenic rule. These findings were all included and described in the Six-Year Review's Occurrence Methodology Document.

EPA has attempted to make its occurrence analysis as clear as possible. In response to the concerns raised by the peer reviewers, a less technical description of the occurrence estimation methodology, aimed at the general reader, was added to the main body of the document. A detailed description of the analysis, intended for readers with technical expertise, including the complete computer code used for model analysis, was incorporated into an appendix of the document. EPA agrees that its estimation methodology is complex, but also believes that it is as transparent as possible while still providing a technically accurate description of the Agency's analysis. The use of simple national occurrence (statistical) assessments is not possible

at this time because there is no national database with a complete collection of regulated contaminant occurrence data. Thus, there is no ideal basis for comparison of national occurrence studies (i.e., the true system contaminant means and national distributions of contaminant occurrence are not, and cannot, be known). The validation approach suggested by the commenter (i.e., basing the model on a portion of the data set and using the remainder to test the model) is intended for a regression-type of model using observed system means to develop a model for system-specific predictions. This approach is not possible for the six-year occurrence assessments, since, to the best of EPA's knowledge, data on the true individual system contaminant mean concentrations and national distributions are not available.

Regarding the other survey studies included in the Occurrence Summary Document, few, if any, provide the quantitative analytical results and national, representative coverage that would enable direct comparison to, or inclusion in, the Six-Year Review estimation analyses conducted with the 16-State cross-section occurrence data.

c. Other Issues Related to the Occurrence Technical Review. One commenter stated that the Agency's current approach to estimate occurrence, employing a conservative methodology and making conservative simplifying assumptions in the absence of definitive data, was appropriate. On the other hand, the commenter argued that it was not appropriate for the Agency to conduct as massive a data collection and analysis project as was undertaken without clear quantitative objectives for the analysis identified a priori. The commenter noted that it was not apparent from either the April 17, 2002, Federal Register or the Occurrence Methodology Document (USEPA, 2002e) that the Agency undertook an effort to set performance objectives for the occurrence estimation.

The commenter felt that the Occurrence Methodology Document does not allow the reader to determine if the data are well apportioned among the categories for which results are reported. They also noted that they were unable to find indications in the support document that such an analysis was undertaken in preparation for constructing the Bayesian model. The commenter stated that the support document does not include actual numeric counts or ranges of detected values and suggested that it would be useful to have this information by contaminant, State, system size category, and water type, as well as an

explicit count of non-detects by this same matrix.

EPA Response: There are several general approaches when undertaking and designing studies that require large amounts of data. As the commenter states, a priori data quality objectives are part of one research approach where study objectives (including technical statistical performance measures) are set, determinations are made on how to meet those objectives, and then the study is designed and implemented accordingly. This ideal was not practical for the national occurrence study conducted for the Six-Year Review because EPA did not have the resources to generate original data, and was thus dependent on the data that could be obtained from the States. The approach taken by the Six-Year Review was to gather a large amount of data that, in aggregate, was expected to be indicative of national contaminant occurrence, develop an occurrence estimation model tbat built upon what has been learned from recent regulatory development work, and then evaluate how good the resulting model estimates are.

As discussed in section IV.A.6.b of today's action, the true national distributions of contaminant occurrence cannot be known. The 16-State national cross-section data set used for the Six-Year Review is the largest compliance monitoring database for drinking water compiled by EPA to date. The database represents approximately 37 percent of the total number of public water systems and 43 percent of the total population served by public water systems in the United States. External peer reviews assessed the approach for developing the national cross-section and its "representativeness" separately under the Chemical Monitoring Reform (CMR) project (in 1998/1999) (USEPA, 1999c) and the Six-Year Review project (USEPA, 2002e), and provided generally favorable comments.

The data management and crosssection development have been described in detail in the support documents for the CMR and the Six-Year Review. Further tabulations of the data have been generated and presented, as the commenter requested, in the final Occurrence Methodology Document (USEPA, 2003d). This information includes the numbers and percentages of analytical detections and nondetections for each contaminant in each of the system size and source water type categories. Generally, because of the large amount of data and the manner in which the Bayesian model handles data, the distribution of observations across the various categories does not significantly affect EPA's estimates. The

number of analytical records differed by contaminant. EPA evaluated 27,648 to 93,062 analytical records for the individual inorganic chemicals, 32,606 to 121,327 records for the synthetic organic chemicals, and 123,229 to 201,235 records for the volatile organic chemicals. Most importantly, the Stage 2 occurrence model also quantifies the uncertainty of the estimates in the different categories of system size and source water type. Hence, the statistical significance of differences in occurrence between the categories can be easily assessed. However, the Agency believes it is more appropriate to consider the universe of potentially affected systems within the 16-State cross-section, rather than individual system categories, when making its revise/not revise decisions as part of the Six-Year Review process.

7. Consideration of Available Economic Information

Some commenters stated that, while the Agency's review of NPDWRs was generally consistent with NDWAC recommendations to EPA (NDWAC, 2000), it is not clear how the Agency took economic factors into account.

EPA Response: An EPA
memorandum, dated March 18, 2002,
describes the Agency's qualitative
evaluation of economic factors (USEPA,
'002c). This memorandum was cited in
the April 17, 2002, Federal Register and

railable in the docket for the Sixr Review (Docket No. OW-2002-0012). It notes that detailed economic analyses were not deemed by the Agency to be necessary to support its decisions of whether or not to revise a particular NPDWR. Rather, a qualitative assessment, based on the extent of occurrence of a contaminant at the MCL, as well as at alternative levels, was undertaken to inform the Agency's judgment about whether possible changes to an MCL offered a meaningful opportunity for health risk reduction and/or cost-savings to public water systems and their customers. EPA has conducted this assessment for 15 of the chemical NPDWRs for which the Agency had determined that a potential health or technological basis may exist for considering a revision to the MCLG/ MCL.2 EPA compared the estimated occurrence and exposure values at the current MCL and at potentially revised regulatory level(s). For 14 of these chemical NPDWRs, the Agency's

assessment showed that the differences were small. In EPA's judgment, these differences are unlikely to provide a meaningful opportunity for health risk reduction or cost-savings to public water systems and their customers. After consideration of these factors, EPA decided that any revision would be a low priority activity for the Agency, and, thus, not appropriate to revise at this time because of: Competing workload priorities; the administrative costs associated with rule making; and the burden on States and the regulated community to implement any regulatory change that resulted. In the case of dichloromethane, the Agency did not have sufficient data to recalculate the PQL to support any potential regulatory revision and thus placed it in the data gaps category.3

B. What Comments or New Information Did EPA Receive on Chemical Contaminant-Specific Issues?

1. Alachlor

One commenter stated that the Office of Pesticide Programs (OPP) found that the chloroacetanilide pesticides (acetochlor, alachlor, and butachlor) should be considered as a group of chemicals having a common mechanism of toxicity due to their ability to cause nasal turbinate tumors. The commenter believes EPA therefore should adopt a strong total chloroacetanilide pesticide standard that would strengthen the current standards.

EPA Response: Butachlor and acetochlor do not presently have an NPDWR and thus, are not included in the Six-Year Review. However, acetochlor is included on the Contaminant Candidate List (CCL) and may in the future be considered as a candidate for regulation. Alachlor is a regulated drinking water contaminant and is included in the Six-Year Review. It is currently undergoing a risk assessment and, therefore, the Agency believes that revision of the NPDWR is not appropriate at this time.

If the Agency decides to regulate either acetochlor or butachlor in the future, EPA may consider regulating them as a group, including alachlor, following a cumulative risk assessment process for pesticides that have a

common mechanism of toxicity. It would be premature to propose a total chloroacetanilide pesticide standard until a cumulative risk assessment is completed because this analysis could impact the Agency's evaluation of specific members of this group, or the group as a whole.

2. Antimony

- a. Health Effects. A number of commenters have suggested that the current MCLG and MCL of 0.006 milligrams per liter (mg/L) for antimony need to be revised. Some of the reasons given were:
- The study used to derive the current MCLG (Schroeder et al.,1970) is not consistent with current good laboratory practice guidelines and there are several newer studies of antimony toxicity that should be considered in deriving a new reference dose (RID).
- Animals used in the Schroeder et al., 1970 study had a viral infection. To compensate for this infection, adjustments were made to the size of the animal groups in an attempt to salvage the data.
- The antimony compound used in the Schroeder et al., 1970 study was potassium antimony tartrate, the most water soluble and toxic form of antimony. Antimony found in drinking water is likely to be in the form of less toxic trivalent and pentavalent antimony species. Therefore, basing the MCLG on the most toxic species of antimony (potassium antimony tartrate) is likely to overestimate the risk posed by antimony in drinking water.

EPA Response: EPA agrees that the MCLG and MCL for antimony may need to be re-evaluated. EPA is in the process of developing a new health risk assessment for antimony, taking into consideration new studies that have become available on the toxicity of antimony. EPA expects to complete the health risk assessment for antimony in the 2003-2004 time frame (68 FR 5870, February 5, 2003 (USEPA, 2003h)). As a result of the ongoing health risk assessment, a revision to the antimony standard is not appropriate at this time, and antimony will be re-evaluated as part of the next Six-Year Review process.

b. Treatment and Implementation Issues. Several commenters questioned the appropriateness of the antimony MCL, and the effectiveness of using the EPA-designated best available technologies (BATs) to meet the antimony MCL. A few small systems in Utah have levels of antimony in water at or above the MCL value of 0.006 mg/L. These systems were granted

²These 15 chemical NPDWRs are: Benzene; beryllium; chlordane; 1,2-dibromo-3chloroproppane dichloromethane; 1,1dichloroethylene; 1,2-dichloropropane; heptachlor, epoxide; hexachlorobenzene; lindane; yl; picloram; toxaphene; and 1,1,2-

The other three NPDWRs in the data gaps category, chromium, fluorida, and lead, were placed there for reasons other than occurrence and economic considerations. Chrominum is in the data gaps category because of the studies being conducted by the National Toxicology Program studies. Fluoride is in the data gaps category pending a National Academy of Sciences update of the health risk assessment and review of the RSC assumptions. Lead is in the data gaps category based on consideration of public comments (see section IV.B.13 of today's action).

exemptions contingent upon testing and installation of treatment by March 2004.

These systems are investigating treatment options for the removal of intimony from their source water. Commenters submitted supporting data documenting the results of their testing and cost analyses. According to commenters, on-site testing indicated that the designated BATs (i.e., reverse osmosis and coagulation/filtration) and most of the other tested treatments were ineffective and/or prohibitively expensive due to: raw water quality concerns; water conservation needs; current costs for water production; and other concerns, such as waste water management. However, commenters did identify treatment options that may be feasible, but these may require further investigation prior to full scale use.

EPA Response: As discussed in the April 17, 2002, Federal Register and as noted in the previous response in section IV.B.2.a, EPA does not believe it is appropriate to consider revisions to the NPDWR for antimony at this time because of the ongoing health risk assessment (67 FR 19030 at 19051

(USEPA, 2002g)).

When EPA initially promulgated the antimony NPDWR in 1992, the Agency estimated that 200 public water systems would be affected (USEPA, 1992). EPA recognizes that implementation of this tandard may present challenges for a

localities. Although the use of the gnated BATs for antimony may not be appropriate in some cases, as long as systems comply with the MCL, they are not limited to these technologies.

EPA believes that the treatment data generated by the commenters may be valuable and may provide insight into potential alternative treatment technologies. The Agency has revised the document, "Water Treatment Technology Feasibility Support Document for Chemical Contaminants; In support of EPA Six-Year Review of National Primary Drinking Water Regulations" (Treatment Feasibility Document) (USEPA, 2003g) to refer to these preliminary test data as they may be applicable to the development of potential new treatment technologies for the removal of antimony and other contaminants.

3. Atrazine

a. Health Effects. Several commenters addressed the EPA decision not to consider revision of the MCL for atrazine at this time. Some of these commenters stated that EPA should use the risk assessment, released by OPP in May 2002, as a basis for reconsidering a drazine NPDWR. One of the nmenters noted that the 2002 risk

assessment is based on reproductive and developmental endpoints which represents a change from the toxicity endpoint that formed the basis of the current MCLG. Two commenters stated that the MCL for atrazine should be revised upward because of the results of the 2002 OPP risk assessment in which the RID increased and the cancer classification changed from "possible human carcinogen" to "not likely to be a human carcinogen." The commenters stated that the change in the cancer assessment implies elimination of the additional 10-fold risk management factor used in 1991 to derive the MCLG/ MCL for atrazine. Another commenter stated that atrazine should be regulated using a non-linear approach which recognizes that there is a level at which no known health effects occur and that these findings must be part of the new

Conversely, another commenter stated that there is substantial new evidence from epidemiological and occupational studies that atrazine poses a serious cancer risk, and that it is an endocrine disruptor at low levels. The commenter believes EPA should adopt a revised atrazine and total triazine standard lower than (i.e., more stringent than) the current 0.003 mg/L standard for atrazine.

A commenter also urged the Agency to:

 Provide a definitive timetable for review of the standard;

 Outline a preliminary scope for its review of the standard; and

 State the underlying premise for the scope of the review.

Other commenters stated that the existing NPDWR only regulates the parent compound atrazine, and that a revised NPDWR should include the chloro-metabolite degradants (i.e., diaminochlorotriazine (DACT), desethyl atrazine (DEA), and desisopropyl atrazine (DIA)). These commenters believe that inclusion of the chlorometabolites would strengthen compliance monitoring programs for public water systems under SDWA and thereby strengthen public health protection. They stated that a regulation for atrazine and the chloro-metabolites should be developed and promulgated within the next 12 to 18 months. Another commenter stated that since the Agency has found that atrazine, simazine, propazine, and the degradants DACT, DEA, and DIA have a common mechanism of toxicity, these should be regulated in a total triazine regulation.

EPA Response: EPA does not believe it is appropriate to consider revisions to the NPDWR for atrazine at this time because the revised risk assessment has

not been finalized. For purposes of the Six-Year Review protocol, EPA considers a risk assessment final when an Interim Reregistration Eligibility Decision (IRED), Reregistration Eligibility Decision (RED), and/or IRIS assessments are complete. 4 Even though an IRED for atrazine was signed on January 31, 2003, an amended IRED is scheduled to be released in October 2003 which will include a Scientific Advisory Panel (SAP) peer review of new data related to health effects. Based upon the outcome of the SAP review, the October 2003 IRED may include additional information that could impact a revise/not revise decision. Therefore, EPA does not believe it is appropriate to consider possible revisions to the NPDWR at this time.

In reviewing the atrazine regulation, EPA will apply an approach consistent with the protocol used for the current review. The Agency will consider the same key elements (i.e., health effects review, technology review, other regulatory revisions review, and, if appropriate, occurrence/exposure analyses and consideration of available economic information) and apply the same basic decision tree for making a revise/not revise decision.

To address the issue of regulating the triazines as a group, the Agency is evaluating the unregulated triazines as part of the CCL process. When the risk assessment is completed for atrazine, the Agency will consider whether or not there are compelling reasons for considering a revision to the atrazine regulation or to wait until the risk assessment for the triazines, which considers issues of cumulative risk, is finalized. EPA will use the CCL regulatory determination process in deciding whether the triazines should be regulated as a group.

b. Costs of Treatment. Commenters stated that the costs associated with not revising the MCL are great. These commenters are concerned that State agencies will be required to develop total maximum daily loads (TMDLs) based on 303(d) 5 listings resulting from

Continued

The IRED is an intermediate decision for an individual pesticide that does not take into account cumulative risk issues for pesticides with a common mode of action. The RED does include cumulative risk. If an IRIS assessment is also in process when the IRED or RED is signed. EPA will make a case-by-case decision on whether to wait for the IRIS assessment before considering possible revisions to the NPDWR.

⁵Section 303(d) of the Clean Water Act and the implementing regulations (40 CFR 130.7) require States to develop TMDLs for waters where required point and nonpoint source pollution controls are not stringent enough to attain or maintain compliance with State water quality standards after the application of technology-based and other

an outdated MCL which creates a burden on State and local government, its citizens, and diverts limited resources away from programs that provide real benefits. Some commenters also stated that the treatment costs to hundreds of community water systems are considerable. One commenter also stated that these are real dollars that would otherwise be available for emergency services, education, nutrition programs, and other vital programs that are the responsibilities of local and State agencies.

EPA Response: As stated in the previous response in section IV.B.3.a, EPA does not believe it is appropriate to revise the NPDWR for atrazine at this time because the risk assessment is not yet final. If EPA decides to revise the NPDWR for atrazine, economic factors, including feasibility and an assessment of costs and benefits, will be taken into consideration for the drinking water program.

4. Beryllium

Two commenters believed that the current drinking water standard for beryllium is more stringent than necessary for the protection of public health and felt that EPA should adopt a higher value for the beryllium standard. These commenters disagreed with EPA on the use of an uncertainty factor of 300 in deriving the 1998 RfD. The

nmenters stated the use of certainty factors of 3 for database uncertainty, 10 for extrapolating data from a dog study to humans, and 10 for intraspecies variation is inappropriate. The commenters stated that EPA has the authority to raise the current drinking water standards for beryllium based on new information that allows for a smaller margin of safety than the one used by EPA. The commenter felt that the current standard for beryllium is "lower than necessary to protect the public from beryllium toxicity and results in clean-up standards that are lower than naturally occurring level of beryllium in water sources and soils." This commenter also expressed concern that the local application of the Federal drinking water standard to private wells in some cases caused undue concerns among users of those wells.

EPA Response: One of the purposes of the Six-Year Review is to determine if the MCL of a chemical should be changed based on a revised RfD or cancer classification. Analytical methods and treatment technologies are

required controls. A TMDL establishes the maximum amount of a pollutant that may be introduced into a waterbody while still ensuring 'ainment and maintenance of water quality indeeds.

considered, as well as occurrence in public water systems. The RfD for beryllium was revised in 1998 based on extensive Agency internal and external reviews, and is unlikely to be revised in the absence of new data. The 1998 assessment also provided separate cancer classification for inhalation and oral exposures (USEPA, 1998). In the revised assessment, the carcinogenicity of beryllium by the inhalation route was described as "likely," while that by the oral route of exposure "cannot be determined." As discussed in the April 17, 2002, Federal Register, the Agency considered the occurrence of beryllium at both potentially higher and lower regulatory levels. EPA concluded that a revision to the NPDWR would not result in a meaningful opportunity for health risk reduction or cost-savings to public water systems and their customers. As a result, revision of this NPDWR is a low priority action for the Agency and is not appropriate at this time.

The goal of drinking water standards is to protect public health. Therefore, it does not matter whether the source of contamination is naturally-occurring or man-made. While EPA appreciates the information on private wells, the SDWA requirements do not apply to private wells (i.e., wells that are not part of a "public water system"). The costs and benefits of a drinking water standard are assessed only with regard to the impacts on public water systems and their customers.

Carbofuran

Some commenters mentioned that the Agency concluded that N-methyl carbamates, including carbofuran, should be considered as a class because they have a common mechanism of toxicity. Therefore, they believe EPA should issue a stronger standard for total N-methyl carbamates, including carbofuran, which would be more stringent than the current carbofuran standard of 0.04 mg/L.

EPA Response: EPA is re-evaluating the toxicity of carbofuran. However, a final assessment has not been issued by EPA. The Agency considers N-methyl carbamate pesticides as a group of chemicals having a common mechanism of toxicity due to their ability to inhibit acetylcholinesterase. However, it is not appropriate to revise the NPDWR for carbofuran at this time because the Agency has not yet completed the final health risk assessment for carbofuran or the other N-methyl carbamates.

6. Chromium

One commenter requested that EPA move quickly in making a revise/not revise determination once the new data

on chromium become available from the National Toxicology Program (NTP) studies of the health effects of chromium VI.

EPA Response: The NTP studies that the commenter refers to should be available before the end of the next Six-Year Review cycle. Meanwhile, EPA is continuing to follow the progress of NTP in conducting subchronic and chronic studies of chromium VI.5 NTP made the data from the subchronic portion of the study available to the public in June 2002 (NTP, 2002). A peer review meeting was held at NTP on July 24, 2002. EPA will examine the peer review report covering the subchronic data once it becomes available. Once the subchronic and chronic studies are completed, the health effects data will be evaluated with regard to their impact on the present RfD and cancer assessment, and integrated with the occurrence and analytical method data before making a new revise/not revise decision.

7. 1,1-Dichloroethylene

In the April 17, 2002, Federal Register, the Agency preliminarily placed 1,1-dichloroethylene in the no revision category because a health risk assessment was pending at the time of publication. Since the publication of the April 17, 2002, Federal Register, the Agency has finalized the risk assessment for 1,1-dichloroethylene. The remaining paragraphs in this section include a brief background discussion about the original promulgation of the 1,1dichloroethylene NPDWR, the results of the appropriate six-year technical reviews and the Agency's revise/not revise decision.

a. Background. EPA published the current NPDWR for 1,1-dichloroethylene on July 8, 1987 (52 FR 25690 (USEPA, 1987)). The NPDWR established an MCLG and an MCL of 0.007 mg/L. The Agency based the MCLG on an RfD of 0.009 milligram per kilogram of body weight per day (mg/kg/day) and a cancer classification of C, possible human carcinogen.

b. Technical Reviews. EPA updated the risk assessment for 1,1-dichloroethylene on August 13, 2002 (USEPA, 2002i). The new risk assessment established an RfD of 0.046 mg/kg/day, based on the same toxicological study as that of the MCLG, but using an uncertainty factor of 100

^o Since NTP is posting its progress on its internet site http://ntp-server.niehs.nih.gov/htdocs/Studies/HexChromium/hexchromiumpg.html, EPA and the public will be able to evaluate the new data relative to the existing EPA assessment for chromium VI as it is released.

instead of 1,000, and using benchmark dose modeling for the dose-response analysis. Under the 1986 cancer guidelines (51 FR 33992, September 24, 1986 (USEPA, 1986)), 1,1dichloroethylene was assigned to Group C, possible human carcinogen. Under the draft revised "Guidelines for Carcinogen Risk Assessment" (USEPA, 1999a), the data for 1,1-dichloroethylene were considered inadequate for an assessment of human carcinogenic potential by the oral route.

Based on the change in RfD for 1,1dichloroethylene, using a 20 percent RSC and a 10-fold risk management factor for possible carcinogenicity, EPA

used 0.03 mg/L as a level for evaluating the occurrence data. Without the use of the 10-fold risk management factor, EPA also used 0.3 mg/L as a level for evaluating the occurrence data.

Analytical or treatment feasibility do not pose any limitations for the current MCL and would not be a limiting factor at the 0.03 mg/L or the 0.3 mg/L level (USEPA, 2002a; USEPA, 2003g). The Agency's review of possible "other regulatory revisions" did not identify any issues that are specific to 1,1dichloroethylene (USEPA, 2003b).

EPA evaluated the results of the occurrence and exposure analyses for 1,1-dichloroethylene to determine

whether possible changes to the standard would be likely to result in a meaningful opportunity for cost-savings to public water systems and their customers (USEPA, 2003d). Table IV-1 shows the results of the detailed occurrence and exposure analysis based on the 16-State cross-section for the current MCL (0.007 mg/L), and for two higher levels (0.03 mg/L and 0.3 mg/L). Based on the detailed analysis, it appears that 1.1-dichloroethylene is unlikely to occur at concentrations above 0.007 mg/L in the States used for the cross-section. BILLING CODE 6560-50-P

	Table IV-1: 1,	1-Dichlor	oethylene O	ccurr	ence ¹	
		Syste				
Level (in mg/L) Upper Level	16-State Cross- Section - Total Systems with Dat	> I	ated # of Syst Level Evaluate Lible intervals	ed	-	mated % of Systems > Level Evaluated credible intervals)3
Evaluated (without 10-fold risk 0.3 management factor)	19,101		0-0)		0.000%	(0.000% - 0.000%)
Lower Level Evaluated (with 10-fold risk 0.03 management factor) ³	19,101	0	(0-0)	0.	000%	(0.000% - 0.000%)
Current MCL 0.007	19,101	3	(1-6)			
	Population		by Systems ²	1 0.0	0144%	(0.00518% - 0.0311%
Level (in mg/L)	16-State Cross- Section - Total Population Served by Systems with Data	Estima Served > Lev	ted Population I by Systems el Evaluated tle intervals)	n	> L	ed % of Population yed by Systems evel Evaluated lible intervals)
Upper Level ivaluated (without 10-fold risk 0.3 management factor) ⁴	106,607,600	0	(0-0)	0.00		(0.000% - 0.000%)
Lower Level Evaluated (with 10-fold risk 0.03 management factor) ⁵	106,607,600	0	(0-0)	0.000	%	(0.000% - 0.000%)
Current MCL 0.007	106,607,600	14,400	(0 12(000)	<u> </u>		
s: Sults are based on the number		, 100	(0 - 136,900)	0.0135	i% (0.0000328% - 0.128%)

Results are based on the number and percent of systems (and the corresponding population served by those systems) with estimated mean concentrations above the specified level of evaluation.

² All percentages are shown to three significant figures. All system values are rounded to the nearest whole system. All population values are rounded to the nearest hundred.

^{3 &}quot;Credible intervals" are generated to quantify the uncertainty around each estimated probability in the Bayesian analysis of the occurrence data. For further explanation of credible intervals and the Bayesian analysis, please see "Occurrence Estimation Methodology and Occurrence Findings Report for the Six-Year Review of Existing National Primary Drinking Water Regulations" Based on the change in the RfD and a 20 percent RSC.

Based on the change in the RfD, a 20 percent RSC, and a risk management factor of 10.

⁶ This value dnes not necessarily reflect the number of systems out of compliance with the current MCL, because these data were collected over the 1993-1997 time period, and because the value represents the estimated mean value over that time period, not the running quarterly average on which compliance is based.

The results of the detailed occurrence and exposure analysis indicate that less than 0.02 percent of the 19,101 systems sampled in the 16-State cross-section, and less than 0.02 percent of the population served by those 19,101 systems might be affected if EPA were to consider levels as high as 0.03 mg/L to 0.3 mg/L. The current BATs and small system compliance technology for 1,1-dichloroethylene have other beneficial effects (e.g., reduction of other co-occurring contaminants, or other common impurities) in addition to 1,1-dichloroethylene removal. Therefore, if EPA were to consider any of these higher levels, the Agency does not know how many of these public water systems that are currently treating to comply with the current MCL of 0.007 mg/L would be likely to discontinue treatment that is already in place (USEPA, 2002c; USEPA, 2003g).

- c. Current Decision. Although there are new health effects data that might support calculation of a less stringent standard for 1,1-dichloroethylene, EPA does not believe a revision to the NPDWR for 1,1-dichloroethylene is appropriate at this time. In making this decision, the Agency considered whether any potential revision to the 1,1-dichloroethylene NPDWR is likely to provide a meaningful opportunity for cost-savings to public water systems and their customers. After consideration of this factor, EPA has decided that any evision to 1,1-dichloroethylene would be a low priority activity for the Agency, and, thus, is not appropriate to revise at this time because of:
- Competing workload priorities;
 The administrative costs associated
- with rulemaking; and
 The burden on States and the regulated community to implement any regulatory change that resulted.

8. Dichloromethane

One commenter stated that it may be difficult to lower the PQL for dichloromethane below the range of 0.001 to 0.002 mg/L since it is required in a number of EPA methods and therefore is a common laboratory contaminant. Because it is a common laboratory contaminant, the commenter stated that using the MDL for 524.2 and 502.2 does not constitute a reasonable basis for assuming that the PQL can be lower. The commenter stated that none of the existing WS studies had spike samples this low and, in addition, the occurrence data may have been compromised due to laboratory contamination.

EPA Response: The basis for EPA indicating that a lower PQL "may exist" as due to the fact that laboratories had ceater than 95 percent laboratory

passing rates using a +/-40 percent acceptance window at "known" spike concentrations close to current MCL of 0.005 mg/L. If laboratory contamination due to dichloromethane were a problem, such high passing rates at this value would not be expected. The MDLs for 524.2 and 502.2 were only used with the 10 times MDL multiplier to estimate what the lower value could be. However, EPA does agree that, at this time, the Agency does not have sufficient data to recalculate the PQL for dichloromethane and for this reason, the Agency placed it in the data gap category

Regarding the occurrence issue, EPA has no data to suggest that high occurrence values were due to false positives from laboratory contamination and the Agency is proceeding on the assumption that State data are accurate unless there is information to the contrary. If laboratory contamination due to dichloromethane does exist, laboratories should be able to identify and discern a contamination issue if they are running laboratory blanks.

9. Di(2-ethylbexyl)adipate (DEHA)

One commenter submitted detailed comments regarding di(2-ethylhexyl)adipate (DEHA). The commenter believed that EPA should consider removing the regulation for DEHA and provided the following reasons:

• The regulation of DEHA in drinking water does not provide any meaningful reduction in the health risk to humans because it is unlikely to cause adverse effects to humans, including reproductive effects, except at very high doses which cannot be attained in drinking water, due to the low water solubility of DEHA

solubility of DEHA.

• The weight of evidence indicates that the peroxisome proliferation mechanism of DEHA rodent carcinogenicity is not relevant to humans. Thus, the MCLG for DEHA should not include an additional 10-fold risk management factor for possible carcinogenicity.

 The legislative history of the 1996 SDWA indicates that Congress envisioned circumstances where relaxation of an MCL would provide the same level of health protection as the existing regulation. Accordingly, if DEHA cannot be deregulated, the commenter believes the MCLG and MCL should be increased.

EPA Response: DEHA was regulated in 1992. Since that time, new studies have become available on the toxicity of DEHA and its metabolites. For this reason, EPA decided to initiate a new health risk assessment of DEHA (67 FR 1212, January 9, 2002 (USEPA, 2002a)).

The assessment will include examination of the studies on which the current NPDWR is based, as well as an evaluation of the data provided by this commenter and new studies that have become available since DEHA was regulated. This health risk assessment is planned for completion in the 2003– 2004 time frame (68 FR 5870, February 5, 2003 (USEPA, 2003h)) and is expected to include development of an RfD for non-cancer health effects, as well as an assessment of potential carcinogenicity from oral exposure. At this time, it is premature to predict the outcome of the Agency's assessment. Thus, as discussed in section IV.A.2.a of today's action, the Agency believes that revision to the NPDWR for DEHA is not appropriate at this time. EPA will determine in the future if revision of the MCLG/MCL is warranted. Any revision to the MCLG/MCL will also take into consideration all the new information, including the water solubility of DEHA under various environmental conditions.

As stated by the commenter, the legislative history of the 1996 SDWA Amendments supports EPA's interpretation that the Agency could increase an MCLG and MCL as long as the relaxed standard does not lessen the level of public health protection. However, EPA does not believe, at the present time, that it can demonstrate that deregulating DEHA would maintain the current level of public health protection (see section IV.A.1.c of today's action).

10. Di(2-ethylhexyl)phthalate (DEHP)

The same commenter who submitted comments on DEHA also submitted detailed comments regarding di(2-ethylhexyl)phthalate (DEHP). The commenter felt that EPA should consider removing the regulation for DEHP for a variety of reasons, including the following:

- The regulation of DEHP in drinking water does not provide any meaningful reduction in the health risk to humans.
- The weight of the evidence indicates that the mode of action through which DEHP causes cancer in rodents is not relevant to humans and, thus, the MCLG for DEHP should not be zero. Any MCLG for DEHP should be based on a threshold endpoint and not on cancer. The commenter cited the February 2000 International Agency for Research on Cancer reclassification of DEHP from Group 2B (possibly carcinogenic to humans) to Group 3 (not classifiable as to its carcinogenicity to humans) as justification for recommending that EPA also reconsider its cancer classification.

 The solubility of DEHP in drinking water is well below any concentrations that would pose a risk to humans.

 If DEHP were to be considered for regulation under the statutory requirements of the 1996 SDWA, it would not be regulated.

• The legislative history of the 1996 SDWA indicates that Congress envisioned circumstances where relaxation of an MCL would provide the same level of health protection as the existing regulation. Accordingly, the commenter believes consideration should be given to increasing the MCLG for DEHP based on the new health effects data.

• Reproductive effects from DEHP as observed in rodents do not appear to be relevant for primates and the doses that are associated with effects in animals are well above those that would be experienced for humans exposed through drinking water because of solubility limitations. The commenter also highlighted the findings of the NTP Center for the Evaluation of Risk to Human Reproduction that there was "minimal concern for reproductive or developmental toxicity for the general population, based on estimates of total exposure to DEHP."

EPA Response: Revision of the NPDWR for DEHP is not appropriate at this time because an Agency health risk issessment is currently in process. The

essment is anticipated to be apleted in the 2003–2004 time frame [68 FR 5870, February 5, 2003 (USEPA, 2003h)). Advances in understanding differences between the primate and rodent response to DEHP and the body of toxicological data that have become available in the past decade motivated the Agency's re-examination of DEHP and will be fully considered in the reassessment.

Once the Agency assessment is completed, EPA will consider the findings and will determine if there is a compelling reason to review the DEHP NPDWR prior to the next Six-Year Review cycle. As discussed in sections IV.A.1.b and IV.A.2.a of today's action, "revise" versus "not revise" decisions under the Six-Year Review take into consideration occurrence, advances in analytical methods, treatment technologies, available economic information, and other factors.

As stated by the commenter, the legislative history of the 1996 SDWA Amendments supports EPA's interpretation that the Agency could increase an MCLG and MCL as long as the relaxed standard does not lessen the level of public health protection.

'Yowever, EPA does not believe, at the esent time, that it can demonstrate

that deregulating DEHP would maintain the current level of public health protection (see section IV.A.1.c of today's action).

11. Fluoride

EPA received three comments on the Agency's decision to place fluoride in the data gap category while the National Academy of Sciences (NAS) examines the toxicological and RSC data published over the last decade. Two of the commenters supported EPA's decision. One of these requested that the NAS concentrate its review on all of the data on the toxicology of fluoride and not just data on the critical skeletal effects. A third commenter requested that EPA not lower the MCL for fluoride from 4 mg/L to 2 mg/L and supported the 1986 EPA decision that dental fluorosis is a cosmetic effect rather than an adverse health effect. The commenter stated that the Public Health Service (PHS) recommended fluoridation level to be used at schools is 3 mg/L. The commenter also stated that if EPA were to lower the MCL, then schools that are currently fluoridating might have a conflict with the PHS recommendations and the EPA MCL.

EPA Response: The National Research Council (NRC) of the NAS has agreed to review the toxicological data on fluoride that have been published since it completed the 1993 study of "Health Effects of Ingested Fluoride" (NRC, 1993), and to examine the data on relative fluoride exposure from drinking water compared to fluoride exposure from the diet and fluoride-containing dental products. Although the Agency indicated in the April 17, 2002, Federal Register that new data on bone effects were a reason for initiating the data review (because bone effects were the basis of the present MCLG), the NAS review will look at the new toxicological data for all endpoints. It is anticipated that the NAS review will take about two years to complete. Because of this pending review, revision of the NPDWR for fluoride is not appropriate at this time.

It is therefore premature to make any judgment regarding the NAS findings and whether or not they may lead to a consideration of a change in the MCL. However, PHS recommendations for school fluoridation programs are designed to provide the benefits of fluoridation without increasing the risk for dental fluorosis. The PHS recommends school water fluoridation only if:

 The school has its own source of water;

 The school is not connected to a community water system; More than 25 percent of students are not served by a public water system that provides water at levels adequate to protect against dental caries; and

 The students served are kindergarten age or greater.

12. Glyphosate

Two commenters made the statement that, despite continued use of glyphosate in pesticide applications, available data and the Agency's occurrence analysis, which includes a prediction of frequency of occurrence at levels below detection, indicate that glyphosate is not observed in compliance monitoring. One of these commenters stated that the occurrence appeared to be rare (less than 0.1 percent) at concentrations 1,000 times lower than the MCL. In addition, according to the commenters, the cost of analyzing for glyphosate is expensive, since it is a single analyte analysis. Accordingly, the commenters wanted EPA to reconsider the glyphosate standard taking costs and benefits into account. The commenters felt that the data may indicate that a glyphosate standard is inappropriate and does not result in any additional public health protection. Therefore, the commenters recommended EPA pursue data gaps that the Agency would need to fill in order to demonstrate that eliminating the glypbosate standard would not lower public health protection.

EPA Response: EPA is conducting an Agency risk assessment for glyphosate that will update the 1993 OPP assessment. As a part of this process, EPA is considering all the data that have been published or submitted to EPA since the completion of the RED in 1993 (USEPA, 1993). Accordingly, revision of the glyphosate NPDWR is not appropriate at this time due to the pending Agency assessment.

EPA recognizes that some utilities feel that the analysis of glyphosate in drinking water is expensive and that this should be taken into consideration with respect to cost and benefits. This will be considered when EPA evaluates glyphosate in the next review cycle (unless there is a compelling reason to evaluate glyphosate on an accelerated schedule). For the reasons stated in section IV.A.1.c of today's action, EPA does not believe it is appropriate to consider deregulation of glyphosate at this time.

13. Lead and Copper

a. Research Needs. Three commenters acknowledged the Agency's January 2000 revisions to the Lead and Copper Rule (LCR) but stated that the Agency should continue to consider how to

make the LCR easier to implement. In particular, they recommended that the following three LCR-related research areas be incorporated into EPA's overall research strategy:

1. How well LCR monitoring results correlate to actual exposure and the effectiveness of the rule in protecting public health.

2. Whether there is a correlation between water quality at indoor and outdoor taps.

 What effect the ban on lead in fixtures has had on lead levels and whether changes need to be made based on this ban.

The commenters explained their rationale for recommending that the Agency determine if a correlation could be established between indoor and outdoor water quality. They stated that a major weakness of the LCR is that sample integrity may be compromised by allowing customers to collect water samples. If the Agency could establish such a correlation, the LCR could be revised to allow water system operators to collect samples from outdoor taps; thereby removing the need for customer-collected sampling.

EPA Response: EPA recognizes that the LCR is a challenging rule that requires difficult solutions to implement, but continues to believe that the public health objective addressed by the rule is as important and essential today as it was when the rule was first

omulgated. Since the Agency promulgated the revisions to the LCR in January 2000 (65 FR 1950, January 12, 2000 (USEPA, 2000)), the Agency has received no significant new information that would support a revision. However, the Agency recognizes that more research would be useful to obtain additional information that could be utilized to address some of the issues associated with the implementation of this rule. For this reason, EPA has revised its rationale for not revising the NPDWR for lead and placed it in the data gaps category. Although the Agency continues to believe that the NPDWR for copper belongs in the risk assessment in process category at the present time, EPA will also consider copper-related risk management and implementation issues as a part of any LCR-related research plans. The Agency is committed to working with stakeholders to support and coordinate identification and prioritization of LCRrelated research needs. Until this research is completed, EPA believes it is premature to consider revisions to the LCR; as a result, revision of the LCR is not appropriate at this time.

The Agency believes that iderstanding the possible correlation

between monitoring results and actual rates of exposure and public health protection is a valid issue. However, EPA recognized during the initial regulatory development of the LCR that a significant effort would be necessary to provide a statistically valid number and frequency of samples for an exposure assessment. The Agency thus adopted an alternative approach which specified a monitoring scheme that sought to "* * * assure that systems are performing 'optimal corrosion control' in part by requiring systems to conduct comprehensive tap sampling at homes specifically targeted for their potential to contain elevated levels of lead and copper" (56 FR 26460 at 26514, June 7, 1991 (USEPA, 1991b)). One issue in assessing exposure reduction resulting from the LCR is a determination of an exposure baseline. EPA does not have a lot of data against which to measure changes in exposure that have occurred as a result of rule implementation. For these reasons, EPA believes that there is still insufficient information to change the basic monitoring approach adopted in the original rule, but recognizes that additional research may be useful.

Research on whether a correlation exists between the water quality at indoor taps and water quality at outdoor taps is a very complex issue. Several variables potentially affect whether a reliable correlation exists between indoor and outdoor taps. These variables include: standing time within the system; contact time with the building plumbing; and the content of the interior plumbing. These variables, coupled with the fact that lead levels from building-to-building can be highly site-specific, make a correlation between indoor and outdoor taps difficult to establish. EPA continues to believe that focusing on the point of delivery to the customer most closely links the data collected to the water quality consumed by the customer.

EPA recognizes the commenter's concerns regarding the integrity of samples collected by drinking water customers. To date, however, the Agency has not been able to identify an acceptable alternative to monitoring at the consumer's tap that can produce results equivalent to those obtained at the point of consumption in terms of ensuring adequate public health protection.

Regarding the commenter's third recommendation, EPA will consider this research need as part of the Agency's overall drinking water research planning process.

b. Relaxing the Monitoring Requirements. Three commenters recommended that water systems be

allowed to conduct water quality parameter (WQP) monitoring in lieu of continued lead and copper tap monitoring. One of these commenters added that this should be allowed once the system has demonstrated that it does not have a lead problem. This commenter also stated that the new requirements to use lead-free solder and plumbing fixtures should preclude problems with lead. Two commenters noted the difficulty that water systems are having maintaining their current sampling pool because homeowners no longer want to participate in the LCR monitoring program. One of these commenters recommended using WQP results to ensure corrosion control treatment is being adequately maintained and to stop lead and copper monitoring after three to five years. The commenter added that once the system ceases lead and copper monitoring, it can use public education to supplement continuing corrosion control, and can use coupons to demonstrate that corrosion rates meet accepted standards.

EPA Response: While EPA is sensitive to the difficulties associated with the monitoring requirements of the LCR, the Agency is also concerned about the implications of reduced or discontinued monitoring. Significant treatment changes or water chemistry disturbances (such as new water sources, major pH/ coagulation changes, disinfectant changes, or seasonal water/treatment changes) can influence the effectiveness of corrosion control, which in turn will require appropriate adjustments of treatment. Current regulations require water systems to continue monitoring lead and copper levels to assure that water quality changes adversely affecting the presence of these contaminants in the drinking water are detected and to assure that appropriate adjustments to maintain optimal corrosion control are made. Proper process control, including water quality and corrosion inhibitor residual concentration monitoring in the distribution system, is the key to making any corrosion control or other treatment work, and assure the continuation of proper water quality. However, EPA recognizes that some changes might be justified in the future based on new, scientifically valid, information and/or research. EPA is considering aspects such as the implications of simultaneous treatment modifications on water quality, including lead and copper control, in its research planning. EPA is not yet able to determine whether the outcome of such research will provide a basis for modifications to the LCR treatment or monitoring

requirements. As stated in the response in section IV.B.13.a. of today's action, EPA has placed the LCR in the data gaps category pending the completion of future research.

c. Corrosion Control Treatment Strategy. Two commenters noted concerns regarding the lead and copper corrosion control strategy. One commenter indicated that the LCR should be revised to allow systems to change corrosion control strategies. The commenter stated that considerable development of the corrosion control market has occurred since systems made their initial assessments and implemented corrosion control programs. The commenter felt that currently, the "LCR locks utilities into a given control strategy," when in some instances limited pilot work and ongoing WQP monitoring would allow a system to re-assesses its treatment and implement an alternative corrosion control inhibitor.

The second commenter indicated that the current corrosion control strategies are marginally effective at preventing particulate lead and copper from entering the water supply. The commenter recommended that EPA consider methods for mitigating the release of insoluble components from plumbing fixtures.

EPA Response: The Agency disagrees at the LCR locks utilities into a given irol strategy, but feels it is necessary demonstrate a sound basis for reassessing and implementing an alternative treatment strategy in the context of the existing regulation. EPA notes that the current regulation provides some flexibility to both States and water systems in the choice of a corrosion control strategy. For example, in response to its own initiative, a request by a water system or other interested party, a State may modify its determination of the optimal corrosion control treatment, among those listed in the Federal regulation, or may modify optimal WQPs if the State determines such changes are necessary to ensure that the system continues to optimize corrosion control treatment (40 CFR 141.82(h)).

The Agency believes that the existing requirements to notify the State when changing a corrosion control strategy remain necessary and appropriate. After they have optimized corrosion control, water systems must notify the State of any treatment changes within 60 days of the change (40 CFR 141.90(a)(3)). The

Agency encourages water systems to notify the State prior to making any changes thus allowing the Primacy Agency to review the changes to reduce the potential for detrimental sideeffects. In the Agency's experience, changes in treatment, such as (but not restricted to) replacement of high pH treatment with corrosion inhibitor, changes in coagulant and coagulation conditions, changes in disinfection, installation of membrane processes, or introduction of chemically different waters into the distribution system provide potential for detrimental sideeffects. Water treatment changes, therefore, should only be done with the greatest care and pilot investigations. While changes to treatment can be made under the existing regulation, systems should conduct additional monitoring (e.g., of lead, copper, and WQPs) until the new treatment is fully implemented and stabilized.

EPA also recognizes that the current LCR may limit flexibility to some extent, particularly in the adoption of new or emerging technologies. The original rule attempted to balance this concern with the need to provide strong public health protection by ensuring that only control strategies of proven effectiveness are adopted. The Agency does not have an adequate basis to revise the treatment requirements at this time but will continue to monitor new developments, including emerging technology. The Agency may consider revisions to the LCR prior to the end of the next Six-Year Review cycle if the Agency receives new, scientifically-valid, information that provides a basis for achieving significant improvement in public health protection or significant cost-savings to utilities and their customers while maintaining current public health protection.

EPA has always recognized that the release of insoluble particulate material containing lead and copper can be an issue in some water systems. While more research may be of interest to improve optimization of corrosion control approaches with respect to this source, EPA expects that evaluations and pilot studies by water systems should include testing and consideration of the relative effectiveness of different treatments towards particulate release in systems for which it is important.

d. Lead Levels in School Drinking Water. One commenter was concerned that the data on lead levels that was analyzed under the Six-Year Review of NPDWR standards may not indicate actual lead contamination of drinking water sources. As an example, the commenter noted that even though

Baltimore City is in compliance for lead levels, ½ of Baltimore schools are using alternative sources of drinking water due to lead contamination. The commenter expressed concern that since data obtained from schools, such as the data from Baltimore, was not considered in the evaluation of lead contamination in drinking water, the most vulnerable population may not be protected from exposure to lead. The commenter stated that it is time for the Agency to reassess how lead levels are evaluated.

EPA Response: The LCR is designed to address system-wide problems with lead and copper contamination. The rule does not specifically target particular structures, such as schools, but rather contains a monitoring protocol designed to ensure that the overall levels of lead and copper system-wide are minimized. Once optimal treatment is implemented, any remaining problems with elevated lead levels in schools may be due to plumbing, coolers, or other materials in the building. These potential sources of lead in schools are of concern and for this reason are explicitly addressed under the provisions of the Lead Contamination Control Act of 1988 (LCCA) (sections 1461 to 1465 of SDWA). The LCCA directed EPA to publish a guidance manual and testing protocol to assist States and schools in identifying sources and determining the extent of lead contamination in school drinking water and, if necessary, in remedying such contamination. In January 1989, the Agency published and distributed the guidance manual, "Lead in School's Drinking Water," to States and schools (USEPA, 1989). In 1994, the Agency updated and revised the guidance manual entitled "Lead in Drinking Water in Schools and Nonresidential Buildings" (USEPA, 1994). A copy of this manual may be obtained from the Safewater website http:// www.epa.gov/safewater/consumer/ leadinschools.html. In addition, the LCCA imposed a ban on the manufacture and sale of water coolers that are not lead free. The LCCA requirements are independent of the NPDWRs and therefore are not addressed under the Six-Year Review process. However, the Agency is continuing to work with schools and States to address problems dealing with lead in school drinking water.

14. Lindane (γ-hexachlorocyclohexane)

In the April 17, 2002, Federal Register, the Agency preliminarily placed lindane in the no revision category because a health risk assessment was pending at the time of publication. One commenter stated that

Treatment changes are categorized as changes to 2 Treatment process, including (but restricted to) disenfection, disinfection by-Luct removal, and corrosion control.

the RED risk assessment for lindane, issued after publication of the April 17, 2002, Federal Register, should be considered in the Agency's review of the NPDWR and expressed concerns regarding the existing regulation. The commenter stated that the current NPDWR is based on an RfD developed in 1988 on the basis of adverse kidney effects and should be revised (USEPA, 1988). The kidney effects were determined to occur through a pathway that is not relevant to human health risk assessment. The commenter stated that the new OPP toxicological assessment has resulted in a significant change to the quantitative dose-response assessment for lindane and that there are no data gaps or uncertainties which would prevent a revision of the NPDWR for lindane at this time.

EPA Response: Since the publication of the April 17, 2002, Federal Register and receipt of the comment regarding lindane, the Agency has finalized the risk assessment for lindane and signed the RED on July 31, 2002. The remaining paragraphs in this section include a brief background discussion about the original promulgation of the lindane NPDWR, the results of the appropriate six-year technical reviews and the Agency's revise/not revise recision.

a. Background. EPA published the current NPDWR for lindane on January 30, 1991 (56 FR 3526 (USEPA, 1991a)). The NPDWR established an MCLG and an MCL of 0.0002 mg/L. The Agency based the MCLG on an RfD of 0.0003 mg/kg/day and a cancer classification of C, possible human carcinogen.

b. Technical Reviews. EPA updated the risk assessment on July 31, 2002 (USEPA, 2002h). The new risk assessment established an RfD of 0.0047 mg/kg/day. The Food Quality Protection Act (FQPA) of 1996 provides for an additional safety factor of up to 10-fold, if necessary, in assessing the risks to infants and children to take into account the potential for pre- and post-natal toxicity, and the completeness of the toxicity and exposure databases. This is referred to as the FQPA safety factor. The Agency concluded that an FQPA safety factor of three was required for lindane since there is evidence for increased susceptibility of the young demonstrated in a developmental neurotoxicity and two-generation reproductive toxicity study in rats. The rationale for using an FQPA safety factor of three is detailed in the RED.

In accordance with the 1999 EPA Draft "Guidelines for Carcinogen Risk Assessment" (USEPA, 1999a), the Agency classified lindane as "suggestive evidence of carcinogenicity, but not sufficient to assess human carcinogenic potential." Based on the RfD for lindane of 0.0047 mg/kg/day, the application of the additional FQPA safety factor of three to this RfD, a 20 percent RSC, and a 10-fold risk management factor of suggested evidence of carcinogenicity, EPA used 0.001 mg/L as a level for evaluating the occurrence data.

Analytical or treatment feasibility do not pose any limitations for the current MCL and would not be a limiting factor at the 0.001 mg/L level (USEPA, 2003a; USEPA 2003g). The Agency's review of possible "other regulatory revisions" did not identify any issues that are specific to lindane (USEPA, 2003b).

EPA evaluated the results of the occurrence and exposure analyses for lindane to determine whether possible changes to the standard would be likely to result in a meaningful opportunity for cost-savings to public water systems and their customers (USEPA, 2003d). Table IV-2 shows the results of the detailed occurrence and exposure analysis based on the 16-State cross-section for concentrations of 0.0002 mg/L (the current MCL), and for 0.001 mg/L. Based on the detailed analysis, it appears that lindane is unlikely to occur at concentrations above 0.0002 mg/L in the States used for the cross-section. BILLING CODE 6560-50-P

	Table IV-	2: Lindane	Occurrence	ı	
		Systems ²			
Level (in mg/L)	16-State Cross- Section - Total Systems with Data	> Level]	# of Systems Evaluated intervals) ^{3,4}	>	nated % of Systems Level Evaluated redible intervals) ³
Level Evaluated 0.001	16,098	0	(0-0)	0.000%	(0.000% - 0.000%)
Ситепt MCL 0.0002	16,098	. 0	(0,0)	-	
	Population	on Served by S	(0-0) Systems ²	0.000%	(0.000% - 0.000%)
Level (in mg/L)	16-State Cross- Section - Total Population Served by Systems with Data	Estimated I Served by > Level Ev (credible in	Population Systems valuated	Ser > L	ted % of Population ved by Systems evel Evaluated dible intervals) ³
Level Evaluated 0.001	99,942,600	0	(0-0)		
Current MCL 0.0002	99,942,600	0	(0.0)	0.000%	(0.000% - 0.000%)
otes:		U	(0-0)	0.000%	(0.000% - 0.000%)

¹ Results are based on the number and percent of systems (and the corresponding population served by those systems) with estimated mean concentrations above the specified level of evaluation.

² All percentages are shown to three significant figures. All system values are rounded to the nearest whole system. All population values are rounded to the nearest hundred.

^{3 &}quot;Credible intervals" are generated to quantify the uncertainty around each estimated probability in the Bayesian analysis of the occurrence data. For further explanation of credible intervals and the Bayesian analysis, please see "Occurrence Estimation Methodology and Occurrence Findings Report for the Six-Year Review of Existing National Primary Drinking Water Regulations" (USEPA, 2003d).

⁴ This value does not necessarily reflect the number of systems out of compliance with the current MCL, because these data were collected over the 1993-1997 time period, and because the value represents the estimated mean value over that time period, not the running quarterly average on which compliance is based.

The results of the detailed occurrence and exposure analysis indicate that few, if any, of the 16,098 systems sampled in the 16-State cross-section might be affected if EPA were to consider levels as high as 0.001 mg/L. The current BATs and small system compliance technology for lindane have other beneficial effects (e.g., reduction of other co-occurring contaminants, or other common impurities) in addition to lindane removal. Therefore, if EPA were to consider a higher level, the Agency does not know how many of these public water systems that are currently treating to comply with the current MCL of 0.0002 mg/L would be likely to discontinue any treatment that is already in place (USEPA, 2002c; USEPA, 2003g).

c. Current Decision. Although there are new health effects data that might support calculation of a less stringent standard for lindane, EPA does not believe a revision to the NPDWR for lindane is appropriate at this time. In making this decision, the Agency considered whether any potential revision to the lindane NPDWR is likely to provide a meaningful opportunity for cost-savings to public water systems and their customers. After consideration of this factor, EPA has decided that any revision to lindane would be a low riority activity for the Agency, and, hus, is not appropriate to revise at this e because of:

- Competing workload priorities;
- The administrative costs associated with rulemaking; and
- The burden on States and the regulated community to implement any regulatory change that resulted.

15. Simazine

One commenter agreed that simazine should be addressed after the risk assessment is completed in 2003 or 2004. The commenter requested that the Office of Water (OW) work closely with the OPP on the risk assessment at that time. The commenter also recommended that OW address the revision of the existing simazine NPDWR before the next review cycle year, scheduled for 2008. The commenter believes the extensive mammalian toxicology database, submitted as part of the Triazine Special Review, can be used in this process.

EPA Response: OW has been coordinating with OPP for the revision of the atrazine and simazine risk assessments. Once the simazine risk assessment is completed, EPA will determine whether a compelling reason

sts to consider review of the simazine 'DWR on an accelerated schedule.

C. What Comments Did EPA Receive Regarding the Review of Implementation-Related Issues for Chemical NPDWRs?

Several commenters recommended that EPA ensure consistent application of rules by making rules more consistent with respect to monitoring frequency, triggers for increased monitoring, criteria for returning to routine monitoring, and criteria for reducing sample requirements. In addition, commenters suggested that the Agency review possible ways for reducing the reporting burden on States, which could free up State resources currently used to implement rules.

One commenter was concerned about monitoring and reporting issues in conjunction with CMR. The commenter felt that EPA should not miss an opportunity to relieve some of the unnecessary confusion that the monitoring requirements of Phase II and V have created. This confusion includes issues such as, what a detection is and what the monitoring requirements are for systems in States without a waiver program. EPA was encouraged to provide this consistency as much as possible, including using the standard monitoring framework to allow States and water systems to more easily understand rule requirements and reduce the need for States to update their data management systems.

One commenter said EPA should ensure consistent application of rules by determining whether or not chronic contaminants should be regulated at non-transient non-community water systems (NTNCWSs), and review existing NPDWRs to ensure that rules are applied consistently. Another commenter recommended that the compliance language for the synthetic organic chemicals (SOCs) and volatile organic chemicals (VOCs) in the Final Arsenic Rule (66 FR 6975, January 22, 2001 (USEPA, 2001)) be adopted for the inorganic chemicals (IOCs), and that systems not be considered in violation of the MCL until it has completed one year of quarterly samples.

EPA Response: The Agency agrees that consistency across regulations is desirable to the extent that it does not jeopardize public health protection or the environment.

As part of the Advanced Notice of Proposed Rulemaking for CMR (62 FR 36100, July 3, 1997 (USEPA, 1997a)), EPA considered some of the issues raised by the commenters. However, during the comment period for the CMR, stakeholders generally indicated that the existing monitoring framework was sufficient. Most State commenters

indicated that it would be too burdensome to adopt CMR. As a result, the Agency decided to take no further action on the CMR. However, the Agency established a standardized monitoring framework which applies to all of the regulated chemical and radiological contaminants (except lead and copper). The new chemical and radiological rules that EPA has promulgated (e.g., arsenic and radionuclides) are coordinated with the standardized monitoring framework. The Agency made special efforts to ensure that the reduced monitoring periods are in line with the 3-year compliance periods in the standardized monitoring framework.

To assist States with understanding rule requirements, the Agency conducted a series of Phase II/V training in 2001. The training provided information to help States make informed decisions about reducing quarterly monitoring requirements. With respect to reduced monitoring, States currently have the flexibility to reduce the frequency of monitoring and/ or to waive sampling requirements for any given contaminant after minimum criteria are met to demonstrate that the system is reliably and consistently below the MCL and/or not vulnerable to contamination.

NTNCWSs are traditionally regulated for chronic contaminants. However, through an alternative mechanism, the Agency is currently evaluating risk and exposure as they pertain to NTNCWS monitoring requirements. This review will not be completed in time for this Six-Year Review process. Until all the issues have been identified and specific options have been formulated, it will not be clear if a revision to regulations is indicated.

EPA intends to consistently implement compliance determination provisions for IOCs, SOCs, and VOCs for all NTNCWSs and community water systems, as described in the preamble to the Final Arsenic Rule (66 FR 6975 at 6990, January 22, 2001 (USEPA, 2001)). The rule makes compliance determinations based on a running annual average. The clarifications to compliance determinations for SOCs, IOCs, and VOCs are based on the average of the initial MCL exceedance and any subsequent State-required confirmation samples. States have the flexibility to require confirmation samples and more frequent monitoring, in addition to required quarterly samples. The average of the exceedance and confirmation sample constitutes the first quarterly sample. Compliance with the MCL is based on the average of the first quarterly sample and three

additional samples over a period of one year, unless any one quarterly sample would cause the running annual average to exceed the MCL. Then the system is out of compliance immediately.

D. What Comments Did EPA Receive on the Total Coliform Rule?

Several commenters addressed the TCR. Several commenters raised several issues relating to monitoring. Some contended that routine monitoring should be focused on critical locations in the distribution system, rather than on the current requirement to monitor all parts of the distribution system. They also urged EPA to allow the use of dedicated sampling taps. Some commenters argued for allowing a finished water storage reservoir as a routine monitoring site. Two commenters urged EPA to focus on E. coli as the measure of water quality in the distribution system, rather than on total coliforms. In addition to routine monitoring, a few commenters addressed the topic of repeat samples after a total coliform-positive sample. One commenter, for example, urged EPA to eliminate the requirement to take upstream and downstream repeat samples after a total coliform-positive sample. Environmental groups urged EPA to strengthen the TCR and other iles that protect against pathogens, and xhorted EPA not to ease the TCR ırden such that public health is impromised.

EPA Response: EPA's announcement in the April 17, 2002, Federal Register was only intended to discuss the Agency's intent to begin the process for revising the TCR. EPA will consider the commenters' suggestions as part of the revision process. As stated in the April 17, 2002, Federal Register, the Agency plans to consider revisions to the TCR with new requirements for ensuring the integrity of distribution systems. The Agency remains committed to obtaining input from stakeholders as part of the rule development process. EPA agrees with the comment that public health should not be compromised, and will consider only those revisions that will assure public health protection.

E. What Comments Did EPA Receive on Research Needs?

Commenters found that EPA's information on potential research resulting from the review of NPDWRs would be better represented by a summary of research needs that were identified by the Agency. Commenters felt that this summary is important to inform future regulatory decisions.

nmenters also suggested additional

identified by EPA in its preliminary review.

EPA Response: EPA agrees that the identification of research needs is an important component of the review of NPDWRs. Research findings may support future reviews and/or revisions to NPDWRs.

The Agency is considering research needs that it identified as part of the review as well as those suggested by commenters. EPA will continue to identify areas where data are lacking. Dialogue with industry and other groups, including those that sponsor or conduct research on priority areas, would be beneficial to the drinking water program. Collaboration in sponsoring studies can provide multiple benefits.

There are two research needs associated with the Six-Year Review that are being addressed through mechanisms external to EPA. The National Research Council of the National Academy of Sciences is conducting an assessment of recent data on fluoride health effects. In addition, the National Toxicology Program is conducting a study on chromium VI toxicity. Both of these research efforts are discussed in the April 17, 2002, Federal Register announcement of EPA's preliminary revise/not revise decisions. The current review identified several general and specific areas of potential research related to treatment. The treatment-related research areas are briefly discussed in the Treatment Feasibility Document (USEPA, 2003g).

EPA is currently in the process of examining whether specific research needs exist within each of the Six-Year Review areas of regulatory consideration (i.e., health effects, analytical methods, treatment, implementation, and occurrence/exposure). Some of the research needs identified during the Six-Year Review effort will be discussed in the context of the Multi-Year Plan (MYP) for drinking water. The MYP describes the EPA Office of Research and Development's fiscal year 2003 to 2010 research program to support the regulatory development activities of the EPA Office of Water. EPA plans to make this document available to the public in

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Dated: July 11, 2003.
Linda J. Fisher,
Acting Administrator.
[FR Doc. 03–18152 Filed 7–17–03; 8:45 am]
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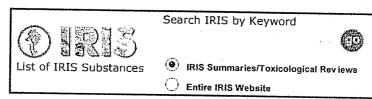
IRIS

IRIS Summaries

1,1-Dichloroethylene (CASRN 75-35-4)

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MAIN CONTENTS



Reference Dose for Chronic Oral Exposure (RfD)



You will need Adobe Reader to view some of the files on this page. See <u>EPA's PDF page</u> to learn more.

Note: A TOXICOLOGICAL REVIEW is available for this chemical in Adobe PDF Format (72 Pages, 531 Kbytes). Similar documents can be found in the List of Available IRIS Toxicological Reviews.

Links to specific pages in the toxicological review are available throughout this summary. To utilize this feature, your Web browser and Adobe program must be configured properly so the PDF displays within the browser window. If your browser and Adobe program need configuration, please go to EPA's PDF page for instructions.

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1,1-Dichloroethylene (1,1-DCE); CASRN 75-35-4; 08/13/02

Health assessment information on a chemical substance is included in IRIS only after a comprehensive review of chronic toxicity data by U.S. EPA health scientists from several Program Offices, Regional Offices, and the Office of Research and Development. The summaries presented in Sections I and II represent a consensus reached in the review process. Background information and explanations of the methods used to derive the values given in IRIS are provided in the Background Documents.

STATUS OF DATA FOR 1,1-Dichloroethylene (1,1-DCE)

File First On-Line 01/30/1987

Category (section)	Status	Last Revised
Oral RfD Assessment (I.A.)	on-line	08/13/2002*
Inhalation RfC Assessment (I.B.)	on-line	08/13/2002*
Carcinogenicity Assessment (II.)	on-line	08/13/2002*

^{*}A comprehensive review of toxicological studies was completed (05/27/05) - please see sections I.A.6., I.B.6., and II.D.2. for more

information.

_I. Chronic Health Hazard Assessments for Noncarcinogenic Effects

_I.A. Reference Dose for Chronic Oral Exposure (RfD)

Substance Name - 1,1-Dichloroethylene (1,1-DCE) CASRN - 75-35-4 Last Revised - 08/13/2002

The oral Reference Dose (RfD) is based on the assumption that thresholds exist for certain toxic effects such as cellular necrosis. It is expressed in units of mg/kg-day. In general, the RfD is an estimate (with uncertainty spanning perhaps an order of magnitude) of a daily exposure to the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a lifetime. Please refer to the Background Document for an elaboration of these concepts. RfDs can also be derived for the noncarcinogenic health effects of substances that are also carcinogens. Therefore, it is essential to refer to other sources of information concerning the carcinogenicity of this substance. If the U.S. EPA has evaluated this substance for potential human carcinogenicity, a summary of that evaluation will be contained in Section II of this file.

This summary replaces the summary dated 04/01/1989. This RfD differs from the previous EPA value of 0.009 mg/kg-day. The previous EPA evaluation used the same study but considered the lowest exposure of 9 mg/kg-day in female rats as a LOAEL for minimal hepatocellular fatty change and minimal hepatocellular swelling and applied a total uncertainty factor (UF) of 1000 (10 for LOAEL-to-NOAEL extrapolation, 10 for interspecies extrapolation, and 10 for human variability). EPA no longer considers hepatocellular swelling, in the absence of other effects such as increased liver enzymes in the serum, as biologically significant in this bioassay. The increased incidence of midzonal fatty change at 9 mg/kg-day in female rats is not statistically significant. The NOAEL in this bioassay is 9 mg/kg-day. In addition, the present evaluation uses benchmark dose (BMD) methodology and calculates a BMDL₁₀ for midzonal fatty change in female rats.

_I.A.1. Oral RfD Summary

Critical Effect	Experimental Doses*	UF	MF	RfD
Liver toxicity (fatty change)	NOAEL: 9 mg/kg-day LOAEL: 14 mg/kg-day			
Rat chronic drinking	THE MISSISSE CONTROL OF THE REAL PROPERTY OF THE PROPERTY OF T			
water study	BMDL ₁₀ : 4.6 mg/kg-day	100	1	5E-2
Quast et al. (1983)		•		mg/kg-day
The state of the s	and the state of t			

^{*}Conversion Factors and Assumptions — The authors provided the exposure data from the bioassay based on measured drinking water consumption.

__I.A.2. Principal and Supporting Studies (Oral RfD)

Quast et al. (1983) conducted a 2-year chronic toxicity and carcinogenicity study of 1,1-DCE in Sprague-Dawley rats (6–7 weeks old). The control group comprised 80 rats of each sex, and each exposed group comprised 48 rats of each sex. The 1,1-DCE was incorporated in the drinking water of the rats at nominal concentrations of 0, 50, 100, or 200 ppm. The time-weighted average exposure

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over the 2-year period was 7, 10, or 20 mg/kg-day for males and 9, 14, or 30 mg/kg-day for females. Rampy et al. (1977) also reported some of the data. Humiston et al. (1978) reported more detailed data. No significant differences were observed among the groups in appearance and demeanor, mortality, body weight, food consumption, water consumption, hematology, urinalysis, clinical chemistry determinations, organ weights, or organ to body weight ratios. After 1 year on study, there was no depletion of the nonprotein sulfhydryl levels in the liver or the kidneys (Rampy et al., 1977).

The only treatment-related effect observed in rats was minimal hepatocellular midzonal fatty change and hepatocellular swelling. At the termination of the study, male rats showed increased incidence of minimal hepatocellular fatty change (control, 14/80; 50 ppm, 5/48; 100 ppm, 13/48; 200 ppm, 19/47) and minimal hepatocellular swelling (control, 0/80; 50 ppm, 1/48; 100 ppm, 2/48; 200 ppm, 3/47). The changes were statistically significant (p<0.05) only in the 200 ppm group. At the termination of the study, female rats showed an increased incidence of minimal hepatocellular fatty change (control, 10/80; 50 ppm, 12/48; 100 ppm, 14/48; 200 ppm, 22/48; statistically significant [p<0.05] at 100 and 200 ppm) and minimal hepatocellular swelling (control, 3/80; 50 ppm, 7/48; 100 ppm, 11/48; 200 ppm, 20/48; statistically significant [p<0.05] in all groups). No exposure-related neoplastic changes occurred at any exposure. No hepatocellular necrosis was evident at any exposure. Based on the minimal nature of the hepatocellular swelling reported by the authors and no change in liver weight, no change in clinical chemistry measurements diagnostic for liver damage, and no other indication of abnormal liver function, the hepatocellular swelling is not considered biologically significant or an adverse effect in this study. The statistically significant hepatocellular midzonal fatty change, however, is considered a minimal adverse effect in this study. Accordingly, the NOAEL in male rats is 10 mg/kg-day and the LOAEL is 20 mg/kg-day; the NOAEL in female rats is 9 mg/kg-day and the LOAEL is 14 mg/kg-day. A BMD analysis was conducted for the results in female rats. In female rats, the ${\rm BMD}_{10}$ is 6.6 mg/kg-day and the ${\rm BMDL}_{10}$ is 4.6 mg/kg-day.

A three-generation study by Nitschke et al. (1983), described in Section I.A.4, corroborated the results of Quast et al. (1983).

The National Toxicology Program conducted 104-week chronic toxicity and carcinogenicity studies of 1,1-DCE in male and female F344 rats (200 of each sex, 9 weeks old) by gavage in corn oil at 0, 1, or 5 mg/kg-day (NTP, 1982). There were no significant differences in survival, clinical signs, or body weight as compared with controls for any group, suggesting that the maximum tolerated dose was not achieved. The results of histopathological examination indicated chronic renal inflammation in male rats (26/50, 24/48, 43/48) and female rats (3/49, 6/49, 9/44). The increase was statistically significant only in males. As this lesion commonly occurs in male rats (Kluwe et al., 1984, 1990), it is not considered biologically significant in this study. The NOAEL in this study 5 is mg/kg-day (the highest exposure tested).

NTP also conducted 104-week chronic toxicity and carcinogenicity studies of 1,1-DCE in male and female B6C3F $_1$ mice (50 of each sex in each group, 9 weeks old) by gavage in corn oil at 0, 2, or 10 mg/kg (NTP, 1982). There were no significant differences in survival, clinical signs, or body weight in any group. The only noncancer effect observed by histopathological examination was necrosis of the liver (male: 1/46; 3/46; 7/49; female: 0/47; 4/49; 1/49). The effect was not statistically significant at either exposure (p=0.6 and 0.06 at the mid- and high-exposure levels in males using a two-tailed test, respectively). In male and female mice the NOAEL is 10 mg/kg-day (the highest exposure tested). The BMD $_{10}$ is 7.8 mg/kg-day and the BMDL $_{10}$ is 4.1 mg/kg-day. This study was not used to derive the RfD because the gavage route of exposure will affect the pharmacokinetics of 1,1-DCE and the exposure-response relationship.

__I.A.3. Uncertainty and Modifying Factors (Oral RfD)

UF - 100

The critical effect is liver toxicity (fatty change) in rats, with a BMDL $_{10}$ of 4.6 mg/kg-day. Although this minimal effect might not be considered adverse—as there is no evidence of a functional change in the liver in rats exposed and glutathione levels are not reduced in this bioassay—the ${\rm BMDL}_{10}$ was used to derive the RfD, because limiting exposure to the BMDL_{10} will protect the liver from more serious damage (fatty liver or necrosis) that could compromise liver function. Individual UFs of 10 each were used for interspecies extrapolation and intraspecies variability because there were no applicable data to justify departure from the default values. Derivation of the RfD from the BMDL_{10} for the minimal fatty change in the liver does not require an effect-level extrapolation. This conclusion is based on the minimal nature of the fatty change and its questionable biological significance because of the absence of any observable functional deficit in the liver. A subchronic-to-chronic extrapolation factor was not applied because the study exposed the animals for 2 years. A database UF is not applied because the database is considered complete. A number of long-term bioassays in rodents by the oral or inhalation route show that liver toxicity is the critical effect. There is no chronic bioassay in a nonrodent mammal. However, there are 90-day bioassays in several species (rats, mice, dogs, guinea pigs, rabbits, and monkeys) that suggest similar exposure-response relationships across species. Therefore, the lack of a chronic bioassay in a nonrodent mammal is not considered a data gap. There are no focused studies on neurotoxicity, but there is no indication from chronic, reproductive, and developmental bioassays in rats and mice by oral or inhalation exposure that neurotoxicity in an important toxic endpoint. No long-term studies have evaluated immunotoxicity in laboratory animals by any route of exposure. The existing bioassays, however, provide no suggestion that immunotoxicity is a critical effect. EPA does not consider these data gaps compelling enough to require application of a database UF.

MF = 1.

_I.A.4. Additional Studies/Comments (Oral RfD)

NTP (1982) conducted a study in male and female F344 rats (10 of each sex, 9 weeks old) administered 1,1-DCE by gavage in corn oil at 0, 5, 15, 40, 100, or 250 mg/kg. Animals were exposed five times per week for 13 weeks. Representative tissues from animals receiving 250 mg/kg and from control animals were examined microscopically. Livers from all groups were examined. Three female rats receiving 250 mg/kg died during the first week of the study. No other rats died. The mean body weight was depressed 13% for male rats receiving 250 mg/kg as compared with controls. Mean body weight in other groups was comparable. Only the liver showed effects attributed to 1,1-DCE. At 250 mg/kg, the three female rats that died showed severe centrilobular necrosis. Minimal to moderate hepatocytomegaly was seen in the rest of the rats at 250 mg/kg. Minimal to mild hepatocytomegaly was seen in 6/10 male rats and 3/10 female rats that received 100 mg/kg. No biologically significant changes were observed in rats that received 40 mg/kg or less. The NOAEL in this study is 40 mg/kg (equivalent to 28.5 mg/kg-day); the LOAEL is 100 mg/kg (equivalent to 71.4 mg/kg-day).

NTP (1982) conducted a study in male and female B6C3F $_1$ mice (10 of each sex, 9 weeks old) administered 1,1-DCE by gavage in corn oil at 0, 5, 15, 40, 100, or 250 mg/kg. Animals were exposed five times per week for 13 weeks. Representative tissues from mice receiving 100 and 250 mg/kg and from control animals were examined microscopically. Livers from all groups were also examined. Survival was 20/20, 19/20, 19/20, 19/20, 15/20, and 1/20 at 0, 5, 15, 40, 100, and 250 mg/kg, respectively. At 100 mg/kg there was a decrease in mean body weight in males (14%) but not in females. No change in mean body weight was observed at lower exposures. Only the liver showed effects attributed to 1,1-DCE. Centrilobular necrosis of the liver was observed in 5/10 males and 5/10 females that received 250 mg/kg and 2/10 males and 2/10 females that received 100 mg/kg. No biologically significant changes in the liver occurred in mice receiving 40 mg/kg or less. The NOAEL in this study is 40 mg/kg (adjusted to a continuous daily exposure of 28.6 mg/kg-day); the LOAEL is 100 mg/kg (adjusted to a continuous daily exposure of 71.4 mg/kg-day).

Quast et al. (1983) conducted a study in beagle dogs (four per group, 8 months old) administered 1,1-DCE by gavage in peanut oil at 0, 6.25, 12.5, or 25 mg/kg-day for 97 days. No significant differences were observed among groups in appearance and demeanor, mortality, body weight, food consumption, hematology, urinalysis, clinical chemistry determinations, organ weights, and organ-to-body-weight ratios. No exposure-related gross or histopathological changes were present in tissues. There was no depletion of the nonprotein sulfhydryl levels in the liver or kidneys. The NOAEL in this study is 25 mg/kg-day (the highest exposure tested).

Reproductive and Developmental Studies

Nitschke et al. (1983) evaluated the reproductive and developmental toxicity of 1,1-DCE in Sprague-Dawley rats. Three generations of the test animals were exposed to drinking water containing nominal 1,1-DCE concentrations of 0 (initially 15 males and 30 females), 50, 100, or 200 ppm (initially 10 males and 20 females at each exposure). The authors provided no information on water consumption. This study was a companion study to Quast et al. (1983) and used the same concentrations of 1,1-DCE in drinking water; in Quast et al. (1983) the average exposure to females was 9, 14, or 30 mg/kg-day. After 100 days of exposure, the rats were mated.

In this three-generation study, there were no biologically significant changes in fertility index, in average number of pups per litter, in average body weight of pups, or in pup survival at any exposure. Neonatal survival was decreased from concurrent control values in the f_2 and f_{3a} litters of dams ingesting 1,1-DCE from drinking water. The survival indices, however, were within the range of control values for this strain of rats in this laboratory. The authors attributed the decreased survival index in f_2 to increased litter size at birth in dams exposed to 1,1-DCE. The apparent effect seen in the f_{3a} litters was not repeated in subsequent matings of the same adults to produce either the f_{3b} or the f_{3c} litters. The authors attributed the decreased survival in the f_{3a} litters as being due to chance.

Histopathological examination of tissues of rats exposed to 1,1-DCE in the drinking water in utero, during lactation, and postweaning revealed slight hepatocellular fatty change and an accentuated hepatic lobular pattern of a reversible nature in the adult rats (data not reported, but the observation is consistent with that reported by Quast et al. [1983] in a chronic bioassay). These effects were observed in the 100 and 200 ppm groups in the F_1 generation and in all groups of the F_2 generation. The authors did not present incidence data and did not report statistical analysis. Exposure to 1,1-DCE in drinking water at concentrations causing mild, dose-related changes in the liver did not affect the reproductive capacity of rats through three generations that produced six sets of litters. The NOAEL for reproductive and developmental toxicity in this study is 200 ppm for exposure to 1,1-DCE in drinking water (the highest exposure tested and about 30 mg/kg-day).

Murray et al. (1979) evaluated the developmental toxicity of 1,1-DCE administered in drinking water at 0 (27 animals) or 200 ppm (26 animals) to pregnant Sprague-Dawley rats (body weight 250 g). Rats were exposed on gestation days 6–15 at 40 mg/kg-day. No teratogenic effects were seen in the embryos using standard techniques for soft and hard tissue examination, and there was no evidence of toxicity to the dams or their offspring. The NOAEL for developmental toxicity in this study is 40 mg/kg-day (the highest exposure tested).

Dawson et al. (1993) evaluated the ability of 1,1-DCE administered in drinking water at 110 ppm or 0.15 ppm to female Sprague-Dawley rats (body weight 250 g) to induce fetal cardiac changes. Rats were administered 110 ppm 1,1-DCE for 61 days before mating or for 48 days before mating and for 20 days during gestation. Other rats were administered 0.15 ppm 1,1-DCE for 82 days before mating or for 56 days before mating and for 20 days during gestation. The dams were killed on gestational day 22 and the gravid uterus was removed and examined. There was no effect on maternal weight gain, average resorption sites (sites where development began but resorption later occurred), or average implantation sites (sites that did not appear to develop beyond implantation and contained a

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metrial gland only). No increase in the incidence of cardiac changes occurred when dams were exposed only before mating. There was, however, a statistically significant increase (p<0.01) in the percent of fetuses with cardiac changes (atrial septal, mitral valve, and aortic valve changes) when the dams were exposed before mating and during gestation. The incidence was control, 7/232 (3%); 0.15 ppm, 14/121 (12%); and 110 ppm, 24/184 (13%).

This statistical analysis was based on total occurrence of affected fetuses. Because the exposure was to the dam and not to individual fetuses, a nested statistical analysis is preferred. Such an analysis takes into account the correlation among fetuses within a litter and the possible nesting of effects within litters. This analysis has not been conducted because all the necessary data are not available. The author provided additional data to resolve typographical errors in the exposure information for each group and to clarify the number of affected litters and number of fetuses per litter affected (letter from Brenda Dawson, University of Auckland, New Zealand, to Robert Benson, U.S. EPA, January 24, 2001). The exposure to dams before and during pregnancy was 0, 0.02, or 18 mg/kg-day in the control, 0.15 ppm, and 110 ppm groups, respectively. The number of affected litters was 5/21 (24%), 8/11 (73%), and 13/17 (76%). The mean number of affected fetuses per litter for affected litters only was 1.40 (13% of the fetuses in the litter), 1.75 (16% of the fetuses in the litter), and 1.85 (17% of the fetuses in the litter). The mean number of affected fetuses per litter for all litters was 0.33 (3% of the fetuses in the litter), 1.27 (12% of the fetuses in the litter), and 1.41 (13% of the fetuses in the litter).

Dawson et al. (1993) did a much more thorough evaluation of alterations in cardiac development than is done in standard developmental toxicity testing protocols. There is no experience with the background rates or the functional significance of such alterations from other studies or laboratories. The incidence of alterations in control fetuses (3% of all fetuses, 24% of all litters, and 1.40 affected fetuses per affected litter) suggests a high background incidence. The authors reported that examinations were done blind to the treatment group, so the data are presumed not to be affected by observer bias.

There is no demonstrated exposure-response relationship in Dawson et al. (1993). A 900-fold increase in exposure did not produce a significant, increase in response in any measure of effect. The cardiac changes are of questionable biological significance, as there were no biologically significant effects reported on growth and survival in the three-generation study (Nitschke et al., 1983). No cardiac effects were reported in a prenatal developmental study (Murray et al., 1979); however, in this study exposure to 1,1-DCE did not occur throughout pregnancy. The pharmacokinetics of 1,1-DCE make it biologically implausible that the cardiac changes were causally associated with exposure to 1,1-DCE. The exposures used in Dawson et al. (1993) are below the level of saturation of CYP2E1 in the rat liver. Essentially all of the 1,1-DCE administered to the dams will be metabolized in the liver and will react with glutathione or macromolecules in the liver. (See the discussion and references in Section 3.) Therefore, it is extremely unlikely that any significant amount of 1,1-DCE or any toxic metabolite will be in the fetal compartment. CYP2E1 is not expressed in fetal liver but begins to be expressed shortly after birth (Cresteil, 1998).

EPA is not aware of any information on the expression of CYP2E1 in fetal cardiac tissue. Cardiac tissue, however, is not generally considered to be a tissue with significant potential for metabolism of xenobiotics. For these reasons EPA cannot conclude that the cardiac changes are caused by exposure to 1,1-DCE. It would be helpful if more definitive studies with a greater range of exposures were conducted to determine the cause and biological significance of the cardiac changes apparently associated with exposure to 1,1-DCE during the period of cardiac organogenesis.

For more detail on Susceptible Populations, exit to the toxicological review, Section 4.7 (PDF).

__I.A.5. Confidence in the Oral RfD

Study — High Database — Medium RfD — Medium

The overall confidence in this RfD assessment is medium. The principal study (Quast, 1983) was well conducted, with an adequate number of animals and appropriate evaluation of a wide variety of endpoints. This study is supported by an additional bioassay in rats (NTP, 1982) and a three-generation reproductive and developmental study showing consistent effects in the liver. A three-generation reproductive study and several bioassays show that reproductive and developmental toxicity are not critical effects. One developmental study, however, shows variations in cardiac morphology that have appear to have little or no physiological consequence. There are no focused studies on neurotoxicity, but there are no indications from chronic, reproductive, or developmental bioassays in rats and mice by oral or inhalation exposure that neurotoxicity is an important toxic endpoint. No long-term studies have evaluated immunotoxicity in laboratory animals by any route of exposure. The existing bioassays, however, provide no suggestion that immunotoxicity is a critical effect. Accordingly, the database is given a medium confidence, but no additional UF is considered necessary.

For more detail on Characterization of Hazard and Dose Response, exit to the toxicological review, Section 6 (PDF).

__I.A.6. EPA Documentation and Review of the Oral RfD

Source Document — Toxicological Review of 1,1-Dichloroethylene (2002)

This assessment was peer reviewed by external scientists. Their comments have been evaluated carefully and incorporated in finalization of this IRIS Summary. A record of these comments is included as an appendix to the Toxicological Review of 1,1-Dichloroethylene. <u>To review this appendix, exit</u> to the toxicological review, <u>Appendix A</u>, <u>Summary of and Response to External Peer Review Comments (PDF)</u>.

Other EPA Documentation — This assessment replaces previous assessments (U.S. EPA, 1985a,b).

Agency Consensus Date — 06/07/2002

A comprehensive review of toxicological studies published through May 2005 was conducted. No new health effects data were identified that would be directly useful in the revision of the existing RfD for 1,1-Dichloroethylene (1,1-DCE) and a change in the RfD is not warranted at this time. For more information, IRIS users may contact the IRIS Hotline at hotline.iris@epa.gov or 202-566-1676.

__I.A.7. EPA Contacts (Oral RfD)

Please contact the IRIS Hotline for all questions concerning this assessment or IRIS, in general, at (202)566-1676 (phone), (202)566-1749 (FAX) or hotline.iris@epa.gov (internet address).

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_I.B. Reference Concentration for Chronic Inhalation Exposure (RfC)

Substance Name -1,1-Dichloroethylene (1,1-DCE)

CASRN — 75-35-4 Last Revised — 08/13/2002

The inhalation Reference Concentration (RfC) is analogous to the oral RfD and is likewise based on the assumption that thresholds exist for certain toxic effects such as cellular necrosis. The inhalation RfC considers toxic effects for both the respiratory system (portal-of-entry) and for effects peripheral to the respiratory system (extrarespiratory effects). It is generally expressed in units of mg/cu.m. In general, the RfC is an estimate (with uncertainty spanning perhaps an order of magnitude) of a daily inhalation exposure of the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a lifetime. Inhalation RfCs were derived according to the Interim Methods for Development of Inhalation Reference Doses (EPA/600/8-88/066F August 1989) and subsequently, according to Methods for Derivation of Inhalation Reference Concentrations and Application of Inhalation Dosimetry (EPA/600/8-90/066F October 1994). RfCs can also be derived for the noncarcinogenic health effects of substances that are carcinogens. Therefore, it is essential to refer to other sources of information concerning the carcinogenicity of this substance. If the U.S. EPA has evaluated this substance for potential human carcinogenicity, a summary of that evaluation will be contained in Section II of this file.

The previous EPA evaluation did not derive an RfC.

__I.B.1. Inhalation RfC Summary

Critical Effect	Experimental Doses*	UF	MF	RfC
Liver toxicity (fatty change)	NOAEL _{HEC} : 17.7 mg/m ³ LOAEL _{HEC} : 53.2 mg/m ³			•
Rat chronic inhalation study Quast et al. (1986)	BMCL _{10HEC} : 6.9 mg/m ³	30	1	2E-1 mg/m ³

*Conversion Factors and Assumptions — The NOAEL from the chronic bioassay is 25 ppm, where the exposure was for 6 hrs/day, 5 days/wk. The conversion factor is 1 ppm = 3.97 mg/m³. The human equivalent concentration (HEC) was calculated using the equation for a category 3 gas (U.S. EPA, 1994). The blood:gas partition coefficient in the rat is 5 (D'Souza and Andersen, 1988). No useable data are available on the blood:gas coefficient in humans. Accordingly the default value of 1 is used for the ratio of

NOAEL $_{\rm HEC}$ = NOAEL $_{\rm adj}$ × $(H_{\rm b/g})_{\rm A}/(H_{\rm b/g})_{\rm H}$ = 25 ppm × 6/24 × 5/7 × 1 × 3.97 = 17.7 mg/m 3 BMCL $_{\rm HEC}$ = BMCL $_{\rm adj}$ × $(H_{\rm b/g})_{\rm A}/(H_{\rm b/g})_{\rm H}$ = 9.8 ppm × 6/24 × 5/7 × 1 × 3.97 = 6.9 mg/m 3

__I.B.2. Principal and Supporting Studies (Inhalation RfC)

Quast et al. (1986) and Rampy et al. (1977) reported results from studies that exposed male and female Sprague-Dawley rats (Spartan substrain, 86 animals/group) to 1,1-DCE by inhalation 6 hrs/day, 5 days/wk, for up to 18 months. Interim sacrifices occurred at 1, 6, and 12 months. Rats were exposed to 1,1-DCE concentrations of 10 ppm and 40 ppm for the first 5 weeks of the study. Because of the absence of observable treatment-related effects among rats sacrificed after 1 month of exposure, the concentrations were increased to 25 and 75 ppm. Exposures were continued at these concentrations through the 18th month of the study. The surviving animals were then held without exposure to 1,1-DCE until 24 months. Cytogenetic evaluations were performed on a separate group of animals (four/sex) exposed to 0, 25, or 75 ppm for 6 months. A separate 90-day study using 20 rats/sex/treatment group was conducted at 0, 25, and 75 ppm, with an interim sacrifice of 8 rats/group at 30 days. There were no exposure-related changes in mortality, appearance and demeanor, body weight, clinical chemistry determinations, hematologic evaluations, urinalysis, or

cytogenetic evaluation of bone marrow preparations.

Minimal hepatocellular fatty change in the midzonal region of the hepatic lobule was observed in both male and female rats in the 25 ppm and 75 ppm groups at the 6-month interim sacrifice (male: control, 0/5; 25 ppm, 1/5; 75 ppm, 4/5; female: control, 0/5; 25 ppm, 2/5; 75 ppm, 4/5). The fatty change was also observed at the 12-month sacrifice, but there was no indication of progression of severity (male: control, 0/5; 25 ppm, 3/5; 75 ppm, 5/5; female: control, 0/5; 25 ppm, 5/5; 75 ppm, 5/5). At the 18-month sacrifice the incidence of this change was no longer increased in male rats (control, 0/27; 25 ppm, 0/25; 75 ppm, 1/27). However, the change persisted in female rats (control, 0/16; 25 ppm, 6/29; 75 ppm, 7/20). The effect was statistically significant (p<0.05) only at the higher exposure. During the last 6 months of the study, after exposure had been discontinued, this effect was no longer discernible (male: control, 0/46; 25 ppm, 1/47; 75 ppm, 0/51; female: control, 0/49; 25 ppm, 0/46; 75 ppm, 1/48).

Although the incidences of several tumors and/or tumor types were found to be statistically increased or decreased compared with controls, none of these differences were judged to be attributable to 1,1-DCE. The tumor incidence data for both control and treated rats in this study were comparable to historical control data for the Sprague-Dawley rats (Spartan substrain) used by this laboratory for several studies of similar design and duration.

Although the minimal hepatocellular midzonal fatty change was reversible and did not result in altered organ weight, clinical chemistry changes diagnostic for liver damage, or any obvious decrement in liver function, the fatty change in liver is considered a minimal adverse effect. Accordingly, the NOAEL in male rats in this study is 75 ppm (the highest exposure tested). The NOAEL for female rats in this study is 25 ppm; the LOAEL is 75 ppm. A benchmark dose analysis was conducted. In female rats the BMC $_{10}$ is 15.1 ppm and the BMCL $_{10}$ is 9.8 ppm, equivalent to 1.8 ppm adjusted for continuous exposure (9.8 ppm × 6/24 × 5/7).

__I.B.3. Uncertainty and Modifying Factors (Inhalation RfC)

UF = 30.

The critical effect is liver toxicity (fatty change) in rats with a BMCL_{10HEC} of 6.9 mg/m³. Although this minimal effect might not be considered adverse—as there is no evidence of a functional change in the liver in rats exposed at this level and glutathione levels are not reduced—it is used to derive the RfC, because limiting exposure to this level will protect the liver from more serious damage (fatty liver or necrosis) that could compromise liver function. The total UF is 30 and the modifying factor is 1. A UF of 3 is used for interspecies extrapolation because a dosimetric adjustment was used.

There is some suggestion that the effects in the kidney of male mice might occur at an exposure lower than the level that produced effects in the liver of rats. Thus, there is some uncertainty as to whether the most sensitive species has been used to derive the RfC. A UF of 10 is used for intraspecies variability because there were no applicable data to depart from the default value. Derivation of the RfD from the BMDL₁₀ for the minimum fatty change in the liver does not require an effect-level extrapolation. This conclusion is based on the minimal nature of the fatty change and its questionable biological significance because of the absence of any observable functional deficit in the liver. Although the animals were exposed for 18 months, rather than the full lifetime, there was no indication that the fatty change was progressing. In contrast, the evidence indicated the fatty change was decreasing in incidence with continued exposure. EPA, therefore, did not apply a subchronic-to-chronic extrapolation factor. A database UF is not applied because the database is considered complete.

A number of long-term bioassays in rodents by the oral or inhalation route show that liver toxicity is the critical effect. There is no chronic bioassay in a nonrodent mammal. However, there are 90-day bioassays in several species (rats, mice, dogs, guinea pigs, rabbits, and monkeys) that suggest similar

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exposure-response relationships across species. Therefore, the lack of a chronic bioassay in a nonrodent mammal is not considered a data gap. There are no focused studies on neurotoxicity, but there are no indications from chronic, reproductive, and developmental bioassays in rats and mice by oral or inhalation exposure that neurotoxicity is an important toxic endpoint. No long-term studies have evaluated immunotoxicity in laboratory animals by any route of exposure. The existing bioassays, however, provide no suggestion that immunotoxicity is a critical effect. EPA does not consider these data gaps compelling enough to require application of a database UF.

MF = 1.

__I.B.4. Additional Studies/Comments (Inhalation RfC)

Prendergast et al. (1967) evaluated the toxicity of 1,1-DCE in Long-Evans or Sprague-Dawley rats, Hartley guinea pigs, beagle dogs, New Zealand albino rabbits, and squirrel monkeys. The test animals (15 rats/group, 15 guinea pigs/group, 3 rabbits/group, 2 dogs/group, or 3 or 9 monkeys/group) were exposed continuously for 90 days to 1,1-DCE vapors at 189 ± 6.2 , 101 ± 4.4 , 61 ± 5.7 , or 20 ± 2.1 mg/m³. The concurrent controls included 304 rats, 314 guinea pigs, 48 rabbits, 34 dogs, and 57 monkeys. The age of the animals was not specified. The exposed animals were evaluated for visible signs of toxicity, mortality, and hematologic, biochemical, pathologic, and body weight changes. There was apparent exposure-related mortality in guinea pigs and monkeys. In guinea pigs the mortality was 2/314, 2/45, 3/15, 3/15, and 7/15 and in monkeys it was 1/57, 1/21, 0/9, 2/3, and 3/9 in the 0, 20, 61, 101, or 189 mg/m³ exposure groups, respectively. The guinea pigs died between days 3 to 9 of exposure; the monkeys died on days 26, 39, 47, 60, and 64 of exposure. There were no visible signs of toxicity in any surviving animals.

At the highest exposure in monkeys, but not in guinea pigs, there was some histopathological evidence of liver damage (see below). In guinea pigs at the highest exposure, there was an increase in serum glutamic-pyruvic transaminase and liver alkaline transaminase (see below). Because visible signs of toxicity were not observed and only minor liver damage is apparent in this study, the mortality data in guinea pigs and monkeys are given no weight.

Varying degrees of growth depression were found in all exposures, but were significant in all species only at 189 mg/m³. The test animals exhibited no significant hematologic alterations, and serum urea nitrogen levels were within control limits in all exposures in which determinations were made. Significant elevations of serum glutamic-pyruvic transaminase and liver alkaline phosphatase activities were found in rats (a threefold and 1.75-fold increase, respectively) and guinea pigs (sevenfold and 2.4-fold increase, respectively) exposed to 189 mg/m³ (other species not tested) but not at 20 mg/m³ (enzyme levels at intermediate exposures not tested). Histopathological examination of liver from dogs, monkeys, and rats revealed damage at 189 mg/m³ (other species not examined). The effects observed included fatty metamorphosis, focal necrosis, hemosiderosis deposition, lymphocytic infiltration, bile duct proliferation, and fibrosis. The changes were most severe in dogs. Sections of kidney from all rats showed nuclear hypertrophy of the tubular epithelium. No detectable liver or kidney damage was observed in any species exposed to 101 mg/m³ or less. The NOAEL in this study is 101 mg/m³ (equivalent to 25 ppm); the LOAEL is 189 mg/m³ (equivalent to 47 ppm).

Short et al. (1977) evaluated developmental toxicity of 1,1-DCE administered by inhalation to pregnant CD-1 rats (Charles River). Animals were exposed to 0 (58 animals), 15 ppm (18 animals), 57 ppm (20 animals), 300 ppm (18 animals), or 449 ppm (18 animals) for 22-23 hours/day on gestation days 6 to 16. Dams were sacrificed on gestation day 20. Maternal toxicity was exhibited as severe maternal weight loss (> 28 grams/dam) at 15 ppm and higher and by maternal mortality at 57 ppm and higher. There was a statistically significant increase in the mean number of fetuses per litter with hydrocephalus at 15 and 57 ppm, with malaligned sternebrae at 15 ppm, and with unossified sternebrae at 57 ppm. Because of the severe maternal toxicity at 15 ppm (60 mg/m³) and higher, this study is not useful for evaluating developmental toxicity.

Short et al. (1977) evaluated developmental toxicity of 1,1-DCE administered by inhalation to pregnant CD-1 mice (Charles River). Animals were exposed to 0 (65 animals), 15 ppm (23 animals), 30 ppm (19 animals), 57 ppm (21 animals), 144 ppm (18 animals), or 300 ppm (15 animals) for 22–23 hrs/day on gestation days 6 to 16. Dams were sacrificed on gestation day 17. At 30 ppm and higher there was maternal toxicity, as shown by statistically significant decreases in maternal weight gain. At 144 and 300 ppm there was an increase in maternal mortality. At 30 ppm and higher there was severe fetal toxicity, with complete early resorption of the litters. At 15 ppm there was no evidence of maternal toxicity, no decrease in fetal body weight, and no decrease in the percentage of viable fetuses. At 15 ppm, there was an increase in the mean number of fetuses per litter with hydrocephalus, occluded nasal passages, micropthalmia, cleft palate, small liver, and hydronephrosis. None of these changes, however, were statistically significant when compared to controls. Also at 15 ppm there was a statistically significant increase in the mean number of fetuses with an unossified incus and with incompletely ossified sternebrae. This study provides evidence of fetal toxicity at 15 ppm, the only exposure without significant maternal toxicity. In this study the LOAEL for developmental toxicity is 15 ppm (60 mg/m³), the lowest exposure tested.

Short et al. (1977) also evaluated developmental neurotoxicity of 1,1-DCE administered by inhalation to CD-1 rats (Charles River). Pregnant rats were exposed to 0 (24 animals), 56 ppm (20 animals), or 283 ppm (19 animals) for 22–23 hrs/day on gestation days 8 to 20. Maternal toxicity was observed at both exposures, as shown by weight loss of 7 g per dam at 56 ppm and 15 grams per dam at 283 ppm. There was complete resorption of three litters at 283 ppm. There was a statistically significant decrease in average pup weight as compared to control at both exposures on post-natal day 1. The difference in pup weight between control and exposed groups decreased with time and disappeared by postnatal day 21. There was no evidence of developmental neurotoxicity at either exposure in pups tested at various times from postnatal day 1 to day 21 in a battery of behavioral tasks, including surface righting, pivoting, auditory startle, bar holding, righting in air, visual placing, swimming ability, physical maturation, and activity. This study shows evidence of maternal and fetal toxicity at both exposures but no evidence of developmental neurotoxicity at either exposure. Accordingly, the NOAEL for developmental neurotoxicity in this study is 283 ppm (1124 mg/m³), the highest exposure tested.

Murray et al. (1979) evaluated developmental toxicity of 1,1-DCE administered by inhalation to pregnant Sprague-Dawley rats (body weight 250 g). Animals were exposed to 0 (20 or 47 animals), 20 ppm (44 animals), 80 ppm (30 animals), or 160 ppm (30 animals) for 7 hrs/day on gestation days 6–15. At 20 ppm there was no maternal toxicity and no effect on embryonal or fetal development. At 80 and 160 ppm, there was toxicity to the dams (statistically significant depression in weight gain at gestation day 6–9, more severe at 160 ppm). At 80 and 160 ppm, there were also statistically significant increased incidences of wavy ribs and delayed ossification of the skull, which are regarded as embryotoxic effects. Both effects were more severe at 160 ppm. No teratogenic effects were seen at any exposure. The NOAEL for developmental toxicity in this study is 20 ppm; the LOAEL is 80 ppm. Under the Guidelines for Developmental Toxicity (U.S. EPA, 1994), these values are not adjusted to continuous exposure.

Murray et al. (1979) evaluated the developmental toxicity of 1,1-DCE administered by inhalation to New Zealand white rabbits (body weight 3.4–4.7 kg). Animals were exposed to 0 (16 animals), 80 ppm (22 animals), or 160 ppm (18 animals) for 7 hrs/day on gestation days 6–18. At 80 ppm there was no maternal toxicity and no effect on embryonal or fetal development. Toxicity to both the dams and their developing embryos was observed at 160 ppm. There was a marked increase in the incidence of resorptions per litter (0.3 ± 0.6 vs. 2.7 ± 3.9). A significant change occurred in the incidence of several minor skeletal variations in their offspring, including an increase in the occurrence of 13 pairs of ribs and a increased incidence of delayed ossification of the fifth sternebra (data not reported). No teratogenic effects were seen at any exposure. The NOAEL for developmental toxicity in this study is 80 ppm; the LOAEL is 160 ppm. Under the Guidelines for Developmental Toxicity (U.S. EPA, 1991), these values are not adjusted to continuous exposure.

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See also studies showing liver toxicity and the reproductive and developmental studies summarized in the RfD section.

For more detail on Susceptible Populations, exit to the toxicological review, Section 4.7 (PDF).

__I.B.5. Confidence in the Inhalation RfC

Study — High Database — Medium RfC — Medium

The overall confidence in this RfC assessment is medium. The principal study (Quast, 1986) was a well-conducted inhalation bioassay with adequate numbers of animals and appropriate evaluation of a wide variety of endpoints. The result is supported by several other 90-day inhalation studies in a variety of species (Prendergast et al. 1967). These inhalation studies are supported by an additional bioassay in rats and a 90-day study in dogs, both by the oral route of exposure showing NOAELs (see the summary of these studies in the RfD section). There is no evidence from the inhalation bioassays that the respiratory tract is a target tissue of low-dose exposure. Several studies by the inhalation route of exposure show that developmental toxicity is not a critical effect. A three-generation reproductive study by the oral route of exposure showed no significant reproductive effects, and several bioassays showed no developmental toxicity. However, one developmental study by the oral route of exposure shows variations in cardiac morphology that appear to have little or no physiological consequence. There are no focused studies on neurotoxicity, but no indication from chronic, reproductive, and developmental bioassays in rats and mice by oral or inhalation exposure that neurotoxicity is an important toxic endpoint. No long-term studies have evaluated immunotoxicity in laboratory animals by any route of exposure. The existing bioassays, however, provide no suggestion that immunotoxicity is a critical effect. Accordingly, the database is given medium confidence, but no additional UF is considered necessary.

For more detail on Characterization of Hazard and Dose Response, exit to the toxicological review, Section 6 (PDF).

___I.B.6. EPA Documentation and Review of the Inhalation RfC

Source Document — Toxicological Review of 1,1-Dichloroethylene (2002)

This assessment was peer reviewed by external scientists. Their comments have been evaluated carefully and incorporated in finalization of this IRIS Summary. A record of these comments is included as an appendix to the Toxicological Review of 1,1-Dichloroethylene. To review this appendix, exit to the toxicological review, Appendix A, Summary of and Response to External Peer Review Comments (PDF).

Other EPA Documentation — None.

Agency Consensus Date — 06/07/2002

A comprehensive review of toxicological studies published through May 2005 was conducted. No new health effects data were identified that would be directly useful in the revision of the existing RfC for 1,1-Dichloroethylene (1,1-DCE) and a change in the RfC is not warranted at this time. For more information, IRIS users may contact the IRIS Hotline at hotline.iris@epa.gov or 202-566-1676.

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__I.B.7. EPA Contacts (Inhalation RfC)

Please contact the IRIS Hotline for all questions concerning this assessment or IRIS, in general, at (202)566-1676 (phone), (202)566-1749 (FAX), or hotline.iris@epa.gov (email address).

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_II. Carcinogenicity Assessment for Lifetime Exposure

Substance Name - 1,1-Dichloroethylene (1,1-DCE) CASRN - 75-35-4 Last Revised - 08/13/2002

Section II provides information on three aspects of the carcinogenic assessment for the substance in question: the weight-of-evidence judgment of the likelihood that the substance is a human carcinogen and quantitative estimates of risk from oral exposure and from inhalation exposure. The quantitative risk estimates are presented in three ways. The slope factor is the result of application of a low-dose extrapolation procedure and is presented as the risk per (mg/kg)/day. The unit risk is the quantitative estimate in terms of either risk per $\mu g/L$ drinking water or risk per $\mu g/cu$.m air breathed. The third form in which risk is presented is a concentration of the chemical in drinking water or air associated with cancer risks of 1 in 10,000, 1 in 100,000, or 1 in 1,000,000. The rationale and methods used to develop the carcinogenicity information in IRIS are described in The Risk Assessment Guidelines of 1986 (EPA/600/8-87/045) and in the IRIS Background Document. IRIS summaries developed since those Guidelines where indicated (Federal Register 61(79):17960-18011, April 23, 1996). Users are referred to Section I of this IRIS file for information on long-term toxic effects other than carcinogenicity.

This IRIS Summary replaces the summary dated 03/31/1987. The assessment of carcinogenicity by the inhalation route of exposure under the draft revised guidelines for carcinogen risk assessment (U.S. EPA, 1999) differs from the previous EPA evaluation (U.S. EPA, 1985a, b). EPA's previous evaluation considered the incidence of kidney adenocarcinomas (Maltoni et al., 1985) as providing sufficient evidence of carcinogenicity to justify deriving an inhalation unit risk for quantifying the potential human cancer risk. As noted in Sections 4.4.3 and 4.6 of the Toxicological Review of 1,1-Dichloroethylene, the new data suggesting that the kidney adenocarcinomas could be a sex- and exposure. Accordingly, the present evaluation does not derive an inhalation unit risk. This conclusion is consistent with the evaluation by the International Agency for Research on Cancer (IARC) (IARC,

In addition, this assessment of carcinogenicity by the oral route of exposure under the draft revised guidelines for carcinogen risk assessment (U.S. EPA, 1999) differs from the previous EPA evaluation (U.S. EPA, 1985a, b). The previous EPA evaluation derived an oral slope factor from the highest of four slope factors calculated from two studies (NTP, 1982; Quast et al., 1983) that did not show statistically significant increases in tumor incidence attributable to oral exposure. The highest slope factor was based on the adrenal pheochromocytomas in male rats (NTP, 1982). Under the 1999 draft revised guidelines for carcinogen risk assessment, EPA emphasizes the importance of using data that show a statistically significant increase in tumor incidence for calculating a slope factor. As there is no statistically or biologically significant increase in tumor incidence at any site in the relevant oral bioassays, the present evaluation characterizes the weight-of-evidence as *inadequate* and accordingly

does not derive an oral slope factor. This conclusion is consistent with the evaluation by IARC (1999).

_II.A. Evidence for Human Carcinogenicity

__II.A.1. Weight-of-Evidence Characterization

Under the 1986 cancer guidelines (U.S. EPA, 1986), 1,1-DCE is assigned to Group C, possible human carcinogen.

Under the draft revised guidelines for carcinogen risk assessment (U.S. EPA, 1999), EPA concludes 1,1-DCE exhibits suggestive evidence of carcinogenicity but not sufficient evidence to assess human carcinogenic potential following inhalation exposure in studies in rodents. Male mice developed kidney tumors at one exposure in a lifetime bioassay, a finding tempered by the absence of similar results in female mice or male or female rats and by the enzymatic differences (i.e., CYP2E1) between male mice and female mice, male and female rats, and human kidney cells. Limited evidence of genotoxicity has been reported in bacterial systems with metabolic activation. The data for 1,1-DCE are inadequate for an assessment of human carcinogenic potential by the oral route, based on the absence of statistically or biologically significant tumors in limited bioassays in rats and mice balanced against the suggestive evidence in male mice in a single bioassay by inhalation and the limited evidence of genotoxicity. The human epidemiological results on the carcinogenicity of 1,1-DCE are too limited to draw useful conclusions. EPA concludes that the results of kidney tumors in one sex and one exposure in a single species of rodents are too limited to support an exposure-response assessment.

Bioassays for cancer by the oral route of exposure have been conducted in rats (Maltoni et al., 1985; NTP, 1982; Ponomarkov and Tomatis, 1980; Quast et al., 1983) mice (NTP, 1982), and trout (Hendricks et al., 1995). Some of these bioassays were conducted at an exposure below the maximum tolerated dose. The bioassay conducted by Maltoni et al. (1985) exposed the animals for only 1 year. The bioassay conducted in rats by Quast et al. (1983) and the bioassay conducted in mice by NTP (1982) were well conducted and both showed some toxicity in the liver at the highest exposure. Neither of these bioassays provides any significant evidence that 1,1-DCE is a carcinogen by the oral route of exposure. The genotoxicity studies are incomplete, but most studies in mammalian cells indicate a lack of genotoxicity.

Bioassays for cancer by the inhalation route of exposure have been conducted in rats (Lee et al., 1977, 1978; Viola and Caputo, 1977; Hong et al., 1981; Maltoni et al., 1985; Quast et al., 1986; Cotti et al., 1988), mice (Lee et al., 1977, 1978; Hong et al., 1981; Maltoni et al., 1985), and hamsters (Maltoni et al., 1985). None of these bioassays was conducted by a protocol that meets current standards. The major defects in most of these bioassays include exposure of the animals for 1 year and exposure at less than the maximum tolerated dose. The only bioassay that showed some evidence of carcinogenicity was the study in Swiss-Webster mice (Maltoni et al., 1985). This study was conducted at or near the maximum tolerated dose, as animals exposed at 50 ppm died after a few exposures. Although the animals were exposed for only 1 year and then observed until natural death, this study showed an increased incidence of kidney adenocarcinomas in male mice at 25 ppm but not at 10 ppm. The incidence of mammary carcinomas in female mice and pulmonary adenomas in male and female mice did not increase with increased exposure. The responses were actually lower at 25 ppm than at 10 ppm, but survival and other toxicities were comparable.

There is evidence that the induction of kidney adenocarcinomas is a sex- and species-specific response related to the expression of CYP2E1 in the kidney of male mice (Speerschneider and Dekant, 1995; Amet et al., 1997; Cummings et al., 2000). The data presented by these researchers, however, are not sufficient to justify a conclusion that the kidney tumors in male mice have no relevance for a human health risk assessment. This conclusion is made with the knowledge that compounds similar in structure to 1,1-DCE (e.g., tetrachloroethylene, trichloroethylene, and 1,2-dichloroethylene) produce

varying degrees of kidney tumors in animal bioassays.

The genotoxicity studies are incomplete, but most studies in mammalian cells indicate a lack of genotoxicity. Accordingly, EPA concludes that the data on the increased incidence of kidney adenocarcinomas in male mice (Maltoni et al., 1985) provide *suggestive evidence* of carcinogenicity by the inhalation route of exposure. EPA also concludes, considering the evidence of a potential sex- and species-specific response, that the results of this bioassay showing an increase in tumors in one sex and one exposure in a single species of rodents are too limited to support an exposure-response assessment.

1,1-DCE causes gene mutations in microorganisms in the presence of an exogenous activation system. Although most tests with mammalian cells show no evidence of genetic toxicity, the test battery is incomplete because it lacks an in vivo test for chromosomal damage in the mouse lymphoma system.

There are a number of uncertainties in the assessment of the carcinogenicity of 1,1-DCE. As noted above, many of the bioassays by the inhalation route of exposure were not conducted at the maximum tolerated dose or for the full lifetime of the animals. EPA has acknowledged this uncertainty in the weight-of-evidence classification. In addition, our knowledge of the metabolic pathways for 1,1-DCE in the human is incomplete. Although it is likely that the initial oxidation of 1,1-DCE in humans occurs via CYP2E1, there could be other CYP isoforms that could activate 1,1-DCE. Thus, there is some potential for a species-specific carcinogenic response in humans similar to the apparent sex- and species-specific response observed by Maltoni et al. (1985) in the kidney of male mice.

For more detail on Characterization of Hazard and Dose Response, exit to the toxicological review, Section 6 (PDF).

For more detail on Susceptible Populations, exit to the toxicological review, Section 4.7 (PDF).

__II.A.2. Human Carcinogenicity Data

Ott et al. (1976) investigated the health records of 138 employees occupationally exposed to 1,1-DCE in processes not involving vinyl chloride. The individuals included in the study had worked in experimental or pilot plant polymerization operations, in a monomer production process as tankcar loaders, or in a production plant that manufactured a monofilament fiber. Time-weighted-average concentrations (8 hours) of 1,1-DCE in the workplace were estimated from job descriptions and the results of industrial hygiene sampling. The subjects were grouped into three exposure categories: less than 10 ppm, 10–;24 ppm, and greater than 25 ppm. The researchers estimated career exposure by taking into account average duration of employment. Results of the most recent health inventory for individuals in the cohort were compared with findings of matched controls. Analysis of mortalities among the cohort indicated no statistically significant findings. Overall, there were no significant differences between the exposed cohort and the controls in hematology and clinical chemistry parameters. Based on power considerations, this study is inadequate for assessing cancer risk in humans.

__II.A.3. Animal Carcinogenicity Data

Oral

Rats. Ponomarkov and Tomatis (1980) treated 24 female BD IV rats by gavage with 1,1-DCE dissolved in olive oil (150 mg/kg body weight) on the 17th day of gestation. Their offspring (81 males and 80 females) were treated weekly with 1,1-DCE at 50 mg/kg body weight by gavage from the time of weaning for 120 weeks or until the animal was moribund. A control group of offspring (49 males

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and 47 females) received only olive oil. Liver and meningeal tumors were more frequently observed in treated than in untreated animals, but the difference was not statistically significant. The total number of tumor-bearing animals was not statistically different between treated and untreated animals.

NTP (1982) conducted chronic toxicity and carcinogenicity studies of 1,1-DCE for 104 weeks in male and female F344 rats (200 of each sex, 9 weeks old) by gavage in corn oil at 0, 1, or 5 mg/kg-day. No significant differences were observed in survival, clinical signs, or body weight as compared with controls for any group, suggesting that the maximum tolerated dose was not achieved. All of the increased tumor incidences that were statistically significant by the Fisher exact test or by the Cochran-Armitage linear trend test (adrenal pheochromocytoma, pancreatic islet cell adenoma or carcinoma, and subcutaneous fibroma in males and pituitary adenoma in females) were not significant when life-table analyses were used. This difference occurs because life table analyses adjust for intercurrent mortality, and thus minimize the impact of animals dying before the onset of late-appearing tumor. This adjustment was particularly critical for the analyses of tumor incidences in male rats, because 12 controls and 10 low-dose animals were accidentally killed during week 82 of the study. Accordingly, NTP concluded that no increased incidence of tumors was found at any site in these bioassays. Under the conditions of this bioassay, 1,1-DCE administered by gavage was not carcinogenic for F344 rats.

Quast et al. (1983) conducted a 2-year chronic toxicity and carcinogenicity study of 1,1-DCE in Sprague-Dawley rats (6–7 weeks old). There were 80 of each sex rats in the control group and 48 rats of each sex in each exposed group. The 1,1-DCE was incorporated in the drinking water of the rats at nominal concentrations of 0, 50, 100, or 200 ppm. The time-weighted-average exposure over the 2-year period was 7, 10, or 20 mg/kg-day for males and 9, 14, or 30 mg/kg-day for females. No significant differences were found among the groups in appearance and demeanor, mortality, body weight, food consumption, water consumption, hematology, urinalysis, clinical chemistry determinations, organ weights, or organ-to-body-weight ratios. The only treatment-related effect observed in rats was a minimal amount of midzonal fatty change and hepatocellular swelling. No exposure-related neoplastic changes occurred at any exposure.

Maltoni et al. (1985) conducted a carcinogenicity and toxicity study of 1,1-DCE in Sprague-Dawley rats. Animals (9 or 10 weeks old) were exposed by gavage in olive oil to 0, 0.5, 5, 10, or 20 mg/kg, 4–5 days/wk for 52 weeks. There were two control groups, one with 150 animals (75 of each sex) and the other with 200 animals (100 of each sex). The exposed groups had 100 animals (50 of each sex). Following the 52-week exposure, animals were observed until spontaneous death (total duration 147 weeks). Body weight was measured every 2 weeks during the 52 week exposure and every 8 weeks thereafter. Full necropsy and histopathological examination were performed. No biologically significant changes were observed in mortality or body weight. There were no biologically significant noncancer or cancer effects in any organ.

Mice. NTP (1982) conducted 104 weeks of chronic toxicity and carcinogenicity studies on 1,1-DCE in male and female B6C3F $_1$ mice (200 of each sex, 9 weeks old) by gavage in corn oil at 0, 2, or 10 mg/kg. No significant differences in survival, clinical signs, or body weight were in any group, suggesting that the maximum tolerated dose was not achieved. The only observed significant increase (p<0.05) in tumor incidence occurred in low-dose females for lymphoma (2/48, 9/49, 6/50) and for lymphoma or leukemia (7/48, 15/49, 7/50). These increases were not considered to be related to 1,1-DCE administration because similar effects were not found in the high-dose females or in males. Under the conditions of this bioassay, 1,1-DCE administered by gavage was not carcinogenic for B6C3F $_1$

Trout. Hendricks et al. (1995) conducted an 18-month carcinogenicity study of 1,1-DCE in rainbow trout (8 weeks old) at 4 mg/kg-day. Tissues examined for neoplasms included liver, kidney, spleen, gill, gonads, thymus, thyroid, heart, stomach, pyloric ceca, duodenum, rectum, pancreas, and swimbladder. 1,1-DCE produced no neoplasms and no increase in liver weight. There was no evidence

of any other chronic toxic effects.

Inhalation

Rats. Lee et al. (1977, 1978) exposed 2-month-old Charles River CD rats (36 males and 35 females) to 55 ppm 1,1-DCE for 6 hrs/day, 5 days/wk, for 12 months. No significant changes were observed in survival, body weight, hematology, clinical blood chemistry, pulmonary macrophage count, cytogenetic analysis of bone marrow, x-ray examination of extremities, collagen contents in liver and lung, serum aminolevulinic acid (ALA) synthetase, urinary ALA level, and serum alpha-fetoprotein. A mild to markedly severe focal, disseminated vacuolization was observed in livers of most of the rats. No hemangiosarcomas were found in the liver or lung. The incidence of hemangiosarcomas in mesenteric lymph node or subcutaneous tissue was 2/36 in males and 0/35 in females.

Viola and Caputo (1977) exposed 2-month-old Sprague-Dawley rats (30 males and 30 females per group) to 0, 75 ppm, or 100 ppm 1,1-DCE for 22–24 months (hours of daily exposure not reported). The incidence of tumors observed at necropsy (males and females combined) was 15/60; 10/36 and 20/60 at 0, 75 ppm, and 100 ppm, respectively. The tumors observed were classified as subcutaneous fibromas or abdominal lymphomas. The histopathological results from this study have not been published. No other data are reported for this study.

Viola and Caputo (1977) also exposed 2-month-old albino Wistar rats (37 males and 37 females) to 1,1-DCE for 4 hrs/day, 5 days/wk, for 12 months. The exposure was at 200 ppm for the first 6 months and at 100 ppm for the rest of the study. A control group of 60 animals received air only. The incidence of tumors (described as reticulum cell sarcomas of a nonsincytial type, primarily in the abdominal cavity) was 15/60 and 17/74 in control and exposed groups, respectively. No other data are reported from this study.

Hong et al. (1981) evaluated mortality and tumor incidence in rats exposed to 1,1-DCE. Groups of 2-month-old CD rats of both sexes were exposed to 0 or 55 ppm 1,1-DCE 6 hrs/day, 4 days/wk for 1 month (four of each sex), 3 months (four of each sex), 6 months (four of each sex), or 10 months (16 of each sex). Following exposure, all animals were observed for an additional 12 months. In rats exposed for 10 months, there was an increase in mortality following the 12-month observation period (67% in exposed, 41% in controls). There was no significant increase in tumors at any site for any exposure period.

Maltoni et al. (1985) conducted a carcinogenicity and toxicity study of 1,1-DCE in Sprague-Dawley rats. Animals (16 weeks old) were exposed by inhalation to 0, 10, 25, 50, 100, or 150 ppm for 4 hrs/day, 4-5 days/wk for 52 weeks. The control group had 200 animals (100 of each sex); the 10, 25, 50, and 100 ppm groups had 60 animals (30 of each sex), and the 150 ppm group had 120 animals (60 of each sex). Following the 52-week exposure, animals were observed until spontaneous death (total duration 137 weeks). Body weight was measured every 2 weeks during the 52-week exposure and every 8 weeks thereafter. Full necropsy and histopathological examination were performed. No biologically significant changes were seen in mortality or body weight. There were no biologically significant noncancer effects in any organ in either sex and no increase in tumors in males at any site. There was a statistically significant increase (p<0.05) in each treatment group as compared with controls in the number of females with mammary fibromas and fibroadenomas. The incidence was 44/56 (78.6%), 24/24 (100%), 20/20 (100%), 21/22 (95.4%), 21/23 (91.3%), and 38/43 (88.4%) in the control, 10, 25, 50, 100, and 150 ppm groups, respectively. The latency time and the number of tumors per tumor-bearing animal were similar among all groups. The incidence of mammary carcinoma in exposed groups was consistently less than that of controls. The incidence was 16/56 (28.6%), 5/24 (20.8%), 4/20 (20%), 1/21 (4.5%), 3/21 (13.0%), and 9/38 (20.9%) in the control, 10, 25, 50, 100, and 150 ppm groups, respectively.

Quast et al. (1986) and Rampy et al. (1977) reported results from studies that exposed male and

female Sprague-Dawley rats (Spartan substrain, 86 animals/group) to 1,1-DCE by inhalation 6 hrs/day, 5 days/wk, for up to 18 months. Interim sacrifices occurred at 1, 6, and 12 months. Rats were exposed to 1,1-DCE concentrations of 10 ppm and 40 ppm for the first 5 weeks of the study. Based on the absence of observable treatment-related effects among rats sacrificed after 1 month of exposure, the concentrations were increased to 25 and 75 ppm. Exposures were continued at these concentrations through the 18th month of the study. The surviving animals were then held without exposure to 1,1-DCE until 24 months. Cytogenetic evaluations were performed on a separate group of animals (four per sex) exposed to 0, 25, or 75 ppm for 6 months. There were no exposure-related changes in mortality, appearance and demeanor, body weight, clinical chemistry determinations, hematologic evaluations, urinalysis, or cytogenetic evaluation of bone marrow preparations. Although the incidences of several tumors and/or tumor types were found to be statistically increased or decreased as compared with controls, none of these differences were judged to be attributable to 1,1-DCE. The tumor incidence data for both control and treated rats in this study were comparable to historical control data for the Sprague-Dawley rats (Spartan substrain) used by this laboratory for several studies of similar design and duration.

Cotti et al. (1988) exposed Sprague-Dawley rats to 1,1-DCE at 0 or 100 ppm for 4–7 hrs/day, 5 days/wk. The exposures were to 13-week-old females for 104 weeks (60 control animals and 54 exposed animals) and to 12-day embryos for 15 or 104 weeks (158 males and 149 females as controls, 60 males and 60 females exposed for 15 weeks, and 62 males and 61 females exposed for 104 weeks). Animals were observed until spontaneous death. In males and females exposed for 104 weeks and in male offspring exposed for 15 weeks, a slight decrease in body weight (data not reported) was observed. An increased percentage of rats bearing malignant tumors (30.9% vs. 17.3 % in controls) and an increased number of malignant tumors per 100 animals (34.1% vs. 17.9% in controls) were observed in male and female offspring exposed for 104 weeks (statistical analysis not presented). An increase in leukemia in offspring, which appeared to be related to length of exposure (4.2% for controls, and 8.3% and 11.4% for exposure of 15 and 104 weeks, respectively), was also observed. Tumors at other sites (total benign and malignant tumors, total benign and malignant mammary tumors, malignant mammary tumors, and pheochromocytomas) showed no change or a decreased incidence. Data from this study are also reported in Maltoni et al. (1985).

Mice. Lee et al. (1977, 1978) exposed 2-month-old CD-1 mice (18 males and 18 females) to 0 or 55 ppm 1,1-DCE for 6 hrs/day, 5 days/wk, for up to 12 months. No deaths occurred in the control or exposed groups. Weight gain was comparable between groups. There was no change in hematology, clinical blood chemistry, cytogenetic analysis of bone marrow, x-ray examination of extremities, or serum alpha-fetoprotein. The livers showed no increase in mitotic figures using ¹⁴C-thymidine incorporation. The incidence of bronchioalveolar adenoma (males and females combined) for 1–3 months exposure, 4–6 months exposure, 7–9 months exposure, and 10–12 months exposure was 0/24, 1/8, 2/10, and 3/28, respectively. The incidence of hemangiosarcomas in liver (males and females combined) for 6 months exposure, 7–9 months exposure, and 10–12 months exposure was 0/16, 1/10, and 2/28, respectively. No hemangiosarcomas were found in other tissues.

Hong et al. (1981) evaluated mortality and tumor incidence rates in mice exposed to 1,1-DCE. Groups of 2-month-old albino CD-1 mice of both sexes were exposed to 0 or 55 ppm for 6 hrs/day, 4 days/wk, for 1 month (8 of each sex), 3 months (8 of each sex), or 6 months (12 of each sex). Following exposure, all animals were observed for an additional 12 months. In mice exposed for 6 months, there was a slight increase in mortality following the 12-month observation period (46% in exposed, 39% in controls). There was no significant increase in tumors at any site for any exposure period.

Maltoni et al. (1985) conducted a carcinogenicity and toxicity study of 1,1-DCE in Swiss mice. Animals (9 or 16 weeks old) were exposed by inhalation to 0, 10, or 25 ppm. Animals were exposed for 4 hrs/day, 4–5 days/wk, for 52 weeks. There were two control groups, one with 180 animals (90 of each sex) and the other with 200 animals (100 of each sex). The 10-ppm group had 60 animals (30 of each sex). Two groups were exposed to 25 ppm: one with 60 animals (30 of each sex) and the other with

240 animals (120 of each sex). Following the 52-week exposure, animals were observed until spontaneous death (total duration 126 weeks). Body weight was measured every 2 weeks during the 52-week exposure and every 8 weeks thereafter. Full necropsy and histopathological examination were performed.

No biologically significant changes occurred in body weight. The exposed animals had a somewhat higher survival than controls. There was a statistically significant increase (p<0.01) as compared with controls in kidney adenocarcinomas in male mice at 25 ppm but not in male mice at 10 ppm or in female mice at either exposure. The incidence was 0/126 (0%), 0/25 (0%), and 28/119 (23.5%) in male mice in the combined controls, 10 ppm, and combined 25 ppm groups, respectively.

There was a statistically significant increase (p<0.01) as compared with controls in mammary carcinomas in female mice at both exposures, but there was no clear exposure-response relationship. The incidence was 3/185 (1.6%), 6/30 (20%), and 16/148 (11%) in females in the combined controls, 10 ppm, and combined 25 ppm groups, respectively. There was also a statistically significant increase (p<0.01) compared with control in pulmonary adenomas in both exposed groups, but there was no clear exposure-response relationship. The incidence was 12/331 (3.6%), 14/58 (24.1%), and 41/288 (14.2%) in male and female mice combined in the combined controls, 10 ppm, and combined 25 ppm groups, respectively. There were no pulmonary carcinomas in any mice. The incidence data are reported as the number of tumor-bearing animals as compared with the number of animals alive when the first tumor was observed in that organ (kidney adenocarcinoma, 55 weeks; mammary tumor, 27 weeks; pulmonary adenoma, 36 weeks)

Hamsters. Maltoni et al. (1985) conducted a carcinogenicity and toxicity study of 1,1-DCE in Chinese hamsters. Animals (28 weeks old) were exposed by inhalation to 0 or 25 ppm. Animals were exposed for 4 hrs/day, 4-5 days/wk, for 52 weeks. The control group had 35 animals (18 male and 17 female); the 25 ppm group had 60 animals (30 of each sex). Following the 52-week exposure, animals were observed until spontaneous death (total duration 157 weeks). Body weight was measured every 2 weeks during the 52-week exposure and every 8 weeks thereafter. Full necropsy and histopathological examination were performed. There were no biologically significant changes in mortality or body weight. No biologically significant noncancer or tumor effects were seen in any organ.

Dermal. Van Duuren et al. (1979) evaluated the carcinogenicity of 1,1-DCE in male and female noninbred Ha:ICR Swiss mice. Carcinogenicity was assessed in three types of tests: a dermal initiation-promotion assay, a repeated dermal application assay, and a subcutaneous injection assay. Vehicle, no-treatment, and positive control groups were included in the tests. In the initiation-promotion assay, 1,1-DCE was tested as a tumor-initiating agent with phorbol myristate acetate as the promoter. Thirty female mice were treated with 121 mg 1,1-DCE. A significant increase (p<0.005) was observed in skin papillomas (nine in eight mice). In the repeated dermal application assay, exposures of 40 and 121 mg/mouse were used. 1,1-DCE was applied to the back of the shaved animals (30 females/dose). No sarcomas were observed at the treatment site. Although 19 mice in the high-dose group and 12 in the low-dose group had lung tumors and 2 mice in the high-dose group had stomach tumors, the tumor incidence at both sites was not significantly different from that of controls (30 lung tumors and 5 stomach tumors). In the subcutaneous injection assay, the test animals were given weekly injections of 2 mg of 1,1-DCE. After 548 days on test, none of the injected animals developed sarcomas at the injection site. 1,1-DCE showed initiating activity in the two-stage carcinogenesis experiments but was inactive as a whole-mouse dermal carcinogen and after subcutaneous injection.

_II.A.4. Supporting Data for Carcinogenicity

Reitz et al. (1980) investigated the ability of 1,1-DCE to cause DNA alkylation, DNA repair, and DNA replication in liver and kidney of rats and mice. Male Sprague-Dawley rats (body weight 200-250~g) and male CD-1 mice (body weight 18-20~g) were exposed by inhalation for 6 hours. There was only a minimal increase in DNA alkylation in both rats and mice at 50 ppm. Similarly, DNA repair in kidneys

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of mice was only minimally increased at 50 ppm. However, tissue damage (kidney nephrosis at 50 ppm, minimal effect at 10 ppm), an increase in DNA replication (sevenfold increase in ³H-thymidine incorporation at 10 ppm, 25-fold increase at 50 ppm), and an increase in mitotic figures occurred. There was no observed histopathological damage or increased DNA replication in the liver of mice at 10 or 50 ppm. In rats there was a small increase in DNA replication (twofold increase in ³H-thymidine incorporation) in the kidney but no increase in liver at 10 ppm.

1,1-DCE induced mutations in *Salmonella typhimurium* and *Escherichia coli* in the presence of an exogenous metabolic system. In *Saccharomyces cerevisiae*, 1,1-DCE induced reverse mutation and mitotic gene conversion in vitro and in a host-mediated assay in mice. In a single study in *Saccharomyces cerevisiae*, it induced aneuploidy in the presence and absence of metabolic activation. In vitro, gene mutations were increased in mouse lymphoma cells but not in Chinese hamster lung cells with or without an exogenous metabolic system. In a single study, 1,1-DCE induced sister chromatid exchanges in Chinese hamster lung cells in the presence of an exogenous metabolic system but not in its absence. In single studies in vivo, 1,1-DCE did not induce micronuclei or chromosomal aberrations in bone marrow or in fetal erythrocytes of mice, nor dominant lethal mutations in mice or rats.

1,1-DCE causes gene mutations in microorganisms in the presence of an exogenous activation system. Although most tests with mammalian cells show no evidence of genetic toxicity, the test battery is incomplete because it lacks an in vivo assessment of chromosomal damage in the mouse lymphoma assay, a test that EPA considers to be an important component of a genotoxicity battery.

Speerschneider and Dekant (1995) investigated the metabolic basis for the species- and sex-specific nephrotoxicity and tumorigenicity of 1,1-DCE. In kidney microsomes from male mice, the rate of oxidation of 1,1-DCE depended on the hormonal status of the animals. Oxidation of 1,1-DCE was decreased by castration and restored when the castrate was supplemented with exogenous testosterone. In kidney microsomes from naive female mice, the rate of oxidation of 1,1-DCE was significantly lower than in males, but it could be increased by administration of exogenous testosterone. Using an antibody to rat liver CYP2E1, the researchers showed expression of a crosseacting protein in male mouse kidney microsomes that was regulated by testosterone and correlated with the ability to oxidize 1,1-DCE and other substrates for CYP2E1 (e.g., p-nitrophenol and chlorozoxazone).

The researchers also showed that different strains of mice express different levels of CYP2E1, and the strains most sensitive to the effects of 1,1-DCE express greater levels of CYP2E1. Nephrotoxicity in Swiss-Webster mice after inhalation of 1,1-DCE was observed in males and in females treated with exogenous testosterone, but not in naive females. In kidney microsomes obtained from both sexes of rats and in six samples of human kidney from male donors, no p-nitrophenol oxidase activity was detected. Other research groups have also reported the absence of detectable CYP2E1 in human kidney tissue (Amet et al., 1997; Cummings et al., 2000).

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_II.B. Quantitative Estimate of Carcinogenic Risk from Oral Exposure

Not applicable. 1,1-DCE shows equivocal evidence of carcinogenicity by the oral route of exposure

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	Appendix A	A185
_II.C. Quantitative Estimate	e of Carcinogenic Risk from In	
Not applied to a pos	o or caremogenic Risk from In	ihalation Exposure
exposure. The weight-of-evidence, h	estive evidence of human carcinogenicit lowever, is not sufficient to justify derivi	y by the inhalation route of
Top of page	, 2-1	my an initiation tillt risk.
The destination of the state of	endaren eta ji nigelija. Parasa kumi isaka dasam kantara menji inteksiona dasam dasam denga kantara menjenda d	
_II.D. EPA Documentation, Assessment)	Review, and Contacts (Carcing	ogenicity
II.D.1. EPA Documentatio	on ·	
Source Document — Toxicological Rev		
as an appendix to Toxicological Povious	y external scientists. Their comments had on of this IRIS Summary. A record of the of 1,1-Dichloroethylene. To review the case of 1,4-Dichloroethylene. To review the case of	lese comments is included
Other EPA Documentation — This asse	essment replaces previous assessments	(U.S. EPA, 1985a,b).
II.D.2. EPA Review (Carcin		
Agency Consensus Date — 06/07/2002	2	
Carcinogenicity assessment for 1.1. Dial	al studies published through May 2005 v would be directly useful in the revision hloroethylene (1,1-DCE) and a change i nation, IRIS users may contact the IRIS	of the existing
II.D.3. EPA Contacts (Carci	nogenicity Assessment)	

Please contact the IRIS Hotline for all questions concerning this assessment or IRIS, in general, at (202)566-1676 (phone), (202)566-1749 (FAX), or hotline.iris@epa.gov (email address).

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_III. [reserved] _IV. [reserved] _V. [reserved]

_VI. Bibliography

Substance Name - 1,1-Dichloroethylene (DCE) CASRN - 75-35-4 Last Revised - 08/13/2002

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_VII. Revision History

Substance Name - 1,1-Dichoroethene (DCE) CASRN - 75-35-4 Primary Synonym - Vinylidene Chloride

Date	Section	Description
03/31/1987	II.	Carcinogenicity Section added
03/01/1988	I.A.1.	Dose conversion clarified
03/01/1988	I.A.7.	Contact changed
03/01/1988	II.A.2.	Text added
03/01/1988	II.B.3.	Text revised
03/01/1988	II.B.4.	Confidence statement revised
03/01/1988	II.C.3.	Text added
03/01/1988	II.C.4.	Confidence statement revised
06/30/1988	I.A.7:	Changed primary contact's telephone number
12/01/1988	II.A.3.	van Durren et al. citation year corrected
04/01/1989	.A.	Oral RfD summary noted as pending change
12/01/1989 1	.B.	Inhalation RfD now under review
)3/01/1990 I	I.	Clarified NTP, 1982 citation
)3/01/1990 \	/I.	Bibliography on-line
)1/01/1991 I	I.	Text edited
1/01/1991 I	I.C.1.	Inhalation slope factor removed (global change)

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02/01/1991	II.C.3.	Information on extrapolation process included
08/01/1991	VI.A.	References clarified
08/01/1991	VI.C.	References clarified
01/01/1992	IV.	Regulatory actions updated
08/01/1995	I.A., I.B.	EPA's RfD/RfC and CRAVE workgroups were discontinued in May, 1995. Chemical substance reviews that were not completed by September 1995 were taken out of IRIS review. The IRIS Pilot Program replaced the workgroup functions beginning in September, 1995.
04/01/1997	III., IV., V.	Drinking Water Health Advisories, EPA Regulatory Actions, and Supplementary Data were removed from IRIS on or before April 1997. IRIS users were directed to the appropriate EPA Program Offices for this information.
01/12/2000	I., II.	This chemical is being reassessed under the IRIS Program.
08/13/2002	I-VIII	New RfD, RfC, and cancer assessment
10/28/2003	I.A.6., I.B.6., II.D.2.	Screening-Level Literature Review Findings message has been added.
06/22/2005	I.A.6., I.B.6., II.D.2.	Screening-Level Literature Review Findings message has been removed and replaced by comprehensive literature review conclusions.

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_VIII. Synonyms

Substance Name — 1,1-Dichloroethylene CASRN — 75-35-4 Last Revised — 08/13/2002

- 1,1-Dichloroethene
- * 1,1-DCE
- Dichloroethene, 1,1-
- * Ethylene, 1,1-dichloro-
- NCI-C54262
- RCRA Waste Number U078
- Sconatex
- UN 1303
- Vinylidene chloride
- Vinylidene dichloride
- Vinylidine chloride
- Chlorure de vinylidene
- ▼ VDC

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Chronic Health Hazards for Non-Carcinogenic Effects

Reference Dose for Chronic Oral Exposure (RfD)

- Oral RfD Summary
- Principal and
- Supporting Studies
- Uncertainty and Modifying Factors
- Additional
 - Studies/Comments
- Confidence in the Oral RfD
- **EPA** Documentation and Review

Reference Concentration for Chronic Inhalation Exposure (RfC)

- Inhalation RfC Summary
- Principal and
- Supporting Studies
- Uncertainty and Modifying Factors
- Additional
- Studies/Comments
- Confidence in the Inhalation RfC
- **EPA** Documentation and Review

Carcinogenicity Assessment for Lifetime Exposure

Evidence for Human Carcinogenicity

- Weight-of-Evidence Characterization
- Human
- Carcinogenicity Data
- Animal Carcinogenicity
- Supporting Data for Carcinogenicity

Quantitative Estimate of Carcinogenic Risk from Oral Exposure

- Summary of Risk Estimates
- Dose-Response Data

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Administrative Decisions Cited

Lancaster v. Dep't of Environment & Natural Resources, 04 EHR 1151.

Lucy, Inc. v. Dep't of Environment and Natural Resources, 01 EHR 1695.

Moore v. Dep't of Environment and Natural Resources, 06 EHR 1479

Sharpe v. Dep't of Environment & Natural Resources Environmental Management Commission, 03 EHR 1776.

Southerland Electric Company v. McLawhorn, 00 EHR 2090.

Wheatley Oil Company, Inc. v. Dep't of Environment & Natural Resources, 03 EHR 0030

SECTION .0200. CLASSIFICATIONS AND GROUNDWATER QUALITY STANDARDS

.0201 GROUNDWATER CLASSIFICATIONS

The classifications which may be assigned to the groundwaters will be those specified in the following series of classifications:

- (1) Class GA groundwaters; usage and occurrence:
 - (a) Best Usage. Existing or potential source of drinking water supply for humans.
- (b) Conditions Related to Best Usage. This class is intended for those groundwaters in which chloride concentrations are equal to or less than 250 mg/l, and which are considered suitable for drinking in their natural state, but which may require treatment to improve quality related to natural conditions.
 - (c) Occurrence. In the saturated zone.
- (2) Class GSA groundwaters; usage and occurrence:
 - (a) Best Usage. Existing or potential source of water supply for potable mineral water and conversion to fresh waters.
- (b) Conditions Related to Best Usage. This class is intended for those groundwaters in which the chloride concentrations due to natural conditions is in excess of 250 mg/l, but which otherwise may be considered suitable for use as potable water after treatment to reduce concentrations of naturally occurring substances.
- (c) Occurrence. In the saturated zone.
- (3) Class GC groundwaters: usage and occurtence:
- ((a) Best Usage. The best usage of GC groundwaters is as a source of water supply for purposes other than drinking, including other domestic uses by humans.

- (b) Conditions Related to Best Usage. This class includes those groundwaters that do not meet the quality criteria for GA or GSA groundwaters and for which efforts to improve groundwater quality would not be technologically feasible, or not in the best interest of the public. Continued consumption of waters of this class by humans could result in adverse health affects.
- (c) Occurrence. Groundwaters of this class may be defined by the Commission pursuant to Section .0300 of this Subchapter on a case by case basis.

History Note

Statutory Authority G.S. 143-214.1; 143B-282(2); Eff. June 10, 1979;

Amended Eff. October 1, 1993; August 1, 1989; September 1, 1984; December 30, 1983.

.0202 GROUNDWATER QUALITY STANDARDS

- (a) The groundwater quality standards for the protection of the groundwaters of the state are those specified in this Rule. They are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of the state, which may be tolerated without creating a threat to human health or which would otherwise render the groundwater unsuitable for its intended best usage.
- (b) The groundwater quality standards for contaminants specified in Paragraphs (g) and (h) of this Rule are as listed, except that:
 - (1) Where the standard for a substance is less than the practical quantitation limit, the detection of that substance at or above the practical quantitation limit constitutes a violation of the standard.
- (2) Where two or more substances exist in combination, the Director shall consider the effects of chemical interactions as determined by the Division of Public Health and may establish maximum concentrations at values less than those established in accordance with Paragraphs (c), (g), or (h) of this Rule. In the absence of information to the contrary, in accordance with Paragraph (d) of this Rule, the carcinogenic risks associated with carcinogens present shall be considered additive and the toxic effects associated with non-carcinogens present shall also be considered additive.
- (3) Where naturally occurring substances exceed the established standard, the standard shall

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DEPARTMENT OF ENVIRONMENT AND NATURAL RESOURCES

be the naturally occurring concentration as determined by the Director.

- (c) Except for tracers used in concentrations which have been determined by the Division of Public Health to be protective of human health, and the use of which has been permitted by the Division, substances which are not naturally occurring and for which no standard is specified shall not be permitted in concentrations at or above the practical quantitation limit in Class GA or Class GSA groundwaters. Any person may petition the Director to establish an interim maximum allowable concentration for a substance for which a standard has not been established under this Rule. The petitioner shall submit relevant toxicological and epidemiological data, study results, and calculations necessary to establish a standard in accordance with Paragraph (d) of this Rule. Within three months after the establishment of an interim maximum allowable concentration for a substance by the Director, the Director shall initiate action to consider adoption of a standard for that substance.
- (d) Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of:
- (1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)];
- (2) Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6;
 - (3) Taste threshold limit value;
 - (4) Odor threshold limit value:
 - (5) Maximum contaminant level; or
- (6) National secondary drinking water stan-
- (e) The following references, in order of preference, shall be used in establishing concentrations of substances which correspond to levels described in Paragraph (d) of this Rule.
 - (1) Integrated Risk Information System (U.S. EPA).
 - (2) Health Advisories (U.S. EPA Office of Drinking Water).
- (3) Other health risk assessment data published by U.S. EPA.
- (4) Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.

- (f) Groundwater quality standards specified in Paragraphs (g) and (h) of this Rule and mierim maximum allowable concentrations established pursuant to Paragraph (c) of this Rule shall be reviewed on a triennial basis. Appropriate modifi cations to established standards shall be made in accordance with the procedure prescribed in Page graph (d) of this Rule where modifications considered appropriate based on data published subsequent to the previous review,
- (g) Class GA Standards. Where not otherwise indicated, the standard refers to the total concentration in micrograms per liter of any constituents in a dissolved, colloidal or particulate form which is mobile in groundwater. This does not apply to sediment or other particulate matter which is preserved in a groundwater sample as a result of well construction or sampling procedures. The Class
 - (1) Acenaphthene: 80;
 - (2) Acenaphthylene: 200;
 - (3) Acetone: 6 mg/L;
 - (4) Acrylamide: 0.008;
 - (5) Anthracene: 2 mg/L;
 - (6) Arsenic: 10:
 - (7) Atrazine and chlorotriazine metabolites: 3;
 - (8) Barium: 700;
 - (9) Benzene: 1;
- (10) Benzo(a)anthracene (benz(a)anthracene): 0.05;
- (11) Benzo(b)fluoranthene: 0.05;
- (12) Benzo(k)fluoranthene: 0.5;
- (13) Benzoic acid: 30 mg/L;
- (14) Benzo(g,h,i,)perylene: 200;
- (15) Benzo(a)pyrene: 0.005;
- (16) Bis(chloroethyl)ether: 0.03;
- (17) Bis(2-ethylhexyl) phthalate (di(2-ethylhexyl) phthalate): 3;
 - (18) Boron: 700;
 - (19) Bromodichloromethane: 0.6;
 - (20) Bromoform (tribromomethane): 4;
 - (21) n-Butylbenzene: 70;
- (22) sec-Butylbenzene: 70;
- (23) tert-Butylbenzene: 70;
- (24) Butylbenzyl phthalate: 1 mg/L;
- (25) Cadmium: 2;
- (26) Caprolactam: 4 mg/L;
- (27) Carbofuran: 40;
- (28) Carbon disulfide: 700;

704/343-2060 Fax 704/444-8739 Benne.Hutson@hmw.com

201 North Tryon Street Charlotte, NC 28202 P.O. Box 31247 (28231) 704.343.2000 f704.343.2300

September 28, 2004

Mr. David Hance
Division of Water Quality
North Carolina Department of Environment and Natural Resources
1636 Mail Service Center
Raleigh, North Carolina 27699-1636

Re: Rhodia Inc./Comments on Proposed Revision to North Carolina Groundwater Quality Standard for 1,1-Dichloroethylene Client-Matter No. 4066081.000602

Dear Mr. Hance:

We represent Rhodia, Inc., a global specialty chemicals manufacturer. The purpose of this letter is to comment on the North Carolina Department of Environment and Natural Resources' (NCDENR's) proposed changes to the 2L groundwater standards, specifically the proposed change and subsequent rescission of the change to the standard for 1,1-dichloroethylene (1,1-DCE).

For almost a decade now, Rhodia has been operating a groundwater remediation system at its former facility in Gastonia. The primary contaminant is 1,1-DCE.

As part of its remediation, Rhodia paid nearly \$1 million to provide municipal water to over 100 residences in the surrounding area, thus significantly reducing the risk posed by this site. The remediation system also prevents contaminated groundwater from discharging to an unnamed tributary on Rhodia's former site. This risk reduction is enhanced as there is no public access to the tributary. The closest access point is downstream of the site where contamination from an off-site, non-Rhodia source not being remediated poses the risk of exposure. The continued operation of the groundwater remediation system continues to further reduce the risk through the ongoing removal of contaminant mass.

The clean up standard Rhodia has been striving to meet for 1,1-DCE has been the historic 2L standard of .007 milligrams per liter (mg/L). Last year, the United States Environmental Protection Agency ("U.S. EPA") published its findings on its reevaluation of various compounds. As to 1,1-DCE, U.S. EPA concluded that the risks had been greatly overstated. In light of U.S. EPA's findings, one would have expected the 2L standard for 1,1-DCE to have been raised. Needless to say,

Mr. David Hance September 28, 2004 Page 2

NCDENR's proposal to tighten the 2L standard from .007 mg/L to 5.8 x 10⁻⁵ mg/L was both surprising and unjustified.

NCDENR has now correctly proposed to rescind this proposed reduction. However, keeping the current standard is not legally sufficient either. Rather, based upon its regulations, NCDENR must raise this standard so it is based on current, accurate toxicological data.

Two regulations govern the establishment of groundwater quality standards in North Carolina. First, under 15A N.C.A.C. 2L.0202(d), the lesser of six different standards is to be used to set a compound's groundwater limit. Second, 15A N.C.A.C. 2L.0202(e) requires NCDENR to establish groundwater standards based upon four sources of toxicity data listed "in order of preference" in the regulation. Although separate, 15A N.C.A.C. 2L.0202(d) and (e) are not independent regulations. Rather they must be read in tandem as the sources for establishing standards listed in 15A N.C.A.C. 2L.0202(e) must "correspond to levels described in Paragraph (d) of this Rule." (Emphasis added.) From this it follows that if a basis for a groundwater quality standard listed in 15A N.C.A.C. 2L.0202(d) does not "correspond" to an accepted reference source listed in 15A N.C.A.C. 2L.0202(e), then the standard in subparagraph (d) cannot be used.

That is the current situation with regard to 1,1-DCE. On its face, the "lesser" of the six identified standards in 15A N.C.A.C. 2L.0202(d) for 1,1-DCE is the maximum contaminant level (MCL) of .007 mg/L. The Safe Drinking Water Act requires that determination of an MCL for a compound must be based in part on the oral reference dose (RfD) for that compound. In its reevaluation of 1,1-DCE in August 2002, U.S. EPA recalculated the RfD for 1,1-DCE, changing it from 0.009mg/kg-day to 0.046mg/kg-day. U.S. EPA published these changes as part of its revision to the file for 1,1-DCE on the Integrated Risk Information System (IRIS). Pursuant to 15A N.C.A.C. 2L.0202(e), IRIS is the most preferred reference source for the establishment of groundwater quality

For reasons of workload, administrative costs and lack of evidence of public cost savings, U.S EPA decided not to change the MCL for 1,1-DCE despite its revision to IRIS. (See National Primary Drinking Water Regulations; Announcement of Completion of EPA's Review of Existing Water Standards, 68 Fed. Reg. 42,908, 42,921 (July 18, 2003)). As a result, for non-scientific reasons, the current MCL for 1,1-DCE no longer "corresponds" to the IRIS evaluation of 1,1-DCE. Thus, under North Carolina regulations, the current MCL cannot be used to establish the 2L groundwater quality

15A N.C.A.C. 2L.0202(d) provides that groundwater quality standards can also be the lesser of the following for a compound:

- Systemic threshold concentration; 1.
- Concentration corresponding to a 1X10-6 incremental lifetime cancer risk; 2. 3.
- Taste threshold limit value;

Mr. David Hance September 28, 2004 Page 3

- 4. Odor threshold limit value; or
- 5. National secondary drinking water standard.

1,1-DCE does not have a taste threshold limit value, odor threshold limit value or national secondary drinking water standard. Since in its IRIS revisions U.S. EPA removed the oral cancer slope factor for 1,1-DCE, there is no cancer risk concentration that can be used to establish a groundwater quality standard.

Thus, the only basis for establishing a groundwater quality standard for 1,1-DCE is the systemic threshold concentration for 1,1-DCE. According to 15A N.C.A.C. 2L.0202(d)(1), the systemic threshold concentration is calculated as follows:

Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution [(.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)]

Since this formula can use U.S. EPA's recalculated reference dose (RfD) for 1,1-DCE, the systemic threshold concentration "corresponds" with IRIS and thus complies with 15A N.C.A.C. 2L.0202(e). Performing the calculation using the recalculated RfD, the systemic threshold concentration for 1,1-DCE is .322 mg/L, nearly 50 times higher than the current 2L standard.

At least at groundwater remediation sites and no doubt more generally, making this change in the 1,1-DCE standard would likely improve environmental quality in North Carolina. Rhodia has been operating a traditional pump-and-treat remediation system at its former Gastonia site for nearly a decade. During that time, it has continually evaluated the feasibility of other remedial options for the site. However, none of these options has proved viable as each option identified would have involved significant cost increases without corresponding reductions in the amount of time needed to reach the current .007 mg/L standard. With a revised standard, clean up can be achieved quicker, making it more likely that Rhodia will consider changes that would result in a faster rate of clean up than is currently occurring. That would clearly be a benefit to North Carolina's environment.

In responding to a previous comment on this proposed change, NCDENR noted that groundwater quality standards can be based on "other appropriate, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data", citing 15A N.C.A.C. 2L.0202(e)(4) as support for this position. While 15A N.C.A.C. 2L.0202(e)(4) allows for other appropriate risk assessment data to be used to establish groundwater quality standards, it is "in order of preference" the least favored reference source. Furthermore, NCDENR has not cited the "other appropriate, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data" on which it is supporting its decision.

IRIS is the most preferred source. If it can be used, it must be used. With regard to establishing a groundwater quality standard for 1,1-DCE, IRIS can be used through the calculation of

Mr. David Hance September 28, 2004 Page 4

the systemic threshold value. As a result, pursuant to 15A N.C.A.C. 2L.0202(d) and (e), NCDENR must raise the groundwater quality standard for 1,1-DCE to .322 mg/L. If it fails to do so, NCDENR is failing to act as required by rule, acting erroneously, and acting arbitrarily and capriciously, all of which expose the agency to a likely successful administrative challenge to its action under N.C. Gen. Stat. § 150B-23(a)(2), (4), and (5). NCDENR should seriously reevaluate its position in light of the comments set forth in this letter.

Sincerely,

HELMS MULLISS & WICKER, PLLC

Benne C. Hutson

cc: Rhodia Management Team

Mr. David Hance Groundwater Section Division of Water Quality Department of Environment and Natural Resources 1636 Mail Service Center Raleigh, NC 27699-1636 of North Carolina, Inc.
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ENVIRONMENTAL

ARCADIS G&M

Subject:

1,1-Dichloroethene

Dear Mr. Hance:

During our meeting on Thursday, August 29, you stated that the comment period for the proposed changes to the 2L Groundwater Standards remained open. The focus of this letter is the substance 1,1-dichloroethene or 1,1-dichloroethylene as it is referenced in the rule. Craig A. Bromby of the law firm of Hunton & Williams has collaborated with me on the preparation of the comments contained herein.

The proposed reduction in the 2L standard from 0.007 milligrams per liter (mg/L) to 5.8 × 10⁻⁵ mg/L is inappropriate. On August 13, 2002, the U.S. Environmental Protection Agency (USEPA) revised the file for 1,1-dichloroethene on the Integrated Risk Information System (IRIS) database (URL: www.epa.gov/iris) (URL for 1,1-dichloroethene: http://www.epa.gov/iris/subst/0039.htm). The reevaluation of 1,1-dichloroethene resulted in the removal of the oral cancer slope factor and the recalculation of the oral reference dose. The result is that if the 2L standard were recalculated using the latest revised toxicity values, it would be much higher than the existing standard, as seen below.

ate:

13 September 2004

Contact:

Shawn Sager

Phone:

Ext. 225

E-mail:

ssager@arcadis-us.com

Our ref:

NC102014.0001

Calculation Of A Groundwater Standard Following 15A NCAC 2L .0202

 Systemic threshold concentration: Oral reference dose revised on August 13, 2002.

RfD = 5×10^{-2} mg/kg-day

Systemic threshold concentration

= RfD \times 70 kg \times 0.2/2 L/day

Systemic threshold concentration

= $(5 \times 10^{-2} \text{ mg/kg-day} \times 70 \text{ kg} \times 0.2)/2 \text{ L/day}$

= 0.350 mg/L

Mr. David Hance 13 September 2004

- 2. Concentration corresponding to an incremental lifetime cancer risk of 1×10^{-6} is not calculated because there is no oral cancer slope factor on IRIS (as of August 13, 2002, when the old value was withdrawn by the USEPA due to a reevaluation of the toxicity information).
- 3. Taste threshold: none found.
- 4. Odor threshold: none found.
- Maximum contaminant level: 0.007 mg/L.
- Secondary maximum contaminant level: none listed.

Based on the literal application of 15A NCAC 2L .0202(e), the standard for 1,1dichloroethene should remain at 0.007 mg/L. However, it is apparent from the revision to the information contained in the IRIS file for this compound that a standard of 0.007 mg/L for 1,1-dichloroethene is inappropriately stringent, since the systemic threshold concentration is calculated as 0.35 mg/L. The changes in the IRIS file for this constituent demonstrate a flaw in the hierarchy used to develop the 2L standards, as discussed below.

It is incumbent on the Division of Water Quality to propose a change to 15A NCAC 2L .0202(e) to require, rather than a rote reliance on the lowest number, notwithstanding its validity, that the best scientific information available be used to achieve an appropriately protective level, which reflects the actual risk posed by the constituent. Proposal of a standard any lower than 0.007 mg/L, given this information, both violates 15A NCAC 2L .0202 and is without valid scientific basis. Allowing the standard to remain at 0.007 mg/L is only marginally more justifiable, and should be corrected now or at the latest in the next biennial review.

We propose that the 15A NCAC 2L .0202(d) be amended to allow the selection of the most scientifically defensible value among the sources listed rather than the lowest value, and that 15A NCAC 2L .0202(e) be amended to provide that the derivation of the appropriate protective level be guided by the references included in 15A NCAC 2L .0202(e) according to the scientific validity of the reference rather than proceeding in a particular order of preference. The data developed for 1,1dichloroethene demonstrate that a more dynamic system is needed in this area.

Groundwater standards are used to set clean-up levels for contaminated sites. Overly stringent levels set orders of magnitude too low can result in the needless expenditure of public and private resources for no justifiable reason. Standards based on outdated information, particularly when the invalidity of the information has been

Mr. David Hance 13 September 2004

acknowledged by the agency on whom the rule presently directs you to rely, cannot be justified and as a matter of sound public policy, should not be allowed to stand.

The case of 1,1-dichloroethene plainly illustrates the flaw in the existing regulation, which the division should seek to correct. The correction will not jeopardize public health or the quality of the environment, as in many, if not most, cases the lowest number may still reflect the most scientifically sound value. Since the groundwater standards do not necessarily apply to public water supplies, relying on a maximum contaminant level (MCL) that has not been reevaluated based on current toxicity information, may be inappropriate. Additionally, it is unsuitable for the EMC to adopt a standard that is not supported by the most recent and reliable data, and relies instead on a historical published artifact.

We urge you to reconsider your position regarding the proposed reduction in the 2L standard. Based on the most up-to-date information presented herein, the standard should be recalculated. If you have any questions, please feel welcome to contact the undersigned or Craig Bromby of Hunton & Williams.

Sincerely,

ARCADIS G&M of North Carolina, Inc.

Shawn L. Sager, Ph.D. Principal Scientist

Copies:
Carl Bailey—NCDENR
Arthur Mouberry—NCDENR
Steve Olp—Celanese
Craig Bromby—Hunton & Williams



North Carolina Department of Health and Human Services Division of Public Health • Epidemiology Section 1912 Mail Service Center • Raleigh, North Carolina 27699-1912 Tel 919-733-3410 • Fax 919-733-9555

Michael F. Easley, Governor

October 1, 2004

Cannen Hooker Odom, Secretary

David Hance 1617 Mail Service Center 512 N. Salisbury Street Raleigh, NC 27604 David.Hance@ncmail.net

Kevin Martin Soil and Environmental Consultants, PA 11010 Raven Ridge Road Raleigh, NC 27614 (Environmental Management Commission Member) kmartin@sandEC.com

Andrew Pitner DENR/DWQ/Aquifer Protection Section Mooresville Regional Office 610 East Center Avenue e 301 oresville, NC -115 Andrew.Pitner@ncmail.net

Dear Mr. Hance, Mr. Martin, and Mr. Pitner:

I am writing in response to the comments received today on thirty-three groundwater quality standards that recently went to public hearing on August 19. The detailed responses to comments are attached. Based on review of the comments, I am recommending changes to cyanide and 1,1-dichloroethylene that were not mentioned by me at the public hearing at the August 19 public hearing.

- For cyanide, I am recommending a change in the descriptive narrative from cyanide (as free cyanide or hydrogen cyanide) to cyanide (as free cyanide).
- For 1,1-dichloroethylene, I am recommending an increase in the standard for 1,1-dichloroethylene from 0.007 mg/L to 0.35 mg/L. Using the new 2002 data provided by US EPA, the recommended groundwater quality standard for 1,1-dichloroethylene is 0.35 mg/L instead of 0.007 mg/L. However, the lesser of the 1-6 criteria specified in 15A NCAC 2L .0202 (d) is 0.007 mg/L, the US EPA maximum contaminant level. The 0.007 mg/L meets the criteria of the rule (as being the lesser of the six criteria) but the 0.35 mg/L is the more scientifically valid level.



North Carolina Department of Health and Human Services Division of Public Health • Epidemiology Section 1912 Mail Service Center • Raleigh, North Carolina 27699-1912 Tel 919-733-3410 • Fax 919-733-9555

Michael F. Easley, Governor

Carmen Hooker Odom, Secretary

October 1, 2004 letter

Mr. Hance

Mr. Martin

Mr. Pitner

Please do not hesitate to call me if you have any questions at 919-715-6429.

Sincerely,

Dr. Luanne K. Williams, Toxicologist Occupational and Environmental Epidemiology Branch NC Division of Public Health

#98

September 30, 2004 Comments from Mr. Charles R. Wakild with Progress Energy – Progress Energy questions why the 0.000058 mg/L was stated as the recommended level in the August 19 public hearing notice instead of the US EPA maximum contaminant level goal and maximum contaminant level of 0.007 mg/L.

September 28, 2004 comments from Benne C. Hutson with Helms Mulliss Wicker – questions were raised as to why the outdated US EPA maximum contaminant level of 0.007 mg/L is being recommended as opposed to the systemic threshold concentration of 0.35 mg/L which was calculated using the revised reference dose.

NC Division of Public Health Response to Comments

Response to Progress Energy Comments

The August 19 public hearing notice did not have the correct or latest NC Division of Public Health recommended groundwater quality standard for 1,1-dichloroethylene of 0.007 mg/L. In 2002, the NC Division of Public Health forwarded new recommendations to the Groundwater Section (see enclosed memorandum) regarding 1,1-dichloroethylene. The new recommended level was 0.007 mg/L which was an increase from the previously recommended level of 0.000058 mg/L made in 1999. The 0.000058 mg/L corresponded to a one in a million cancer risk based on the US EPA recommended oral slope factor of 0.6 (mg/kg-day)⁻¹. The 0.007 mg/L level is the lesser of the six criteria used to calculate the standards as mentioned in 15A NCAC 2L .0202(d).

Response to Helms Mulliss Wicker Comments

in August 2002, US EPA completed a review of the noncancer and carcinogenicity studies which resulted in revision to the oral reference dose and carcinogenic assessment but not the maximum contaminant level. The previous chronic oral reference dose was 0.009 mg/kg-day based on hepatic lesions in rats observed in the 1983 Quast et al. study. This reference dose served as the basis for the calculation of the existing maximum contaminant level as follows (see 1995 IRIS):

0.009 mg/kg-day x 70 kg x 1 day/2 Liters x 0.10 safety factor account for carcinogenicity x 0.20 relative source contribution = 0.0063 mg/L or rounded up to 0.007 mg/L

According to US EPA's 2002 assessment, there is no statistically or biologically significant increase in tumor incidence at any site in the relevant oral bioassays. Therefore, a safety factor of 10 to account for carcinogenicity is no longer needed. US EPA also conducted additional review of the 1983 Quast et al. study and changed the reference dose based on this review from 0.009 mg/kg-day to 0.05 mg/kg-day. Using this revised chronic oral reference dose, a systemic threshold concentration can be calculated as follows:

A systemic threshold concentration for 1,1-dichloroethylene can be calculated as shown (US Environmental Protection Agency, Integrated Risk Information System (IRIS) Online Computer Database 2004 http://www.epa.gov/iris):

0.05 mg/kg/day x 70 kg x 1day / 2L x 0.20(RSC) = 0.35 mg/L RSC = Relative Source Contribution

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Note: The oral reference dose of 0.05 mg/kg-day is based upon a rat chronic drinking water study where liver toxicity was the critical effect reported.

Using the new 2002 data provided by US EPA, the recommended groundwater quality standard for 1,1-dichloroethylene is 0.35 mg/L instead of 0.007 mg/L. However, the lesser of the 1-6 criteria specified in 15A NCAC 2L .0202 (d) is 0.007 mg/L, the US EPA maximum contaminant level. The 0.007 mg/L meets the criteria of the rule but the 0.35 mg/L is the more scientifically valid level. It is recommended to use the more scientifically valid level of 0.35 mg/L which is calculated as follows:

Criteria Specified in 15A NCAC 2L .0202

1. A systemic threshold concentration for 1,1-dichloroethylene can be calculated as shown (US Environmental Protection Agency, Integrated Risk Information System (IRIS) On-line Computer Database 2004 http://www.epa.gov/iris):

0.05 mg/kg/day x 70 kg x 1 day / 2L x 0.20(RSC) = 0.35 mg/L

RSC = Relative Source Contribution

Note: The oral reference dose of 0.05 mg/kg-day is based upon a rat chronic drinking water study where liver toxicity was the critical effect reported.

- According to US EPA, there is no statistically or biologically significant increase in tumor incidence at any site in the relevant oral bioassays. Using the draft revised guidelines for carcinogenic risk assessment, US EPA characterizes the carcinogenic weight of evidence as inadequate and accordingly chose not to derive an oral slope factor (US Environmental Protection Agency, Integrated Risk Information System (IRIS) On-line Computer Database 2004 http://www.epa.gov/iris). The 2002 (8/13/02) IRIS summary replaces the 1985 summary where a slope factor was generated from two studies that did not show statistically significant increases in tumor incidence attributable to oral exposure. Under the 1999 draft revised guidelines for carcinogen risk assessment, EPA emphasizes the importance of using data that show a statistically significant increase in tumor incidence for calculating a slope factor. According to US EPA, the data for 1,1-dichloroethylene are inadequate for an assessment of human carcinogenic potential by the oral route, based on the absence of statistically or biologically significant tumors in limited bioassays in rats and mice balanced against the suggestive evidence in male mice in a single bioassay by inhalation and the limited evidence of genotoxicity. The human epidemiological results on the carcinogenicity of 1,1-dichloroethylene are too limited to draw useful conclusions. EPA concludes that the results of kidney tumors in one sex and one exposure in a single species of rodents are too limited to support an exposure-response assessment.
- 3. Currently, the maximum contaminant level for 1,1-dichloroethylene is 0.007 mg/L (U.S. Environmental Protection Agency Office of Ground Water and Drinking Water http://www.epa.gov/safewater/mcl.html 2004). The previous chronic oral reference dose available on IRIS was 0.009 mg/kg-day which was based on hepatic lesions in rats observed in the 1983 Quast et al. study. This reference dose served as the basis for the calculation of the existing maximum contaminant level as follows (see the 1995 IRIS):
- 0.009 mg/kg-day x 70 kg x 1 day/2 Liters x 0.10 safety factor account for carcinogenicity x 0.20 relative source contribution = 0.0063 mg/L or rounded up to 0.007 mg/L

This level is outdated. The 1983 Quast et al. study was reviewed by US EPA in 2002 and based on this review, US EPA recommended a change in the reference dose to 0.05 mg/kg-day. Also, according to US EPA, there is no statistically or biologically significant increase in tumor incidence at any site in the relevant oral bioassays and a safety factor of 10 is no longer needed to account for carcinogenicity. If US EPA used the same equation above to calculate a maximum contaminant level but removed the safety factor of 10 (which is no longer needed to account for carcinogenicity) and

inserted the new reference dose of 0.05 mg/kg-day, then the maximum contaminant level may be calculated to be 0.35 mg/L.

- 4. Currently, there is no national secondary drinking water standard for 1,1-dichloroethylene (U.S. Environmental Protection Agency Office of Ground Water and Drinking Water http://www.epa.gov/safewater/mcl.html 2004).
- 5. A taste threshold value for 1,1-dichloroethylene was not found in the literature.
- 6. An odor threshold value for 1,1-dichloroethylene was not found in the literature.

The recommended groundwater quality standard for 1,1-dichloroethylene is 0.35 mg/L and not 0.007 mg/L.

Benne C. Hutson 704,343,2060 Fax 704,444,8739 benne.hutson@hmw.com

201 North Tryon Street Charlotte, NC 28202 P.O. Box 31247 (28231) 704.343.2000 f 704.343.2300

January 18, 2005

VIA STANDARD MAIL

Mr. David H. Moreau Chairman Dpt. City & Planning, UNC-CH CB3140 Chapel Hill, NC 27599

Re: Proposed Groundwater Standard for 1,1-dichloroethylene (1,1-DCE). Client-Matter No. 4066081-602

Dear Mr. Moreau:

We represent Rhodia, Inc., a global specialty chemicals manufacturer. For almost a decade now, Rhodia has been operating a groundwater remediation system at its former manufacturing plant in Gastonia. The primary constituent of concern is 1,1-dichloroethylene (1,1-DCE).

We are writing regarding the proposed revision to the groundwater quality standard for 1,1-DCE that the EMC will consider at its February meeting. The regulatory situation regarding 1,1-DCE is different from all of the other chemicals you will consider in February. The current groundwater quality standard for 1,1-DCE is .007 mg/L. Based on U.S. EPA's most current scientific and toxicological evidence, DENR's toxicologist and the two hearing officers all agree that the standard should be .350 mg/L. (A copy of our September 28, 2004 letter explaining why the standard should be .350 mg/L and copies of Dr. Luanne K. Williams' letter and the portions of the hearing officers' report agreeing with this conclusion are enclosed.)

However, the hearing officers concluded that DENR's current regulations would have to be amended to make this change. As we will explain in this letter, we believe that the EMC can make this change under the current regulations and we ask that you do so at your February meeting.

Two regulations govern the establishment of groundwater quality standards.

- 1. 15A N.C.A.C. 2L .0202(d) provides that the lesser of six different standards should be used to set a compound's groundwater limit.
- 2. 15A N.C.A.C. 2L.0202(e) mandates that four sources of toxicity data listed "in order of preference shall be used in establishing" groundwater standards.

January 18, 2005 Page 2

Although separate, these two regulations are not independent of each other. Rather they must be read together as the sources listed in 15A N.C.A.C. 2L.0202(e) must "correspond to levels described in Paragraph (d) of this Rule." From this it follows that if a basis for a groundwater quality standard listed in 15A N.C.A.C. 2L.0202(d) does not "correspond" to an accepted reference source listed in 15A N.C.A.C. 2L.0202(e), then the standard in subparagraph (d) cannot be used.

That is the situation with regard to 1,1-DCE. On its face, the "lesser" of the six identified standards in 15A N.C.A.C. 2L.0202(d) for 1,1-DCE is the maximum contaminant level (MCL) of .007 mg/L. The Safe Drinking Water Act requires that determination of a compound's MCL be based in part on the oral reference dose (RfD) for that compound. In its reevaluation of 1,1-DCE in August 2002, U.S. EPA recalculated the RfD for 1,1-DCE, changing it from 0.009 mg/kg-day to 0.046 mg/kg-day. U.S. EPA published these changes as part of its revision to the file for 1,1-DCE on the Integrated Risk Information System (IRIS). Pursuant to 15A N.C.A.C. 2L.0202(e), IRIS is the most preferred reference source for the establishment of groundwater quality standards.

Even though it made changes to the IRIS file for 1,1-DCE, for reasons of workload, administrative costs and lack of evidence of public cost savings, U.S. EPA decided not to change the MCL for 1,1-DCE despite its revision to IRIS. (See enclosed copy of portions of National Primary Drinking Water Regulations; Announcement of Completion of EPA's Review of Existing Water Standards, 68 Fed. Reg. 42,908, 42,921 (July 18, 2003)). For non-scientific reasons, the current MCL for 1,1-DCE no longer "corresponds" to the IRIS evaluation of 1,1-DCE. Thus, under North Carolina regulations, the current MCL cannot be used to establish the 2L groundwater quality standard for 1,1-DCE.

15A N.C.A.C. 2L.0202(d) provides that groundwater quality standards can also be the lesser of the following for a compound:

- 1. Systemic threshold concentration;
- 2. Concentration corresponding to a 1X10⁻⁶ incremental lifetime cancer risk;
- Taste threshold limit value;
- 4. Odor threshold limit value; or
- 5. National secondary drinking water standard.

1,1-DCE does not have a taste threshold limit value, odor threshold limit value or national secondary drinking water standard. In its reevaluation, U.S. EPA concluded that 1,1-DCE was not a carcinogen and removed the oral cancer slope factor for 1,1-DCE. Thus, there is no cancer risk concentration that can be used to establish a groundwater quality standard.

January 18, 2005 Page 3

The only basis for establishing a groundwater quality standard for 1,1-DCE is the systemic threshold concentration for 1,1-DCE. According to 15A N.C.A.C. 2L.0202(d)(1), the systemic threshold concentration is calculated as follows:

Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)]

Since this formula can use U.S. EPA's recalculated reference dose (RfD) for 1,1-DCE, the systemic threshold concentration "corresponds" with IRIS and thus complies with 15A N.C.A.C. 2L.0202(e). Performing the calculation using the recalculated reference dose, the systemic threshold concentration for 1,1-DCE is .350 mg/L.

Under the proposal before you in February, the groundwater quality standard for 1,1-DCE would continue to be the maximum contaminant level of .007 mg/L. From a scientific and toxicological basis, everyone agrees that this is not correct – rather, the groundwater quality standard for 1,1-DCE should be .350 mg/L. The express language of DENR's regulations for establishing groundwater quality standards mandates the same result – the groundwater quality standard for 1,1-DCE must be .350 mg/L. Please take appropriate action at your meeting in February to adopt this standard.

Sincerely,

HELMS MULLISS & WICKER, PLLC

Benne C. Hutson

Enclosures

cc: Robynn Moraites (w/o encls.)(via electronic mail)
Rhodia Management Team (w/o encls.)(via electronic mail)
Environmental Management Commission Members
Kevin Martin
Andrew Pitner
Frank Crawley
Dr. Luanne K. Williams
Alan Clark
Jeff Manning
David Hance

Benne C. Hutson Direct: 704.343.2060

McGuireWoods

bhutson@mcguirewoods.com Direct Fax: 704.444.8739

May 28, 2009

VIA OVERNIGHT MAIL AND ELECTRONIC MAIL

Sandra Moore
North Carolina Department of
Natural and Environmental Resources
Division of Water Quality
Planning Section
1617 Mail Service Center
Raleigh, North Carolina 27699-1617

Re:

Rhodia Inc., Radiator Specialty Company and Ashland Inc. / Comments on

Proposed Amendments to North Carolina Groundwater Quality Standards – Legal

Requirement to Amend Standard for 1,1-Dichloroethylene

Dear Ms. Moore:

We represent Rhodia Inc., Radiator Specialty Company and Ashland Inc. The purpose of this letter is to provide comments on the proposed changes to the North Carolina groundwater quality standards established by 15A N.C.A.C. 2L.0202(g). These comments are being provided on or before June 1, 2009, the deadline established for the submission of comments in the Notice of the Proposed Rule Amendments as published in the April 1, 2009 edition of the North Carolina Register.

The specific comment being submitted by Rhodia, Radiator Specialty and Ashland is that there is no proposed change to the groundwater quality standard for 1,1-dichloroethylene ("1,1-DCE"). Based upon applicable North Carolina statutes and regulations, including but not limited to 15A N.C.A.C. 2L.0202(d) and (e), the Environmental Management Commission ("EMC") has a mandatory legal obligation to amend the groundwater quality standard for 1,1-DCE from the current standard of 7 micrograms per liter (μ g/L) to 350 μ g/L as the current standard does not correspond to the United States Environmental Protection Agency's revised health risk assessment standard. A toxicologist with the Occupational Environmental Epidemiology Branch of the North Carolina Division of Public Health previously considered this matter in 2004 and recommended that the standard for 1,1-DCE be changed to 350 μ g/L.

The full basis for this comment is set forth in the remainder of this letter.

I. <u>Background</u>

A. Rhodia

Rhodia formerly operated a specialty chemical manufacturing facility in Gastonia. For over a decade now, Rhodia has been operating a groundwater remediation system at this former facility. The primary contaminant of concern is 1,1-DCE.

As part of its remediation, Rhodia paid over \$1 million to provide municipal water to over 100 residents in the surrounding area, thus significantly reducing the risks posed by this site. Operation of the remediation system has prevented and continues to prevent contaminated groundwater from reaching an unnamed tributary on Rhodia's former site. This risk reduction is enhanced as there is no public access to the tributary. The ongoing operation of the groundwater remediation system continues to further reduce the risks through the removal of contaminant mass.

Since 1997, Rhodia has been discharging its treated groundwater effluent to this unnamed tributary pursuant to the terms of an NPDES permit. In 2002, Rhodia received a determination from the Hazardous Waste Section of the Division of Waste Management that the treated effluent would not be characterized as a hazardous waste under the provisions of the Resource Conservation and Recovery Act ("RCRA"). This determination was based upon the finding, supported by years of sampling data, that the treated effluent did not contain levels of contaminants, including 1,1-DCE, that exceeded the lowest known health-based standards for those contaminants. As a result of this determination, the treated effluent now is primarily reused in the manufacturing operations of the current owner of Rhodia's former facility with any amounts that cannot be used in the manufacturing process being discharged to the unnamed tributary pursuant to the terms of the NPDES permit.

B. Radiator Specialty

Radiator Specialty Company operates a manufacturing facility in Indian Trail, North Carolina. Since 1972, Radiator Specialty Company has manufactured and packaged various aerosol and liquid automotive and general purpose chemicals at this facility. In 1987, Radiator Specialty closed two former surface impoundments at the facility. In conjuction with these closure activities, Radiator Specialty initiated the process of applying for a post-closure permit to manage these closed impoundments in accordance with the requirements of RCRA. The initial post-closure permit was issued in 1996 and is currently going through the renewal process.

In 1994, prior to issuance of the permit, Radiator Specialty voluntarily designed and began operation of a groundwater remediation system. That system has been in continuous operation since that time with the treated effluent being discharged to a Union County publicly

owned treatment works. One of the contaminants of concern is 1,1-DCE.

In 2003, Radiator Specialty received a determination from the Hazardous Waste Section of the Division of Waste Management that the treated effluent would not be characterized as a hazardous waste under the provisions of RCRA. The determination was based upon the finding, supported by years of sampling data, that the treated effluent did not contain levels of contaminants, including 1,1-DCE, that exceeded the lowest known health-based standards for those contaminants.

C. Ashland

Ashland has two former facilities in Greensboro and Raleigh where it has been voluntarily operating groundwater remediation systems for several years. In Greensboro, Ashland operated a distribution center for bulk industrial chemicals and solvents from 1968 to 2001. The Raleigh facility operated as a package warehouse and bulk distribution center from the late 1950s until bulk operations ceased in 1988. From that time, the Raleigh facility was used as a distribution warehouse until operations ceased in 2006. Both facilities have been sold to third parties. However, Ashland retains contractual responsibility for the historic contamination at each site.

At the Raleigh facility, Ashland has voluntarily been conducting investigation and remediation activities since 1989. This has included the operation of a groundwater extraction and treatment and a soil vapor extraction system. Similar work has been done at the Greensboro facility. 1,1-DCE is a contaminant of concern at both sites. Ashland is currently negotiating administrative orders on consent with the Hazardous Waste Section of the Division of Waste Management to govern future actions at both sites.

II. The History of the Standard for 1,1-DCE.

The cleanup standard each company has been striving to meet for 1,1-DCE in groundwater at each site has been the historic 2L standard of 7 μg/L. In 2003, the United States Environmental Protection Agency ("U.S. EPA") published its findings on the re-evaluation of risks posed by various compounds. As to 1,1-DCE, U.S. EPA concluded that the risk had been greatly overstated. As part of its re-evaluation, U.S. EPA recalculated the oral reference dose (RfD) for 1,1-DCE, changing it from 0.009 mg/kg/day to 0.050 mg/kg/day. U.S. EPA published these changes as part of its revision to the file for 1,1-DCE in the Integrated Risk Information System ("IRIS"). U.S. EPA subsequently reviewed the IRIS standard for 1,1-DCE on October 28, 2003 and June 22, 2005 and determined that these changes were still appropriate and no further revisions were necessary. (See Section VII. of U.S. EPA's current IRIS file for 1,1-DCE attached as Exhibit A.)

III. North Carolina Laws and Regulations Require that the Groundwater Standard for 1,1-DCE Be Changed from 7 μg/L to 350 μg/L.

In its proposed revisions to the 2L groundwater quality standards, the EMC and NCDENR have not proposed to make any change to the current standard for 1,1-DCE. Rather, the standard for 1,1-DCE would remain at 7 μ g/L. However, keeping the 1,1-DCE standard the same violates North Carolina laws and regulations applicable to the establishment of groundwater quality standards. To comply with such statutory and regulatory requirements, the EMC and NCDENR must base the standard for 1,1-DCE on the most current and accurate toxicological data and, based upon that data, must increase the groundwater quality standard for 1,1-DCE to 350 μ g/L.

A. 15A N.C.A.C. 2L.0202(d) and (e) – The North Carolina Regulations Governing the Establishment of Groundwater Quality Standards.

Two regulations govern the establishment of groundwater quality standards in North Carolina. First, under 15A N.C.A.C. 2L.0202(d), the lesser of six different standards is to be used to set a compound's groundwater limit. Second, 15A N.C.A.C. 2L.0202(e) requires the EMC and NCDENR to establish groundwater standards based upon four sources of toxicity data listed "in order of preference" in the regulation. Although separate, 15A N.C.A.C. 2L.0202(d) and (e) are not independent regulations. Rather, they must be read in tandem because the sources of toxicity data for establishing standards listed in 15A N.C.A.C. 2L.0202(e) must "correspond to levels described in Paragraph (d) of this Rule". (Emphasis added.) From this it follows that if a basis for a groundwater quality standard listed in 15A N.C.A.C. 2L.0202(d) does not "correspond" to an accepted reference source listed in 15A N.C.A.C. 2L.0202(e), then the standard in subparagraph (d) cannot be used. (For ease of reference, copies of these two regulations are enclosed with this letter as Exhibit B.)

This is the current situation with regard to 1,1-DCE. On its face, the "lesser" of the six identified standards in 15A N.C.A.C. 2L.0202(d) is the Federal Safe Drinking Water Act's maximum contaminant level ("MCL") of 7 µg/L. The Safe Drinking Water Act requires that determination of an MCL for a compound must be based in part on the oral reference dose (RfD) for that compound. In its re-evaluation of 1,1-DCE in 2003, U.S. EPA recalculated the RfD for 1,1-DCE, changing it from 0.009 mg/kg/day to 0.050 mg/kg/day. U.S. EPA published these changes as part of its revision to the file for 1,1-DCE in IRIS. (See Section I.A.1. of EPA's current IRIS file for 1,1-DCE attached as Exhibit A.) Pursuant to 15A N.C.A.C. 2L.0202(e), IRIS is the most preferred reference source for the establishment of groundwater quality standards.

For reasons of competing workload priorities, administrative costs and lack of evidence of public cost savings, U.S. EPA in 2002 decided not to change the MCL for 1,1-DCE despite its revision to IRIS. (See National Primary Drinking Water Regulations; Announcement of Completion of EPA's Review of Existing Water Standards, 68 Fed. Reg. 42908, 42921 (July 18, 2003) attached as Exhibit C). That situation remains the same today. As a result, for non-

Page 5

scientific and non-toxicological reasons, the current MCL for 1,1-DCE no longer "corresponds" to the IRIS evaluation of 1,1-DCE. Thus, under North Carolina regulations, the current MCL cannot be used to establish the 2L groundwater quality standard for 1,1-DCE.

B. Under 15A N.C.A.C. 2L.0202(d) and (e), the Groundwater Quality Standard for 1,1-DCE Must be 350 μg/L.

15A N.C.A.C. 2L.0202(d) provides that besides an MCL, groundwater quality standards can also be the lesser of the following for a compound:

- 1. Systemic threshold concentration;
- 2. Concentration corresponding to a 1 X 10⁻⁶ incremental lifetime cancer risk;
- 3. Taste threshold limit value;
- 4. Odor threshold limit value; or
- 5. National secondary drinking water standard.

1,1-DCE does not have a taste threshold limit value, odor threshold limit value or national secondary drinking water standard. In its IRIS revisions, U.S. EPA removed the oral cancer slope factor for 1,1-DCE and concluded that the compound was only a possible human carcinogen. As a result, there is no cancer risk concentration that can be used to establish a groundwater quality standard.

The only basis for establishing a groundwater quality standard for 1,1-DCE is the systemic threshold concentration for 1,1-DCE. According to 15A N.C.A.C. 2L.0202(d)(1), the systemic threshold concentration is calculated as follows:

[Reference dose mg/kg/day x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganic; .20 for organics)] / [2 liters/day (avg. water consumption)]

Since this formula can use U.S. EPA's recalculated RfD for 1,1-DCE, the systemic threshold concentration "corresponds" with IRIS and thus complies with 15A N.C.A.C. 2L.0202(e). Performing the calculation using the recalculated RfD of 0.050 and a relative source contribution of .20 as 1,1-DCE is an organic compound, the systemic threshold concentration for 1,1-DCE is $350~\mu g/L$, 50~times higher than the current 2L standard.

IV. Rhodia Made These Same Comments in 2004 and the State's Toxicologist Agreed that the Standard for 1,1-DCE Should Be 350 μg/L.

Rhodia brought these same points to the attention of the EMC and NCDENR in September 4, 2004 when the EMC last considered revisions to North Carolina's groundwater quality standards. Those comments were set forth in my letter to David Hance of the Division of Water Quality dated September 28, 2004. (A copy of this letter is attached as Exhibit D.) Similar comments had also been submitted by Dr. Shawn L. Sager, a principal scientist with Arcadis G&M of North Carolina, Inc. in a letter dated September 13, 2004. (A copy of this letter is included as Exhibit E.) These letters were reviewed by Dr. Luanne K. Williams, a toxicologist with the Occupational Environmental Epidemiology Branch of the North Carolina Division of Public Health. In comments enclosed with a letter dated October 1, 2004, Dr. Williams concluded that based upon the information provided in Rhodia's public comments, "the recommended groundwater quality standard for 1,1-dichloroethylene is 350 µg/L and not 7 µg/L." (Emphasis added.) A copy of Dr. Williams' letter is included as Exhibit F.

Despite the uncontroverted information presented in Rhodia's and Dr. Sager's 2004 public comments and the uncontroverted recommendation of Dr. Williams, the State's toxicologist, the 2004 hearing officers recommended that the standard for 1,1-DCE remain the same, erroneously concluding that the groundwater quality regulations in 15 N.C.A.C. Subchapter 2L would have to be amended in order for the 1,1-DCE standards to be changed. To the contrary, the regulations as they existed at that time required (as previously explained in this letter) that the standard for 1,1-DCE had to be changed to 350 μ g/L. Rhodia presented this position to Dr. David H. Moreau, who was then chairman of the EMC, in a January 18, 2005 letter. (A copy of that letter is attached as Exhibit G.) Even though North Carolina's toxicologist agreed and North Carolina law and regulations required that the groundwater quality standard for 1,1-DCE had to be set at 350 μ g/L, the EMC did not make that change but rather kept the standard at 7 μ g/L.

V. <u>U.S. EPA Has Changed the National Recommended Water Quality Criteria for 1,1-DCE Based on the Change in the RfD.</u>

As previously stated, for non-scientific reasons U.S. EPA has not changed the drinking water standard for 1,1-DCE. However, U.S. EPA, based upon its 2003 revised risk assessment, has changed the recommended surface water quality standard for 1,1-DCE. On December 31, 2003, U.S. EPA published changes to the national recommended water quality criteria for the protection of human health for 15 compounds including 1,1-DCE. 68 Fed. Reg. 75507 (December 31, 2003). These criteria are numeric values that describe ambient water concentrations that protect human health from the harmful effects of pollutants. According to U.S. EPA, "these criteria are based solely on data and scientific judgments about the relationship between pollutant concentrations and environmental and human health effects." 68 Fed. Reg. 75509.

At the time these changes were published, the water quality criteria for 1,1-DCE was 0.057 μ g/L. In the revisions to the criteria for 1,1-DCE, U.S. EPA "incorporated into the recalculations . . . a new reference dose (RfD)" of 0.050 mg/kg/day. 68 Fed. Reg. 75510. This was the revised RfD that EPA published in its re-evaluation of the risk posed by 1,1-DCE. Based on the use of this revised RfD, the national recommended water quality criteria for 1,1-DCE went from 0.057 μ g/L to 330 μ g/L, an increase of nearly 6,000 times the previous standard. That revised standard remains in effect today. (A copy of the December 31, 2003 revisions to the national recommended water quality criteria is attached as Exhibit H.)

VI. Conclusion

The legal and regulatory situation regarding 1,1-DCE remains the same today as it did the last time the EMC considered revisions to North Carolina's groundwater quality standards. Under applicable North Carolina laws and regulations, specifically 15A N.C.A.C. 2L.0202(d) and (e), the EMC and NCDENR must change the groundwater quality standard for 1,1-DCE to 350 μ g/L. If they fail to do so, the EMC and NCDENR are failing to act as required by law and are acting erroneously. Furthermore, since they had the same information in 2004 and failed then to take the legally required action and since they are being presented with the same information and legal standards today, if the EMC and NCDENR do not make this change to the standard this time, they will be acting arbitrarily and capriciously. All of these form a basis which expose the EMC and NCDENR to a successful administrative challenge to their failure to act under the North Carolina Administrative Procedure Act. To avoid such a challenge, the EMC and NCDENR must change the groundwater quality standard for 1,1-DCE from 7 μ g/L to 350 μ g/L.

Sincerely,

McGuireWoods LLP

Benne C. Hutson

Enclosures

cc: Rhodia Management Team (w/encls.)

Radiator Specialty Management Team (w/encls.)

Ashland Management Team (w/encls.)

STATE OF NORTH CAROLINA DEPARTMENT OF ENVIRONMENT AND NATURAL RESOURCES

REPORT OF PROCEEDINGS TO THE NORTH CAROLINA ENVIRONMENTAL MANAGEMENT COMMISSION FOR THE PROPOSED REVISIONS TO THE 15A NCAC 02L .0202 GROUNDWATER QUALITY STANDARDS

PUBLIC HEARINGS

- April 21, 6:30 PM Western Piedmont Community College Moore Hall, 1001 Burkemont Avenue, Morganton
- April 23, 6:30 PM Bladen Community College Multipurpose Auditorium Building, 7418 Highway 41 West, Dublin
- April 30, 6:30 PM Archdale Building, Ground Floor Hearing Room, 512 N. Salisbury Street, Raleigh

Hearing Officers' Report Triennial Review of Groundwater Standards September 10, 2009

Noting that this information could be of great importance to clearly identifying the human risk, staff agrees that maintaining the current standard would be adequately protective of public health

1,1 DICLOROETHYLENE (1,1-DCE)

Proposed Standard: 7 ug/L Current Standard: 7 ug/L

26. Comments received in regard to the 1,1,-dichloroethylene standard from McGuireWoods LLP (Benne C. Hutson) representing Rhodia Inc., Radiator Specialty

There is no proposed change to the groundwater quality standard for 1,1dichloroethylene. Based upon applicable North Carolina statutes and regulations, including but not limited to 15A N.C.A.C. 2L.0202 (d) and (e), the Environmental Management Commission ("EMC") has a mandatory legal obligation to amend the groundwater quality standard for 1,1-DCE from the current standard of 7 micrograms per liter (ug/L) to 350 ug/L as the current standard does not correspond to the United States Environmental Protection Agency's revised health risk assessment standard. A toxicologist with the Occupational Environmental Epidemiology Branch of the North Carolina Division of Public Health previously considered this matter in 2004 and recommended that the standard for 1,1-DCE be changed to 350 ug/L.(See letter for further detail)

15A NCAC 2L .0202(d) and (e) although separate are not independent regulations. The sources of toxicity data for establishing standards listed in .0202(e) must "correspond to levels described in paragraph (d) of this Rule". The USEPA has not changed the MCL due to competing workload priorities, administrative cost and lack of evidence of pubic cost savings despite its revision of the oral reference dose from 0.009 mg/kg/day to 0.050 mg/kg/day in the IRIS database. The IRIS database is listed in .0202(e) as the preferred

By not amending the standard to the lowest known health based-standard, the EMC and DENR are failing to act as required by law and are acting erroneously. Further, they are acting arbitrarily and capriciously and may expose themselves to a successful administrative challenge to their failure to act under the NCAPA.

Response: DENR regulations require that groundwater standards be established in accordance with 15A NCAC 021 .0202 (d) -- the least of the six criteria.

The first two criteria, a systemic threshold concentration and a concentration corresponding to an incremental lifetime cancer risk of one-in-a-million, are concentrations calculated by DENR using the references cited in .0202 (e). The remaining four criteria used to establish a groundwater standard--taste threshold, odor threshold, the federal maximum contaminant level and the National secondary drinking water standard -- are concentrations that have been

Hearing Officers' Report Triennial Review of Groundwater Standards September 10, 2009

EPA establishes the Federal Primary Drinking Water Standards and Secondary Drinking Water Standards according to federal regulations. The references set out in 15A NCAC 02L .0202 (e) are intended for DENR use when calculating the criteria in accordance with 15A NCAC 02L .0202 (d)(1) and (d)(2).

This issue led to a Groundwater Stakeholder Work Group, made up of industry, environmental groups and DENR agencies that met between July 14, 2005 and December 1, 2005. The Stakeholder Group, led by Carl Bailey, DWQ, and Kevin Martin, EMC, discussed needed changes to the groundwater rules to update groundwater standards so that the most up-to-date toxicity information was being incorporated. In the end, the EMC Groundwater Committee decided not to propose any changes to the groundwater rules stating that the 1,1,-DCE issue could be dealt with using the variance process (Attachment A-20).

VIII. RECOMMENDATIONS

Following a careful and comprehensive review of all of the submitted written and oral comments, supporting data, and attachments to this record, the Hearing Officers recommend that the North Carolina Environmental Management Commission adopt the changes to the groundwater quality standards as proposed with the following modifications:

 Based on information obtained from the US EPA (Office of Water), comments, concerns and implementation issues noted during the public hearing process, the Hearing Officers (HOs) and staff recommend adoption of an arsenic standard at 10 ppb rather than the 0.02 ppb proposed. This recommendation is based upon the following information.

According to the US EPA chemical manager for the inorganic arsenic assessment in the Office of Water, the final draft Toxicological Review for Inorganic Arsenic has been completed recently and is scheduled for public release in the near future. This latest assessment contains toxicity indices based on lung and bladder cancers. These internal cancers also served as endpoints for the Arsenic Rule in 2001 which established the Federal Drinking Water Standard of 10 ug/L. The available assessment in IRIS is based solely on skin cancer and does not address the known lung and bladder cancer risk.

The Federal Drinking Water Standard of 10 ppb is considered by EPA to be protective of human health in drinking water sources. The HOs recommend that under 15A NCAC 02L .0103, establishment of a standard for arsenic at 10 ppb meets the Commission's policy for protection of the groundwater as a drinking water source and establishes a level that is protective of public health.

The HOs make note here that a substantial number of comments were received from concerned parties with respect to the proposed arsenic level. The overwhelming majority disapproved of the proposed level due to the significant financial burden compliance would place on them at this time. Concerned parties also noted issues with naturally-occurring background levels that range from non-detect (practical quantitation limit at 1 ppb) to 190 ppb across the state. Of note, laboratories routinely detect and quantitate arsenic at approximately 1 - 2 ppb.

Hearing Officers' Report Triennial Review of Groundwater Standards September 10, 2009

A-20 EMC Groundwater Committee 1,1-DCE Recommendation

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CAMENO SOURCE SERVED

1996 to Present

Date

Agenda Title

Recommendation that the EMC use existing Subchapter 2L Rules to Resolve conflicts between Criteria in 15A NCAC 2L.0202 (d) and (e)

Type

GWC Action

The GWC unamimoulsy accepts & concurs with the Division recommendation that in individual site specific cases where there is a difference between 2L.0202 (d)(1) requirements in the 2L.0202 Groundwater Standards and the Federal MCLs, a variance under 2L.0113 could be approved that would allow less restrictive Groundwater Quality Standards while providing the site specific . requirements necessary to protect Public Water Supplies

Incident Number

Location

711D

Date

5/9/2007

Agenda Title

INFORMATION--Results from the Groundwater Standards Stakeholders

Process- Stakeholders Recommendations regarding the use of Federal MCLs as

Type

NC groundwater standards & the modification of the Relative Source

Contribution component in 2L.0202(d)(1)

GWC Action

Incident Number

Location

711D

Table 8			
Summary of Remedial Technologies			
207 Telegraph Drive, Gastonia, North Card	nlina		
201 Telegraph 21110, Gastonia, Trottin Gast			
	Containment		
	Containment		
-	Hydraulic Five groundwater extraction wells currently operating; providing plume containment and mass reduction.		
Physical Barriers	Not practical to install slurry walls as deep as needed. Grouting is possible but		
(Slurry wall, grouting)	not likely to be completely effective. Would address migration through		
	bedrock fractures.		
Permeable Reactive Barriers	Depth is too great and would be difficult to install into the bedrock.		
(Zero Valent Iron, bio, e-barriers)			
Phytoremediation	Depth is too great.		
,			
Source Zo	ne Restoration/Partial Mass Removal		
Electrical Resistive Heating	Installation of electrodes into the bedrock paired with a vapor extraction		
(three-phase or six-phase heating)	system. Resistive heating would "boil off" the VOCs, which would then be		
	captured and treated. High potential for removing a significant mass of		
	DNAPL. Installation into the bedrock would be technically challenging and		
	very expensive. Potential concern with enhancing migration of possible DNAPL as the matrix is heated. May have to heat first from below the possible		
	DNAPL as the matrix is heated. May have to hear first from below the possible DNAPL zone ("hot floor" method - not readily applicable to bedrock).		
	Divid B Zone (not noor incurous incurous incurous applicable to occitock).		
Steam Flushing	Contact with contaminants not likely to be complete in fractured bedrock.		
In-Situ Oxidation	Injection of oxidizing chemicals such as hydrogen peroxide or potassium		
	permanganate. Difficult to inject the chemicals into the bedrock fractures where		
	possible DNAPL exists.		
In-Situ Reduction	Injection of reducing chemicals such as a slurry of zero-valent iron. Difficult to		
	locate and inject the chemicals into the bedrock fractures where possible		
	DNAPL exists.		
Dual-Phase Extraction	Application of a high vacuum to deep wells to extract mobile DNAPL,		
	contaminated groundwater, and vapors. If it is possible to dewater the		
	formation, soil vapor extraction can be performed. Even if possible to implement this technology, it will still leave a mass of possible DNAPL.		
Enhanced Reductive Dechlorination	Injection of vegetable oil or other electron donor to stimulate reductive dechlorination. Difficult to locate and inject the chemicals into the bedrock		
·	fractures where possible DNAPL exists.		

Generated by: Daniel Brown

Checked by: Jessica High

Appendix B: Petition Summary and Background Information

On March 16, 2011, McGuireWoods, on behalf of Rhodia Inc., filed a petition for rulemaking to the Division of Water Quality Director (DWQ), Coleen Sullins. The petition requests an amendment of the groundwater quality standard for 1,1-dichloroethylene (1,1-DCE) contained in 15A NCAC 2L .0202(g)(59) from 7 ug/L to 350 ug/L.

By regulation, groundwater standards are established as the lower of the six criteria contained in 15A NCAC 2L .0202(d) (1) – (6). Based on these criteria, the current standard for 1,1-DCE is the maximum contaminant level (MCL) of 7 ug/L. MCLs are federal drinking water standards established by the USEPA Office of Water and are applicable to public water supply systems regulated under the Federal Safe Drinking Water Act.

The petitioner seeks to amend the groundwater standard because the federal MCL for 1,1-DCE was calculated using outdated health effects data. The DWQ and the USEPA acknowledge that updated health effects data support calculation of a less stringent MCL. However, EPA does not plan to update the MCL because any potential revision is not likely to provide a meaningful opportunity for cost-savings or health risk reduction to public water systems and their customers. A revised standard of 350 ug/L would reduce cleanup costs for Rhodia and other sites.

The petitioner provided a legal opinion that 15A NCAC 2L .0202(d) and (e), in tandem, are sufficiently broad to establish the groundwater standard at 350 ug/L under 2L .0202(d)(1), based on the current toxicity data published in the USEPA's Integrated Risk Information System (IRIS) database, rather than the federal MCL, which was calculated prior to the updated toxicity data being published.

The 1,1-DCE standard issue was first brought forth by Rhodia and others during the Groundwater Triennial Review (GWTR) that ended in 2005 and again during the GWTR that ended in 2010. Both times the Environmental Management Committee (EMC) approved a 1,1,DCE groundwater standard of 7 ug/L. After consultation with its legal counsel, Frank Crawley, the EMC determined that 2L .0202(d) and (e) were not sufficiently broad to allow a change to the 1,1-DCE standard from 7 ug/L to 350 ug/L as requested.

In May 2005, the EMC Groundwater Committee (GWC) directed the DWQ to establish a Groundwater Stakeholder Group (GWSG) to discuss ways to amend the groundwater rules to ensure the use of the most recent toxicity information when developing groundwater standards. The GWSG consisted of representatives from various stakeholder groups, such as, Department of Environment and Natural Resources (DENR) regulatory agencies, city and county governments, major industries, environmental groups, agricultural interests, and public health. After much discussion, the GWSG could not reach consensus on how to amend the groundwater regulations and ultimately, the DWQ recommended to the GWC that the issue be dealt with by the variance process in 2L .0113. At its July 12, 2006 meeting, the GWC unanimously accepted and concurred with the DWQ recommendation that in individual site-specific cases, a variance under 2L .0113 could be approved that would allow less restrictive Groundwater Quality Standards while providing the site specific requirements necessary to protect public water supplies.

In November 2010, Rhodia submitted a site-specific variance request for a 1,1-DCE standard of 350 ug/L. The Division of Waste Management (DWM), the regulatory authority over the site, reviewed the request and determined it to be incomplete based on the requirements in 2L .0113. Rhodia withdrew the variance request, stating that the variance approach was not an appropriate mechanism for seeking relief from a standard that was not based on current health effects information. Subsequently, Rhodia submitted the rulemaking petition.

At its May 2011 meeting, the GWC heard presentations on the rulemaking petition from the petitioner and DWQ staff. DWQ staff recommended that the petition be denied and that Rhodia work with DWM staff to address the deficiencies identified in their variance request.

After discussion, the GWC passed a motion to recommend that the full EMC proceed with rulemaking as proposed by the petitioner to amend the 1,1-DCE standard from 7 ug/L to 350 ug/L. The Committee acknowledged that, according to legal counsel, rule language is needed to allow deviation from 2L .0202(d), which requires that the groundwater standard be established at the lowest of the six criteria.

On July 14, 2011, the EMC approved Rhodia's Rulemaking Petition and directed the DWQ to initiate rulemaking to amend the groundwater standard for 1,1-DCE from 7 ug/L to 350 ug/L.

15A NCAC 02L .0202 is proposed for amendment as follows: (Option 1)

15A NCAC 02L .0202 GROUNDWATER QUALITY STANDARDS

- (a) The groundwater quality standards for the protection of the groundwaters of the state are those specified in this Rule. They are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of the state, which may be tolerated without creating a threat to human health or which would otherwise render the groundwater unsuitable for its intended best usage.
- (b) The groundwater quality standards for contaminants specified in Paragraphs (g) and (h) of this Rule are as listed, except that:
 - (1) Where the standard for a substance is less than the practical quantitation limit, the detection of that substance at or above the practical quantitation limit constitutes a violation of the standard.
 - (2) Where two or more substances exist in combination, the Director shall consider the effects of chemical interactions as determined by the Division of Public Health and may establish maximum concentrations at values less than those established in accordance with Paragraphs (c), (g), or (h) of this Rule. In the absence of information to the contrary, in accordance with Paragraph (d) of this Rule, the carcinogenic risks associated with carcinogens present shall be considered additive and the toxic effects associated with non-carcinogens present shall also be considered additive.
 - Where naturally occurring substances exceed the established standard, the standard shall be the naturally occurring concentration as determined by the Director.
- (c) Except for tracers used in concentrations which have been determined by the Division of Public Health to be protective of human health, and the use of which has been permitted by the Division, substances which are not naturally occurring and for which no standard is specified shall not be permitted in concentrations at or above the practical quantitation limit in Class GA or Class GSA groundwaters. Any person may petition the Director to establish an interim maximum allowable concentration for a substance for which a standard has not been established under this Rule. The petitioner shall submit relevant toxicological and epidemiological data, study results, and calculations necessary to establish a standard in accordance with Paragraph (d) of this Rule. Within three months after the establishment of an interim maximum allowable concentration for a substance by the Director, the Director shall initiate action to consider adoption of a standard for that substance.
- (d) Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of:
 - (1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)] / [2 liters/day (avg. water consumption)];
 - (2) Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6;
 - (3) Taste threshold limit value:
 - (4) Odor threshold limit value;
 - (5) Maximum contaminant level; or
 - (6) National secondary drinking water standard.
- (e) The following references, in order of preference, shall be used in establishing concentrations of substances which correspond to levels described in Paragraph (d) of this Rule.
 - (1) Integrated Risk Information System (U.S. EPA).
 - (2) Health Advisories (U.S. EPA Office of Drinking Water).
 - (3) Other health risk assessment data published by U.S. EPA.
 - (4) Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.
- (f) Groundwater quality standards specified in Paragraphs (g) and (h) of this Rule and interim maximum allowable concentrations established pursuant to Paragraph (c) of this Rule shall be reviewed on a triennial basis. Appropriate modifications to established standards shall be made in accordance with the procedure prescribed in Paragraph (d) of this Rule where modifications are considered appropriate based on data published subsequent to the previous review.
- (g) Class GA Standards. Where not otherwise indicated, the standard refers to the total concentration in micrograms per liter of any constituent in a dissolved, colloidal or particulate form which is mobile in groundwater. This does not apply to sediment or other particulate matter which is preserved in a groundwater sample as a result of well construction or sampling procedures. The Class GA standards are:
 - (1) Acenaphthene: 80;
 - (2) Acenaphthylene: 200;

```
1
               (3)
                        Acetone: 6 mg/L;
 2
               (4)
                       Acrylamide: 0.008;
 3
               (5)
                        Anthracene: 2 mg/L;
 4
               (6)
                        Arsenic: 10;
 5
               (7)
                        Atrazine and chlorotriazine metabolites: 3;
 6
                        Barium: 700;
               (8)
 7
               (9)
                        Benzene: 1;
 8
               (10)
                        Benzo(a)anthracene (benz(a)anthracene): 0.05;
 9
               (11)
                       Benzo(b)fluoranthene: 0.05;
10
               (12)
                        Benzo(k)fluoranthene: 0.5;
                       Benzoic acid: 30 mg/L;
11
               (13)
12
               (14)
                       Benzo(g,h,i,)perylene: 200;
13
               (15)
                        Benzo(a)pyrene: 0.005;
14
               (16)
                        Bis(chloroethyl)ether: 0.03;
15
               (17)
                        Bis(2-ethylhexyl) phthalate (di(2-ethylhexyl) phthalate): 3;
               (18)
16
                       Boron: 700;
               (19)
                       Bromodichloromethane: 0.6;
17
18
               (20)
                       Bromoform (tribromomethane): 4;
19
               (21)
                       n-Butylbenzene: 70;
20
               (22)
                       sec-Butylbenzene: 70;
21
                       tert-Butylbenzene: 70;
               (23)
22
               (24)
                       Butylbenzyl phthalate: 1 mg/L;
23
               (25)
                        Cadmium: 2;
24
               (26)
                        Caprolactam: 4 mg/L;
25
                        Carbofuran: 40;
               (27)
26
               (28)
                        Carbon disulfide: 700;
27
               (29)
                       Carbon tetrachloride: 0.3;
28
                        Chlordane: 0.1;
               (30)
29
               (31)
                       Chloride: 250 mg/L;
30
               (32)
                        Chlorobenzene: 50;
31
               (33)
                       Chloroethane: 3,000;
32
               (34)
                        Chloroform (trichloromethane): 70;
33
                       Chloromethane (methyl chloride): 3;
               (35)
                       2-Chlorophenol: 0.4;
34
               (36)
35
               (37)
                       2-Chlorotoluene (o-chlorotoluene): 100;
36
               (38)
                       Chromium: 10;
37
               (39)
                        Chrysene: 5;
38
               (40)
                       Coliform organisms (total): 1 per 100 milliliters;
39
               (41)
                        Color: 15 color units;
40
               (42)
                       Copper: 1 mg/L;
41
               (43)
                        Cyanide (free cyanide): 70;
42
               (44)
                        2, 4-D (2,4-dichlorophenoxy acetic acid): 70;
43
                       DDD: 0.1;
               (45)
44
                       DDT: 0.1;
               (46)
45
               (47)
                       Dibenz(a,h)anthracene: 0.005;
46
               (48)
                       Dibromochloromethane: 0.4;
47
               (49)
                        1,2-Dibromo-3-chloropropane: 0.04;
48
               (50)
                        Dibutyl (or di-n-butyl) phthalate: 700;
49
               (51)
                        1,2-Dichlorobenzene (orthodichlorobenzene): 20;
50
               (52)
                        1,3-Dichlorobenzene (metadichlorobenzene): 200;
51
               (53)
                        1,4-Dichlorobenzene (paradichlorobenzene): 6;
52
               (54)
                        Dichlorodifluoromethane (Freon-12; Halon): 1 mg/L;
53
               (55)
                        1,1-Dichloroethane: 6;
54
                        1,2-Dichloroethane (ethylene dichloride): 0.4;
               (56)
55
                        1,2-Dichloroethene (cis): 70;
               (57)
```

```
1
               (58)
                        1,2-Dichloroethene (trans): 100;
 2
               (59)
                       1,1-Dichloroethylene (vinylidene chloride): 7350;
 3
               (60)
                        1,2-Dichloropropane: 0.6;
 4
               (61)
                        1,3-Dichloropropene (cis and trans isomers): 0.4;
 5
               (62)
                       Dieldrin: 0.002;
 6
                        Diethylphthalate: 6 mg/L;
               (63)
 7
               (64)
                        2,4-Dimethylphenol (m-xylenol): 100;
 8
               (65)
                       Di-n-octyl phthalate: 100;
 9
               (66)
                        1,4-Dioxane (p-dioxane): 3;
10
               (67)
                        Dioxin (2,3,7,8-TCDD): 0.0002 ng/L;
                        1,1- Diphenyl (1,1,-biphenyl): 400;
11
               (68)
12
               (69)
                       Dissolved solids (total): 500 mg/L;
13
               (70)
                        Disulfoton: 0.3;
14
                       Diundecyl phthalate (Santicizer 711): 100;
               (71)
15
               (72)
                        Endosulfan: 40;
16
               (73)
                        Endrin, total: (includes endrin, endrin aldehyde and endrin ketone): 2;
                        Epichlorohydrin: 4;
17
               (74)
18
               (75)
                       Ethyl acetate: 3 mg/L;
19
               (76)
                       Ethylbenzene: 600;
20
               (77)
                       Ethylene dibromide (1,2-dibromoethane): 0.02;
21
               (78)
                       Ethylene glycol: 10 mg/L;
22
               (79)
                       Fluoranthene: 300;
23
               (80)
                       Fluorene: 300;
24
               (81)
                       Fluoride: 2 mg/L;
25
               (82)
                       Foaming agents: 500;
26
               (83)
                        Formaldehyde: 600;
27
               (84)
                        Gross alpha (adjusted) particle activity (excluding radium-226 and uranium): 15 pCi/L;
28
                        Heptachlor: 0.008;
               (85)
29
               (86)
                       Heptachlor epoxide: 0.004;
30
               (87)
                       Heptane: 400;
31
               (88)
                       Hexachlorobenzene (perchlorobenzene): 0.02;
32
               (89)
                       Hexachlorobutadiene: 0.4;
33
               (90)
                       Hexachlorocyclohexane isomers (technical grade): 0.02;
                       n-Hexane: 400;
34
               (91)
35
               (92)
                        Indeno(1,2,3-cd)pyrene: 0.05;
               (93)
                       Iron: 300;
36
37
               (94)
                        Isophorone: 40;
38
               (95)
                       Isopropylbenzene: 70;
39
               (96)
                       Isopropyl ether: 70;
40
               (97)
                       Lead: 15;
41
               (98)
                        Lindane (gamma hexachlorocyclohexane): 0.03;
               (99)
42
                       Manganese: 50;
                       Mercury: 1;
43
               (100)
44
               (101)
                       Methanol: 4 mg/L;
                        Methoxychlor: 40;
45
               (102)
46
               (103)
                        Methylene chloride (dichloromethane): 5;
47
                        Methyl ethyl ketone (2-butanone): 4 mg/L;
               (104)
48
               (105)
                        2-Methylnaphthalene: 30;
49
                        3-Methylphenol (m-cresol): 400;
               (106)
50
                        4-Methylphenol (p-cresol): 40;
               (107)
51
               (108)
                       Methyl tert-butyl ether (MTBE): 20;
52
               (109)
                       Naphthalene: 6;
53
               (110)
                       Nickel: 100;
54
                       Nitrate: (as N) 10 mg/L;
               (111)
55
               (112)
                       Nitrite: (as N) 1 mg/L;
```

```
1
               (113)
                        N-nitrosodimethylamine: 0.0007;
 2
               (114)
                        Oxamvl: 200:
 3
                        Pentachlorophenol: 0.3;
               (115)
 4
               (116)
                        Petroleum aliphatic carbon fraction class (C5 - C8): 400:
 5
               (117)
                        Petroleum aliphatic carbon fraction class (C9 - C18): 700;
 6
                        Petroleum aliphatic carbon fraction class (C19 - C36): 10 mg/L;
               (118)
 7
               (119)
                        Petroleum aromatics carbon fraction class (C9 - C22): 200;
 8
               (120)
                        pH: 6.5 - 8.5;
 9
               (121)
                        Phenanthrene: 200:
10
                        Phenol: 30;
               (122)
                        Phorate: 1;
11
               (123)
                        n-Propylbenzene: 70;
12
               (124)
13
               (125)
                        Pyrene: 200;
14
                        Selenium: 20;
               (126)
                        Silver: 20;
15
               (127)
16
               (128)
                        Simazine: 4;
                        Styrene: 70;
17
               (129)
18
               (130)
                        Sulfate: 250 mg/L;
19
               (131)
                        1,1,2,2-Tetrachloroethane: 0.2;
                        Tetrachloroethylene (perchloroethylene; PCE): 0.7;
20
               (132)
                        2,3,4,6-Tetrachlorophenol: 200;
21
               (133)
22
                        Toluene: 600;
               (134)
23
                        Toxaphene: 0.03;
               (135)
24
               (136)
                        2, 4, 5,-TP (Silvex): 50;
25
                        1,2,4-Trichlorobenzene: 70;
               (137)
26
               (138)
                        1,1,1-Trichloroethane: 200;
27
               (139)
                        Trichloroethylene (TCE): 3;
28
                        Trichlorofluoromethane: 2 mg/L;
               (140)
29
               (141)
                        1,2,3-Trichloropropane: 0.005;
30
               (142)
                        1,2,4-Trimethylbenzene: 400;
31
               (143)
                        1,3,5-Trimethylbenzene: 400;
32
                        1,1,2-Trichloro-1,2,2-trifluoroethane (CFC-113): 200 mg/L;
               (144)
33
                        Vinyl chloride: 0.03;
               (145)
                        Xylenes (o-, m-, and p-): 500; and
34
               (146)
35
               (147)
                        Zinc: 1 mg/L.
36
      (h) Class GSA Standards. The standards for this class are the same as those for Class GA except as follows:
37
               (1)
                        chloride: allowable increase not to exceed 100 percent of the natural quality concentration; and
38
               (2)
                        total dissolved solids: 1000 mg/l.
39
      (i) Class GC Waters.
40
                        The concentrations of substances which, at the time of classification, exceed the standards applicable
               (1)
41
                        to Class GA or GSA groundwaters shall not be caused to increase, nor shall the concentrations of other
42
                        substances be caused to exceed the GA or GSA standards as a result of further disposal of
43
                        contaminants to or beneath the surface of the land within the boundary of the area classified GC.
44
                        The concentrations of substances which, at the time of classification, exceed the standards applicable
               (2)
45
                        to GA or GSA groundwaters shall not be caused to migrate as a result of activities within the boundary
46
                        of the GC classification, so as to violate the groundwater or surface water quality standards in
47
                        adjoining waters of a different class.
48
               (3)
                        Concentrations of specific substances, which exceed the established standard at the time of
49
                        classification, are listed in Section .0300 of this Subchapter.
50
51
                        Authority G.S. 143-214.1; 143B-282(a)(2);
      History Note:
52
                        Eff. June 10, 1979;
53
                        Amended Eff. November 1, 1994; October 1, 1993; September 1, 1992; August 1, 1989;
54
                        Temporary Amendment Eff. June 30, 2002;
55
                        Amended Eff. August 1, 2002;
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1 Temporary Amendment Expired February 9, 2003; 2 Amended Eff. November 2012; January 1, 2010; April 1, 2005. 3 15A NCAC 02L .0202 is proposed for amendment as follows: (Option 2)

1 2 3

GROUNDWATER QUALITY STANDARDS 15A NCAC 02L .0202

(a) The groundwater quality standards for the protection of the groundwaters of the state are those specified in this Rule. They are the maximum allowable concentrations resulting from any discharge of contaminants to the land or waters of the state, which may be tolerated without creating a threat to human health or which would otherwise render the groundwater unsuitable for its intended best usage. (b) The groundwater quality standards for contaminants specified in Paragraphs (g) and (h) of this Rule are as listed,

except that:

Where the standard for a substance is less than the practical quantitation limit, the detection of that (1)

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14

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9

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substance at or above the practical quantitation limit constitutes a violation of the standard. Where two or more substances exist in combination, the Director shall consider the effects of chemical (2) interactions as determined by the Division of Public Health and may establish maximum concentrations at values less than those established in accordance with Paragraphs (c), (g), or (h) of this Rule. In the absence of information to the contrary, in accordance with Paragraph (d) of this Rule, the carcinogenic risks associated with carcinogens present shall be considered additive and the toxic effects associated with non-carcinogens present shall also be considered additive.

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Where naturally occurring substances exceed the established standard, the standard shall be the (3) naturally occurring concentration as determined by the Director.

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21

22

23

24

(c) Except for tracers used in concentrations which have been determined by the Division of Public Health to be protective of human health, and the use of which has been permitted by the Division, substances which are not naturally occurring and for which no standard is specified shall not be permitted in concentrations at or above the practical quantitation limit in Class GA or Class GSA groundwaters. Any person may petition the Director to establish an interim maximum allowable concentration for a substance for which a standard has not been established under this Rule. The petitioner shall submit relevant toxicological and epidemiological data, study results, and calculations necessary to establish a standard in accordance with Paragraph (d) of this Rule. Within three months after the establishment of an interim maximum allowable concentration for a substance by the Director, the Director shall initiate action to consider adoption of a standard for that substance.

(d) Except as provided in Paragraph (f), groundwater Groundwater quality standards for substances in Class GA and Class GSA groundwaters are established as the least of:

31 32

30

(1) Systemic threshold concentration calculated as follows: [Reference Dose (mg/kg/day) x 70 kg (adult body weight) x Relative Source Contribution (.10 for inorganics; .20 for organics)]/[2 liters/day (avg. water consumption)];

33

(2) Concentration which corresponds to an incremental lifetime cancer risk of 1x10-6;

34 35

(3) Taste threshold limit value;

36 37

(4) Odor threshold limit value; (5) Maximum contaminant level; or

38 39

(6) National secondary drinking water standard.

40 41

(e) The following references, in order of preference, shall be used in establishing concentrations of substances which correspond to levels described in Paragraph (d) of this Rule.

42

Integrated Risk Information System (U.S. EPA). (1)

43 44

Health Advisories (U.S. EPA Office of Drinking Water). (2) Other health risk assessment data published by U.S. EPA. (3)

45

(4) Other relevant, published health risk assessment data, and scientifically valid peer-reviewed published toxicological data.

46

(f) The Commission may establish groundwater standards less stringent than existing maximum contaminant levels or national secondary drinking water standards if it finds, after public notice and opportunity for hearing, that

47 48 49

(1) more recent data published in any of the EPA health references listed in paragraph (e) results in a standard which is protective of public health, taste threshold, or odor threshold,

50 51

(2) such a standard will not endanger the public health and safety, including health and environmental effects from exposure to groundwater contaminants, and

52 53 (3) compliance with a standard based on the maximum contaminant level or national secondary drinking water standard would produce serious hardship without equal or greater public benefit.

54 55

(flg) Groundwater quality standards specified in Paragraphs (g)(h) and (h)(i) of this Rule and interim maximum

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2
      allowable concentrations established pursuant to Paragraph (c) of this Rule shall be reviewed on a triennial basis.
 3
      Appropriate modifications to established standards shall be made in accordance with the procedure prescribed in
 4
      Paragraph (d) of this Rule where modifications are considered appropriate based on data published subsequent to the
 5
      previous review.
 6
      (g)(h) Class GA Standards. Where not otherwise indicated, the standard refers to the total concentration in micrograms
 7
      per liter of any constituent in a dissolved, colloidal or particulate form which is mobile in groundwater. This does not
 8
      apply to sediment or other particulate matter which is preserved in a groundwater sample as a result of well construction
      or sampling procedures. The Class GA standards are:
 9
10
                        Acenaphthene: 80;
               (1)
11
                        Acenaphthylene: 200;
               (2)
12
                        Acetone: 6 mg/L;
               (3)
13
               (4)
                        Acrylamide: 0.008;
14
                        Anthracene: 2 mg/L;
               (5)
15
               (6)
                        Arsenic: 10;
16
               (7)
                        Atrazine and chlorotriazine metabolites: 3;
                        Barium: 700;
17
               (8)
18
               (9)
                        Benzene: 1;
19
               (10)
                        Benzo(a)anthracene (benz(a)anthracene): 0.05;
                        Benzo(b)fluoranthene: 0.05;
20
               (11)
21
                        Benzo(k)fluoranthene: 0.5;
               (12)
22
               (13)
                        Benzoic acid: 30 mg/L;
23
               (14)
                        Benzo(g,h,i,)perylene: 200;
24
               (15)
                        Benzo(a)pyrene: 0.005;
25
                        Bis(chloroethyl)ether: 0.03;
               (16)
26
               (17)
                        Bis(2-ethylhexyl) phthalate (di(2-ethylhexyl) phthalate): 3:
27
               (18)
                        Boron: 700;
28
               (19)
                        Bromodichloromethane: 0.6;
29
               (20)
                        Bromoform (tribromomethane): 4;
30
               (21)
                        n-Butylbenzene: 70;
                        sec-Butylbenzene: 70;
31
               (22)
32
                        tert-Butylbenzene: 70;
               (23)
33
                        Butylbenzyl phthalate: 1 mg/L;
               (24)
34
               (25)
                        Cadmium: 2;
35
               (26)
                        Caprolactam: 4 mg/L;
                        Carbofuran: 40;
36
               (27)
37
               (28)
                        Carbon disulfide: 700;
38
               (29)
                        Carbon tetrachloride: 0.3;
39
               (30)
                        Chlordane: 0.1;
40
               (31)
                        Chloride: 250 mg/L;
41
               (32)
                        Chlorobenzene: 50;
42
               (33)
                        Chloroethane: 3,000;
43
               (34)
                        Chloroform (trichloromethane): 70;
44
                        Chloromethane (methyl chloride): 3;
               (35)
45
               (36)
                        2-Chlorophenol: 0.4;
46
               (37)
                        2-Chlorotoluene (o-chlorotoluene): 100;
47
               (38)
                        Chromium: 10;
48
               (39)
                        Chrysene: 5;
49
               (40)
                        Coliform organisms (total): 1 per 100 milliliters;
50
               (41)
                        Color: 15 color units;
51
               (42)
                        Copper: 1 mg/L;
52
               (43)
                        Cyanide (free cyanide): 70;
53
               (44)
                        2, 4-D (2,4-dichlorophenoxy acetic acid): 70;
54
               (45)
                        DDD: 0.1;
55
                        DDT: 0.1;
               (46)
```

1

```
1
               (47)
                        Dibenz(a,h)anthracene: 0.005;
 2
               (48)
                        Dibromochloromethane: 0.4;
 3
               (49)
                        1,2-Dibromo-3-chloropropane: 0.04;
 4
               (50)
                        Dibutyl (or di-n-butyl) phthalate: 700;
 5
               (51)
                        1,2-Dichlorobenzene (orthodichlorobenzene): 20;
 6
                        1,3-Dichlorobenzene (metadichlorobenzene): 200;
               (52)
 7
               (53)
                        1,4-Dichlorobenzene (paradichlorobenzene): 6;
 8
               (54)
                        Dichlorodifluoromethane (Freon-12; Halon): 1 mg/L;
 9
               (55)
                        1,1-Dichloroethane: 6;
10
                        1,2-Dichloroethane (ethylene dichloride): 0.4;
               (56)
               (57)
                        1,2-Dichloroethene (cis): 70;
11
12
               (58)
                        1,2-Dichloroethene (trans): 100;
13
               (59)
                        1,1-Dichloroethylene (vinylidene chloride): 7;
14
                        1,2-Dichloropropane: 0.6;
               (60)
                        1,3-Dichloropropene (cis and trans isomers): 0.4;
15
               (61)
16
               (62)
                        Dieldrin: 0.002;
17
               (63)
                        Diethylphthalate: 6 mg/L;
18
               (64)
                       2,4-Dimethylphenol (m-xylenol): 100;
19
               (65)
                        Di-n-octyl phthalate: 100;
20
               (66)
                        1,4-Dioxane (p-dioxane): 3;
21
                       Dioxin (2,3,7,8-TCDD): 0.0002 ng/L;
               (67)
22
                        1,1- Diphenyl (1,1,-biphenyl): 400;
               (68)
23
               (69)
                        Dissolved solids (total): 500 mg/L;
24
               (70)
                        Disulfoton: 0.3;
25
                       Diundecyl phthalate (Santicizer 711): 100;
               (71)
26
               (72)
                        Endosulfan: 40;
27
               (73)
                        Endrin, total: (includes endrin, endrin aldehyde and endrin ketone): 2;
28
                        Epichlorohydrin: 4;
               (74)
29
               (75)
                       Ethyl acetate: 3 mg/L;
30
               (76)
                        Ethylbenzene: 600;
31
               (77)
                       Ethylene dibromide (1,2-dibromoethane): 0.02;
32
               (78)
                       Ethylene glycol: 10 mg/L;
33
               (79)
                       Fluoranthene: 300;
34
               (80)
                       Fluorene: 300;
35
               (81)
                        Fluoride: 2 mg/L;
                       Foaming agents: 500;
36
               (82)
37
               (83)
                        Formaldehyde: 600;
38
               (84)
                        Gross alpha (adjusted) particle activity (excluding radium-226 and uranium): 15 pCi/L;
39
               (85)
                       Heptachlor: 0.008;
40
               (86)
                       Heptachlor epoxide: 0.004;
41
               (87)
                       Heptane: 400;
                       Hexachlorobenzene (perchlorobenzene): 0.02;
42
               (88)
43
               (89)
                       Hexachlorobutadiene: 0.4;
               (90)
                       Hexachlorocyclohexane isomers (technical grade): 0.02;
44
               (91)
45
                       n-Hexane: 400;
46
               (92)
                        Indeno(1,2,3\text{-cd})pyrene: 0.05;
47
                       Iron: 300;
               (93)
48
               (94)
                        Isophorone: 40;
49
               (95)
                       Isopropylbenzene: 70;
                        Isopropyl ether: 70;
50
               (96)
51
               (97)
                       Lead: 15;
52
               (98)
                       Lindane (gamma hexachlorocyclohexane): 0.03;
53
               (99)
                       Manganese: 50;
54
               (100)
                       Mercury: 1;
                       Methanol: 4 mg/L;
55
               (101)
```

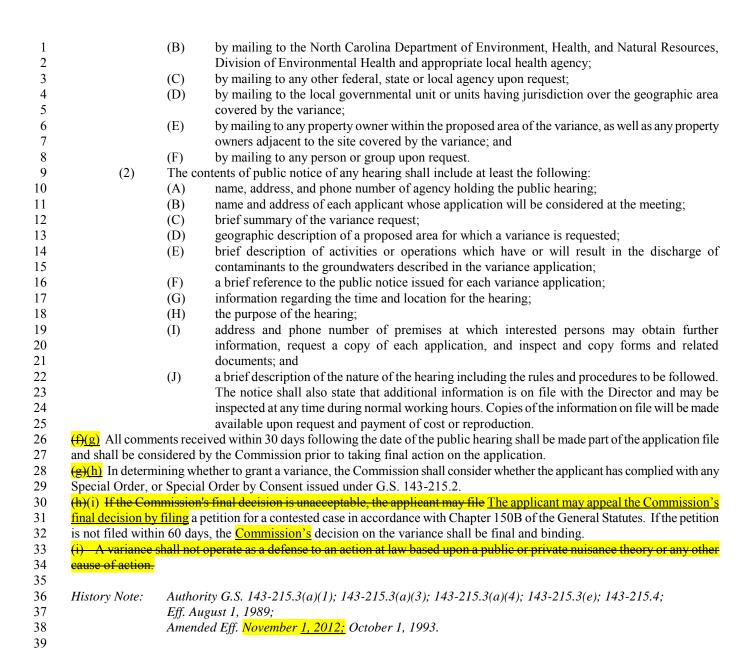
```
1
               (102)
                        Methoxychlor: 40;
 2
               (103)
                        Methylene chloride (dichloromethane): 5;
 3
                        Methyl ethyl ketone (2-butanone): 4 mg/L;
               (104)
 4
               (105)
                        2-Methylnaphthalene: 30;
 5
               (106)
                        3-Methylphenol (m-cresol): 400;
 6
                        4-Methylphenol (p-cresol): 40;
               (107)
 7
               (108)
                        Methyl tert-butyl ether (MTBE): 20;
 8
               (109)
                        Naphthalene: 6;
 9
               (110)
                        Nickel: 100;
10
                        Nitrate: (as N) 10 mg/L;
               (111)
                        Nitrite: (as N) 1 mg/L;
11
               (112)
                        N-nitrosodimethylamine: 0.0007;
12
               (113)
13
               (114)
                        Oxamyl: 200;
14
               (115)
                        Pentachlorophenol: 0.3;
                        Petroleum aliphatic carbon fraction class (C5 - C8): 400;
15
               (116)
16
               (117)
                        Petroleum aliphatic carbon fraction class (C9 - C18): 700;
                        Petroleum aliphatic carbon fraction class (C19 - C36): 10 mg/L;
17
               (118)
18
               (119)
                        Petroleum aromatics carbon fraction class (C9 - C22): 200;
19
               (120)
                        pH: 6.5 - 8.5;
20
               (121)
                        Phenanthrene: 200;
21
                        Phenol: 30;
               (122)
22
                        Phorate: 1;
               (123)
23
               (124)
                        n-Propylbenzene: 70;
24
               (125)
                        Pyrene: 200;
25
                        Selenium: 20;
               (126)
26
               (127)
                        Silver: 20;
27
               (128)
                        Simazine: 4;
28
                        Styrene: 70;
               (129)
                        Sulfate: 250 mg/L;
29
               (130)
30
               (131)
                        1,1,2,2-Tetrachloroethane: 0.2;
31
               (132)
                        Tetrachloroethylene (perchloroethylene; PCE): 0.7;
32
                        2,3,4,6-Tetrachlorophenol: 200;
               (133)
33
                        Toluene: 600;
               (134)
34
               (135)
                        Toxaphene: 0.03;
35
               (136)
                        2, 4, 5,-TP (Silvex): 50;
                        1,2,4-Trichlorobenzene: 70;
36
               (137)
37
               (138)
                        1,1,1-Trichloroethane: 200;
38
               (139)
                        Trichloroethylene (TCE): 3;
39
                        Trichlorofluoromethane: 2 mg/L;
               (140)
40
               (141)
                        1,2,3-Trichloropropane: 0.005;
41
               (142)
                        1,2,4-Trimethylbenzene: 400;
                        1,3,5-Trimethylbenzene: 400;
42
               (143)
                        1,1,2-Trichloro-1,2,2-trifluoroethane (CFC-113): 200 mg/L;
43
               (144)
44
                        Vinyl chloride: 0.03;
               (145)
45
               (146)
                        Xylenes (o-, m-, and p-): 500; and
46
               (147)
                        Zinc: 1 mg/L.
47
      (h)(i) Class GSA Standards. The standards for this class are the same as those for Class GA except as follows:
48
               (1)
                        chloride: allowable increase not to exceed 100 percent of the natural quality concentration; and
49
               (2)
                        total dissolved solids: 1000 mg/l.
50
      (i)(i) Class GC Waters.
51
                        The concentrations of substances which, at the time of classification, exceed the standards applicable
               (1)
52
                        to Class GA or GSA groundwaters shall not be caused to increase, nor shall the concentrations of other
53
                        substances be caused to exceed the GA or GSA standards as a result of further disposal of
54
                        contaminants to or beneath the surface of the land within the boundary of the area classified GC.
```

1 2 3 4	(2)	The concentrations of substances which, at the time of classification, exceed the standards applicable to GA or GSA groundwaters shall not be caused to migrate as a result of activities within the boundary of the GC classification, so as to violate the groundwater or surface water quality standards in adjoining waters of a different class.
5	(3)	Concentrations of specific substances, which exceed the established standard at the time of
6		classification, are listed in Section .0300 of this Subchapter.
7		
8	History Note:	Authority G.S. 143-214.1; 143B-282(a)(2);
9		Eff. June 10, 1979;
10		Amended Eff. November 1, 1994; October 1, 1993; September 1, 1992; August 1, 1989;
11		Temporary Amendment Eff. June 30, 2002;
12		Amended Eff. August 1, 2002;
13		Temporary Amendment Expired February 9, 2003;
14		Amended Eff. November 1, 2012; January 1, 2010; April 1, 2005.
15		<u> </u>

15A NCAC 02L .0113 is proposed for amendment as follows: (Option 3)

15A NCAC 02L .0113 VARIANCE

- (a) The Commission, on its own initiative or pursuant to a request under G.S. 143-215.3(e), may grant variances to the rules of this Subchapter.
- (b) Requests for variances are filed by letter from the applicant to the Environmental Management Commission. The application shall be mailed to the chairman of the Commission in care of the Director, Division of Environmental Management, Post Office Box 29535, Raleigh, N.C. 27626 0535. Water Quality, 1617 Mail Service Center, Raleigh, N.C. 27699-1617.
- (c) For site-specific variances, the The application shall contain the following information:
 - (1) Applications filed by counties or municipalities must include a resolution of the County Board of Commissioners or the governing board of the municipality requesting the variance.
 - (2) A description of the past, existing or proposed activities or operations that have or would result in a discharge of contaminants to the groundwaters.
 - (3) Description of the proposed area for which a variance is requested. A detailed location map, showing the orientation of the facility, potential for groundwater contaminant migration, as well as the area covered by the variance request, with reference to at least two geographic references (numbered roads, named streams/rivers, etc.) must be included.
 - (4) Supporting information to establish that the variance will not endanger the public health and safety, including health and environmental effects from exposure to groundwater contaminants. (Location of wells and other water supply sources including details of well construction within 1/2 mile of site must be shown on a map).
 - (5) Supporting information to establish that requirements of this Rule cannot be achieved by providing the best available technology economically reasonable. This information must identify specific technology considered, and the costs of implementing the technology and the impact of the costs on the applicant.
 - (6) Supporting information to establish that compliance would produce serious financial hardship on the applicant.
 - (7) Supporting information that compliance would produce serious financial hardship without equal or greater public benefit.
 - (8) A copy of any Special Order that was issued in connection with contaminants in the proposed area and supporting information that applicant has complied with the Special Order.
 - (9) A list of the names and addresses of any property owners within the proposed area of the variance as well as any property owners adjacent to the site covered by the variance.
- (d) For state-wide variances to groundwater standards established in Section .0202, the application shall contain the following information:
 - (1) Supporting information to establish that the variance will not endanger the public health and safety, including health and environmental effects from exposure to groundwater at the proposed constituent levels. This should include information obtained from the following references.
 - (a) Integrated risk Information System (U.S. EPA).
 - (b) Health Advisories (U.S. EPA Office of Drinking Waters).
 - (c) Other health risk assessment data published by U.S. EPA.
 - (d) Other relevant, published health and ecological risk assessment data, and scientifically valid peer-reviewed published toxicological data.
 - (2) A list of all known potentially affected sites, to include permitted sites and incident sites. For each site listed, a map for each site with the location of wells and other water supply sources within ½ mile of the affected site must be provided.
 - (3) A list of increased costs for treatment for any of the wells or water supply sources listed in Paragraph (2) above due to the proposed variance to Section .0202.
- (d)(e) Upon receipt of the application, the Director will review it for completeness and request additional information if necessary. When the application is complete, the Director shall give public notice of the application and schedule the matter for a public hearing in accordance with G.S. 143-215.4(b) and the procedures set out in Paragraph (e)(f) of this Rule.
 - (1) Notice of public hearing on any variance application shall be circulated in the geographical areas of the proposed variance by the Director at least 30 days prior to the date of the hearing:
 - (A) by publishing the notice one time in a newspaper having general circulation in said county;



Appendix F: Summary of Proposed Rule Changes

Rule Citation	Proposed Revision	Potential Economic Impact
15A NCAC 02L	(g)(59) Amends the 1,1-	-Compliance cost savings for
.0202(g)(59)	dichloroethylene groundwater	facilities with releases of 1,1-
(Rhodia Option 1)	standard from 7 ug/L to 350 ug/L.	dichloroethylene (1,1-DCE).
		-Compliance cost savings to
		permitted facilities (for example, non-
		discharge, pretreatment, landfills).
		-Compliance cost to public water
		supply systems using 1,1-DCE-
		contaminated groundwater above the
		federal maximum contaminant level
		(MCL) of 7 ug/L.
15A NCAC 02L	(d) Allows an exception to the	-Compliance cost savings for
.0202(d) & (f)	criteria for establishing	facilities with releases of 1,1-
(DWQ Option 2)	groundwater standards.	dichloroethylene (1,1-DCE).
	(f) Allows the Environmental	-Compliance cost savings to
	Management Commission to	permitted facilities (for example, non-
	establish groundwater standards	discharge, pretreatment, landfills).
	less stringent than existing	-Compliance cost to public water
	Maximum Contaminant Levels	supply systems using 1,1-DCE-
	and secondary drinking water	contaminated groundwater above the
	standards under certain	federal maximum contaminant level
	circumstances.	(MCL) of 7 ug/L.
15A NCAC 02L	(b) Updates the Division of Water	-No cost or cost savings expected.
.0113 (b) - (i)	Quality mailing address.	
(DWQ Option 3)	(d) Adds a state-wide variance	
	option and application	
	requirements.	
	Note: original paragraphs (d)-(i)	
	are re-alphabetized (e)-(i).	
	(h) & (i) adds and deletes text for	
	clarity.	

Appendix G



RADIATOR SPECIALTY COMPANY

600 Radiator Road, Indian Trail, NC 28079-5225

VIA OVERNIGHT AND ELECTRONIC MAIL

Environmental Management Commission c/o Lois Thomas, EMC Recording Clerk Director's Office Division of Water Quality 512 North Salisbury Street Raleigh, North Carolina 27604

Re: Letter of Support for Rhodia Inc.'s Petition for Rulemaking to Change Groundwater Quality Standard for 1,1-Dichloroethylene (1,1-DCE)

Dear Members of the Environmental Management Commission:

Radiator Specialty Company is writing this letter to express its support of Rhodia Inc.'s rulemaking petition to change the groundwater quality standard for 1,1-DCE to 350 μ g/L.

Radiator Specialty has been headquartered in North Carolina since its founding in 1924. Today, from its corporate headquarters and 400,000 square foot manufacturing plant in Indian Trail, Radiator Specialty develops, manufactures and markets high performance products for auto, motorcycle, plumbing, hardware and industrial applications for more than 1,600 customers in 81 countries. Radiator Specialty is committed to conducting all of its operations in an environmentally responsible manner.

Just as the State expects us to operate in compliance with environmental laws and regulations, Radiator Specialty expects the State to do so as well. During the 2008. Triennial Review, we along with Rhodia Inc. and Ashland Inc. submitted a written public comment that the groundwater quality standard for 1,1-DCE should be changed to 350 μ g/L. The basis for that request was the same as the basis for Rhodia's rulemaking petition. Radiator Specialty believes that North Carolina's regulations require that this change be made and we ask you to approve Rhodia's petition at your July 14 meeting.

We would request that a copy of this letter be provided to each member of the EMC and that it also be placed on the EMC's website as part of the materials related to the Rhodia petition.

Sincerely,

Stuart A. Kerkhoff

Environmental Manager

Radiator Specialty Company

Appendix H

Dear Ms. Thomas,

I am President of Duncklee & Dunham, P.C., a Geological and Engineering Environmental Consulting firm located in Cary, North Carolina. I understand the July 14, 2011 EMC meeting will have an agenda item relating to considering establishing a new groundwater standard for 1,1-dichloroethene (1,1-DCE) at 350 \Box g/L, up from the current level of 7 \Box g/L. I have been following this case with interest, as it makes sense that North Carolina groundwater quality standards are based on the most current scientific and toxicological data available.

I understand the current 1,1-DCE level of $7 \square g/L$ is based on the lowest of the standards the Division of Water Quality considers for comparison, in this case the federal maximum concentration limit (MCL). I understand that the oral reference dose was changed for 1,1-DCE in 2003, but EPA did not change the MCL for this constituent, and has no plans to do so because of non-scientific reasons, e.g. competing workload priorities and administrative costs associated with the rulemaking. I do not believe it was the legislative intent for 2L standards to use the least value if that value was not calculated based on the most current toxicological information.

Even though the state \square s toxicologist calculated the groundwater quality standard should be 350 \square g/L in 2004 based on the updated reference dose information, I understand the Division of Water Quality refused to change the 1,1-DCE standards because of the way their own 2L rules were written. I am optimistic the EMC ruling will be based on logic, not adherence to a standard that no longer has the basis to be set at that level.

Duncklee & Dunham has had many clients affected by the changes/additions in groundwater standards and IMACs, often requiring much more money to be spent to clean up groundwater to meet these standards. It appears that the regulated community may have to bear considerable additional costs to meet a standard that has no scientific basis.

On behalf of the professional geological and engineering staff of Duncklee & Dunham, P.C. please consider this letter of support for the Rhodia Petition for the amendment of the current 1,1-DCE groundwater standard to be raised from $7 \Box g/L$ to $350 \Box g/L$.

Sincerely,

David L. Duncklee, PG, RSM

Senior Hydrogeologist Mobile: 919-417-9923 Office: 919-858-9898 x201 dave@dunckleedunham.com

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15.0	Solar Age	t .	

A Professional Geologic and Engineering Corporation

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Appendix I: NCDENR Outreach via Email to Industry/ Private Sector

Manufacturers and Chemical Industry Council of North Carolina (MCIC)

Preston Howard, President 620 N. West Street, Suite 101

Raleigh, NC 27603 Telephone: 919-834-9459

E-Mail: <u>preston.howard@mcicnc.org</u>

jim.kuszaj jim@mcienc.org

Web Site: http://www.mcicnc.org/index.html

Member Companies:

Member Companies.	
Air Products and Chemicals, Inc.	GlaxoSmithKline
Alcoa	Hexion
Archer Daniels Midland Co	International Paper
BASF Corporation	Invista
Blue Ridge Paper Products	Kao Specialties Americas LLC
Cargill	Kapstone Kraft Paper Corp.
Carolina Stalite	Kimberly Clark
Caterpillar, Inc	Lord Corporation
Celgard	MeadWestvaco
Chemtura	MOEN Incorporated
Ciba Specialty Chemicals	National Starch & Chemical Company
Clariant Corporation	Novozymes North America
Cognis	Oak-Bark Corp.
Corn Products International, Inc.	PCS Phosphate Company, Inc.
Corning Incorporated	Pressure Chemical Co.
Covidien/Mallinckrodt	RJ Reynolds Tobacco Company
DAK Americas	Resinall Corporation
Domtar Paper Company, LLC	Rohm & Haas
Dow Corning	Shurtape Technologies, Inc.
DuPont Company	Surry Chemicals, Inc.
Elementis Chromium	Syngenta Crop Protection
Evonik Stockhausen.	Trinity Manufacturing, Inc.
ExxonMobile Chemical	Unilin US MDF
FMC Corporation	Univar
Fortron Industries	Vertellus Performance Materials
General Electric	Weyerhaeuser Company
Georgia-PacificCorporation	
· · · · · · · · · · · · · · · · · · ·	

MCIC Business Partners

Arcadis

EI, Inc.

ENSR Consulting and Engineering (NC), Inc.

Hunton & Williams

Kilpartick Stockton LLP

McNair Law Firm, P.A.

Stearns & Wheler, PLLC

Trinity Consultants

URS Corporation - North Carolina

Womble, Carlyle, Sandridge & Rice, PLLC

North Carolina Chamber (formerly NC Citizens for Business & Industry)

S. Lewis Ebert, President and CEO Raleigh Corporate Center 701 Corporate Center Drive, Suite 400 Raleigh, N.C. 27607 919-836-1407

info@ncchamber.net

<u>lebert@ncchamber.net</u>

Wesley Gappens, Communications Specialist

wgappens@ncchamber.net

919-836-1418

Web Site: http://www.ncchamber.net/mx/hm.asp?id=home

N.C. Rural Economic Development Center

Thomas W. Lambeth, Chairman Billy Ray Hall, President 4021 Carya Drive, Raleigh, NC 27610 info@ncruralcenter.org

http://www.ncruralcenter.org/ Telephone: (919) 250-4314

Garnet Bass, Director of communications

gbass@ncruralcenter.org

Progress Energy Service Company, LLC

Mick Greeson 410 S. Wilmington Street Raleigh, NC 27601 919-546-6129

mick.greeson@pgnmail.com

Web Site: https://www.progress-energy.com/

Duke Energy Corporation

George T. Everett, vice president, Environmental and Public Policy 526 South Church Street Charlotte, NC 28202

Everett, George T" GTEverett@duke-energy.com

Web Site: http://www.duke-energy.com/north-carolina.asp

Treated Wood Council, Inc.

Jeff Miller, Executive Director 1111 19th St., NW, Ste. 800 Washington, DC 20036 Phone: 202-463-2045

E-mail: jeff miller@treated-wood.org

Web Site: http://www.treated-wood.org/home.html

North Carolina Pork Council

Tommy Stevens, Director of Environmental Services

tommy@ncpork.org 2300 Rexwoods Drive Suite 340 Raleigh, NC 27607

Phone: 919-781-0361

Web Site: http://www.ncpork.org/index.jsp

NC Association of County Commissioners

David F. Thompson, Exec. Director

david.thompson@ncacc.org

215 N Dawson Street

Raleigh, NC 27603

Albert Coates Local Government Center

919-715-2893

Web Site: http://www.ncacc.org/

Email: ncacc@ncacc.org

Todd McGee, Communications Director (919) 715-7336, or todd.mcgee@ncacc.org

Kevin.leonard@ncacc.org

NC League of Municipalities

Erin Wynia

ewynia@nclm.org

215 N Dawson Street

Raleigh, NC 27603

PO Box 3069 (27602-3069)

(919) 715-4000

Web Site: http://www.nclm.org/Pages/default.aspx

Professional Engineers of North Carolina

1015 Wade Ave, Suite A

Raleigh, NC 27605

(919) 834-1144 (phone)

E-Mail: exec@penc.org Web Site: www.penc.org

Marc R. Worth, President

PSNC Energy

4077 Haywood Rd

Mills River, NC 28759

Phone: 828-890-7554

Email: mworth@scana.com

(in the future send directly to Betsy Bailey at bbailey@penc.org.)

National Federation of Independent Businesses - North Carolina

150 Fayetteville Street

Suite 1110

Raleigh, NC 27601

919-833-9480

Gregg Thompson, NFIB/North Carolina State Director

Gregg.Thompson@NFIB.org

Web Site: http://www.nfib.com/north-carolina

NC Council of Governments (NCCOGS):

Southwestern Commission, Land-of-Sky Regional Council, Isothermal Planning and Development Commission, High Country COG, Western Piedmont COG, Centralina COG, Piedmont Triad COG, Northwest Piedmont COG, Triangle J COG, Kerr-Tar Regional COG, Upper Coastal Plain COG, Mid-Carolina COG, Lumber River COG, Cape Fear COG, Eastern Carolina COG, Mid-East Commission, Albemarle Commission, Regional Associate.

NCCOGS Contact Information:

Southwestern Commission

(Region A)

Bill Gibson, Executive Director

Main Street Federal Building, Room 202

P.O. Drawer 850 Bryson City, NC 28713

Tel: (828) 488-9211 or Fax: (828) 488-3950

Email: <u>bill@regiona.org</u>
Website: <u>www.regiona.org</u>

Members: Cherokee, Clay, Graham, Haywood, Jackson,

Macon and Swain Counties

Land-of-Sky Regional Council

(Region B)

Joe McKinney, Executive Director

339 New Leicester Hwy Asheville, NC 28806

Tel: (828) 251-6622 or Fax: (828) 251-6353

jmckinney@landofsky.org Email: info@landofsky.org Website: www.landofsky.org

Members: Buncombe, Madison, Henderson,

and Transylvania Counties

Isothermal Planning and Dev. Commission

(Region C)

Jim Edwards, Executive Director

jedwards@regionc.org 111 West Court Street

P.O. Box 841

Rutherfordton, NC 28139

Tel: (828) 287-2281 or Fax: (828) 287-2735

Website: www.regionc.org

Members: Cleveland, McDowell, Polk, and

Rutherford Counties

High Country Council of Governments

(Region D)

Rick Herndon, Executive Director

Executive Arts Building, 155 Furman Road

P.O. Box 1820 Boone, NC 28607

Tel: (828) 265-5434 or Fax: (828) 265-5439

Email: rherndon@regiond.org
Website: www.regiond.org

Members: Alleghany, Ashe, Avery, Mitchell, Watauga, Wilkes, and Yancey Counties

Western Piedmont Council of Govt's

(Region E)

H. DeWitt Blackwell, Jr. AICP

Executive Director

Western Piedmont Council of Governments

P. O. Box 9026 Hickory, NC 28603 or 736 Fourth Street SW Hickory, NC 28602

dee.blackwell@wpcog.org

828 485-4230 voicemail 828 455-3477 cell

828 322-5991 Fax

http://www.wpcog.org/

Members: Alexander, Burke, Caldwell, and

Catawba Counties

Centralina Council of Governments

(Region F), Executive Director

Jim Prosser

Executive Director 525 North Tryon Street

12th Floor

Charlotte, NC 28202

iprosser@centralina.org

Tel: (704) 372-2416 or Fax: (704) 347-4710

Website: www.centralina.org

Member: Anson, Cabarrus, Gaston, Iredell,

Lincoln, Mecklenburg, Rowan, & Stanly Counties

Piedmont Triad Council of Governments

(Region G)

Ginger Booker, *Interim* Executive Director 2216 W. Meadowview Road, Suite 201

Wilmington Building Greensboro, NC 27407-3408

Tel: (336) 294-4950 or Fax: (336) 632-0457 (Cv Stober, Water Resources Manager

cstober@ptcog.org

Website: www.ptcog.org

Members: Alamance, Caswell, Davidson,

Guilford, Montgomery, Randolph, & Rockingham

Counties

Northwest Piedmont Council of Govt's

(Region I)

Matthew L. Dolge, Executive Director 400 West Fourth Street, Suite 400 Winston-Salem, NC 27101

Tel: (336) 761-2111 or Fax: (336) 761-2112

TDD: (336) 761-2110 Email: regioni@nwpcog.org Website: www.nwpcog.org

Members: Davie, Forsyth, Stokes, Surry, and Yadkin

Counties

Triangle J Council of Governments

(Region J)

Kirby Bowers, Executive Director

kbowers@ticog.org

4307 Emperor Blvd., Suite 110, Durham, 27703

Mailing address: P.O. Box 12276 Research Triangle Park, NC 27709

Tel: (919) 549-0551 or Fax: (919) 558-9393

Email: tjcog@tjcog.org Website: www.tjcog.org

Members: Chatham, Durham, Johnston, Lee, Moore,

Orange, and Wake Counties

Kerr-Tar Regional Council of Governments

(Region K)

Timothy Baynes

tbaynes@kerrtarcog.org, Executive Director

(timmy@kerrtarcog.org) 510 Dabney Drive

P.O. Box 709

Henderson, NC 27536

Tel: (252) 436-2040 or Fax: (252) 436-2055

Email:

Website: www.kerrtarcog.org

Members: Franklin, Granville, Person, Vance, and

Warren Counties

Upper Coastal Plain Council of Govt's

(Region L)

Greg T. Godard, Executive Director 1309 S. Wesleyan Blvd. (37803)

P.O. Box 9

Wilson, NC 27894 Tel: (252) 234-5952

Email: ggodard@ucpcog.org Website: www.ucpcog.org

Members: Edgecombe, Halifax, Northampton, Nash, and Members: Cumberland, Harnett, and

Wilson Counties

Mid-Carolina Council of Governments

(Region M)

James Caldwell, Executive Director

130 Gillespie Street P.O. Drawer 1510 Fayetteville, NC 28302

Tel: (910) 323-4191 or Fax: (910) 323-9330

Email: <u>icaldwell@mccog.org</u> Website: www.mccog.org

Sampson Counties

Lumber River Council of Govt's

(Region N)

James Perry, Executive Director james.perry@lumberrivercog.org

Lumber River Council of Governments

30 CJ Walker Road COMtech Park Pembroke, NC 28372 Tel: (910) 618-5533 or

Website: http://www.lumberrivercog.org/

Email: lrcog@lrcog.org

Members: Bladen, Hoke, Richmond, Robeson, and

Scotland Counties

Eastern Carolina Council of Govt's

(Region P)

Larry Moolenaar, Executive Director

lmoolenaar@eccog.org

233 Middle Street, 3rd floor, O'Marks Bldg.

P.O. Box 1717

New Bern, NC 28563-1717

Tel: (252) 638-3185 or Fax: (252) 638-3187

Email: <u>eccog@eccog.org</u>
Website: www.eccog.org

Members: Carteret, Craven, Duplin, Greene, Jones,

Lenoir, Onslow, Pamlico, Wayne Counties

Albemarle Commission

(Region R)

Mr. Lambert, Executive Director

512 S. Church Street

P.O. Box 646

Hertford, NC 27944 Tel: (252) 426-5753 or Fax: (252) 426-8482

slambert@albemarlecommission.org

Email: bkr27944@yahoo.com (? Did not send email)

http://www.albemarlecommission.org/

Members: Camden, Chowan, Currituck, Dare, Gates, Hyde, Pasquotank, Perquimans, Tyrrell, and Washington

Counties

Cape Fear Council of Govt's

(Region O)

Christopher D. May, Executive Director

1480 Harbour Drive Wilmington, NC 28401 Tel: (910) 395-4553 or Fax: (910) 395-2684

Email: cmay@capefearcog.org
Website: www.capefearcog.org

Contact: Don Eggert, Local Gov't Services Director

deggert@capefearcog.org

Members: Brunswick, Columbus, New Hanover, and

Pender Counties

Mid-East Commission

(Region Q)

Tim Ware, Executive Director 1385 John Small Avenue

P.O. Box 1787

Washington, NC 27889 Tel: (252) 974-1825

Email: tware@mideastcom.org

Website: http://www.mideastcom.org/

Phone: 252.946.8043 **Toll Free:** 1.800.799.9194

Members: Beaufort, Bertie, Hertford, Martin,

and Pitt Counties

Regional Associate

William A. McNeil, AICP Cell: (919) 622-3303

Office: (919) 715-9550

Email: <u>ncregions@mindspring.com</u>

NC Association of Launderers & Cleaners

Sto Fox, Executive Director

stofox@ncalc.org

336-549-5486

1403-A Sunset Drive

Greensboro, NC 27408

336-389-9011

Web Site: http://www.ncalc.org/

Email: ncalc@ncalc.org

Charlie Smith, District 2 Director

charlie@crsassociates.org

North Carolina Farm Bureau

Mailing Address:

PO Box 27766

Raleigh, NC 27611

Physical Address:

5301 Glenwood Ave.

Raleigh, NC 27612

919 782-1705

http://www.ncfb.org/

Contact: Anne Coan

Anne.coan@ncfb.org

919-788-1005

Newaterworks operators association

http://www.ncwoa.com/

Leslie Carreiro, President

City of Asheville – North Fork WTP

3374 North Fork-Left Fork Rd, Black Mountain, NC 28711

Phone: 828-271-6105 Fax: 828-271-6102

lcarreiro@ashevillenc.gov

Cindy Gall, Administrator NCWOA

PO Box 4519

Emerald Isle, NC 28594 Phone: (252) 764-2094

Fax: (252) 764-2095 Email: ncwoa@intrex.net

North Carolina Poultry Federation

Kendall Casey, President 4020 Barrett Drive

Suite 102

Raleigh, NC 27609 Phone: (919) 783-8218

http://www.ncpoultry.org/ rlford@ncpoultry.org

Email Outreach to State Departments and Programs

North Carolina Department of Agriculture and Consumer Services (NCDACS)

Mr. Dewitt Hardee
Environmental Programs Manager
1001 Mail Service Center
Raleigh, NC 27699-1001
Dewitt.hardee@ncagr.gov

http://www.ncagr.gov/environmentalprograms/staff.htm

NC Department of Transportation

Ken D. Pace, State Environmental Operations Engineer Roadside Environmental Unit Environmental Operations & Rest Area 1557 Mail Service Center, Raleigh, NC 27699 (Mail) 1 South Wilmington Street, Raleigh, NC (Delivery) Transportation Building - 5th Floor Annex kpace@ncdot.gov 919 733-2920 x70

T. C. Niver, PG, CHMM - 1-888-457-0512

tniver@dot.state.nc.us

Roadside Environmental Unit Environmental Operations Section Environmental Operations Engineer III 1566 Mail Service Center (Mail) Raleigh, NC 27699 4809 Beryl Road (office trailer)- Delivery Raleigh, NC 27606

Robin Maycock
Environmental Operations Engineer II
1558 Mail Service Center (Mail)
Raleigh, NC 27699
1425 Rock Quarry Road, Suite 106 (Delivery)
Raleigh, NC 27610
rmaycock@ncdot.gov
919 861-3781

John Kirby jkirby@ncdot.gov 508-1816

NCDENR Division of Environmental Health

Terry Pierce, Director 2728 Capital Blvd. 1630 MSC Raleigh NC 27699-1630 Phone: (919) 733-2870

Fax: (919) 715-3242 Terry.pierce@ncdenr.gov

NC Division of Water Resources, Tom Reeder, Director

Tom.reeder@ncdenr.gov

919-715-3045

Public Water Supply Section (now under DWR)

Jessica Godreau, P.E., CPM, Section Chief

(919) 715-3232

Jessica.godreau@ncdenr.gov

PWSS Web site: http://www.ncwater.org/pws/

North Carolina Division of Air Quality

Sheila Holman, Director Sheila.holman@ncdenr.gov 1641 Mail Service Center Raleigh, NC 27699-1641 (919) 715-0971

North Carolina Division of Waste Management

Dexter Matthews, Director Linda Culpepper, Deputy Director 1646 Mail Service Center, Raleigh, NC 27699-1646 (919)508-8400

NC DWQ Aquifer Protection Section

Ted Bush, Chief 919-715-6172 Betty Wilcox 715-6169

NC DHHS Division of Public Health

Mina W. Shehee, Ph.D.
Medical Evaluation & Risk Assessment Unit
Occupational & Environmental Epidemiology Branch
Division of Public Health
North Carolina Department of Health & Human Services
1912 Mail Service Center
Raleigh NC 27699-1912
(919) 707-5920
mina.shehee@dhhs.nc.gov
ken.rudo@dhhs.nc.gov

Email Outreach to Environmental Groups:

Clean Water for North Carolina

Hope Taylor-Guevara, Executive Director Durham Office 2009 Chapel Hill Road Durham, NC 27707 hope@cwfnc.org (919) 401-9600

NC Conservation Network

Grady McCallie, Policy Director 19 E. Martin St., Suite 300 Raleigh, NC 27601 919.857.4699 grady@ncconservationnetwork.org Dan Conrad dan@ncconservationnetwork.org

Environmental Defense Fund

Sam Pearsall SPearsall@edf.org Maggie Clary mclary@edf.org

Southern Environmental Law

Mary Maclean Asbill mmasbill@gmail.com Julie Youngman jyoungman@selcnc.org Geoff Gisler ggisler@selcnc.org NC/SC Office 200 West Franklin St., Suite 330 Chapel Hill, NC 27516-2559 (919) 967-1450 Fax (919) 929-9421

Note: sent information in an email through their web site "send us an email" link. Could not find an email address for any staff online.

American Rivers

Peter Raabe

praabe@americanrivers.org

331 West Main Street

Suite 504

Durham NC 27701

919-682-3500

AWWA (NC American Waterworks Association)

Nbanks@ncsafewater.org Lindsay Roberts, Executive Director <u>lrobertrs@ncsafewater.org</u>

NC League of Conservation Voters (formerly Conservation Council of North Carolina)

Dan Crawford dan@nclcv.org

http://www.nclcv.org/

NC Sierra Club

http://nc.sierraclub.org/about/contact.html info@sierraclub-nc.org

Molly Diggins, State Director

Western NC Alliance (WNCA)

Julie Mayfield, Executive Director www.wnca.org julie@wnca.org

Waterkeepers

Hannah Connor

hconnor@waterkeeper.org

Others contacted as a result of outreach:

Drycleaning and Laundry Institute

Mary Scalco <u>mscalco@ifi.org</u> 1-800-638-2627 704-216-8593

S&ME, Inc.

Connel Ware Senior Project Manager 3718 Old Battleground Road Greensboro, NC 27410 336-288-8980 cware@smeinc.com

ARCADIS U.S., Inc. 801 Corporate Center Drive, Suite 300 Raleigh, NC 27607 919-854-1282 Shawn Sager, Ph.D. Principal Scientist shawn.sager@arcadis-us.com www.arcadis-us.com

Rhodia, Inc. 207 Telegraph Drive Gastonia, NC Benne Hutson bhutson@mcguirewoods.com

Utilities Inc. (aka Carolina Water) Mary Rollins 704-319-0519 MFRollins@uiwater.com

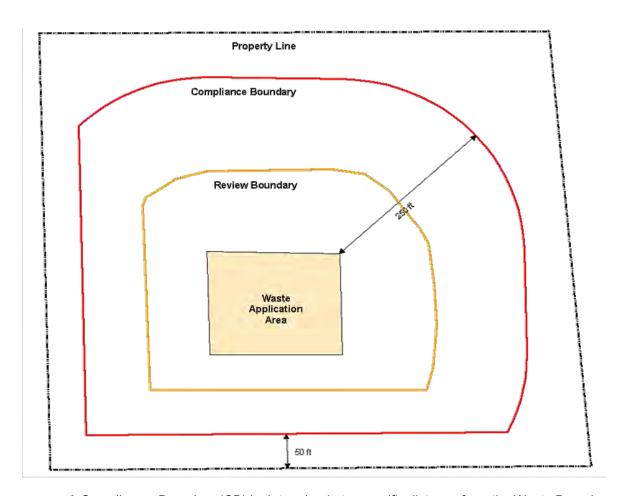
American Truetzschler John Guglielmetti JGuglielmetti@am-trutzschler.com

Excel Civil & Environmental Associates, PLLC Mike Stanforth mstanforth@excelengr.com

Middlesex Public Water Supply Tony Arnold, Director of Client Services Envirolink, Inc.

Phone: 252-235-4900 Cell: 252-236-8168

Email: tarnold@envirolinkinc.com



APPENDIX J: Boundaries at a Typical DWQ Permitted Waste Site

- A Compliance Boundary (CB) is determined at a specific distance from the Waste Boundary
- The Review Boundary (RB) is midway between the CB and the RB
- Exceedances of the groundwater quality standard at the Review Boundary (RB) require
 preventive action/ modeling; exceedances at the Compliance Boundary (CB) require corrective
 action.

As shown: "For disposal systems individually permitted on or after December 30, 1983, a compliance boundary shall be established 250 feet from the waste boundary, or 50 feet within the property boundary, whichever point is closer to the source." 15A NCAC 02L.0107(b)

"For disposal systems individually permitted prior to December 30, 1983, the compliance boundary is established at a horizontal distance of 500 feet from the waste boundary or at the property boundary, whichever is closer to the source." 15A NCAC 02L.0107(a)

"A review boundary is established around any disposal system midway between the compliance boundary and the waste boundary." 15A NCAC 02L.0108

*0-10 Years*Costs and Benefits Associated with Proposed Rule Changes to 15A NCAC 02L .0202 Groundwater Quality Standards

With Two Percent Inflation

Appendix K

	2011-	2012-	2013-	2014-	2015-	2016-	2017-	2018-	2019-	2020-	2021-
Fiscal Year	12	13	14	15	16	17	18	19	20	21	22
Year Number	0	1	2	3	4	5	6	7	8	9	10
Costs											
Private Company Well Closure Costs	\$0	\$5,834	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Total Costs	\$0	\$5,834	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Benefits											
State Benefits											
NCDOT Reduced Reporting	\$0	\$3,672	\$3,537	\$3,396	\$3,247	\$3,091	\$2,928	\$2,757	\$2,578	\$2,390	\$2,194
NCDENR Opportunity Cost Savings	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Private Company Benefits											
Monitoring Cost Savings to Private											
Companies	\$0	\$5,969	\$6,088	\$6,210	\$6,334	\$6,461	\$6,590	\$6,722	\$6,857	\$6,994	\$7,134
Well Closure Savings	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Operations and Maintenance Cost Savings		\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Total Benefits	\$0	\$9,641	\$9,626	\$9,606	\$9,582	\$9,553	\$9,518	\$9,479	\$9,434	\$9,384	\$9,328
Net Impact (benefits-costs)	\$0	\$3,807	\$9,626	\$9,606	\$9,582	\$9,553	\$9,518	\$9,479	\$9,434	\$9,384	\$9,328
Total Impact (benefits+costs)	\$0	\$15,475	\$9,626	\$9,606	\$9,582	\$9,553	\$9,518	\$9,479	\$9,434	\$9,384	\$9,328
30-year Net Present Value (@7% discount rate)	\$960,152				•						

Appendix K

11-20 Years

Costs and Benefits Associated with Proposed Rule Changes to 15A NCAC 02L .0202 Groundwater Quality Standards

With Two Percent Inflation

	2022-	2023-	2024-	2025-	2026-	2027-	2028-	2029-	2030-	2031-
Fiscal Year	23	24	25	26	27	28	29	30	31	32
Year Number	11	12	13	14	15	16	17	18	19	20
Costs										
Private Company Well Closure Costs	\$0	\$0	\$0	\$0	\$0	\$27,126	\$0	\$0	\$0	\$0
Total Costs	\$0	\$0	\$0	\$0	\$0	\$27,126	\$0	\$0	\$0	\$0
Benefits										
State Benefits										
NCDOT Reduced Reporting	\$1,989	\$1,776	\$1,552	\$1,319	\$1,077	\$824	\$560	\$286	\$0	\$0
NCDENR Opportunity Cost Savings	\$0	\$0	\$0	\$0	\$0	\$1,145	\$1,168	\$1,191	\$1,215	\$1,239
Private Company Benefits										
Monitoring Cost Savings to Private										
Companies	\$7,276	\$7,422	\$7,570	\$7,722	\$7,876	\$16,432	\$16,761	\$17,096	\$17,438	\$17,787
Well Closure Savings	\$0	\$0	\$0	\$0	\$7,698	\$0	\$0	\$0	\$0	\$0
Operations and Maintenance Cost										
Savings	\$0	\$0	\$0	\$0	\$0	\$219,646	\$224,039	\$228,519	\$233,090	\$237,752
Total Benefits	\$9,266	\$9,197	\$9,123	\$9,041	\$16,651	\$238,047	\$242,527	\$247,092	\$251,743	\$256,778
Net Impact (benefits-costs)	\$9,266	\$9,197	\$9,123	\$9,041	\$16,651	\$210,920	\$242,527	\$247,092	\$251,743	\$256,778
Total Impact (benefits+costs)	\$9,266	\$9,197	\$9,123	\$9,041	\$16,651	\$265,173	\$242,527	\$247,092	\$251,743	\$256,778

Appendix K
21-30 Years

Costs and Benefits Associated with Proposed Rule Changes to 15A NCAC 02L .0202 Groundwater Quality Standards

With Two Percent Inflation

	2032-	2033-	2034-	2035-	2036-	2037-	2038-	2039-	2040-	2041-
Fiscal Year	33	34	35	36	37	38	39	40	41	42
Year Number	21	22	23	24	25	26	27	28	29	30
Costs										
Private Company Well										
Closure Costs	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Total Costs	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Benefits										
State Benefits										
NCDOT Reduced Reporting	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
NCDENR Opportunity Cost										
Savings	\$1,264	\$1,289	\$1,315	\$1,341	\$1,368	\$1,396	\$1,424	\$1,452	\$1,481	\$1,511
Private Company Benefits										
Monitoring Cost Savings to										
Private Companies	\$18,143	\$18,505	\$18,875	\$19,253	\$19,638	\$20,031	\$20,431	\$20,840	\$21,257	\$21,682
Well Closure Savings	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	35,793
Operations and Maintenance										
Cost Savings	\$242,507	\$247,357	\$252,304	\$257,350	\$262,497	\$267,747	\$273,102	\$278,564	\$284,135	\$289,818
Total Benefits	\$261,913	\$267,151	\$272,495	\$277,944	\$283,503	\$289,173	\$294,957	\$300,856	\$306,873	\$348,803
Net Impact (benefits-costs)	\$261,913	\$267,151	\$272,495	\$277,944	\$283,503	\$289,173	\$294,957	\$300,856	\$306,873	\$348,803
Total Impact (benefits+costs)	\$261,913	\$267,151	\$272,495	\$277,944	\$283,503	\$289,173	\$294,957	\$300,856	\$306,873	\$348,803

Assumptions:

- It will take 30 years to remediate Rhodia's 1,1-DCE groundwater contamination to the current standard of 7 ug/L using the existing pump-and-treat technology.
- It will take 15 years to remediate Rhodia's 1,1-DCE groundwater contamination to the proposed standard of 350 ug/L using the existing pump-and-treat technology.
- Eleven monitoring wells that are sampled twice a year can be closed immediately because the 1,1-DCE concentration is below 350 ug/L, otherwise they would have been closed in 15 years.
- Thirty-eight monitoring wells, seven of which are monitored twice a year and thirty-one of which are monitored once a year, can be closed in fifteen years, the amount of time it is estimated to take Rhodia to cleanup 1,1-DCE groundwater contamination to the proposed standard of 350 ug/L (otherwise, these wells would have been closed in 30 years).
- Pump-and-treat operation and maintenance costs will be reduced by fifteen years if the 1,1-DCE groundwater standard is changed to 350 ug/L.
- It will cost \$520 per well (49 wells) to close monitoring well s in accordance with Title 15A NCAC 02C .0113 regulations.
- NCDENR oversight of Rhodia will be reduced by 15 years.
- Inflation will increase annually at a rate of 2%.

Appendix L

0 – 10 years

Private Industry Costs and Benefits with the Proposed Rule Change

With Two Percent Inflation

	2011-	2012-	2013-	2014-	2015-	2016-	2017-	2018-	2019-	2020-	2021-
Fiscal Year	12	13	14	15	16	17	18	19	20	21	22
Year Number	0	1	2	3	4	5	6	7	8	9	10
Costs											
Well Closure Costs	\$0	\$5,834	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Total Costs	\$0	\$5,834	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Benefits											
Monitoring Cost Savings	\$0	\$5,969	\$6,088	\$6,210	\$6,334	\$6,461	\$6,590	\$6,722	\$6,857	\$6,994	\$7,134
Well Closure Savings	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Operations and Maintenance Cost Savings	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Total Benefits	\$0	\$5,969	\$6,088	\$6,210	\$6,334	\$6,461	\$6,590	\$6,722	\$6,857	\$6,994	\$7,134
Net Impact (benefits-costs)	\$0	\$135	\$6,088	\$6,210	\$6,334	\$6,461	\$6,590	\$6,722	\$6,857	\$6,994	\$7,134
Total Impact (benefits+costs)	\$0	\$11,803	\$6,088	\$6,210	\$6,334	\$6,461	\$6,590	\$6,722	\$6,857	\$6,994	\$7,134
30-year Net Present Value			•	•	•	•	•		•	•	•
(@7% discount rate)	\$930,522										

Appendix L

11 – 20 years

Private Industry Costs and Benefits with the Proposed Rule Change

With Two Percent Inflation

	2022-	2023-	2024-	2025-	2026-	2027-	2028-	2029-	2030-	2031-
Fiscal Year	23	24	25	26	27	28	29	30	31	32
Year Number	11	12	13	14	15	16	17	18	19	20
Costs										
Well Closure Costs	\$0	\$0	\$0	\$0	\$0	\$27,126	\$0	\$0	\$0	\$0
Total Costs	\$0	\$0	\$0	\$0	\$0	\$27,126	\$0	\$0	\$0	\$0
Benefits										
Monitoring Cost Savings	\$7,276	\$7,422	\$7,570	\$7,722	\$7,876	\$16,432	\$16,761	\$17,096	\$17,438	\$17,787
Well Closure Savings	\$0	\$0	\$0	\$0	\$7,698	\$0	\$0	\$0	\$0	\$0
Operations and Maintenance										
Cost Savings	\$0	\$0	\$0	\$0	\$0	\$219,646	\$224,039	\$228,519	\$233,090	\$237,752
Total Benefits	\$7,276	\$7,422	\$7,570	\$7,722	\$15,574	\$236,078	\$240,800	\$245,616	\$250,528	\$255,538
Net Impact (benefits-costs)	\$7,276	\$7,422	\$7,570	\$7,722	\$15,574	\$208,952	\$240,800	\$245,616	\$250,528	\$255,538
Total Impact (benefits+costs)	\$7,276	\$7,422	\$7,570	\$7,722	\$15,574	\$263,204	\$240,800	\$245,616	\$250,528	\$255,538

Appendix L

21 - 30 years

Private Industry Costs and Benefits with the Proposed Rule Change

With Two Percent Inflation

	2032-	2033-	2034-	2035-	2036-	2037-	2038-	2039-	2040-	2041-
Fiscal Year	33	34	35	36	37	38	39	40	41	42
Year Number	21	22	23	24	25	26	27	28	29	30
Costs										
Well Closure Costs	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Total Costs	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0
Benefits										
Monitoring Cost Savings	\$18,143	\$18,505	\$18,875	\$19,253	\$19,638	\$20,031	\$20,431	\$20,840	\$21,257	\$21,682
Well Closure Savings	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$35,793
Operations and Maintenance										
Cost Savings	\$242,507	\$247,357	\$252,304	\$257,350	\$262,497	\$267,747	\$273,102	\$278,564	\$284,135	\$289,818
Total Benefits	\$260,649	\$265,862	\$271,179	\$276,603	\$282,135	\$287,778	\$293,533	\$299,404	\$305,392	\$347,292
Net Impact (benefits-costs)	\$260,649	\$265,862	\$271,179	\$276,603	\$282,135	\$287,778	\$293,533	\$299,404	\$305,392	\$347,292
Total Impact (benefits+costs)	\$260,649	\$265,862	\$271,179	\$276,603	\$282,135	\$287,778	\$293,533	\$299,404	\$305,392	\$347,292

Appendix M *0-20 Years*

Public Water Supply Systems Costs with the Proposed Change

Unadjusted For Future Inflation

Fiscal Year	2011-12	2012-13	2013-14	2014-15	2015-16	2016-17	2017-18	2018-19	2019-20	2020-21	2021-22
Year Number	0	1	2	3	4	5	6	7	8	9	10
Costs											
Estimated Capital Expenditure											
(Inflated to 2012)	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$203,750
Annual Operation & Maintenance											
(Inflated to 2012)	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$68,460
Annual Monitoring	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$600
Total Costs	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$272,810
20 Year Net Present Value (@7%											
discount rate)	(\$455,756)										

Fiscal Year	2022-23	2023-24	2024-25	2025-26	2026-27	2027-28	2028-29	2029-30	2030-31	2031-32
Year Number	11	12	13	14	15	16	17	18	19	20
Costs										
Estimated Capital Expenditure	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$203,750
Annual Operation &										
Maintenance (Inflated to 2012)	\$68,460	\$68,460	\$68,460	\$68,460	\$68,460	\$68,460	\$68,460	\$68,460	\$68,460	\$136,920
Annual Monitoring	\$600	\$600	\$600	\$600	\$600	\$600	\$600	\$600	\$600	\$1,200
Total Costs	\$69,060	\$69,060	\$69,060	\$69,060	\$69,060	\$69,060	\$69,060	\$69,060	\$69,060	\$341,870

Appendix M *0-20 Years*

Public Water Supply Systems Costs with the Proposed Change

Adjusted For Future Two Percent Inflation

Fiscal Year	2011-12	2012-13	2013-14	2014-15	2015-16	2016-17	2017-18	2018-19	2019-20	2020-21	2021-22
Year Number	0	1	2	3	4	5	6	7	8	9	10
Costs											
Estimated Capital Expenditure		\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$248,370
Annual Operation &											
Maintenance		\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$83,452
Annual Monitoring		\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$731
Total Costs	\$0	\$0	\$0	\$0	<i>\$0</i>	<i>\$0</i>	<i>\$0</i>	<i>\$0</i>	<i>\$0</i>	\$0	\$332,554
20 Year Net Present Value											
(@7% discount rate)	(\$605,840)										

Fiscal Year	2011-12	2022-23	2023-24	2024-25	2025-26	2026-27	2027-28	2028-29	2029-30	2030-31	2031-32
Year Number	0	11	12	13	14	15	16	17	18	19	20
Costs											
		\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$0	\$302,762
Estimated Capital Expenditure											
Annual Operation &		\$85,121	\$86,824	\$88,560	\$90,332	\$92,138	\$93,981	\$95,861	\$97,778	\$99,733	\$203,456
Maintenance											
		\$746	\$761	\$776	\$792	\$808	\$824	\$840	\$857	\$874	\$1,783
Annual Monitoring						-	-	-			
		\$85,867	\$87,585	\$89,336	\$91,123	\$92,946	\$94,805	\$96,701	\$98,635	\$100,607	\$508,001
Total Costs	\$0										

Appendix N: Summary of Division of Waste Management Site Information

Name & Address	DWM Program with Oversight *	Is 1,1- dichloroethylene (1,1-DCE) present in groundwater above 7 ug/L?	Is 1,1- dichloroethylene (1,1- DCE) present in groundwater above 350 u/L?	What constituent(s) are driving site assessment, remediation, and other action?
Rhodia Inc. 207 Telegraph Drive Gastonia, NC	IHSB	Yes	Yes	1,1-DCE
Radiator Specialty 100 Radiator Road Indian Trail, NC	HWS	Yes	Yes	1,4-dioxane, PCE, TCE, 1,1-DCA, 1,2-DCA
Suttle Avenue, LLC Formerly Radiator Specialty Intersection of Wilkinson Blvd and Suttle Ave. Charlotte, NC	BFs 06016-02-60 HWS NCD003149663 Also in IHSB Inventory (1900 Wilkinson Ave)	No Maximum on site was 8.1 ug/L and it is now below 7 ug/L	No	PCBs and chlorinated solvents (PCE, TCE, vinyl chloride)
Intersection of Pearson's Turnpike and Chicksaw Road Gastonia Township, NC	BFs "MTE Hydraulics"	No	No	Known petroleum & chlorinated Solvent groundwater Contamination has been remediation to standard
The Park Ministries, Inc. 2500 Independence Blvd Charlotte, NC	BFs 10038-06-60 Know as "Merchandise Mart"	Yes 230 ug/L sampled on 02/02/06	No	No active remediation, only Land Use Restrictions. Risk drivers are chlorinated solvents 1,1-DCE, PCE & TCE. All above 2L standard
Camden Square: Design Center Carolinas, LLC 118-120 West Worthington Ave. Charlotte, NC	BFs	Yes 480 μg/L in MW-1 1,200 μg/L in MW-3	?	Chlorinated solvents
Intersection of Highway 321 and Rankin Lake Road Gastonia, NC	BFs 10063-06-36 Know as "Former ATS Manufacturing"	No Detected in three wells at 52 ppb, 49 ppb & 20 ppb but now below 7 ug/L	No	No active remediation, only Land Use Restrictions. Chlorinated solvents (1,1-DCE, PCE, 1,1,1-TCA) above 2L required a BFA.
Former Ashland, Inc. 1415 South Bloodworth St Raleigh, NC	HWS	Yes	No	PCE & degradation products (TCE, cis-1,2- DCE, vinyl chloride). Also present in significantly lower levels: 1,1- DCE,carbon tetrachloride, chloroform, 1,1-DCA, 1,1,1-TCA and methylene chloride
Former Ashland, Inc. 2802 Patterson Street Greensboro, NC	HWS	Yes	Yes	PCE, TCE and numerous other VOCs and SVOCs present above 2L.

^{*(}Inactive Hazardous Sites Branch (IHSB), Hazardous Waste Section (HWS), Brownfields Program(BFs)

McGuireWoods LLP 201 North Tryon Street P.O. Box 31247 (28231) Charlotte, NC 28202 Phone: 704.343.2000 Fax: 704.343.2300 www.mcguirewoods.com

Benne C. Hutson Direct: 704.343.2060 **McGUIREWOODS**

bhutson@mcguirewoods.com Direct Fax: 704.444.8739

RECEIVED

February 1, 2011

FEB 2 2011

NCDENR MRO IHSB

George P. Adams, PG & EI
Environmental Engineer II
North Carolina Department of
Environment & Natural Resources
Division of Waste Management
Superfund Section – Inactive Hazardous
Sites Branch
610 East Center Avenue, Suite 301
Mooresville, NC 28115

Re:

Rhodia (Former Rhone – Poulenc Facility)

207 Telegraph Road

Gastonia, Gaston County, North Carolina

IHSB # NONCD 0002395-Former APS No. 8919

Dear George:

I have enclosed with this letter the 2010 Annual Groundwater Sampling and Surface Water Sampling Results and Annual Groundwater Extraction System Performance Report for this site.

If you have any questions regarding this report, please let me know.

Sincerely

Benne C. Hutson

BCH Enclosure

cc: Rhodia Management Team (w/enclosure)



FEB 2 2011

NCDENR MRO IHSB

207 Telegraph Drive Gastonia, North Carolina Incident No. 8919

Prepared for:

Rhodia Inc.

Prepared by:



Charlotte, North Carolina January 2011

2010 Annual Groundwater and Surface Water Sampling Results RECEIVED

FEB 2 2011

and

NCDENR MRO IHSB

2010 Annual Groundwater Extraction System Performance Report

207 Telegraph Drive Gaston County, North Carolina Incident No. 8919

Prepared for:

Rhodia Inc.

Prepared by:



Charlotte, North Carolina

January 2011

Daniel Brown Staff Engineer

Tegwyn Williams, L.G. Project Manager NC Licensed Geologist License No. 1501

Introduction

This report presents the results of semi-annual groundwater and surface water sampling performed by CH2M HILL at the site located at 207 Telegraph Drive in Gaston County¹. This periodic sampling was performed in accordance with the February 1998 groundwater remediation system Performance Monitoring Plan (PMP) and subsequent modifications to the PMP required by the North Carolina Department of Environment and Natural Resources (NCDENR) in a letter to Rhodia dated May 3, 2001. The PMP, as modified, addresses the long-term monitoring of the effectiveness of the groundwater extraction component of the remediation system. The PMP includes requirements for: (1) quarterly measurements of water levels in all site monitoring wells, and (2) semi-annual and annual sampling of various monitoring wells, extraction wells, and surface water sampling locations at the project site as listed in Table 1.

The semi-annual monitoring event was conducted in March 2010, and the annual monitoring event was conducted in September 2010. The September 2010 monitoring event documented in this report included all of the monitoring wells, extraction wells, and surface water sampling locations listed in Table 1.

NCDENR also requires submission of an annual groundwater extraction system performance report. Since the September 2010 monitoring event represents the most comprehensive round of sampling for the year, it is the basis for the annual assessment of extraction system performance. Therefore, this report also serves as the 2010 Annual Groundwater Extraction System Performance Report.

The groundwater remediation system operating at the site is a pump-and-treat system consisting of five groundwater extraction wells (CP-1, EW-2, EW-3, EW-4, and EW-5) four of which are located on the site and, a low-profile air stripper, and a control system located on neighboring property owned by Rhodia. The locations of the groundwater extraction wells are shown in **Figure 1**.

In July 2007, Rhodia installed a water transfer system to pump treated groundwater from the remediation system on this neighboring property to a storage tank located on the site, which is currently owned by Lubrizol Advanced Materials (LAM), where it is used as process water. Implementation of this system followed receipt of a determination from NCDENR's Hazardous Waste Section that the treated effluent would not be classified as a hazardous waste under RCRA. Whenever LAM's storage tank is full the treated groundwater is discharged to the unnamed creek at Rhodia's NPDES outfall.

1

¹ The site was formerly owned and operated by Rhone-Poulenc Inc. On July 29, 1994, Rhone-Poulenc sold this site to the B.F. Goodrich Company, Performance Resins and Emulsions Division under an agreement that provided that Rhone-Poulenc maintained responsibility for the groundwater and surface water remediation and sampling required pursuant to the October 19, 1993 Notice of Violation (NOV) and the November 22, 1993 Notice of Requirement (NOR). Effective January 1, 1998, Rhone-Poulenc restructured its worldwide businesses by separating its life sciences and chemicals businesses, which restructuring included the transfer of substantially all of Rhone-Poulenc's chemical businesses and assets to a newly formed corporation named Rhodia Inc. Rhodia is not a successor to Rhone-Poulenc Inc. (now known as Bayer CropScience Inc.) Rhodia has also never owned or operated this site. Rather, Rhodia has performed work at this site pursuant to a contractual obligation to facilitate Rhone-Poulenc's compliance with the NOV and NOR. Rhodia has been performing this contractual obligation since its formation on January 1, 1998.

The remainder of this report is organized into the following sections:

- "Background" summarizes the conceptual model of site hydrogeologic conditions and reiterates the extraction system remedial performance objectives presented in the PMP.
- "Hydraulic Gradient Monitoring" summarizes the water level measurement data collected during performance monitoring activities and evaluates the performance of the extraction system with respect to maintaining hydraulic containment of groundwater contamination.
- "Groundwater and Surface Water Sampling" presents the results of the September 2010 groundwater and surface water sampling activities and evaluates the current spatial distribution of the major groundwater contaminant plumes.
- "Groundwater Contaminant Mass Removal" discusses extraction well sampling
 results and discharge rates, as well as calculations of the estimated mass of volatile
 organic compounds (VOCs) removed from groundwater by the remediation system
 during the preceding 12 months. "Remedial Alternatives" summarizes the on-going
 evaluation of alternative remedial technologies and their potential applicability to
 this site.
- "Conclusions" summarizes the conclusions reached based on the data and information presented in this report.

Appendix A includes tables summarizing extraction well and monitoring well construction data. **Appendix B** contains the laboratory analytical data reports for the 2010 groundwater and surface water sampling.

Background

Site Conceptual Model

A conceptual model of geologic and hydrogeologic conditions at the site has been developed from data generated as part of the Comprehensive Site Assessment (CSA).

Two aquifers have been identified in the vicinity of the site: the *water table aquifer* and the *deep bedrock aquifer*. These aquifers are distinguished by their independent hydraulic behavior.

Within the water table aquifer, there are two distinct geologic units: (1) a layer of saprolite within which the water table occurs over the vast majority of the site, and (2) a relatively shallow portion of the underlying fractured bedrock, which is referred to as the bedrock unit of the water table aquifer. As the hydraulic interconnection between these two geologic units is high, they have been grouped into one aquifer called the water table aquifer.

The *deep bedrock aquifer* consists of the fractured bedrock underlying the water table aquifer. The CSA data indicate that the *deep bedrock aquifer* is not well connected hydraulically to the *water table aquifer* in the vicinity of the site.

Remedial Objectives of the Groundwater Extraction System

As presented in the PMP, the remedial objectives of the groundwater extraction system are as follows:

- Primary Objective: Hydraulically contain and control the movement of groundwater contaminant plumes (namely 1,1-dichloroethene [1,1-DCE], and petroleum-related compounds) in the water table aquifer to prevent further migration. This is being achieved by pumping the extraction wells to produce and maintain hydraulic gradients that are directed inward (toward the extraction wells) from a containment zone boundary defined along portions of the site's property boundaries. The inward hydraulic gradients are intended to prevent migration of the contaminant plumes in the water table aquifer beyond the containment zone boundary. The containment zone boundary is shown in Figure 1.
- <u>Secondary Objective</u>: Reduce the concentration and mass of dissolved contaminants in the water table aquifer.

With respect to these remedial objectives, a portion of the groundwater extraction system (extraction wells CP-1, EW-2, and EW-3) was designed as part of an interim remedial measure required by NCDENR to hydraulically contain and control the plume of 1,1-DCE in the *bedrock unit* of the *water table aquifer*. The containment and control of the petroleum compound plume was the objective of the additional extraction wells (EW-4 and EW-5) that became operational in late 1998.

Hydraulic Gradient Monitoring

Groundwater levels were measured in monitoring wells and extraction wells on December 22, 2009, March 22, 2010, June 25, 2010, and September 10, 2010. The four water level gauging events were conducted while the groundwater remediation system was operating. These measurements are summarized in Table 2. Table 2 also presents the calculated water level elevations for each well based on the surveyed reference points from which the water level depths were measured. Figure 2 shows the locations of all monitoring wells and extraction wells. Well construction data for all monitoring and extraction wells, including screen interval depths, are summarized in Appendix A.

The September 10, 2010 water level data were used to produce two potentiometric surface contour maps (Figures 3 and 4) of the *water table aquifer*. Figure 3 depicts the potentiometric surface contours of the shallowest portion of the *water table aquifer*. The monitoring well data used to prepare this contour map were primarily from wells screened in saprolitic materials, except in the northwest corner of the site where the water table initially occurs in the bedrock unit. Figure 4 depicts the potentiometric surface contours of the *bedrock unit* of the *water table aquifer*.

The information portrayed in Figures 3 and 4 was used to create Figure 5, which depicts the minimum spatial extent of the remediation system zone of capture in the *water table aquifer*. As shown in Figure 5, the zone of capture within the saprolite unit of the *water table aquifer* is slightly less extensive than that in the *bedrock unit* of the *water table aquifer*.

Groundwater and Surface Water Sampling

Groundwater Sampling

Monitoring well and extraction well sampling was performed during the periods of March 23-25, 2010 and September 27- October 1, 2010. The samples were analyzed for selected VOCs either by Environmental Protection Agency (EPA) Method 601 (purgeable halocarbons) or by both EPA Methods 601 and 602 (purgeable aromatics), as indicated in **Table 1**. Prism Laboratories, Inc. of Charlotte, North Carolina performed the laboratory analyses.

The monitoring wells were purged and sampled using a variety of sampling equipment, including QED Well Wizard dedicated environmental sampling pumps, a Grundfos Redi-Flo2™ portable environmental sampling pump, and disposable bailers. In order to document the well purging process, water quality parameters (specific conductance, pH, and temperature) were monitored until stable readings were achieved.

Extraction wells CP-1, EW-2, EW-3, EW-4 and EW-5 were sampled directly from sampling ports located at each well head.

The results of the 2010 groundwater monitoring events are summarized in Tables 3 and 4. Table 3 summarizes the results for samples analyzed for purgeable halocarbons by EPA Method 601 (selected chlorinated VOCs). Table 4 summarizes the results for samples analyzed for purgeable aromatics by EPA Method 602 (selected petroleum-related compounds). Tables 3 and 4 also include the results from previous samples collected from each well. The 2010 sampling results are differentiated by shaded rows in the tables.

Table 5 summarizes the field parameters measured at the conclusion of monitoring well purging during the September 2010 monitoring event. Copies of the laboratory data sheets for the 2010 groundwater monitoring events are provided in **Appendix B**.

1,1-Dichloroethene (1,1-DCE)

Figures 6 and 7 depict the spatial distribution of 1,1-DCE in September 2010 within the saprolitic and bedrock units of the *water table aquifer*, respectively. In both figures, interpolated isoconcentration contours have been added to illustrate the distribution of 1,1-DCE.

Figure 6 shows that the areal extent of 1,1-DCE within the *water table unit* is similar to the previous monitoring event, and exhibits a general northwest-southeast orientation in response to groundwater extraction activities. Review of the historical analytical data presented in **Table 3** also indicates that the source area concentrations, represented by MW-16A, continue to decrease over time.

The distribution of 1,1-DCE in the *bedrock unit* of the *water table aquifer* depicted by Figure 7 is similar to the 2009 monitoring event. However, the concentrations detected in groundwater samples collected from several wells represent the lowest levels reported in the last 10 years, including source area well MW-16B and nearby wells MW-15C and MW-17C, and extraction well EW-3.

The groundwater sample collected from MW-38 was reported to contain a concentration of 1,1,-DCE of 12 μ g/L, consistent with the 2009 monitoring event, and slightly exceeding the current North Carolina Groundwater Quality Standard (7 μ g/L).

Trichloroethene (TCE)

Figures 8 and **9** depict the spatial distribution of TCE in groundwater within the saprolitic and bedrock units of the *water table aquifer*, respectively. The detected concentrations of TCE have generally remained constant or decreased since the previous monitoring event.

Benzene

Figures 10 and **11** depict the spatial distribution of benzene in groundwater within the saprolitic and bedrock units of the *water table aquifer*, respectively. Based upon the September 2010 monitoring results, the spatial extent of benzene in the *water table aquifer* is generally representative of all detected petroleum-related compounds. Overall, benzene concentrations were consistent with those reported in 2009.

Surface Water Monitoring

Two surface water samples (SW-1 and SW-8) were collected during the March and September 2010 monitoring events.

Table 6 summarizes the results of the 2010 surface water sampling for both purgeable halocarbons by EPA Method 601 (selected chlorinated VOCs) and purgeable aromatics by EPA Method 602 (selected petroleum-related compounds). In March, ethylbenzene was detected at a concentration of 1.9 μ g/l in the surface water sample collected from SW-1. No other target analytes were detected in the surface water samples.

Groundwater Contaminant Mass Removal

The total mass of VOCs removed by the groundwater extraction and treatment system is directly related to extraction well contaminant concentrations and discharge rates. A discussion of these items is provided in this section along with a calculation of the total mass of VOCs removed by the groundwater remediation system both during the last 12 months, and since the remediation system began operating over 13 years ago in September 1997.

The sampling results for the groundwater extraction wells are included in **Table 3** (chlorinated compounds) and **Table 4** (petroleum-related compounds), near the end of each table. The majority of the chlorinated VOC mass is now being recovered by extraction wells CP-1, EW-2, and EW-3, with the two remaining extraction wells maintaining hydraulic control over the petroleum plume.

The target discharge rates for groundwater extraction wells CP-1, EW-2, EW-3, EW-4, and EW-5 are 5, 12, 17, 20, and 1.5 gallons per minute (gpm), respectively. During the current reporting period, from September 2009 through September 2010, the average groundwater extraction system discharge rate was approximately 51.5 gpm.

The estimated total mass of 1,1-DCE and toluene removed from groundwater by the extraction and treatment system is presented in Table 7. The mass removal estimates provided in Table 7 are the result of mass balance calculations based on treatment system

influent and effluent concentrations as well as the treatment system influent flow rate. Based on the mass balance calculations, approximately 169 pounds of total VOCs were recovered from groundwater by the remediation system during the 12 months ending September 7, 2010. Since remediation efforts began in September 1997, the groundwater treatment system has removed an estimated mass of 7,192 pounds of total VOCs. Figure 12 illustrates the daily VOC removal rate and the cumulative mass of VOCs removed.

Remedial Alternatives

Several remedial technologies have been developed to contain, remove, or degrade chlorinated solvents in groundwater. However, few, if any, are effective in fractured crystalline bedrock settings such as exist at this site. Rhodia routinely evaluates existing and emerging remedial technologies to assess their potential applicability at the site. Table 8 summarizes the remedial technologies that have been considered for implementation at this site.

The primary impediments to groundwater remediation at this site are:

- The fractured crystalline bedrock setting
- Possible DNAPL present at depth within fractured bedrock
- The primary site contaminant (1,1-DCE) has been assigned a very low groundwater quality standard (7.0 μ g/L) even though, as described below, NCDENR staff have agreed that the standard should be 350 μ g/L.
- Rhodia does not own the site and relies upon the consent of the current property owner for site access not only to operate and monitor the existing remediation system, but also to make any changes to the remedial approach currently being used, all of which must be done within the constraints imposed by the current property owner's business operations.

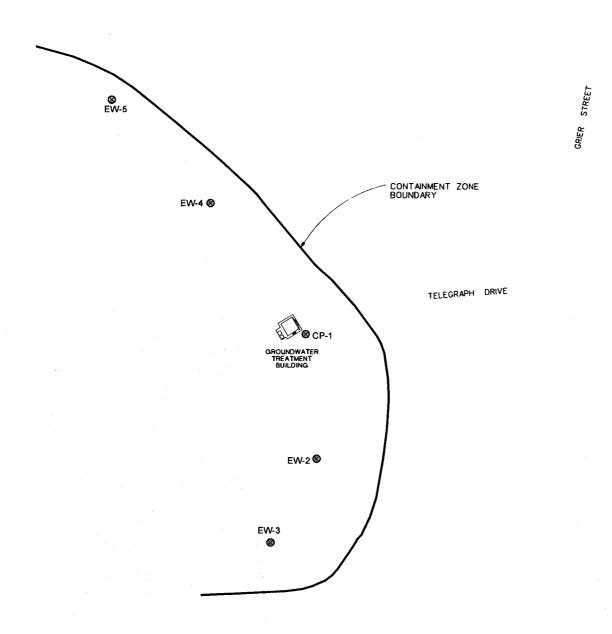
Combined, these factors render most alternative remedial technologies unfeasible, cost prohibitive, or both. However, recent studies by the United States Environmental Protection Agency have concluded that the risks associated with 1,1-DCE have been greatly overstated. In response to two rule-making proceedings by the North Carolina Environmental Management Commission (EMC), Rhodia has twice proposed that based on current scientific evidence and applicable laws and regulations, the groundwater quality standard for 1,1-DCE should be changed to 350 μ g/L, a position that NCDENR's toxicologist and staff person responsible for the second rule-making proceeding have both agreed with. Despite these positions held by NCDENR staff, the EMC has concluded in both rule making proceedings that it would not make this change. According to the September 2010 monitoring data, levels of 1,1-DCE exceeded the current standard of 7.0 μ g/L in samples collected from 21 monitoring wells. If the standard was 350 μ g/L, there would only be exceedances in 10 wells. If the groundwater quality standard for 1,1-DCE is changed to 350 μ g/L, alternate remedial technologies may become more feasible and cost effective.

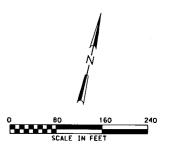
Conclusions

Based on the data and information presented in this report, the following conclusions have been reached:

- Samples collected from several monitoring wells and two extraction wells in or near the source area were reported to contain the lowest concentrations of 1,1-DCE in over 10 years of monitoring.
- The groundwater remediation system continues to meet the objectives of hydraulic containment and source reduction.
- During the 12-month period ending September 7, 2010, the groundwater remediation system removed an estimated 169 pounds of VOCs from the site groundwater. The remediation system has removed a total of approximately 7,192 pounds of VOCs from the groundwater since activation in 1997.
- If the groundwater quality standard for 1,1-DCE is changed to 350 μ g/L, alternate remedial technologies may become more feasible and cost effective.







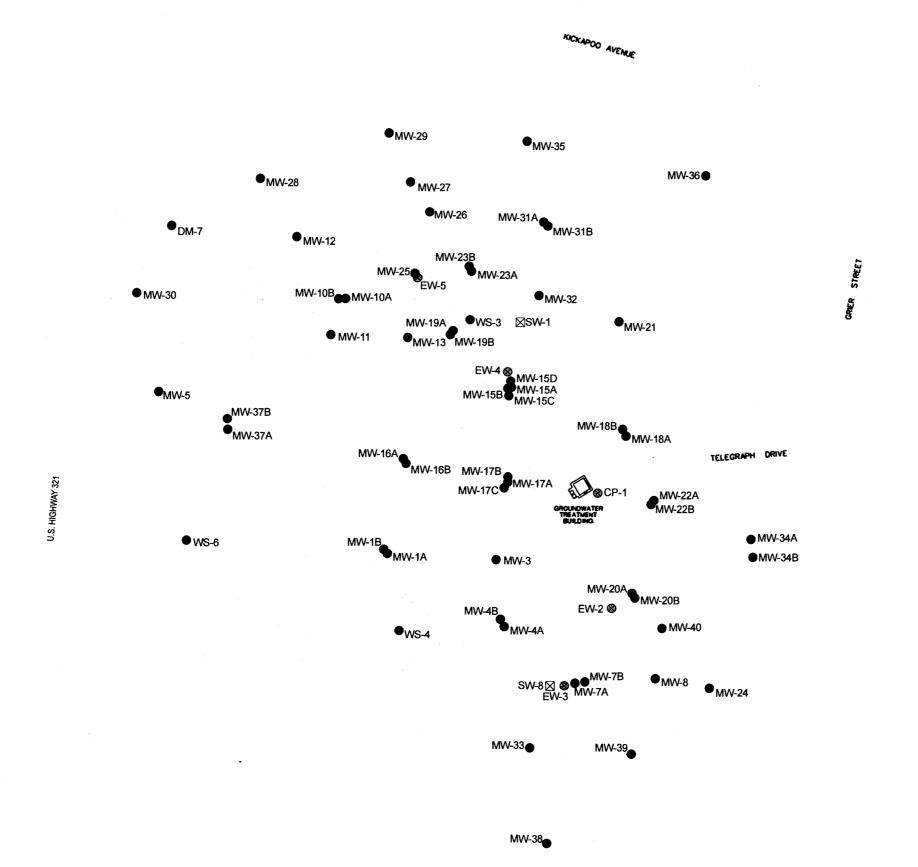
<u>Legend</u>

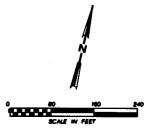
*** * * * Chain Link Fence

Stream

Figure 1 Extraction Well Location Map 207 Telegraph Drive Gastonia, North Carolina







Legend
EW-2

Groundwater Extraction Well

MW-13

Groundwater Monitoring Well

WS-4

Water Supply Well

SW-8

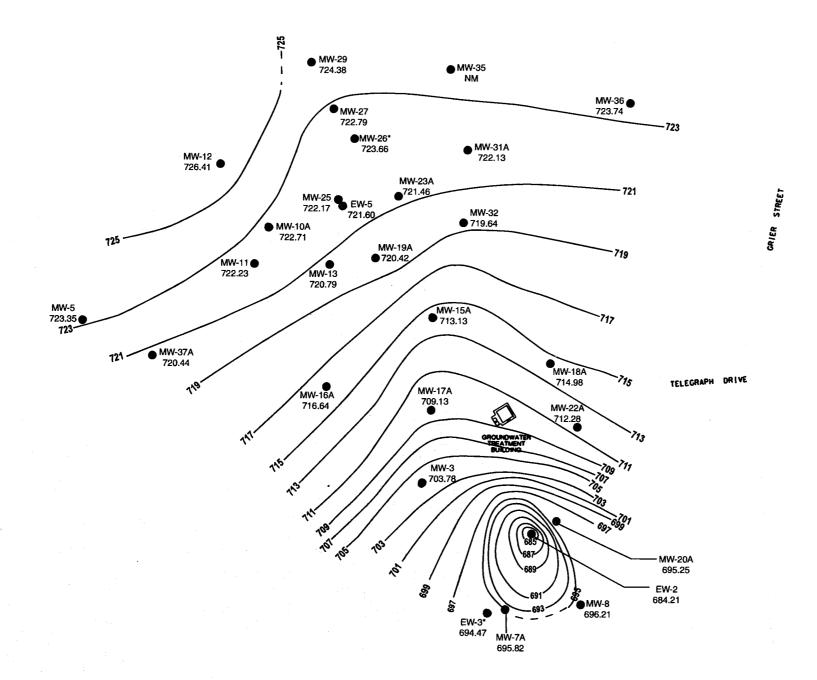
Surface Water Sampling Location

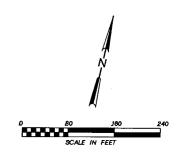
Chain Link Fence

Stream

Figure 2
Monitoring, Water Supply, and Extraction Wells
(And Surface Water Sampling Locations)
207 Telegraph Drive
Gastonia, North Carolina

KICKAPOO AVENUE





Legend

MW-13 Groundwater Monitoring Well

EW-3 ● Groundwater Extraction Well

× × × × Chain Link Fence

Stream

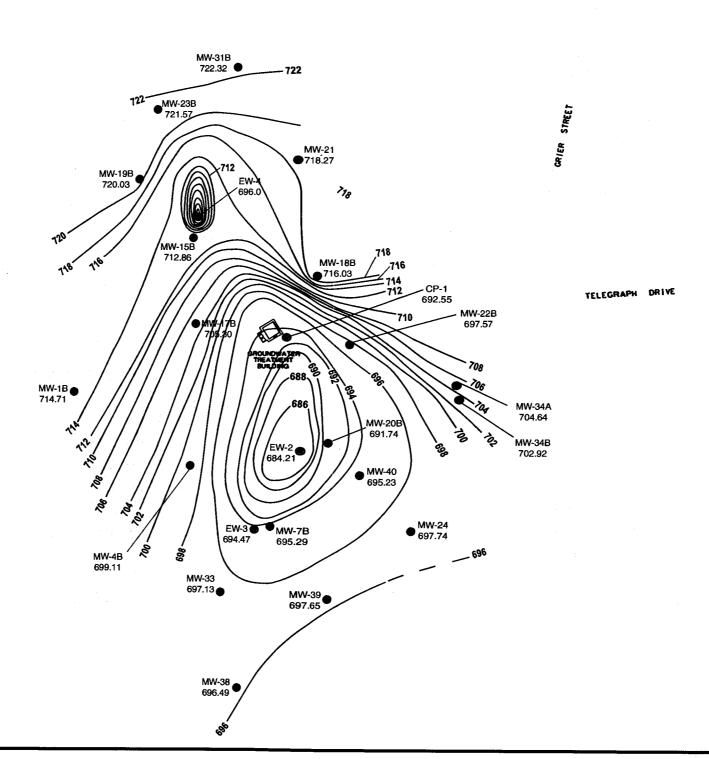
Potentiometric Contours (dashed where inferred)

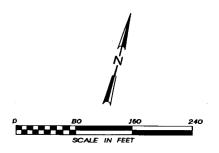
NM Not Measured

- Potentiometric surface contours have been interpolated between monitoring well locations. Actual conditions may differ from those shown here.
- Datat from MW-26* was not used to estimate the potentiometric surface.

Figure 3
Potentiometric Surface Map of the
Saprolitic Unit of the Water Table Aquifer
September 2010
207 Telegraph Drive
Gastonia, North Carolina

KICKAPOO AVEMUE





<u>Legend</u>

Stream

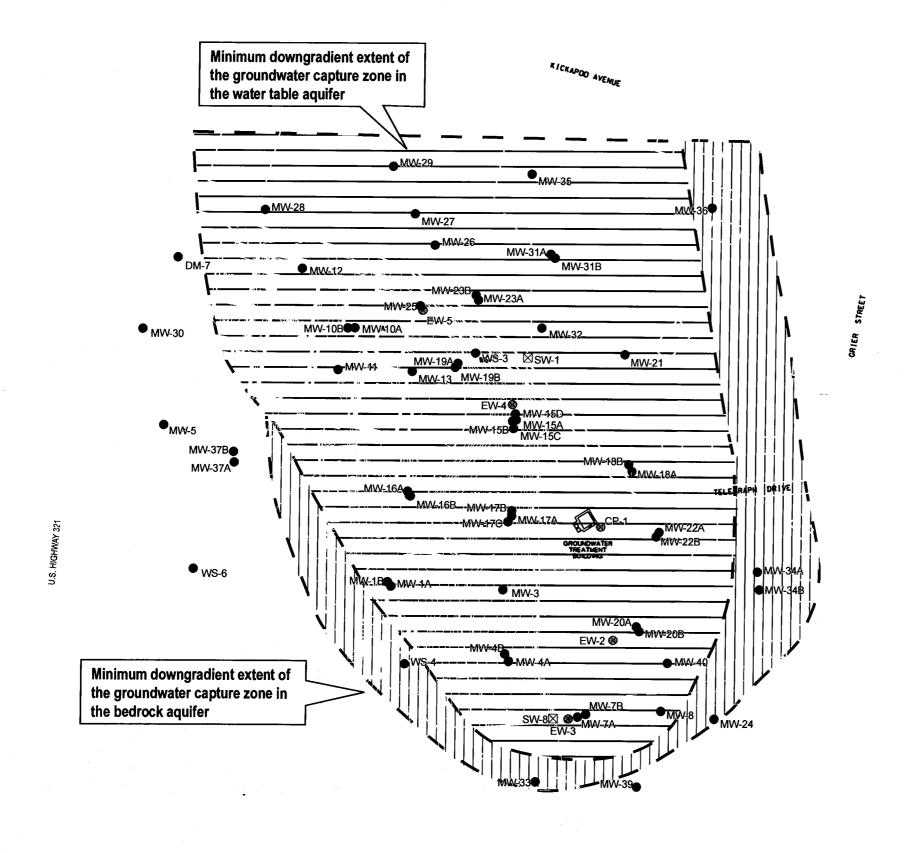
Potentiometric Contours (dashed where inferred)

NM Not Measured

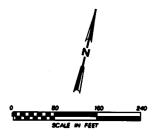
Notes:

 Potentiometric surface contou have been interpolated betwe monitoring well locations. Actu conditions may differ from tho shown here.

Figure 4
Potentiometric Surface Map of the
Bedrock Unit of the Water Table Aquife
September 2010
207 Telegraph Drive
Gastonia, North Carolina



MW-38



<u>Legend</u>

EW-2 Groundwater Extraction Well

MW-13 Groundwater Monitoring Well

WS-4 Water Supply Well

SW-2 Surface Water Sampling Location

××× Chain Link Fence

Stream

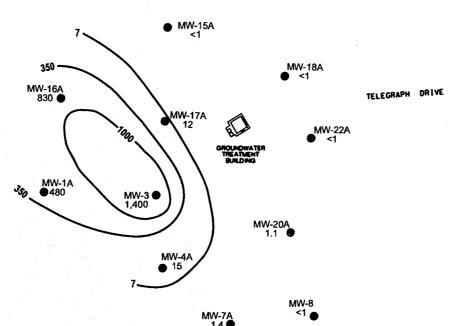
Figure 5
Capture Zones of Groundwater
Remediation Systems
September 2010
207 Telegraph Drive
Gastonia, North Carolina

KICKAPOO AVERUE

●MW-35 <1

MW-25 EW-5

MW-13 • MW-19A 3.1



0 80 160 24
SCALE IN FEET

<u>Legend</u>

MW-13 ● Groundwater Monitoring Well
<1 1,1 DCE Concentration in µg/L

1,7 DCE Concentration in p

EW-5 Groundwater Extraction Well
<1 1,1 DCE Concentration in µg/L

× × × × × Chain Link Fence

Stream

 1,1-DCE Isoconcentration Contour in µg/L

(dashed where inferred)

Figure 6
1,1-DCE Concentrations in the
Saprolitic Unit of the Water Table Aquifer
September 2010
207 Telegraph Drive
Gastonia, North Carolina

U.S. H



MW-29 ● <1

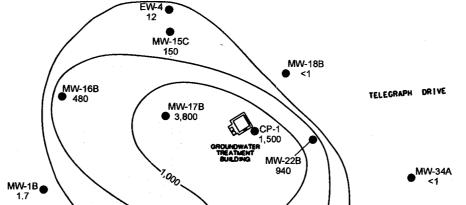
MW-27 ● <1

MW-31B ● <1

MW-23B ● <1

MW-21 ● <1 MW-19B <1 ●

MW-4B



EW-2 ● 960 MW-7B 740 ● ● MW-24 <1

MW-20B 460

MW-39/ MW-33 ● 210

1,500

Legend

→ → → Chain Link Fence

MW-15C ● Groundwater Monitoring Well 150 1,1 DCE Concentration in µg/L

CP-1 Groundwater Extraction Well

1,1 DCE Concentration in µg/L

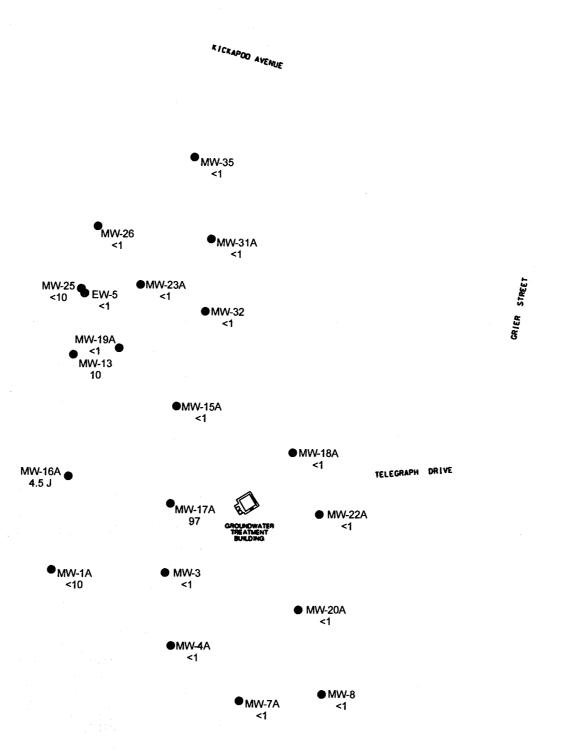
1,1-DCE Isoconcentration

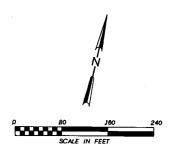
(dashed where inferred)

Contour in µg/L

Figure 7 1,1-DCE Concentrations in Bedrock Unit of the Water Table Aquifer September 2010 207 Telegraph Drive Gastonia, North Carolina





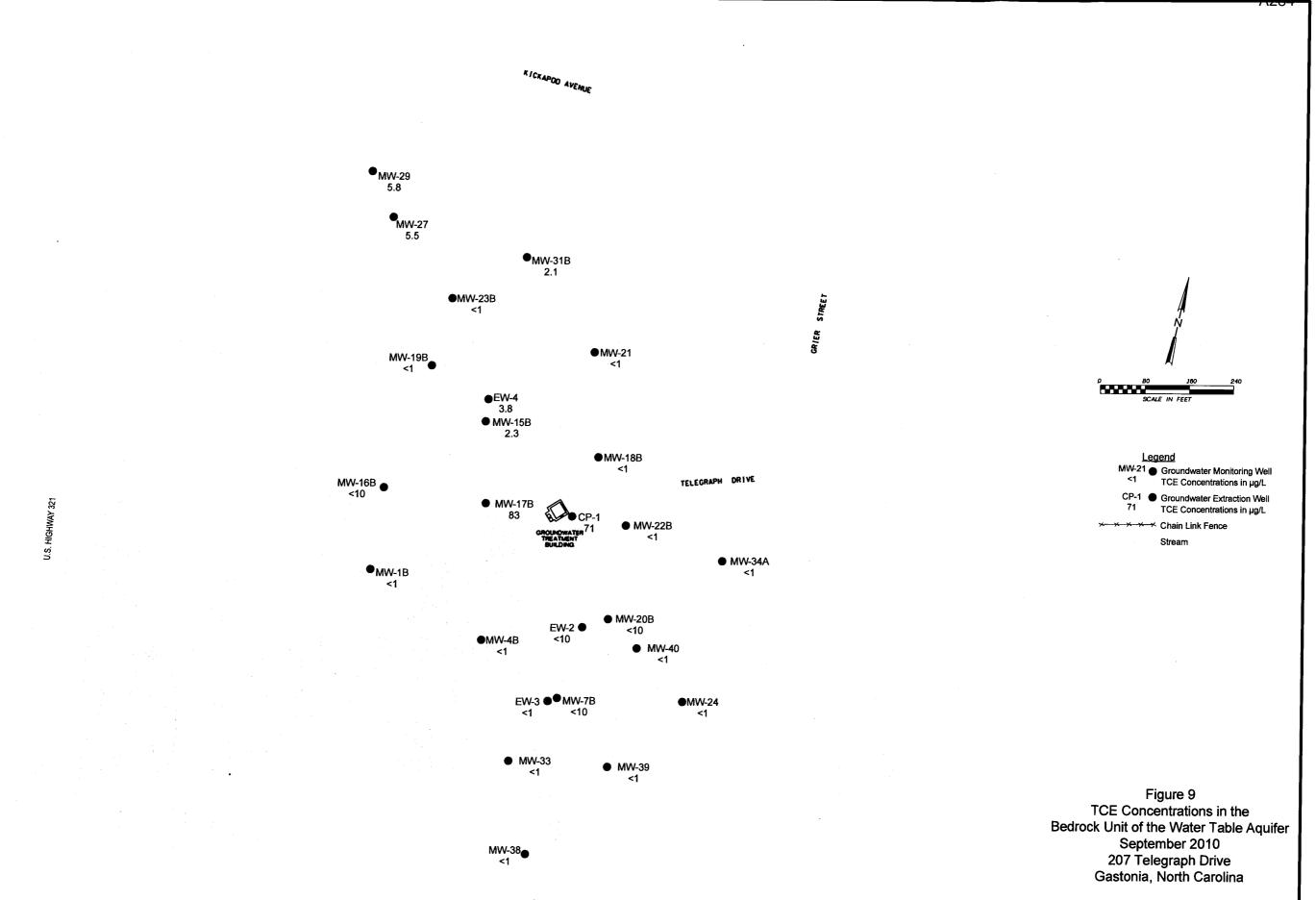


Legend

MW-32 ● Groundwater Monitoring Well

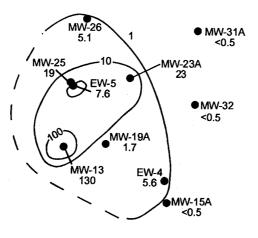
<1 TCE Concentration in μg/L</p>
EW-5 ● Groundwater Extraction Well
<1 TCE Concentration in μg/L</p>
× × × × ✓
Chain Link Fence
Stream
J Estimated Value

Figure 8
TCE Concentration in the
Saprolitic Unit of the Water Table Aquifer
September 2010
207 Telegraph Drive
Gastonia, North Carolina



KICKAPOO AVENUE

MW-35 ● <0.5



STREET

● MW-18A <0.5

TELEGRAPH DRIVE



BO 160 240 SCALE IN FEET

Legend

MW-13 Groundwater Monitoring Well Benzene Concentrations in μg/L

EW-5 Groundwater Extraction Well 7.6 Benzene Concentrations in μg/L

× × × × Chain Link Fence

Stream

Benzene Isoconcentration
 Contour in µg/L.
 (dashed where inferred)

Figure 10
Benzene Concentrations in the
Saprolitic Unit of the Water Table Aquifer
September 2010
207 Telegraph Drive
Gastonia, North Carolina

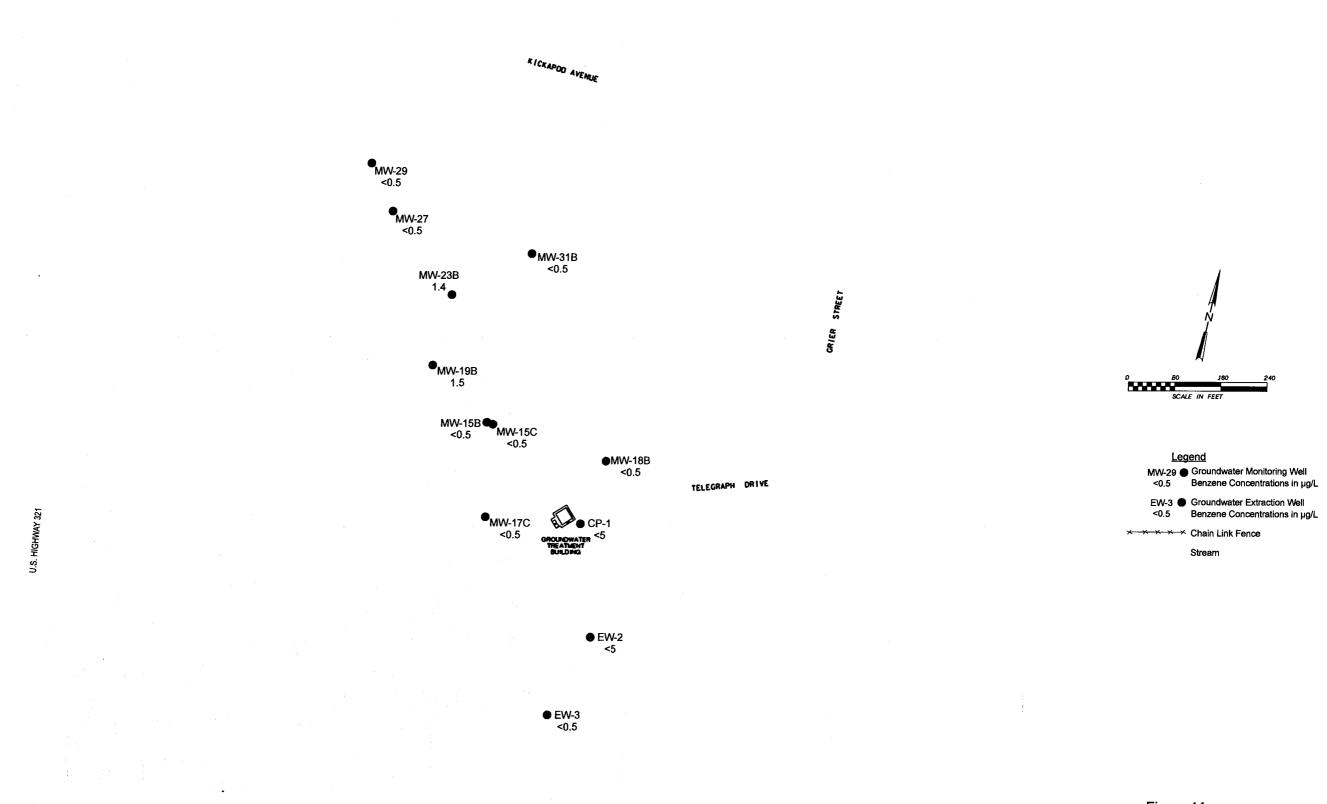
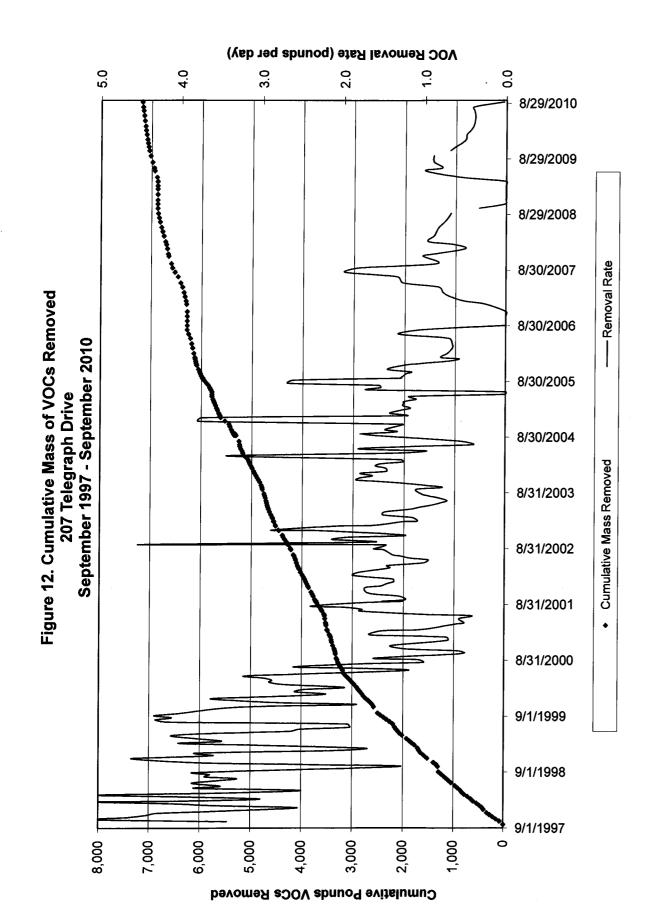
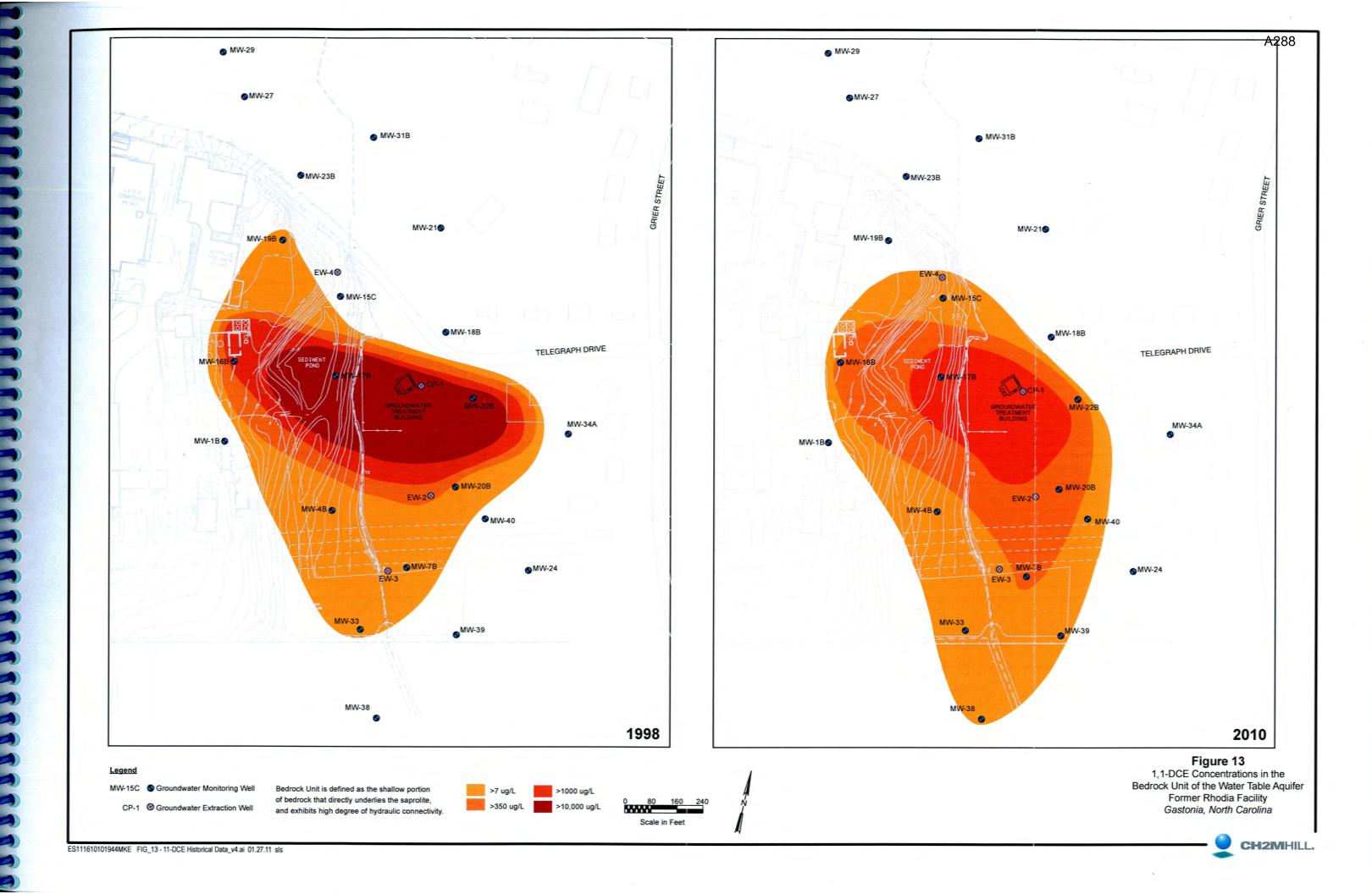


Figure 11
Benzene Concentrations in Bedrock Unit
Monitoring Wells
September 2010
207 Telegraph Drive
Gastonia, North Carolina







Tables

Schedule for Groundwater Quality Monitoring 207 Telegraph Drive, Gastonia, North Carolina	
Sampling Locations and Analyses	Sampling Schedule
EPA Method 601 compounds only: MW-4A, MW-17B, MW-20A, MW-24, MW-34A, MW-38, MW-39, MW-40	Semi-annually (1st and 3rd Quarters)
EPA Method 601 and 602 compounds: MW-17C, MW-19B, MW-31A, MW-31B, MW-32, extraction wells CP-1, EW-2, EW-3, EW-4, EW-5, surface water sampling locations SW-1 and SW-8	Semi-annually (1st and 3rd Quarters)
EPA Method 601 compounds only: MW-1A, MW-1B, MW-3, MW-4B, MW-7A, MW-7B, MW-8, MW-16A, MW-16B, MW-17A, MW-20B, MW-21, MW-22A, MW-22B, MW-33, MW-34B	Annually (3rd Quarter)
EPA Method 601 and 602 compounds: MW-13, MW-15A, MW-15B, MW-15C, MW-15D, MW-18A, MW-18B, MW-19A, MW-23A, MW-23B, MW-25, MW-26, MW-27, MW-29, MW-35	Annually (3rd Quarter)

Table 2 Water Level Measurements

207 Telegraph Drive, Gastonia, North Carolina Water-Level Elevation Water-Level Depth Measurement (feet above MSL) Point Elevation (feet) (feet above MSL) 12/22/2009 3/22/2010 6/25/2010 9/10/2010 12/22/2009 3/22/2010 6/25/2010 9/10/2010 735.17 NM 23.20 22.45 711.97 711.52 712.72 MW-1A 23,65 20.78 18.78 715.38 714.71 MW-1B 20.94 20.14 20.11 714 55 715.35 703.78 706.71 706.85 MW-3 722 56 15.71 15.85 17.04 714.32 10,10 12.05 13.77 DRY 704.22 702 27 700 55 MW-4A DRY 702.36 700.98 699.11 702.35 MW-4B 716.38 14.03 14.02 15.40 17.27 762.55 39.93 38.53 38.40 39.20 722.62 724.02 724 15 723.35 MW-5 697.80 695.82 15.00 15.99 699.45 MW-7A 712.80 13.72 13.35 16.98 699 08 14.40 17.94 698.83 697.24 695.29 698.53 MW-7B 713.23 14.70 25.44 24.79 26.38 28.31 400 US 699 73 698 14 696.21 MW-8 722.71 723.14 724.20 35.32 35.77 36.81 MW-10A 759.52 36.38 MW-10B 759.15 41.38 38.40 40.16 42.40 717.77 720.75 718.99 716.75 723.30 723.27 722.23 35.50 MW-11 757.73 35.72 34 43 34 46 722.01 33.24 33.95 728.31 727,29 726.41 32.05 33.07 727.12 MW-12 760.36 MW-13 743.36 22.17 21.23 21.71 22.57 721.19 722.13 721,65 720.79 12.10 714.12 713.13 11.47 MW-15A 725 23 11.11 10.86 11.52 MW-15B 725.71 713.72 714.19 713.50 712.86 714.29 713.53 712.84 724.25 725.37 MW-15C 10.52 9.96 10 72 11.41 713.73 717.40 717.50 715.25 716.64 6.50 7.97 10.12 718.87 8.15 MW-15D 717.67 MW-16A 735.42 18.22 17.75 17.92 18.78 717.20 704.88 703.20 32.05 705.30 29.20 30.37 MW-16B 735.25 29.95 14.83 18.81 MW-17A 723.96 12.41 17.57 711.30 711.55 706.39 709.13 707.78 707.72 710.41 705.30 MW-17B 724.11 16 33 16 39 13.70 20.59 NM 706.04 704 83 703 11 18.45 MW-17C 723.70 716.83 714.98 718.83 MW-18A 729.13 11.95 10,30 12.30 14.15 717.18 9.37 10.91 12.42 NM 717.54 716.03 10.84 MW-18B 728.45 722.07 720.42 MW-19A 742.49 21.17 20.42 21.30 22.07 721 32 721.19 722.24 22.51 720.03 MW-19B MW-20A 21.20 721.07 742 54 21 47 20.30 32.20 33.70 35.67 698.37 698.72 697 22 695 25 730.92 32.55 693.71 691.74 695.24 MW-20B 730.96 36.05 35.72 37.25 39.22 694.91 20.96 18.60 20.82 722.19 719.97 718.27 740.79 MW-21 18.80 33.79 716.08 714,30 712.28 MW-22A 734.88 20.16 20.58 22 60 714.72 37.20 700.98 699,38 697.57 35.39 700.85 MW-22B 734 77 33 92 8.82 9.81 723.01 723 87 722.45 721 46 731.27 8.26 MW-23A 723.70 722.54 732.03 732.54 10.46 722.74 MW-23B 9 29 8.33 9.49 34.80 21.95 701.19 723.65 31.99 31.35 32.88 700.55 699.66 697.74 MW-24 723.42 722.17 744.12 20.61 20.47 20.70 723.51 19.53 727.51 725.60 723.66 17.62 MW-26 745.13 19.24 MW-27 754.6 29.69 27.90 29.57 31.81 724.91 726 70 725.03 722.79 729.80 729,38 729.20 41.62 41.80 728.81 MW-28 771.00 42.19 41.20 759.44 33.18 31.30 35.06 726.26 728.14 726.44 724.38 MW-29 719.76 717.72 721.06 46.10 5.25 49.44 719.36 MW-30 767.16 47 80 47.40 722.13 722.32 6.62 723.52 724.63 723.26 7.75 MW-31A 729.88 6.36 730.68 6.93 5.70 7.11 8.36 723.75 724,98 723.57 MW-31B 720.56 719.64 8.22 720.78 721.86 MW-32 727.86 7.08 6.00 7,30 695.45 708.16 MW-33 713.73 13.64 18.28 14.80 16.60 700.09 698.93 697.13 706.71 704.64 37.22 706.81 MW-34A 741.86 35.05 33.70 35.15 706.31 727.84 34.98 36.45 704.84 702.92 MW-34B 741.29 36.13 4.41 17.85 726.43 MW-35 730.84 3.00 NM 726.21 19.67 725.56 723.74 724.86 15.72 MW-36 743.41 18.55 753.41 33.61 32.26 32.97 719.80 720.95 721.15 720.44 MW-37A 719.36 719.84 718.74 37.03 716.59 MW-37B 755.77 39.18 35 93 36.41 699.04 701.04 697.85 699.48 696.49 697.65 11.87 13.23 698.50 10,68 11.22 MW-38 709.72 MW-39 13.59 13.11 14.67 16.50 700.56 33.95 35.93 698.27 697.21 695.23 MW-40 DM-7 731.16 32.89 32.44 42.50 24.32 725.62 768.12 43.16 40,28 41.25 724.96 727.84 726.87 694.32 692.55 696.16 22.55 696.11 CP-1 716.87 20.76 20.71 36.90 38.88 687.54 687.41 686.19 684 21 35.55 35.68 EW-2 723.09 697.99 697.46 694.47 697.94 EW-3 709.26 11.32 11.27 11.80 14.79 696.58 722.78 28.02 28.60 697.74 698.14 696.00 26.46 EW-4 724.60 26.86 721.60 723.95 EW-5 742.10 18,85 18.15 19.32 20.50 723 25 15.23 722.58 717.38 721.78 WS-3 730.58 739.52 13.20 8.00 8.80 707.45 717.94 706.14 718.11 32.07 33.38 35.03 706.54 704 49 32.98 WS-4 716.66 41.90 40.62 40.45 41.17 WS-6

BP/DB

NM = Not measured.

MSL = Mean Sea Level

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

	Drive, Gastonia, N	CarOil	74									_
Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CASN 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CASI 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CASS 71-55-6 ppb	1,2-Dichloro- propane CASI 78-87-5 ppb	Trichloro- ethene (TCE) CASN 79-01-6 ppb	Tetrachloro- ethene (PCE) CASW 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASI 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CASW 156-60-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb	Chloroform CASH 67-66-3 ppb
NC Groundwate	r Quality Standard		7	6	200	0.6	3	0,7	70	100	0.3	70
MW-1A	12/15/1993	1	17	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0,5	< 0.5	< 0.5
	2/8/1994	5	18	< 5	240	< 5	< 5	< 5	< 5	< 5	< 5	< 5
	2/8/1994	1	18	<1	220	<1	<1	<1	<1	<1	< 1	<1
	5/25/1994	1	17	<1	100	<1	<1	<1	<1	<1	<1	<1
	7/25/1994	1	38	< 0.5	130	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
	9/19/1996	1	67.4	<1	10.8	<1	<1	< 1	<1	<1	< 1	<1
	8/11/1997	6	300	< 6	27	< 6	< 6	< 6	< 6	< 6	< 6	< 6
	9/2/1998	1	660	< 1	14	<1	<1	<1	<1	<1	<1	<1
	9/20/1999	1	350	<1	20	<1	<1	<1	<1	<1	<1	<l< td=""></l<>
	9/11/2000	1	560	<l< td=""><td>6.3</td><td><<u>l</u></td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><l< td=""></l<></td></l<>	6.3	< <u>l</u>	<1	<1	<1	<1	<1	<l< td=""></l<>
	8/27/2001	11	790	<1	8.8	<1	<1	<1	<1	4.3	<1	<1
	10/1/2002	10	3,800	<10	<10	<10	<10	<10	<10	<10	<10	<10
ļ	9/29/2003	.1	2,500	</td <td>2.5</td> <td><1</td> <td><1</td> <td><!--</td--><td><1</td><td><1</td><td><1</td><td><1</td></td>	2.5	<1	<1	</td <td><1</td> <td><1</td> <td><1</td> <td><1</td>	<1	<1	<1	<1
	10/4/2004	10	410	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/28/2005	10	430	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10
	10/20/2006	10	530	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/21/2007	1/10	800	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/23/2008	1/10	180	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	<1	<1
	9/16/2009	1/10	630	<1	<1	<1	<1	<1	<1	<1	< <u>l</u>	<1
	9/30/2010	10	480	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10
MW-1B	5/25/1994	1	9	< 1	< 1	< 1	<1	<1	<1	<1	<1	<1
	7/27/1994	1	1	< 1	<1	< 1	<1	< 1	<1	<1	< 1	<1
	11/1/1994	1	<1	< 1	<1	< 1	<1	<1	<1	<1	< 1	<1
	9/18/1996	1	< 1	<1	<1	<1	<1	< 1	<1	<1	<1	<1
	8/11/1997	1	<1	<1	<1	<1	<1	< 1	<1	<1	< 1	<1
	9/18/1998	1	<1	<1	<1	< 1	<1	<1	<1	<1	<1	<1
	9/30/1999	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/18/2000	1	<1	<1	<1	< <u>l</u>	<1	<1	<1	<1	<1	1.2
	8/27/2001		<1	<1	<1	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><<u>l</u></td></i<>	<1	<1	< <u>l</u>
	10/3/2002	1	2.5	<1	<1	<1	<1	<1	<1	<1	<1	< <u>l</u>
	9/29/2003	1	<1	</td <td><1</td> <td><1</td> <td><1</td> <td><<u>1</u></td> <td><1</td> <td><1</td> <td><1</td> <td><1</td>	<1	<1	<1	< <u>1</u>	<1	<1	<1	<1
-	9/30/2004	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
-	9/29/2005	1	1.7	<1	< i	<1	<1	<1	<1	. <1	< 1	<1
	10/20/2006	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
}	9/21/2007	1	<1	<1	<1	<u> </u>	<1	<1	<1	<1	<1	<1
}	9/25/2008	1	<1	<1	<1	<1	<1	< <u>l</u>	<1	<1	<1	<1
F	9/14/2009	1	<1	<1	<1	<1	<1	<1	<1	<u><1</u>	<1	<1
	25.4.06.2	1	1.7	<1	<1	<1	<1	<1	<1	<1	<1	<1
MW-3	2/8/1994	1	7	<1	<1	<1	<1	<1	<1	<1	<1	<1
}	5/24/1994	1	26	<1	<1	<1	<1	< 1	<1	<1	<1	<1
-	7/26/1994	1	100	<1	<1	<1	<1	<1	<1	<1	<1	<1
	7/26/1994	-2-	98	<2	< 2	<2	< 2	< 2	< 2	< 2	< 2	< 2
ŀ	9/18/1996	1	10	<1	< 1	<1	<1	<1	<1	<1	<1	<1
	8/12/1997	500	15,000	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500
-	9/2/1998	1	37,000	<1	<1	< 1 <1000	7	< 1 <1000	<1	<1 <1000	<1	<1
-	8/30/2001 9/29/2003	1000	11,000	<1000	<1000		<1000		<1000		<1000	< 1000
H		1	2,000	<10	<1	<1	<1	<1	<1	<1	<1	<1
ŀ	10/1/2004	10	3,100	<10	<10	<10	<10	<10	<10	<10	<10	<10
·	9/30/2005	10	5,900	< 10	< 10 <10	< 10	< 10 <10	< 10	<10	< 10	< 10 <10	< 10
L	10/17/2006 9/20/2007	100	1,000	<10		<10		<10	<10	<10		<10
		100	1,300	<1	<1	<1	<1	<1	<1	<1	< <u> </u>	<u> <1</u>
-	9/16/2009	1/100	940	<1	<1	<1	<1	<1	<1	<1	<1	<1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbona (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

207 Telegrapi	Drive, Gastonia, l	orth Caroli	ina	-								
Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CAS# 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAST 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CASN 71-55-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CASN 79-01-6 ppb	Tetrachloro- ethene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASN 136-39-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAST 156-69-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb 0.3	Chloroform CAS# 67-66-3 ppb
		Т.							· · · · · · · · · · · · · · · · · · ·		T	
MW-4A	12/15/1993	1	< 0,5	< 0.5	< 0,5	< 0.5	< 0.5	< 0.5	< 0.5	< 0,5	< 0.5	< 0.5
ĺ	2/7/1994 5/24/1994	1	<1 <1	<1 <1	<1 <1	<1 <1	<1	<1	<1	<1	<1 <1	<1 <1
	7/25/1994	1	< 0.5	< 0.5	< 0.5	<0.5	< 0.5	< 0.5	<0.5	<0.5	< 0.5	< 0.5
	9/18/1996	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
i	8/11/1997	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/18/1998	i	1.2	<1	3.9	<1	<1	<1	<1	<1	<1 <1	<1
	9/2/1998	i	14	<1	<1	<1	<1	<1	<1	<1	<1	<1
Ĭ	3/22/1999	1	19	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/20/1999	1	66	<1	2.2	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1	<1
	9/12/2000	1	27	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/1/2001	1	6.0	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><l< td=""></l<></td></l<>	<1	<1	<1	<1	<1	<1	<1	<l< td=""></l<>
	8/27/2001	11	60	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1
	3/28/2002	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/26/2003	11	1.2	<1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1
	9/25/2003	1	16	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1	<1	<1
	9/29/2004		4	<1	<1	< <u>l</u>	<1	<1	<1	<1	<1	<1
	1/14/2005	1	3,1	<1	<1	<1	<1	<1	<1	<1	<1	· <1
	6/22/2005	1	42	<1	<1	<1	< <u>i</u>	<1	<1	<1	<1	< 1
	9/28/2005	1	110	<1	< l	<1	<1	<1	1.1	<1	<1	<1
	3/8/2006	. 1	11	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/17/2006	1	35	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/7/2007	1	11	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/20/2007	1	59	<1	<1	<1	<1	<1	<1	<1	<1	<u><1</u>
	3/12/2008 5/1/2009	1	8.8	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1
	9/16/2009	1	8.6 24	<1	<1	<1	<1	<1	√ √	<1	<1	<1
	3/21/2010	1	5.9	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/27/2010	1	15	<1	<1	<1	<1	<1	<1	<1	<1	<1
MW-4B	5/24/1994	1	12	<1	38	<1	<1	<1	<1	<1	<1	<1
12	7/26/1994	i	48	<1	<1	<1	<1	<1	<1	<1	<1	<1
	11/1/1994	1	85	<1	48	<1	<1	<1	<1	<1	2	< 10
	9/18/1996	10	220	< 10	52	< 10	< 10	< 10	< 10	< 10	< 10	< 6.3
	8/12/1997	6.25	250	< 6.3	59	< 6.3	< 6.3	< 6.3	< 6.3	< 6.3	< 6.3	<1
	9/10/1998	1	200	<1	22	<1	<1	<1	<1	<1	<1	<1
	9/29/1999	1	190	<1	7.4	<1	<1	<1	<1	<1	<1	<1
	9/12/2000	1	26	< <u>1</u>	2.9	<1	<1	<1	<1	<1	<1	<1
	8/27/2001	1	240	<1	7.5	<1	<1	<1	<1	<1	<1	<1
	10/3/2002	-1	330	< <u>l</u>	4.4	<1	<1	<1	<1	<1	<1	<1
	9/26/2003	1	910	<i< td=""><td>2.1</td><td><1</td><td>2.1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><10</td></i<>	2.1	<1	2.1	<1	<1	<1	<1	<10
	10/4/2004	10	690	<10	<10	<10	<10	<10	<10	<10	<10	< 10
	9/29/2005	10	440	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	<1
	10/19/2006	10	530	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1
	9/20/2007	10	320 82	<1	<1	<1	<1	<1	<1	<1	<1	<1 <1
	9/16/2009	1/10	730	<1	<1	<1	<1	<1	<1 <1	<i>1</i>	<1	<1
	9710/2009	1/10	250	<1	<1	<1	<1	<1	<1	<1	<1	<1
MW-5	2/8/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	5/25/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	7/27/1994	1	<1	<1	< i	<1	<1	< 1	<1	<1	< l	< 0,5
MW-6	12/15/1993	1	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	<1
	2/8/1994	1	24	<1	< 1	<1	<1	<1	<1	<1	<1	<1
	5/24/1994	1	<1	<1	< 1	· <1	< i	<1	<1	<1	<1	<1
	5/24/1994	1	<1	<1	< 1	<1	<1	<1	<1	<1	<1	< 0.5
	7/25/1994	1	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	<1
	9/19/1996	1	222	<1	<1	<1	<1	<1	<1	<1	<1	<1
	8/1/1997	1	34	<1	<1	<1	<1	<1	<1	<1	<1	<1
	8/11/1997	1	17	<1	<1	<1	<1	<1	< 1	<1	<1	<1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
immary of Detected Purgeable Halocarbons (Chlorinated Compounds)
07 Telegraph Drive, Gastonia, North Carolina

Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CAS# 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAS# 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CAS# 71-55-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CAS# 79-01-6 ppb	Tetrachloro- ethene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASI 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAS# 156-60-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb	Chloroform CAS# 67-66-3 ppb 70
		T .									<1	<1
MW-7A	2/8/1994	1	<1	<1	<1	<1	<1	<1	<1 <1	<1 <1	2	<1
	5/24/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	7/26/1994	1	<1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1	<1	<1	<1
	9/17/1996 8/12/1997	1	7.6 3.3	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/10/1998	î	<1	<1	<1	<1	<1	<1	<1	<1	<1	1.1
	9/27/1999	1	49	<1	<1	<1	<1	<1	<1	<1	<1	1.2
	9/11/2000	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<l< td=""></l<>
	8/27/2001	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	8/27/2001	1	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	<1	<1	<1
	10/1/2002	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/29/2003	1	<1	<1	<1	<1	<1	⊲	<1	<1	<1	<1
	10/1/2004	1	7.1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/29/2005	1	24	< 1	<1	< 1	<1	<1	<1	<1	< 1	< 1
	10/20/2006	1	5	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/20/2007	1	2.2	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/22/2008	11	4.9	<1	<1	<1	<1	<1	<1	<1	<1	_<1
	9/15/2009	1	<t< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></t<>	<1	<1	<1	<1	<1	<1	<1	<1	<1
	47/07/00	1	1.4	<1	<1	<1	<1	<1	<1	<1	<1	<1
MW-7B	7/26/1994	1	10	<1	<1	<1	< 1	<1	<1	<1	2	<1
	11/2/1994	1	46	<1	.4	<1	< 1	<1	< 1	< 1	<1	<1
	1/17/1995	1	69	< 1	3	< i	<1	<1	< 1	<1	<1	< 50
	9/17/1996	50	740	< 50	< 50	< 50	< 50	< 50	< 50	< 50	< 50	< 25
	8/12/1997	25	800	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 25	<u> </u>
	9/10/1998	1	340	<1	20	< 1	<1	<1	<1	<1	<1	<10
	9/29/1999	10	200	<10	<10	<10	<10	<10	<10	<10	<10	<1
	9/11/2000	1	9.9	<1	<110	<1	< <u>l</u>	<1	<1	<1	<1	<1
	8/27/2001	1	90	<1	8.5	<1	<1	<1	<1	<1	<1	<1
	10/4/2002	1	400	<1	5,5	<1	<1	<1	<1	<1	<1	<1
	9/25/2003	1	1600	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/1/2004	10	880	<1	<1	<1	<1	<1	, <1	<1	<1	< 10
	9/28/2005	10	890	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	<1
	10/19/2006	10	770	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/24/2007	1/100	2,200	<1	<1	< <u>l</u>	<1	<1	1.2	<1	<1	0,92 J
	9/23/2008	10	960	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/15/2009	10/100	1,400	<10	<10	<10	<10	<10	<10	<10 <10	<10 <10	<10 <10
	1040000	10	740	<10	<10	<10	<10	<10	<10		<1	<1
MW-8	2/8/1994		<1	<1	<1	<1	<1	<1	<1	<1 <1	<1	<1
	5/24/1994	1	<1	<1 <1	<u><1</u>	<1 <1	<1 <1	<1 <1	<1	<1	<1	2
	7/26/1994	1	<1 <1	<1	<1	<1	<1	<1	<1	<1	<1	17
	9/17/1996 8/13/1997	1	<1	<1	<1	<1	<1 <1	<1	<1	. <1	<1	23
	9/10/1998	,	<1	<1	<1	<1	<1	<1	<1	<1	<1	9.4
		1	<1	<1	<1	<1	<1	<1	<1	<1	<1	3.1
	9/20/1999	1	<1	<1	<1	<1	<i>1</i>	<1	<1	<1	<1	1.9
	9/12/2000 8/28/2001	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	NS
	9/29/2003	1	<1	<1	<1	< <u>1</u>	<1	<1	<1	<1	<1	1,9
	10/1/2004	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	1.2
	9/30/2005	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< 2
	10/16/2006	1	6.6	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/20/2007	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/22/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	1.1
	9/16/2009	1	32	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/10/2009	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1

Table 3	
Groundwater Analytical Data	
Monitoring Wells, Extraction Wells, and Inactive Water Supply We	lls
Summary of Detected Purgeable Halocarbons (Chlorinated Compo-	nds)
207 Telegraph Drive, Gastonia, North Carolina	.,

20/ telegraph	Telegraph Drive, Gastonia, North Carolina										1	
Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CASV 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAS# 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CASV 71-55-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CASN 79-01-6 ppb	Tetrachloro- ethene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASN 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAS# 156-60-5 ppb	Carbon Tetrachloride CASS 56-23-5 ppb	Chloroform CAS# 67-66-3 ppb
NC Groundwate	r Quality Standard		7	6	200	0,6	3	0.7	70	100	0.3	70
MW-10A	12/15/1993	2	5	7	29	< 2	120	3	NA	<2	< 2	<1
	2/9/1994	1	2	2	9	< 1	51	< i	15	<1	<1	< 2
	5/24/1994	. 2	. 7	6	25	< 2	120	< 2	35	< 2	< 2	< 2
	7/27/1994	2	4	5	< 2	< 2	85	3	22	< 2	< 2	1
	11/3/1994	1	2	3	11	<1	60	2	13	<1	<1	<1
	1/17/1995	1	1	4	9	<1	61	<1	12	<1	<1	<1
	4/17/1995	1	1.3	3.3	10	< 1	43	2	13	<1	<1	<1
	7/20/1995	1	< 1	2.8	7.8	<1	37	2.4	9,3	<1	<1	< 1
	10/18/1995	1	< 1	1.9	4.3	< 1	25	6.6	4.9	<1	<1	<1
	1/25/1996	1	1.1	3.1	11	<1	49	7	7.9	<1	< 1	< 10
	9/18/1996	10	< 10	< 10	< 10	< 10	23,7	8.6 J	< 10	< 10	< 10	<1
1	3/24/1997	1	<1	2.5	4	< 1	14	19	3.2	<1	<1	<1
	6/26/1997	1	<1	3.2	4,6	<1	20	21	3.9	<1	<1	<1
	9/30/1997	1	₹	1.7	4	<1	15	24	2.4	<1	<1	<1
	11/7/1997	1	<1	1.7	4.3	<1	14	<1	2.4	<1	<1	< 1
MW-10B	9/9/1994	1	<1	<1	<1	< 1	<1	<1	<1	<1	< 1	< 1
	11/4/1994	1	<1	3	<1	<1	2	<1	<1	<1	<1	<1
	1/18/1995	1	< 1	2	1	< 1	2	< 1	<1	<1	<1	< 1
MW-11	2/9/1994	1	<1	<1	3	4	22	<1	12	<1	< 1	<1
[2/9/1994	1	< 1	<1	5	< 1	31	<1	13	<1	<1	<1
	5/25/1994	1	< 1	< 1	6	< 1	21	<1	9	<1	<1	< 1
	7/27/1994	1	<1	<1	< 1	<1	16	<1	. 5	<1	<1	<1
[11/3/1994	1	< 1	< 1	4	<1	14	<1	6	<1	<1	<1
[1/18/1995	1	<1	<1	3	< 1	14	<1	4	<1	<1	4,6
	11/27/1995	1	<1	<1	< 1	<1	16	<1	2.7	<1	<1	<1
	1/25/1996	1	< 1	< 1	1.9	<1	19	< 1	9.3	<1	<1	3,1
[9/18/1996	1	<1	< 1	<1	< 1	7.9	<1	1.7	<1	<1	5.1
	3/24/1997	1	< 1	<1	1.3	<1	8.2	<1	2.2	<1	< 1	3.8
	6/26/1997	1	<1	<1	<1	<1	12	<1	2.4	<1	<1	3.6
	9/30/1997	1	1.1	1.5	4.3	<1	14	<1	2.4	<1	<1	2.6
	11/7/1997	1	<1	<1	<1	<1	7.6	<1	1.8	<1	<1	<1
MW-12	2/10/1994	1	<1	<1	< 1	<1	<1	<1	<1	<1	<1	<1
	5/25/1994	1	<1	<1	3	3	2	<1	< 1	< 1	2	<1
	7/28/1994	_11	2	<1	<1	< i	<1	< l	<1	<1	<1	< 1
	11/4/1994	1	<1	<1	<1	<1	<1	<1	<1 .	<1	<1	<1
	1/18/1995	1	<1	<1	2	<1	<1	<1	<1	<1	<1	<1
L	4/17/1995	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	7/20/1995	1	<1	<1	1.9	<1	1,9	<1	<1	<1	<1	<1
	10/18/1995	1	<1	<1	1,8	<1	<1	<1	<1	<1	<1	< l
	1/25/1996	1	<1	<1	<1	<1	< 1	<1	<1	<1	<1	<1
	9/18/1996	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
L	3/24/1997	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	6/26/1997	1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><l< td=""><td><1</td></l<></td></i<>	<1	<1	<1	<1	<1	<l< td=""><td><1</td></l<>	<1
	9/30/1997	1	<1	<1	<1	<1	<1	1.4	<1	<1	<1	<l< td=""></l<>
	11/7/1997	1	<1	<1	<1	<1	<l< td=""><td>1.3</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	1.3	<1	<1	<1	<1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonis, North Carolina

Sumple Date Date Date Peter Peter Peter Company		Drive, Gastonia, N				1	T	,			 	1	
	Samula	Date of	Dibution	ethene (1,1-DCE)	ethane (1,1-DCA)	ethane (1,1,1-TCA)	propane	ethene (TCE)	ethene (PCE)	ethene (cis-1,2-DCE)	ethene (trans-1,2-DCE)	Tetrachloride	Chloroform
No.													ppb
	NC Groundwat	er Quality Standard		7	6	200	0.6	3	0.7	70	100	0.3	70
	MW-13	2/9/1994	1	<1	29	8	<1	28	2	32	<1	<1	<1
		5/25/1994	1	<1	20	14	<1	37	4	40	<1	<1	<1
			1										<1
													<1
1000 1000													<1
1,000/1995 1												i	<1
1021/1979													<1
10010096 1												<1	<1
Main		1/25/1996	1	1.1	16	3.9	<1	32	1.2	29	1.4	<1	1.6
MANIAN		9/18/1996	_1_	<1	5.6	3,7	<1	31.4	1,6	20	<1	<1	<1
\$\frac{900097}{90009}		3/24/1997	1										<2
11/11/997													<2
92/1998													<2
990/999													1.1
98/2000 1			-										<u> </u>
MW-14 12191999 1													<1
191/2002												1	<1
MW-14 10 C10			1	-	<1	<1		7.6	,	1.4	<1	<1	<10
109/2005 10		9/29/2003	10	<10	<10	<10	<10	12	<10	<10	<10	<10	<10
NW-14 10/17/2006 10 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 < 0 <			10										< 10
971/2007 10 18 < <10 <10 <10 <10 <10 <10 <10 <10 <10 <													<1
9727008 10													
MW-14													
MW-14													<1
29/1994		200001-9-27 4 34-20-0-0-0-0-1-1-1-1-1-10-00-0-0-0-0-0-0-											</td
777/1994	MW-14	12/15/1993	1	<1	1	3	<1	4	<1	<1	<1	<1	< 1
11/1/1994 1		2/9/1994	1	<1	<1	4	3	8	<1	33	<1	<1	<1
1/17/1995		7/27/1994	1	<1	<1	<1	<1		<1				<1
MW-15A 1 1 1 2 2.6 1 3.3 1.2 1													<1
1/20/1995													
11/27/1995 1													
37/1/996		1				1							
9/18/1996 1 <1 <1 <1 <1 <1 <1 <1			$\overline{}$										<1
9/30/1997 1												<1	<1
MW-15A 27/1994 1 3 5 3 3 41 9 41 2 41 41 41 41 41 41		6/26/1997	1	<1	2.6	<1	<1	1	24	<1	<1	<1	<1
MW-15A 27/1994 1 3 5 3 < 1 9 <1 2 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1		9/30/1997	1	<1	<1	T T							<2
5/25/1994 1 7 15 6 <1			$\overline{}$			1							<1
7/2/11994 1 11 29 4 <1	MW-15A												<1
9/18/1996 5 11 24 < 5 < 5 71 < 5 54 < 5 < 5 < 1 8/12/1997 1 7.5 13 < 1						1							
8/12/1997 1 7.5 13 <1							_		_				
9/18/1998 1 6 13 <1													
9/24/1999 1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <									ľ		i i		<1
9/11/2000 1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <													<1
10/1/2002 1 <1							<1		<1	<1	<1	<1	< <u>I</u>
9/29/2003 1 <1		8/28/2001	1	<1	<1	<1	1						<1
10/1/2004 1 <1			1										<1
9/30/2005 1 22 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <										- 1			<1
10/20/2006 1 7.7 3.2 <1 <1 26 <1 50 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1										18			<1
9/21/2007 1 12 10 <1 <1 52 <2 75 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <													< 5
9/22/2008 1 <1													
9/15/2009 1 <1 <1 <1 <1 <1 <1 <1 <1 <1			-										
TO STATE OF THE ST													<1
		# 15/2009	1	<1	<1	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	<1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

ZOT TEIERLAD	a Drive, Gastonia, N	TOTTE CATOL	I BX							<u> </u>		
Sample Location	Date of Sampling	Dilution Factor	I,1-Dichloro- ethene (I,1-DCE) CASN 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAS# 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CAS# 71-55-6 ppb	1,2-Dichloro- propane CASN 78-87-5 ppb	Trichloro- ethene (TCE) CAS# 79-01-6 ppb	Tetrachloro- ethene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CAS# 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CASW 156-60-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb	Chloroform CASN 67-66-3 ppb
NC Groundwat	ter Quality Standard		7	6	200	0.6	3	0.7	70	100	0.3	70
MW-15B	2/7/1994	5	14	35	13	< 5	110	< 5	61	< 5	< 5	< 5
	2/7/1994	5	15	60	< 5	<5	120	<5	62	< 5	1	<1
	5/25/1994	1	4	9	5	<1					< 5	
	5/25/1994	1	3	7	5	1	37	<1	29	<1	<1	<1
	7/27/1994	5			i	<1	29	<1	22	<1	<1	< 5
			25	60	13	< 5	230	< 5	160	< 5	< 5	< 5
	9/18/1996	20	20	30	< 20	< 20	190	< 20	190	< 20	< 20	< 4
	8/12/1997	4	13	31	<4	<4	120	<4	110	<4	<4	<1
	9/18/1998	1	5	20	<1	<1	97	<1	76	<1	<1	<1
	9/24/1999	1	10	16	< <u>l</u>	<1	210	<1	220	<1	<1	2,1
	9/18/2000	1	0.5	16	<1	<1	120	<1	150	8	<i< td=""><td><1</td></i<>	<1
	8/28/2001	1	9,3	11	1.8	<1	68	<1	110	<1	<1	3.6
	10/1/2002	1	7.6	15	1.2	4.8	84	<1	140	<1	<1	<1
	9/29/2003	. 1	11	14	<1	<1	130	<1	240	<1	<1	<1
	9/30/2004	1	11	13	<1	<1	95	<1	200	<1	<1	< 1
	10/18/2005	1	7.1	5,3	<1	<1	37	<1	90	<1	< 1	<1
	10/20/2006	10	28	16	<1	<1	120	<1	240	<1	<1	<1
	9/21/2007	10	23	8,6	<1	<i< td=""><td>85</td><td><1</td><td>110</td><td><1</td><td><1</td><td>3</td></i<>	85	<1	110	<1	<1	3
	9/23/2008	1	19	2.3	<1	<1	26	<1	50	<1	<1	2.6
	9/15/2009	1	12	1,4	<1	<1	9.9	<u> </u>	18	<1	<1	<i< td=""></i<>
	9/28/2010	1	5.4	0.27 J	<1	<1	2,3	<1	4.5	<1	<1	<1
MW-15C	5/25/1994	1	4	7	9	<1	59	<1	27	<1	< 1	<1
	7/27/1994	<u>t</u>	3	. 7	5	<1	53	< 1	29	<1	<1	< 1
	11/2/1994	1	4	6	6	< l	56	<1	26	<1	< 1	< 2
	9/18/1996	2	1.6	1.9	2.1	< 2	25	< 2	8.9	< 2	< 2	<1
	8/12/1997	1	1.8	1.2	<1	< 1	16	<1	5	<1	<1	<1
	9/18/1998	1	< <u>l</u>	<1	<1	<l< td=""><td>11</td><td><1</td><td>_1.</td><td><1</td><td><1</td><td><1</td></l<>	11	<1	_1.	<1	<1	<1
	9/24/1999	1	<1	53	<u> </u>	<1	1.4	<1	<1	<1	<1	1.1
	9/18/2000	1	150	<1	<1	<1	1.9	<1	1.7	<1	<l< td=""><td><10</td></l<>	<10
	8/28/2001	10	180	<10	<10	<10	<10	<10	<10	<10	·<10	<1
	10/3/2002	1	4,100	<1	<1	<1	6,2	<1	2.7	<1	<l< td=""><td><1</td></l<>	<1
	9/26/2003		690	<1	<1	<1	< <u>l</u>	<1	<1	<1	<1	<1
	9/30/2004	1	1,700	<1	<1	<l< td=""><td>1,1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	1,1	<1	<1	<1	<1	<1
	10/3/2005	1/100	1,300	<1	<1	<1	1.5	<1	<1	<1	<1	<1
	10/18/2006	1/100	1,200	<1	<1	<1	4.1	<1	6,1	<1	<1	1.8
	9/24/2007	1/10	290	<1	<1	<1	0.71 J	<1	1.0	<1	<1	2.6
	9/25/2008	1/10	610	<1	<1	<1	1.4	4.5	0.98 J	<1	<1	< <u>1</u>
	9/15/2009 9/28/2010	1/10	210	<1	<1	<1	<1	<1	<1	<1	<l< td=""><td><1</td></l<>	<1
MOVIED		1/10	150	<1	<1	<1	<1	<1	<1	<1	<1	<1
MW-15D	11/4/1994 11/4/1994	1	61	<1	<1	<1	<1	<1	<1	<1	<u> </u>	<1
	1/17/1995	1	62	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/18/1996	2	36 16	<1	<1 <2	<1	<1	<1	<1	<1	<1	< 2
ŀ	8/12/1997	1	13	<1	<1	<1	<1	<1	<2 <1	< 2 < 1	< 2	<1
ŀ	9/22/1998	1	12	<1	<1	<1	<1	<1	<1	<1	<1 <1	<1
	9/24/1999	1	4.6	<1	<1					. 1	. "	<1
İ	9/18/2000	1	3	<1	<1	<1	<1	<1	<1	<1 <1	<1 <1	<1 <1
ļ	8/28/2001	1	5.6	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/3/2002	1	6.5	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/26/2003	1	7.2	<1	<1	<1	<1	<1	<1	<1	<1	<1 <1
ļ	9/30/2004	1	4.2	<1	<1	<1	<1	<1	<1 <1	<1	<1 <1	<1
	10/3/2005	1	3,0	<1	<1	<1	<1	<1	<1	<1	<1	< 500
ľ	10/20/2006	i	3.4	<1	<1	<1 <1	<1	<1 <1	<1 <1	<1	<1	< 500 <1
Ì	9/25/2007	i	<1	<1	<1	<1	<1	<1	<1 <1	<1	<1	<1
1	9/26/2008	1	2.6	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	<1	<1	<1	<1
ļ	9/18/2009	1	1.5	<1	<1	<1	<1	<1	<1	<1	<1	_ <1
Ī	9/10/2009 (4/29/20/20/00/20	1	2.5	<1	<1	<1	<1	<1	<1	<1	<1	<1
	CONTRACTOR OF THE PARTY OF THE	•	4.0	`*	~1	`1	`1	-1	~1	~1		

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

	1											
	1		1,1-Dichloro-	1,1-Dichloro-	1,1,1-Trichloro-	La Dieble	Trichloro-	Tetrachloro-	cis-1,2-Dichloro- ethene	trans-1,2-Dichloro- ethene	Carbon]
]		ethene (1,1-DCE)	ethane (1,1-DCA)	ethane (1,1,1-TCA)	1,2-Dichloro- propane	ethene (TCE)	ethene (PCE)	(cis-1,2-DCE)	(trans-1,2-DCE)	Tetrachloride	Chlorofor
Sample	Date of	Dilution	CAS# 75-35-4	CAS# 75-34-3	CAS# 71-55-6	CAS# 78-87-5	CAS# 79-01-6	CAS# 127-18-4	CAS# 156-59-2	CAS# 156-60-5	CAS# 56-23-5	CASE 67-66
Location	Sampling	Factor	ppb	ppb	ppb	ppb	ppb	ppb	ppb	ppb	ppb	ppb
C Groundwate	er Quality Standard	T T	77	6	200	0.6	3	0.7	70	100	0,3	70
MW-16A	5/25/1994	500	18,000	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 1,000
	7/27/1994	1,000	14,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 100
	7/27/1994	100	7,300	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 250
	11/1/1994	250	13,000	< 250	< 250	< 250	< 250	< 250	< 250	< 250	< 250	< 100
	9/19/1996	100	132,000	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 500
	11/12/1996	500	10,000	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500
	3/20/1997	500	18,000	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500
	8/11/1997	500	18,000	< 500	< 500	< 500	< 500	< 500	< 500	< 500	< 500	<1
	9/2/1998	1	20,000	<1	< 1	<1	2	<1	< 1	<1	<1	<1
	9/27/1999	1	16,000	<1	<1	<1	1.2	<1	<1	<1	<1	<100
	9/12/2000	100	6,500	<100	<100	<100	<100	<100	<100	<100	<100	<100
	8/29/2001	100	15,000	<100	<100	<100	<100	<100	<100	<100	<100	<100
	10/2/2002	100	1,400	<100	<100	<100	<100	<100	<100	<100	<100	4.4
	9/26/2003	1	9,200	4	<1	<1	18	<1	11	<1	<100	<10
	10/4/2004	10	1,900	<10	<10	<10	46	<10	33	<10	<10	< 10
	9/28/2005	10/100	1,500	< 10	< 10	< 10	23	< 10	15	< 10	< 10	<1
	10/20/2006	10	780	<10	<10	<10	29	<10	22	<10	<10	<10
	9/25/2007	10/100	1,200	<10	<10	<10	29	<10	26	<10	<10	<10
	9/24/2008	10/100	1,600	<10	<10	<10	22	<10	24	<10	<10	<10
	9/14/2009	10/100	1,200	<10	<10	<10	15	<10	18	<10	<10	<10
	997700 O	10	830	<10	<10	<10	4.5 J	<10	9,7 J	<10	<10	<10
MW-16B	5/25/1994	, i	7	<1	<1	<1	<1	<1	<1	<1	< 1	<1
	7/27/1994	i	10	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/31/1994	- 	62	<1	<1	<1	<1	<1	<1	<1	<1	<1
	1/18/1995	î	99	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/19/1996	-,	1,380	<1	<1	<1	<1	<1	<1	<1	<1	< 80
	8/11/1997	80		< 80	< 80	< 80	< 80	< 80	< 80	< 80	< 80	<2
			2,300		<2	<2	<2	<2	<2	<2	<2	<1
	9/14/1998	- 2	3,500	<2			< <u>1</u>	<1	<1	<1	<1	<100
	9/27/1999	1 100	1,200	<1	<1	<100		<100	<100	<100	<100	<40
	9/12/2000	100	1,200	<100	<100	<100	<100 <40	- 1	<40	<40	<40	<100
	8/30/2001	40	1,700	<40	<40	<40		<40	1	- 1	<100	<100
	10/4/2002	100	2,200	<100	<100	<100	<100	<100	<100	<100		<10
	9/26/2003	10	1,900	<10	<10	<10	<10	<10	<10	<10	<10	
	10/1/2004	10	880	<1010	<10	<10	<10	<10	<10	<10	<10	< 10
	9/29/2005	10	510	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	<1
	10/20/2006	10	990	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/25/2007	10	880	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/24/2008	10	890	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/14/2009	10	550	<10	<10	<10	<10	<10	<10	<10	<10	<10
	21-31r/000	10	480	<10	<10	<10	<10	<10	<10	<10	<10	<10
MW-17A	2/7/1994		72	2	4	<1	4	<1	<1	<1	<1	< 5
	5/26/1994	5	230	< 5	< 5	< 5	< 5	< 5	< 5	< 5	< 5	<1
	7/27/1994	1	98	<1	4	<1	4	<1	<1	< 1	<1	<1
	9/19/1996	1	1,170	2	2,1	<1	4	<1	<1	<1	<1	<1
	8/12/1997	1	4	20	4.2	<1	8,3	<1	14	<1	<1	2
	9/18/1998	11	9	<1	<1	<1	6	<1	4	<1	<1	2.6
	9/29/1999	1	11	6.8	2.9	<1	14	<1	11	<1	<1	2.8
	9/18/2000	1	25	14	<1	<1	43	<l< td=""><td>30</td><td>7.1</td><td><1</td><td><<u>l</u></td></l<>	30	7.1	<1	< <u>l</u>
[8/29/2001	1	36	32	6.1	<1	100	<1	81	<1	<l< td=""><td>2.4</td></l<>	2.4
[10/3/2002	1	27	13	<1	<1	36	<1	24	<1	<1	<1
Ī	9/26/2003	1	7	6.8	<1	<1	26	<1	24	<1	<1	<1
	9/30/2004	1	<1	2.4	<1	<1	10	<1	12	<1	<1	1.5
1	9/28/2005	1	7,9	4.0	<1	<1	25	<1	38	<1	<1	< 100
ļ	10/20/2006	1	0.98	<1	<1	<1	5	<1	6	<1	<1	<1
ļ	9/24/2007	1	0.5	<1	<1	<1	<1	<1	<1	<1	<1	<1
ŀ	9/23/2008	1	19	18	<1	<1	150	<1	270	<1	<1	<1
ł	9/23/2008	1	4.1	2,9	<1	<1	35	<1	59	<1	<1	<1
1	9/1//2009	1	12	5.6	<1	<1	97	· <1	190	3.5	<1	<1

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Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells Summary of Detected Pargeable Halocarbons (Chlorinated Compound ected Purgeable Halocarbons (Chlorinated Compounds) 207 Telegraph Drive, Gastonia, North Carolina 1,1-Dichloro 1,1-Dichloro 1,1,1-Trichloro Trichloro-Tetrachloro cis-1,2-Dichloro trans-1,2-Dichloro ethene ethane ethane 1.2-Dichloro-Carbon Tetrachlorid (1,1-DCE) (1,1-DCA) (1,1,1-TCA) (PCE) (cis-1,2-DCE) (trans-1,2-DCE) propane (TCE) Dilution Sample Date of CAS# 75-35-4 CAS# 75-34-3 CAS# 71-55-6 CAS# 78-87-5 CASN 79-01-CAS# 127-18-CASN 156-59-2 CAS# 156-60-5 CAS# 56-23-5 CAS# 67-66-3 ppb C Groundwater Quality Standard 200 0.6 0.7 70 100 70 MW-17B 2/7/1994 100 2,300 < 100 < 100 < 100 < 100 < 100 < 100 < 100 < 100 < 100 5/26/1994 100 7.000 < 100 < 100 < 100 < 100 < 100 < 100 < 100 < 100 < 100 5/26/1994 200 4,700 < 100 70 J < 200 58 J 48 J < 100 < 100 < 100 < 100 7/27/1994 100 6,400 < 100 < 100 < 100 < 100 < 100 < 100 < 100 < 100 < 500 9/19/1996 500 161,000 < 500 < 500 < 500 < 500 < 500 < 500 < 500 < 500 < 500 9/19/1996 500 < 500 < 500 104,000 < 500 < 500 < 500 < 500 < 500 < 500 < 2,000 11/12/1996 2,000 53,000 < 2,000 < 2,000 < 2,000 < 2.000 < 2.000 < 2.000 < 2,000 < 2,000 < 1.300 3/20/1997 1,250 60,000 < 1,300 < 1,300 < 1,300 < 1,300 < 1,300 < 1,300 < 1,300 < 1,300 < 2,000 8/12/1997 2,000 70,000 < 2,000 < 2,000 < 2,000 <2,500 < 2,000 < 2,000 < 2,000 < 2,000 < 2,000 3/27/1998 2,500 43,000 <2,500 <2,500 <25,00 <2,500 <2,500 <2,500 <2,500 <2.500 < 100 9/18/1998 100 66,000 < 100 < 100 < 100 < 100 < 100 < 100 < 100 < 100 <400 3/24/1999 400 21,000 <400 <400 <400 <400 <400 <400 <400 <400 <1 9/16/1999 43,000 24 13 16 160 <1 96 <1 <1 <1000 24,000 3/23/2000 1,000 <1000 <1000 <1000 <1000 <1000 <1000 <1000 <1000 <1000 9/11/2000 1,000 23,000 <1000 <1000 <1000 <1000 <1000 <1000 <1000 <1000 <400 2/28/2001 400 35,000 <400 <400 <400 <400 <400 <400 <400 <400 <1000 8/30/2001 1,000 34,000 <1000 <1000 <1000 <1000 <1000 <1000 <1000 <1000 <100 3/28/2002 100 8,200 <100 <100 <100 110 <100 <100 <100 <100 <100 9/30/2002 100 18,000 <100 <100 <100 380 <100 410 <100 <100 <100 3/25/2003 100 7,700 <100 <100 <100 280 <100 240 <100 <100 <100 9/25/2003 100 12,000 <100 <100 <100 290 <100 350 <100 <100 <100 9/29/2004 100 3,300 <100 <100 <100 <100 380 <100 <100 <10 1/14/2005 100 8,000 <100 <100 <100 360 <100 420 <100 <100 <100 6/22/2005 10/100 7,700 16 <10 <10 200 <10 270 <10 <10 <1 9/28/2005 100 2,300 < 100 < 100 < 100 290 < 100 410 < 100 < 100 < 1 100 4,100 3/10/2006 <100 <100 <100 340 <100 450 <100 <100 <100 10/18/2006 100 2,500 <100 <100 <100 280 <100 370 <100 <100 <100 3/8/2007 100 7,700 < 100 < 100 < 100 190 < 100 180 < 100 < 100 < 100 9/21/2007 100 10,000 <100 <100 <100 <100 <100 110 <100 <100 <100 3/11/2008 10/100 14 < 10 < 10 < 10 170 < 10 300 < 10 < 10 < 10 9/23/2008 10/100 4,100 24 <10 <10 370 210 <10 <10 <10 <10 5/1/2009 10/100 2,400 12 <10 <10 110 <10 270 <10 <10 <10 9/15/2009 10/100 13 2,900 <10 <10 140 <10 240 <10 <10 <10 9/24/2010 10/100 <10 3,300 <10 <10 110 <10 180 <10 <10 <10 +973070010 ± 10/100 3,800 <10 <10 <10 83 <10 170 <10 <10 <10 MW-17C 5/26/1994 1 50 <1 < 1 <1 <1 < 1 <1 < 1 < 1 < 10 7/27/1994 36 < 1 < 1 < 1 < 1 < 1 < 1 < I < 1 < 25 11/2/1994 10 250 < 10 < 10 < 10 < 10 < 10 < 10 < 10 < 10 < 1 1/17/1995 25 800 < 25 < 25 < 25 < 25 < 25 < 25 < 25 < 25 < 50 9/19/1996 2,650 < 1 < 1 < 1 <1 <1 < 1 1.8 <1 <50 50 1,600 < 50 < 50 < 50 < 50 < 50 8/12/1997 < 50 < 50 < 50 < 1 3/27/1998 50 700 <50 <50 <50 <50 <50 <50 <50 <50 <! 9/14/1998 820 < 1 < 1 < 1 < 1 < i <1 < 1 < 1 <1 3/24/1999 77 <1 <! <1 <1 <1 <1 <40 <1 <1 9/16/1999 170 <1 <1 <1 <1 <! <1 <1 <! <10 3/23/2000 40 <40 <40 <40 <40 <40 <40 <40 <40 160 <1 10 <10 9/11/2000 270 <10 <10 <10 <10 <10 <10 <10 <1 2/28/2001 180 <1 <1 <1 <1 <1 <1 <1 <1 <1 8/30/2001 <1 <1 <1 <1 <1 <1 <1 <1 160 <1 3/28/2002 190 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/30/2002 320 <1 <1 <1 <1 <1 <1 <1 <1 <1 140 <1 <1 3/25/2003 1 <1 <1 <! <1 <1 <1 <1 9/25/2003 280 <1 <1 <1 <1 <1 <1 <1 <1 <10 9/29/2004 290 <1 <1 <1 <1 <1 < 1 1/14/2005 320 <t <1 <1 <1 <1 <1 <1 <1 <1 6/22/2005 10 140 <10 <10 <10 <10 <10 <10 <10 <10 <1 9/28/2005 1/10 280 < 10 < 1 < 1 < 1 < 1 < 1 <1 3/10/2006 1/10 210 <1 <1 <1 <i <1 <1 **<**1 <1 <1 10/18/2006 1/10 230 <1 <1 <1 <1 <1 <1 <1 <1 <1 3/8/2007 1/10 130 <1 < 1 < 1 < 1 < 1 < 1 < 1 <1 < 1 9/21/2007 1/10 190 <1 <1 <1 <1 <1 <1 <1 <1 <1 3/12/2008 1/10 150 <I <1 <1 <1 <1 <1 <1 <1 <1 1/10 130 9/22/2008 <1 <1 <1 <1 <1 <1 <1 <1 <1 4/30/2009 1/10 120 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/16/2009 1/10 <1 <1 <1 110 <1 <1 <1 <i <1 <1 1/10 100 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 ۲۱ <1 <1 **~1**

Table 3

roundwater Analytical Data

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

	Drive, Gastonia, N	The Caron	0.4		**:	1	T		I	T**	r	
Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CAS# 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAS# 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CASF 71-55-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CASW 79-01-6 ppb	Tetrachloro- ethene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASN 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CASW 156-60-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb	Chloroform CAS# 67-66-3 ppb
NC Groundwater	Quality Standard		7	6	200	0.6	3	0.7	70	100	0.3	70
MW-18A	2/9/1994	1	<1	<1	3	<1	<1	<1	<1	<1	<1	<1
	5/26/1994	1	< 1	<1	<1	<1	< 1	<1	<1	<1	<1	<1
	7/28/1994	1	3	< 1	< 1	<1	<1	<1	<1	<1	<1	<1
	9/17/1996	. 1	<1	<1	<1	< 1	<1	<1	<1	<1	<u><1</u>	<1
	8/13/1997	1	< 1	< 1	<1	<1	<1	<1	<1	<1	<1	<1
	9/5/1998	1	<1	< 1	<1	< 1	<1	<1	<1	<1	<1	<1
	9/28/1999	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/12/2000	_1	<1	<1	< <u>l</u>	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1
	8/29/2001	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
[9/30/2002	11	<1	<1	≤1	<i< td=""><td><1</td><td><1</td><td><1</td><td><l< td=""><td><1</td><td><1</td></l<></td></i<>	<1	<1	<1	<l< td=""><td><1</td><td><1</td></l<>	<1	<1
	9/29/2003	1	<1	<1	<]	<1	<1	<1	<1	<1	<1	<1
	9/28/2004	11	<1	<1	<1	<1	<1	<1	<1	<u> </u>	<1	<1
[9/27/2005	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/19/2006	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/26/2007	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/25/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
L	9/17/2009	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	(9/29/2010	1	<1	<1	<1	<1	<1	<1	<1	< <u>l</u>	<1	<1
MW-18B	5/26/1994	1	<1	<1	< 1	<1	<1	<1	<1	<1	< 1	<1
L	7/28/1994	1	3	<1	< 1	<1	<1	<1	<1	<1	<1	<1
Į.	11/4/1994		<1	<1	3	<1	<1	<1	<1	<1	<1	<1
1	9/17/1996	1	9.4	< 1	< 1	<1	< 1	<1	<1	<1	<1	<1
1	8/13/1997	1	<1	<1	<1	<1	<1	< 1	<1	<1	<1	<1
Į.	9/5/1998	1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><l< td=""></l<></td></l<>	<1	<1	<1	<1	<1	<1	<1	<1	<l< td=""></l<>
Ļ	9/28/1999	1	<1	<1	<1	<1	<1	< <u>l</u>	<1	<1	<1	<1
Ĺ	9/12/2000	1	<1	<l< td=""><td><1</td><td><1</td><td>·<1</td><td><1</td><td><1</td><td><1</td><td><1</td><td>1.3</td></l<>	<1	<1	·<1	<1	<1	<1	<1	1.3
Ļ	8/29/2001	1	16	<1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1
Ļ	10/4/2002	_11	4	<1	<1	<1	<1	<1	<1	<1	<1	<1
L	9/25/2003	1	<1	<1	<1	< <u>l</u>	< <u>i</u>	<1	<1	<1	<1	<1
ļ.	9/28/2004	1	<1	<1	<1	<1	<1	<1	<1	<u> </u>	<1	< 2
	9/26/2005		11	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/19/2006	1	0.82	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1	<1
	9/26/2007		<1	<1	<1	<1	<1	< <u>l</u>	<1	<1	<1	<1
ļ	9/25/2008	1	<u> </u>	<1	<1	<1	<1	<1	<1	<1	<1 <1	<1 <1
	200000	1	<1	<1	<1	<1	<1	<1	<1		< 2	2
MW-19A	2/7/1994	2	13	7	20	<2	90 72	16 18	31 27	< 2 < 1	<1	<10
-	5/25/1994	1	. 8	6	18 11	<1 <1	62	19	<1	<1	<1	<10
-	7/26/1994	1	6	4	<10	<10	<10	<10	<10	<10	<10	<10
	9/14/1998	10	<10	<10 <10	<10	<10	14	11	<10	<10	<10	<10
-	9/20/1999 9/8/2000	10	32 30	<10	<10	<10	<10	<10	<10	<10	<10	<1
<u> </u>	8/28/2000 8/28/2001	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<1
<u></u>	10/1/2002	10	1	<1	<1	<1	1	<1	<1	<1	<1	<1
ŀ	9/29/2003	1	15	<1	<1	<1	<1	<1	<1	<1	<1	<1
F	10/1/2004	1	11	<1	<1	<1	<1	<1	1.4	<1	<1	<1
ŀ	10/3/2005		11	<1	<1	<1	<1	<1	1,2	<1	<1	<1
 	10/3/2005	$\overrightarrow{}$		<1	<1	<1	<1	<1	<1	<1	<1	<1
H	9/26/2007	+	6.3	<1	<1	<1	2	<1	<1	<1	<1	<i< td=""></i<>
ŀ	1		1.9	- <u><1</u>	<1	<1	<1	<1	<1	<1	<1	1
}	9/22/2008	+ +	4.2	<1	<1 <1	<1	<1	<1	<1	<1	<1	<1
Į.	9/17/2009	1	3.1	<1	<1	<1	<1	<1	<1	<1	<1	1.5

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

C Groundwater MW-19B	r Quality Standard 2/8/1994 5/25/1994 5/25/1994 7/27/1994 3/26/1998 9/14/1998 3/24/1999 9/20/1999 3/24/2000 9/8/2000 3/1/2001 3/3/8/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005 9/3/10066		7	6 2 3 3 3 4 4 4 4 4 4 4	200 2 5 <1 <1 <1 <1 <1 <1 <1 <1 <1	0.6 2	3 4 2 3 <1 <1 <1 <1 <1 <1 <1 <1	0.7 2 3 2 < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < < <	3 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <	100 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	0.3	70
MW-19B	5/25/1994 7/27/1994 3/26/1998 9/14/1998 3/24/1999 9/20/1999 3/24/2000 9/8/2000 3/1/2001 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2003 9/25/2003 9/25/2003 9/25/2003		3 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	3 3 3 <1 2 1.1 1.1 <1 <1 <1 <1	5 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	4 4 4 4 4 4 4 4	2 3 <1 <1 <1 1.1 <1	3 2 <1 <1 <1 <1	<1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <	<1 <1 <1 <1 <1	<1 <1 <1 <1 <1 <1	<1 <1 <1 <1 <1 <1 <1 <1 <1
	7/27/1994 3/26/1998 9/14/1998 3/24/1999 9/20/1999 9/20/1999 3/24/2000 9/8/2000 3/1/2001 3/12/2001 3/12/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005		<1 4 13 4	3	<1 <1 <1 <1 <1 <1 <1 <1 <1 <1	্বা ব্ ব্ ব্ ব্ ব্ ব্	3 <1 <1 <1 <1 1.1 <1	2 <1 <1 <1 <1 <1 <1	<1 <1 <1 <1 <1	<1 <1 <1 <1	<1 <1 <1 <1 <1	<1 <1 <1 <1
	3/26/1998 9/14/1998 3/24/1999 9/20/1999 3/24/2000 3/1/2001 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005	1 1 1 1 1 1 1 1 1 1 1 1	4 13 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	< < < < < < < <	्य ्य ्य ्य ्य ्य ्य ्य ्य ्य ्य ्य ्य	ব ব ব ব ব ব	<1 <1 <1 1.1 <1	<1 <1 <1 <1	<1 <1 <1 <1	<1 <1 <1	<1 <1 <1 <1	<1 <1 <1
	9/14/1998 3/24/1999 9/20/1999 3/24/2000 9/8/2000 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005	1 1 1 1 1 1 1 1 1 1 1	13 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	\(\) 2 1.1 1.1 \(\) < < < < < < < < <	<1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <	্ব ব্য ব্য ব্য ব্য	<1 <1 1.1 <1	<1 <1 <1	<1 <1 <1	<1 <1	<1 <1 <1	<1 <1
	3/24/1999 9/20/1999 3/24/2000 9/8/2000 3/1/2001 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005	1 1 1 1 1 1 1 1 1 1	4 4 4 4 4 4 4 4 4	2 1.1 1.1 <1 <1 <1 <1 <1	<1 <1 <1 <1 <1 <1	<1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <	<1 1.1 <1	<1 <1	<1	<1 <1	<1 <1	<1
	9/20/1999 3/24/2000 9/8/2000 3/1/2001 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005	1 1 1 1 1 1 1 1 1	4 4 4 4 4 4 4	1.1 1.1 <1 <1 <1 <1	<1 <1 <1 <1 <1	্ব ব ব ব	1,1	<1	<1	<1	<1	
	3/24/2000 9/8/2000 3/1/2001 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005	1 1 1 1 1 1 1 1	रा रा रा रा रा रा रा रा	1.1 <1 <1 <1 <1 <1	<1 <1 <1 <1	<1 <1 <1	<1					
	9/8/2000 3/1/2001 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/25/2004 1/14/2005 6/23/2005	1 1 1 1 1 1 1	ব ব ব ব ব ব ব	<1 <1 <1 <1 <1	<1 <1 <1	<1 <1		<1	 	<1		
	3/1/2001 8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/29/2004 1/14/2005 6/23/2005	1 1 1 1 1 1	ব ব ব ব ব	্ব ব ব ব	<1 <1	< <u>1</u>	<1				<1	<u><1</u>
-	8/31/2001 3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/29/2004 1/14/2005 6/23/2005 9/28/2005	1 1 1 1 1 1 1 1 1 1	<1 <1 <1 <1 <1	<1 <1	<1			<1	<1	<1	<1	<1
	3/28/2002 10/1/2002 3/26/2003 9/25/2003 9/29/2004 1/14/2005 6/23/2005 9/28/2005	1 1 1 1 1	ব ব ব	<1 <1			<1	<1	<1	<1	<1	<1
-	10/1/2002 3/26/2003 9/25/2003 9/29/2004 1/14/2005 6/23/2005 9/28/2005	1 1 1 1	<1 <1 <1	<1	٧,	<1	<1	<1	<1	<1	<1	<1
-	3/26/2003 9/25/2003 9/29/2004 1/14/2005 6/23/2005 9/28/2005	1 1 1	<1 <1		_,	<1	<1	<1	<1	<1	<1	<1
-	9/25/2003 9/29/2004 1/14/2005 6/23/2005 9/28/2005	1 1	<1		<1	<1	<1	<1	<1	<1	<1	<1
-	9/29/2004 1/14/2005 6/23/2005 9/28/2005	1		<1	<1 <1	<1 <1	<1 <1	<1 <1	<1	<1	< <u>1</u> <1	<1 <1
- - - - -	1/14/2005 6/23/2005 9/28/2005	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
- - - -	6/23/2005 9/28/2005		<1	<1	<1	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1
 - -	9/28/2005	1 1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
 - -		1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
		1	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><t< td=""><td><1</td><td><1</td><td><1</td></t<></td></i<>	<1	<1	<t< td=""><td><1</td><td><1</td><td><1</td></t<>	<1	<1	<1
	10/17/2006	1	<l< td=""><td><1</td><td><1</td><td><l< td=""><td>0.94</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<></td></l<>	<1	<1	<l< td=""><td>0.94</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	0.94	<1	<1	<1	<1	<1
-	3/8/2007	1	<1	<1	<1	<1	1.4	<1	<1	<1	<1	<1
1	9/21/2007	1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1
Γ	3/12/2008	. 1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/23/2008	1	<1	<1	<1	<1	1.4	<1	<1	<1	<1	<1
	4/30/2009	1	<1	<1	<1	<1	0.81 J	<1	<1	<1	<1	<l< td=""></l<>
L	9/15/2009	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	5/5/24/2010	1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1	<1	<1	<1
	1:9/30/2010	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
MW-20A	5/24/1994	1	<1	<1	<1	< 1	<1	<1	<1	<1	3	< 1
L	7/26/1994	1	< 1	<1	< 1	<1	<1	<1	<1	<1	<1	<1
L	11/1/1994	1	2	<1	<1	<1	<1	<1	<1	<1	<1	<1
F	9/17/1996	11	7	<1	< 1	< 1	<1	<1	<1	<1	<1	<1
-	8/12/1997	1	<1	<1	<1	< 1	<1	<1	<1	<1	<1	<1
-	3/26/1998	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/10/1998	1	<1	<1	<1	<1	< <u>l</u>	<1	<1	<1	<1	<1
-	3/22/1999	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
-	9/15/1999	-!-	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
-	3/24/2000	1	<1	<1	<1	<1	<1	<1	<1	<1<1	<1	<1
	9/11/2000	1	<1	<1	<1	<1	<1 <1	< <u>1</u>	<1	<1	<1	<1
	3/1/2001 8/30/2001	1	<1 <1	<1	<1 <1	< <u>1</u>	<1	<1 <1	<1	< <u> </u>	<1 <1	<1 <1
	3/29/2002	1	<1	<1	<1	<1	< <u>1</u>	<1	<1	<1	<1	<1
	9/30/2002	1	<1	<1	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	<1
	3/25/2003	1	<1	<1	<i>i</i>	<1	<1	<1	<1	<1	<1	<i< td=""></i<>
r	9/25/2003	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/30/2004	1	<1	<1	<1	- <1	<1	<1	<1	<1	<1	<1
	1/17/2005	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	5/27/2005	1	<1	<1	<1	<1	<1	<1	<1	<1	<l< td=""><td><1</td></l<>	<1
	9/27/2005	.1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/10/2006	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/19/2006	1	24	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><!--</td--><td><1</td></td></l<>	<1	<1	</td <td><1</td>	<1
	3/7/2007	1	6.5	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/21/2007	1	<1	< <u>l</u>	<1	<1	<1	<1	<1	<1	<1	< <u> </u>
	3/11/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
L	9/22/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
L	4/30/2009	1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><l< td=""><td><1<1</td></l<></td></i<>	<1	<1	<1	<1	<1	<l< td=""><td><1<1</td></l<>	<1<1
<u> </u>	9/15/2009	1	3,3	<1	<1	<1	<1	<1	<1	<1	<1	< <u>I</u>
	popular.	1	1.7	<1	<1	<1	<1	<1	</td <td><1</td> <td><1</td> <td><1</td>	<1	<1	<1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Sammary of Detected Pargeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CASN 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAS# 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CASN 71-35-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CASW 79-01-6 ppb	Tetrachloro- ethene (PCE) CASN 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASN 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAS# 156-60-5 ppb	Carbon Tetrachloride CASY 56-23-5 ppb 0.3	Chloroform CASN 67-66-3 ppb
	r Quality Standard		7	6	200	0.6	3	0.7	70			
MW-20B	5/24/1994	1	92	<1	<1	<1	<1	<1	<1	<1	3	< 5
	7/26/1994	1	10	<1	<1	<1	<1	<1	<1	1>	2	< 20
	11/1/1994	5	200	< 5	< 5	< 5	< 5	< 5	< 5	< 5	< 5	<1
	9/17/1996	20	190	< 20	< 20	< 20	< 20	28	< 20	< 20	< 20	<10
	3/26/1997	2	64	<1	<1	< 1	<1	<1	<1	<1	<1	<25
	5/29/1997	10	270	<10	<10	<u> </u>	<10	<10	<10	<1	<1	< 25
	7/7/1997	25	520	<25	<25	<1	<25	<25	<25	<1	<1	<50
	8/12/1997	25	620	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 25	<1
	11/13/1997	50	530	<50	<50	<1	<50	<50	<50	<1	<1	<1
	9/10/1998	l	310	<1	<1	<1	<1	< <u>l</u>	<1	<1	<1	<10
	9/28/1999	1	560	<u> <1</u>	<1	<1	<l< td=""><td><l< td=""><td><1</td><td><1</td><td><1</td><td><1</td></l<></td></l<>	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1
	9/19/2000	10	420	<10	<10	<10	<10	<10	<10	<10	<10	<100
	8/29/2001	1	470	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><10</td></l<>	<1	<1	<1	<1	<10
	10/4/2002	100	1,700	<100	<100	<100	<100	<100	<100	<100	<100	<10
	9/25/2003	10	1,000	<10	<10	<10	<10	<10	<10	<10	<10	< 10
	10/4/2004	10	1,500	<10	<10	<10	<10	<10	<10	<10	<10	<1
	9/28/2005	10	1,500	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 1
,	10/19/2006	10/100	11,000	<10	<10	<10	29	<10	38	<10	<10	<10
	9/25/2007	10	710	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/24/2008	10	650	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/16/2009	10/100	1,400	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9002000	10	460	<10	<10	<10	<10	<10	<10	<10	<10	<10
MW-21	2/9/1994	1	<1	< 1	5	<1	<1	<1	< l	<1	<1	<1
]	5/26/1994	1	< 1	< 1	5	< 1	<1	<u> </u>	<1	<1	<1	<1
ļ	7/28/1994	1	44	<1	4	< 1	<1	<1	<1	<1	<1	<1
	9/16/1996	_1	1.1	<1	<1	< l	<1	<1	<1	<1	<1	<1
,	8/13/1997	1	11	< 1	<1	<1	1.2	<1	<1	<1	<1	<1
	9/5/1998		<1	<1	< 1	<1	< 1	< 1	<1	<1	<1	<1
	9/29/1999	1	<1	<1	<1	<1	< <u>i</u>	<1	<1	<1	<1	<1
1	9/13/2000	1	<1	<1	<1	<1	</td <td><1</td> <td><1</td> <td><1</td> <td><l< td=""><td><1</td></l<></td>	<1	<1	<1	<l< td=""><td><1</td></l<>	<1
ļ	8/29/2001	1	<1	<1	<i< td=""><td><<u>l</u></td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i<>	< <u>l</u>	<1	<1	<1	<1	<1	<1
	10/3/2002	1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1	<1	<1
	9/24/2003	1	6.2	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1	<1
	9/28/2004	1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><<u>l</u></td><td><1</td></l<>	<1	<1	<1	<1	<1	<1	< <u>l</u>	<1
,	9/27/2005	1	<1	<1	<1	< 1	<1	<1	<1	<1	<1	<1
,	10/19/2006		<1	<1	<1	<1	<1	<1	<1	< <u>l</u>	<1	<u><1</u>
ļ	9/26/2007	1	<1	<1	< <u>l</u>	<1	0.75 J	<1	<1	<l< td=""><td><1</td><td><1</td></l<>	<1	<1
	9/26/2008	1	<1	<1	<1	<l< td=""><td><1</td><td>1.7</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	1.7	<1	<1	<1	<1
	9/17/2009	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<u> <1</u>
	\$ 950,000 E	1	<1	<1	<l< td=""><td><1</td><td><1</td><td>1.8</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	1.8	<1	<1	<1	<1
MW-22A	5/26/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
ļ.	7/28/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
- 1	11/3/1994	1	<1	<1	6	<1	<1	<1	<1	<1	<1	<1
}	9/16/1996		5.7	<1	4	<1	<1	<1	<1	<1	<1	<1
	8/13/1997	1	8,6	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/14/1998	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/29/1999	1	4.8	<u><1</u>	<1	<u><1</u>	<1	<1	<u> </u>	<1	<1	<1
	9/13/2000	1	<1	<1	<1	<1	<1	< <u>l</u>	<1	<1	<u> </u>	<1
-	8/29/2001	1	<1	<1	<1	<1	<u><1</u>	<1	<u><1</u>	<1	<1	<1
}	10/4/2002	1	5.7	<1	<1	<1	<1	<1	<1	<1	<1	<1
ļ.	9/24/2003	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
ļ	9/30/2004	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< 25
L	9/26/2005	1	<1	<1	<1	<1	<1	< <u>l</u>	<1	<1	<1	< 25
Ĺ	10/19/2006	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
Į.	9/26/2007	1	<1	<1	<1	<1	<1	<i< td=""><td>⊴</td><td><1</td><td><1</td><td><1</td></i<>	⊴	<1	<1	<1
L	9/25/2008	1	5.5	<1	<1	<1	<1	<1	<1	<1	<1	<u><1</u>
Ĺ	9/17/2009	1	<1	<1	<1	<1	<1	<1	<1	<1	< <u>1</u>	<1
8	9/00/5100	1	< 1	<1	<1	<1	<1	<1	<1	<1	<1	<1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

	Drive, Gastonia, N	T Caron	102	,	<u> </u>							· ·
Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CAS# 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAS# 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CAS# 71-55-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CASN 79-01-6 ppb	Tetrachloru- ethene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASW 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAS# 156-60-5 ppb	Carbon Tetrachloride CASI 56-23-5 ppb	Chloroform CAS# 67-66-3 ppb
NC Groundwate	r Quality Standard		7	6	200	0.6	3	0.7	70	100	0.3	70
MW-22B	5/26/1994	25	770	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 25
	7/28/1994	25	460	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 200
	11/3/1994	25	810	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 1,000
	9/16/1996	200	2,400	< 200	< 200	< 200	< 200	< 200	< 200	< 200	< 200	<1
	8/13/1997	1,000	42,000	< 1,000	< 1,000	< 1,000	<1,000	< 1,000	< 1,000	< 1,000	< 1,000	<1,000
	9/10/1998	1	56,000	<1	<1	5	<1	<1	<1	<1	< 1	<100
	9/29/1999	1,000	15,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1000
	9/13/2000	100	38,000	<100	<100	<100	<100	<100	<100	<100	<100	<100
	8/29/2001	1,000	12,000	<1000	<1000	<1000	<1000	<1000	<1000	<1000	<1000	<10
	10/4/2002	100	2,700	<100	<100	<100	<100	<100	<100	<100	<100	<10
	9/24/2003	10	370	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/30/2004	10	1,400	<10	<10	<10	11	<10	22	<10	<10	<1
	9/26/2005	1	510	<10	<10	<10	<10	<10	<10	<10	<10	<1
	10/19/2006	10/100	1,200	<10	<10	<10	17	<10	23	<10	<10	<10
	9/26/2007	10	320	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/25/2008	1/10	820	<1	<l< td=""><td><1</td><td>0,74 J</td><td><1</td><td>0.95 J</td><td><1</td><td><1</td><td><1</td></l<>	<1	0,74 J	<1	0.95 J	<1	<1	<1
	9/17/2009	1/10	380	<1	<1	<1	2.9	<1	2.3	<1	<1	<1
	9/28/2010	1/10	940	< <u>1</u>	<1	<1	<1	<1	3.7	<1	<1	<1
MW-23A	5/26/1994	_1_	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	7/28/1994	1	3	<1	<1	< 1	<1	<1	<1	<1	< 1	< 1
	11/4/1994		<1	<1	<1	<1	< 1	<1	<1	<1	< 1	<1
	9/18/1998	1	<1	<1	<1	<1	<1	<1	<1	<1	< 1	<1
	9/28/1999	1	<1	<1	<1	<1	4.8	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1
	9/13/2000	1	<1	<1	<1	<1	2.3	<1	<1	<1	<1	<1
	8/28/2001	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<100
	9/30/2002	1	<1	<1	<u><l< u=""></l<></u>	<u> </u>	<1	<1	<1	<1	<1	<1
	9/29/2003	100	<100	<100	<100	<100	<100	<100	<100	<100	<100	< 1
	9/29/2004	1	<1	<1	<1	< <u>i</u>	7.1	<1	<1	<1	<1	<1
	9/30/2005	1	< 1	< 1	< 1	< 1	5	<1	<1	<1	<1	<1
}	10/16/2006	1	<1	<1	<1	<1	5,3	<t< td=""><td><1</td><td><1</td><td><1</td><td><1</td></t<>	<1	<1	<1	<1
	9/26/2007	1	<1	<1	<1	<1	5,2	<1	3.2	<1	<1	< <u>l</u>
	9/25/2008	1	<1	<1	<1	<1	3.1	<1	<1	<1	<1	<1
ļ	9/18/2009	1/10	<1	<1	<1	<1	1.6	<1	<l< td=""><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1
	10/1/2010	1/10	<1	<1	<1	<1	<1	<1	<1	<1	<1	< <u>i</u>
MW-23B	5/26/1994	1	<1	<1	6	< <u>1</u>	3	<1	<1	<1	< 1	<1
	7/28/1994	1	2	<1	<1	<1	4	<1	<1	<1	<1	<1
,	11/4/1994	1	<1	<1	3	<1	4	<1	< 1	<1	< l	<1
- 1	9/21/1998	1	<1	<1	<1	<1	9	<1	<1	<1	<1	<1
}	9/28/1999	1	<1	<1	<1	<1	2.9	<1	<1	<1	<1	<l< td=""></l<>
}	9/12/2000	1	<1	<1	<1	<1	2.3	<1	<1	<1	< <u>1</u>	<1
ŀ	8/28/2001	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
ŀ	9/30/2002	1	<1	<1	<1	<1	<1	<1	<1	<u><1</u>	<1	<1
}	9/29/2003	1	2	<1	<1	<1	<1	<1	<1	<1	<1	< 1
	9/29/2004	1	<1	<1	<1	<1	3,3	<1	<1	<1	<1	< I
}	9/27/2005	1	<1	<1	<1	<1	6.1	<1	<1	<1	<1	<1
ļ	10/17/2006	-1	<1	<1	<1	<1	7.9	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1
ŀ	9/26/2007	1	<1	<1	<1	<1	6.4	<1	<1	<1	<1	·<1
-	9/25/2008	1	<1	<1	< <u>l</u>	<1	3.9	<1	<1	<1	< <u>l</u>	<1
9	9/17/2009	1	<i< td=""><td><1</td><td><1</td><td><1</td><td>2.1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	2.1	<1	<1	<1	<1	<1
	1.0080000	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonis, North Carolina

Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CAS# 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CASW 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CASN 71-55-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CASI 79-01-6 ppb	Tetrachloro- ethene (PCE) CASI 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CAS# 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAS# 156-60-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb	Chloroform CAS# 67-66-3 ppb
	er Quality Standard		7	6	200	0,6	3	0.7	70	100	0.3	70
MW-24	7/26/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	2	<1
	11/1/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	1/17/1995	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< 1
	9/17/1996	1	<1	<1	<1	<1	< 1	<1	<1	<1	<1	<1
	9/17/1996	1	<1	<1	<1	<1	<1	<1	<1	<1	<1 <1	<u><1</u> <1
	3/26/1997	1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1 <1	<1	<1
	5/29/1997 7/7/1997	1 1	<1	<1	<1 <1	<u> </u>	<1 <1	<1	<1	<1	<1	<1
	8/13/1997	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	2,2
	11/13/1997	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/26/1998	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/10/1998	_11	<1	<1	<1	<1	< 1	<1	<1	<1	<1	3,1
	3/24/1999	1	<1	<5	<1	<1	<1	<1	<1	<1	<1	3.6
	9/16/1999	1	<1	<1	<1	<1	<1	<1	<1	<1	<1 <1	2.6 4.1
	3/23/2000	1	< <u> </u>	<1 <1	<1 <1	<1 <1	<1 <1	< <u>1</u>	<1 <1	<1 <1	<1 <1	4.1 <1
	9/11/2000 3/1/2001	1	<1 <1	<1 <1	<1	<1	<1	<1	<1	<1	<1	<1
	8/30/2001	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	3.3
	3/29/2002	i	<1	<1	<1	<1	<1	<1	<1	<1	<1	1,8
	9/30/2002	1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td>1.2</td></l<>	<1	<1	<1	<1	1.2
	3/25/2003	1	<1	<1	<1	<1	<1	<1	<1	<1	< <u>1</u>	< <u>l</u>
	9/25/2003	1	< <u>l</u>	<1	<1	<1	<1	< <u>I</u>	<1	<1	<1	<u><1</u>
	9/29/2004	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	1/17/2005 5/27/2005	-1	0.64	<1 <1	<1 <1	<1 <1	<1 <1	0.27 <1	< <u>1</u>	<1 <1	<1 <1	0.54 < 1
	9/27/2005	1 1	<1 <1	<1	<1 <1	<1	<1	<1	<1		۹ ا	<1
	3/10/2006	1	<1	<1	<1	<1	<1	<1	<1	<1	<	<1
i	10/18/2006	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/7/2007	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/21/2007	1	<1	<1	<1	<1	<1	<1	<1	<1	<l< td=""><td>< 5</td></l<>	< 5
	3/11/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< 5
	9/22/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<u><1</u>	<1
	4/29/2009	1	1.1	<1	<1	<1	<1	<1	<u> </u>	<1 <1	<1 <1	<1 <1
	9/15/2009	1	1.2 <1	<1	<1 <1	<u><1</u> <1	<1 <1	<1	<1 <1	<1	<1	<1
	e granda na kalendara ka	1	<1	<1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	<1	<1	<1
MW-25	7/28/1994	1	<1	<1	3	<1	2	<1	<1	<1	<1	<400
	11/1/1994	1	1	<1	4	<1	3	<1	<1	<1	<1	<10
	9/2/1998	400	<400	<400	<400	<400	<400	<400	<400	<400	<400	<1
	9/30/1999	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<40
	9/19/2000	1	<1	<1	<1	<1	3.5	<1	<1	<1 <40	<1 <40	<100 <40
	8/29/2001 10/1/2002	100	<40 <100	<40 <100	<40 <100	<40 <100	<40 <100	<40 <100	<40 <100	<100	<100	<40
	9/29/2003	40	<100 <40	<40	<40	<40	<40	<40	<40	<40	<40	< 40
	10/1/2004	40	<40	<40	<40_	<40	<40	<40	<40	<40	<40	<1
	9/30/2005	40	< 40	< 40	< 40	< 40	< 40	< 40	< 40	< 40_	< 40	6
	10/18/2006	40	<40	<40	<40	<40	<40	<40	<40	<40	<40	<40
	9/26/2007	40	<40	<40	<40	<40	<40	<40	<40	<40	<40	<40
	9/26/2008	10	<10	<10	<10	<10	<10	<10	<10	<10	<10 <10	<10 <10
	9/17/2009	10	<10 <10	<10 <10	<10 <10	<10 <10	<10 <10	<10 <10	<10 <10	<10 <10	<10	<10 <10
MW-26	7/28/1994	1	<10	<10	<1	<1	2	<1	<1	<1	<1	< 100
.** 17 -40	11/4/1994	1	<1	<1	3	<1	2	<1	<1	<1	<1	<10
	9/19/1998	100	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 100	<10
	9/30/1999	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/13/2000	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	8/28/2001	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/30/2002	10	<10	<10	<10	<10	<10	<10	<10	<10	<10 <10	<1 <1
	9/29/2003 10/4/2004	10	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10 <1	<10	<1
	9/30/2005	1	<1	<1	<1	<1	7.4	<1	<1	<1	<1	<1
	10/17/2006	1	<1	<1	<1	<1	8,6	<1	<1 _	<1	<1	<1
ľ	9/25/2007	ī	<1	<1	<1	<1	6.2	<1	<1	<1	<1	<1
į	9/25/2008	1/10	<1	<1	<1	<1	4.8	<1	<1	<1	<1	<1
[9/17/2009	1/10	<1	<1	<1	<1	0.77 J	<1	<1	<1	<1	<1
	3001 003	1/10	<1	<1	<1	<1	<1	<l< td=""><td><l< td=""><td><1</td><td><1</td><td><1</td></l<></td></l<>	<l< td=""><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1

Table 3 A305 roundwater Analytical Data Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells ummary of Detected Purgeable Halocarbons (Chlorinated Compounds) 207 Telegraph Drive, Gastonia, North Carolina 1,1,1-Trichloro 1 I-Dichlom-1.1-Dichloro Trichloro Tetrachloro cis-1,2-Dichloro trans-1,2-Dichloro 1,2-Dichloro-Carbon ethene ethane ethene ethene propane CAS# 78-87-5 (1,1-DCE) (1,1-DCA) (1,1,1-TCA) (TCE) (PCE) (cis-1,2-DCE) (trans-1,2-DCE) Tetrachloride Chloroform Sample Date of Dilution CAS# 75-35-4 CAS# 75-34-3 CAS# 56-23-5 CAS# 67-66-3 CAS# 71-55-6 CAS# 79-01-0 CAS# 127-18-CAS# 156-59-2 CAS# 156-60-5 Location Sampling Factor ppb ppb ppb NC Groundwater Quality Standard 200 100 70 MW-27 7/28/1994 <1 < 1 < 1 <1 11 <1 < 1 7/28/1994 <1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 11/3/1994 <1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <1 9/9/1998 <1 <1 <1 < 1 19 < 1 < 1 <1 <1 <1 9/27/1999 <1 <1 <1 <1 11 <1 <1 <1 <1 <1 9/12/2000 <1 <1 <1 <1 12 <1 <1 <1 <1 <1 8/28/2001 1 <1 <1 <1 <1 3,9 <1 <1 <1 <1 <1 9/30/2002 <1 <1 <1 <1 4.6 <1 <1 <! <I <1 9/29/2003 <1 <1 <1 <I 9.4 <! <1 <1 <1 < 1 9/29/2004 <1 <1 <1 <1 7.6 <1 <1 <1 <1 9/28/2005 <1 < 1 < 1 < 1 7.7 < 1 < 1 <1 < 1 11 10/16/2006 <1 <1 <1 9/25/2007 1 <1 <1 <1 <1 3.8 <1 <1 <1 <1 <1 9/26/2008 <1 <1 <1 <1 6,3 <1 <1 <1 <1 <1 9/17/2009 <1 <1 <1 4.2 <1 <1 <1 <1 <1 10/1/2010 <1 <1 <1 <1 5.5 <1 <1 <1 <1 <1 MW-29 7/28/1994 <1 <1 <1 < 1 < 1 < 1 < 1 <1 < 1 11/3/1994 < 1 < 1 < 1 < 1 < 1 < 1 <1 <1 9/9/1998 < 1 < 1 < 1 < 1 24 < 1 < 1 < 1 < 1 <1 9/27/1999 <1 <1 <1 <1 <1 <1 <1 8,3 <1 <1 9/13/2000 <1 <1 <1 <1 <1 <1 <1 <1 <1 2.5 8/28/2001 <1 <1 <1 <1 1.6 <1 <1 <1 <1 <1 9/30/2002 <1 <1 <1 <1 3.5 <1 <1 <1 <1 <1 9/29/2003 <1 <1 <1 <1 11 <1 <1 <1 <1 < 1 9/29/2004 <1 <1 <1 <1 <! <1 <<u>l</u> <1 < 1 9/28/2005 <<u>l</u> < 1 < 1 < 1 < 1 < 1 <1 < 1 10 <1 10/16/2006 <1 <i <1 <1 <1 <1 <1 <1 <1 9/25/2007 <1 <1 <1 <1 4.9 <1 <1 <1 <1 <1 9/26/2008 <1 <1 <1 <1 <1 <1 <1 4.2 <1 <1 9/17/2009 <1 <1 <1 <1 4,3 <1 <1 <1 <1 <<u>l</u> 10/1/2010 1 <1 <1 <1 5.8 <1 <1 <1 **~1** <1 MW-30 9/9/1994 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 11/2/1994 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <1 1/19/1995 < 1 < 1 < 1 < 1 < 1 <1 1 1/19/1995 < 1 2 < 1 < t < 1 < 1 < 1 MW-31A 11/4/1994 < 1 <1 <1 <1 < 1 <1 <1 < 1 < 1 11/4/1994 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <1 1/19/1995 < I < 1 < I < 1 < 1 < 1 < 1 < 1 <1 3/18/1998 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 5/27/1998 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Pargeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastoula, North Carolina

Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CASN 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CASW 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CASN 71-55-6 ppb	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CASW 79-01-6 ppb	Tetrachloro- ethene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASN 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAS# 156-60-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb	Chlorofori CAS# 67-66- ppb
Groundwate	r Quality Standard		7	6	200	0,6	3	0.7	70	100	0.3	70
MW-31B	11/3/1994	1	<1	<1	<1	<1	1	<1	<1	<1	<1	<1
	1/19/1995	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/26/1998	1	<1	<1	<1	<1	4.2	<1	<1	<1	<1	. <1
	5/29/1998	1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td>4</td><td><1 <1</td></l<>	<1	<1	<1	<1	4	<1 <1
	9/5/1998	1	<1	<u> </u>	<1	<1	<1	<1	<1	<1<1	<1 <1	<1
	12/31/1998	1	<1	<1	<1	<1	3 <1	<1 <1	<u> </u>	<1	<1	<1
	3/17/1999 5/20/1999	1	<1	<1	< <u>1</u>	<1 <1	2	< <u> </u>	<1	<1	<l< td=""><td><1</td></l<>	<1
	9/28/1999	1	< <u>1</u>	<1	<1	<1	3	<1	<1	<1	<1	<1
	12/16/1999	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/23/2000	1	<1	<1	<1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td></l<>	<1	<1
	6/16/2000	1	<1	<1	<1	<1	1.2	<l< td=""><td><1</td><td><1</td><td><<u>l</u></td><td><1</td></l<>	<1	<1	< <u>l</u>	<1
	9/7/2000	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< <u>l</u>
	12/28/2000	1	<1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1
	2/28/2001	1	<1	<1	<1	<1	1,2	<1	<1	<1	<1	<1_
	6/7/2001	Į.	<1	<1	<1	<1	1.1	<1	<1	<1	<1	<1
	8/31/2001	1	<1	<1	<1	< <u>l</u>	<1	<1	<1	<1	<1	<1
	3/15/2002	1_	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1
	10/1/2002	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/26/2003	1	<1	<1	<1	<1	<1	<1	<1	<1	<1 <1	< <u>1</u>
	9/24/2003	1	<1	<1	<1	<1	<1	<1	<1	< <u>1</u>	<1 <1	<1
	9/28/2004	1	<1	<1	<1	<1 <1	1.7 2,4	<u><1</u> <1	<1 <1	<1	<1	<1
	1/17/2005	1	<1	<1	<1		1.7	<1	<1	<1	<1	<1
	6/23/2005	1	<1	<1	<1	<1 <1	<1.7 <1	<1	<1	<1	<1	<1
	9/26/2005	1	<1	<1 <1	<1 <1		<1	<1	<1	<1	<1	<1
	3/10/2006	-	< <u>1</u>	<1	<1	<1	1.5	<1	<1	<1	<1	<1
	10/16/2006	1 1	<1	<1	<1	<1	1.7	<1	<1	<1	<1	<1
	3/8/2007 9/24/2007	- 	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/10/2008	1	<1	<1	<1	<1	1.4	<1	<1	<1	<1	<1
	9/23/2008	i	<1	<1	<1	<1	1.6	<1	<1	<1	<1	<1
	4/30/2009	i	<1	<1	<1	<1	1.3	<1	<1	<1	<1	<1
	9/16/2009	î	<1	<1	<l< td=""><td><1</td><td>1.2</td><td><<u>l</u></td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	1.2	< <u>l</u>	<1	<1	<1	<1
	9/28/2010	1	<1	<1	<1	<1	2.1	<1	<1	<1	<1	<1
	100	1	<1	<1	<1	<1	1.3	<1	<1	<1	<1	<1
	Service Const	1	<1	<1	<1	<1	2,1	<1	<1	<1	<1	<1
MW-32	11/4/1994	1	<1	<1	<1	<1	<1	<1	<1	≤1	<1	< <u>l</u>
	1/19/1995	1	<1	<1	< 1	<1	<1	<1	<1	<1	<1	<l< td=""></l<>
	3/18/1998	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< <u>l</u>
	9/5/1998	1	<1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td><td><<u>l</u></td><td><<u>l</u></td></l<>	<1	<1	< <u>l</u>	< <u>l</u>
	3/19/1999	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/29/1999	1	<1	<1	<1	<1	<1	<1	<1	<1	<u><1</u>	<l< td=""></l<>
	3/27/2000	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< <u>l</u>
	9/12/2000	. 1	<1	<1	<1	<1	<1	<1	<1	<1 <1	<1 <1	<1 <1
	2/28/2001	1	<1	< <u>i</u>	<1	<1	<1	<1	<1	<1	<1 <1	< <u> </u> <
	8/31/2001	1	<1	<1	<1	<1	<1 <1	<1 <1	<1 <1	<1	<1	<1
	3/15/2002	1	<1	<u><1</u>	<1	<1 <1	<1	<1	<1	<1	<1	<l< td=""></l<>
	10/1/2002	1	<1	<1	<1 <1	<i <i<="" td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i>	<1	<1	<1	<1	<1	<1
	3/26/2003 9/24/2003	1.	<1	<1 <1	<1	<u><1</u>	<1	<1	<1	<1	<1	<1
	9/24/2003 9/28/2004	1	<1 <1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	1/17/2005	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	6/23/2005		<1	<1	<1	<1	0.97	<1	<1	<1	<1	2
	9/26/2005	<u>-</u>	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/10/2006	i	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	10/17/2006	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/8/2007	1	<1	<1	<1	<1	<1	<1	<1	<1	< 1	<1
	9/24/2007	i	<1	<1	<1	<1	<1	< <u>l</u>	⊲	<1	<1	<1
	3/11/2008	1	<1	<1	<1	<1	<1	<1	<1	_<1	<1	<1
	9/23/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	4/30/2009	1 _	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/16/2009	1	<1	<1	<1	<1	<1	<1	<1	<1	· <1	<1
	9/28/2010	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
		1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<u> </u>
		1	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></i<>	<1	<1	<1	<1	<1	<1	<1	<1

ater Analytical Data Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells nmary of Detected Purgeable Halocarbons (Chlorinated Compounds) 207 Telegraph Drive, Gastonia, North Carolina 1,1,1-Trichloro 1 1-Dichloro-1.1-Dichloro Trichloro Tetrachloro cis-1,2-Dichloro trans-1,2-Dichloro-1,2-Dichloroethane ethane ethene ethene ethene Carbon (1,1-DCA) CAS# 75-34-3 (1,1,1-TCA) CASI 71-55-6 propane CASI 78-87-5 (1,1-DCE) (TCE) (PCE) (cis-1.2-DCE) (trans-1.2-DCE) Tetrachloride Chloroform Sample Date of Dilution CAS# 75-35-4 CAS# 67-66-3 CAS# 79-01-6 CAS# 127-18-CAS# 156-59-2 CAS# 156-60-5 CAS# 56-23-5 Sampling Factor ppb ppb ppb NC Groundwater Quality Standard 200 0,6 70 100 70 0.3 MW-33 9/27/1994 < 1 <1 < 1 <1 <1 <1 <1 < 1 11/2/1994 < 1 < 1 6 < 1 < 1 < 1 < 1 < 1 < 1 < 1 11/2/1994 <1 <1 < 1 <1 <1 < 1 < 1 < 1 2.4 < 1 1/17/1995 1 5 <1 <1 <1 < 1 <1 <1 < 1 <1 9/19/1996 36.4 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <1 3/26/1997 11 < 1 <1 < 1 < 1 < 1 < i < 1 <1 1.1 5/29/1997 17 <1 <1 <1 <1 <1 <1 <1 <1 1.7 7/7/1997 18 <1 <1 <1 <1 <1 <1 <1 <1 <1 8/13/1997 14 < 1 < 1 < 1 < 1 < 1 <1 < 1 < 1 <1 11/13/1997 ı 13 <l <1 <1 <1 <1 <1 <1 <1 <1 9/2/1998 15 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/28/1999 2.9 <1 <1 <1 <1 <1 9/18/2000 1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 8/27/2001 <1 <1 <1 <1 <1 <1 <l <1 <1 <1 10/4/2002 1.9 <<u>1</u> <1 <1 <1 <1 <1 <1 9/25/2003 <1 <1 <1 <1 <1 <1 <1 <1 <1 < 1 10/1/2004 <1 <1 <1 <1 <1 <i <1 <! <1 < 1 9/28/2005 < 1 < 1 < 1 < 1 < 1 < 1 <1 < 1 < 1 < 1 10/19/2006 1/10 440 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/26/2007 1/10 710 <1 <1 <1 <l <1 <1 <1 <1 <1 1/10 110 9/25/2008 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/16/2009 1/10 420 <1 <1 <1 <1 <1 <1 <1 <1 <19/28/201b 1/10 210 <1 <1 <1 <1 <1 <1 <1 <1 <1 MW-34A 1/20/1995 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 9/16/1996 < 1 < 1 < 1 < 1 < 1 < <u>1</u> <1 <1 8/13/1997 <1 < 1 <1 < 1 <1 < 1 < 1 <1 <1 1 <1 3/26/1998 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 5/29/1998 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/2/1998 <1 <1 <1 <1 <1 <1 <1 <1 <i <1 12/31/1998 <1 <1 <1 <1 <1 <1 <1 <1 3/17/1999 <Į <1 <1 <1 <1 1 5/19/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/28/1999 <<u>l</u> <1 <1 <1 <1 <1 <1 <1 <1 <1 12/16/1999 <1 <1 <1 <1 ≤1 <1 <1 <1 <1 <1 3/27/2000 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 6/15/2000 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/8/2000 <1 <1 <1 <1 <1 <1 <1 <1 <i <1 12/28/2000 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 2/28/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 6/7/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 8/30/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 3/29/2002 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 10/1/2002 <] <1 <1 <1 <1 <1 <1 <1 <1 <1 3/26/2003 <1 <1 <1 <1

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Table 3

9/24/2003

9/28/2004

1/17/2005

5/27/2005

9/27/2005

3/10/2006

10/18/2006

3/8/2007

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3/12/2008

9/23/2008

4/29/2009

9/14/2009

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Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells ummary of Detected Purgeable Halocarbons (Chlorinated Compounds) 207 Telegraph Drive, Gastonia, North Carolina 1,1-Dichloro-1,1-Dichloro-1,1,1-Trichloro-Trichloro Tetrachloro cis-1,2-Dichlorotrans-1,2-Dichloroethene 1.2-Dichloro ethene ethene ethene Carbon (cis-1,2-DCE) (trans-1,2-DCE) Tetrachloride Chloroform (1.1-DCE) (1,1,1-TCA) (TCE) (PCE) (1,1-DCA) propane Dilution CAS# 71-55-6 CAS# 78-87-3 CAS# 79-01-6 CAS# 127-18 CAS# 156-59-2 CAS# 156-60-5 CAS# 56-23-5 CAS# 67-66-3 Sample ppb Sampling ppb ppb ppb ppb ppb 0.7 70 100 0.3 NC Groundwater Quality Standard 200 0.6 <1 <1 <1 < 1 < 1 < 1 MW-34B 6/28/1995 < 1 < 1 < 1 9/16/1996 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <1 < 1 < 1 <1 < 1 <1 < 1 <1 3/25/1997 3.4 < 1 <1 5/30/1997 <1 <1 <1 <1 <! <1 <1 <1 <1 <1 <1 <1 <1 <1 1.2 7/7/1997 1.4 <1 < 1 <1 < 1 < 1 < 1 < i <1 8/14/1997 2.7 < 1 <1 11/13/1997 1.7 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/2/1998 <1 <i <1 <1 <1 9/29/1999 <1 <1 <1 <1 <1 <1 1.5 9/13/2000 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 8/30/2001 <1 <1 <1 <1 10/4/2002 2.9 <1 <I <1 <1 <1 9/24/2003 1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <l <1 <1 <1 9/28/2004 <1 <! <1 <1 9/26/2005 <1 <1 <1 <1 <1 10/18/2006 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/26/2007 3.5 <1 <1 <1 <1 9/26/2008 6.6 <1 <1 <1 <1 9/18/2009 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <I <1 <1 <1 #3/pv/pi0107* 5.1 <1 <1 <1 <1 <1 MW-35 9/19/1998 <1 <1 <1 <1 <1 <1 9/28/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/13/2000 <1 <1 <1 <1 8/29/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 10/1/2002 <1 <1 <1 <1 <1 9/29/2003 <1 <1 <1 <1 <1 < 1 9/29/2004 <1 <1 <1 <1 <! <1 <1 <1 <1 <1 <1 < 1 < 1 < 1 < 1 9/27/2005 < 1 <1 <1 <1 10/16/2006 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/26/2007 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/26/2008 <1 <1 <1 9/18/2009 <1 <1 <1 <I <1 <1 <1 <1 <i <1 <1 <1 <1 <1 <1 <1 <1 <1 1090-0100 <1 < 1 <1 < <u>ļ</u> MW-37A < 1 3/7/1996 < 1 < 1 < 1 < 1 < 1 MW-37B 3/7/1996 < 1 < 1 < 1 < 1 < 1 < 1 <1 < <u>1</u> <1 <1 <1 < 1 < 1 < 1 <1 MW-38 7/25/1997 < 1 <1 8/13/1997 < 1 < 1 < 1 < 1 < 1 < 1 <1 3/27/1998 <1 <1 <1 <<u>1</u> <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 5/29/1998 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/2/1998 <1 <1 <1 <1 <1 <1 <1 <1 <1 12/29/1998 <1 <1 <1 3/22/1999 <1 <1 <I <1 <1 <1 <1 5/20/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 1.3 9/16/1999 <1 <1 <1 12/20/1999 <1 <1 <1 <1 2.4 2.5 <1 <1 <1 <1 <1 <1 3,6 4.2 <1 <1 <1 <l <<u>1</u> <1 3/24/2000 1 <1 <1 1.2 <1 6/15/2000 <1 1.7 <1 9/8/2000 <1 <1 <1 <1 <1 <I <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <u><1</u> <1 12/28/2000 <1 <1 <1 <1 <1 <1 3/1/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 6/7/2001 <1 <1 <1 <1 <1 <i <1 <1 8/27/2001 <1 <1 <1 <1 <1 <1 <I <1 <1 <1 <1 <1 3/28/2002 <1 <1 <1 3.7 <1 <1 <1 <1 <1 9/30/2002 <1 <<u>1</u> <1 <1 <1 <1 3/25/2003 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/24/2003 <1 <1 <1 <1 <1 <1 <1 <1 9/29/2004 <1 <1 1.6 1/14/2005 <1 <1 <1 <1 1.3 <1 <1 <1 <1 <1 <1 <1 <1 <1 < 1 <1 <1 <<u>1</u> 6/22/2005 <1 <1 <1 <1 <1 <1 9/27/2005 <1 <1 <1 3/10/2006 <1 <1 <<u>l</u> <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 10/19/2006 <1 < 1 < 1 1.8 3/7/2007 4.1 < 1 < 1 < 1 < 1 < 1 <1 <1 <1 <<u>i</u> <1 <1 <1 <1 2.0 9/20/2007 <1 <1 <1 <1 <1 1,0 <1 3/11/2008 7.0 <1 <1 4.4 <1 <1 <1 <1 <1 <1 <1 9/22/2008 <1 <i <1 <1 <1 0.98 J <1 <1 11.0 4/29/2009 <1 <1 <1 <1 1.2 9/15/2009 12.0 <1 <1 <1 <1

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207 Telegraph Drive, Gastonia, North Carolina 1,1-Dichloro-1,1-Dichloro 1,1,1-Trichloro-Trichloro-Tetrachloro cis-1,2-Dichlorotrans-1,2-Dichloroethene ethane ethane 1.2-Dichloroethene Carbon (1,1-DCE) (1,1-DCA) (1,1,1-TCA) (TCE) (cis-1,2-DCE) propane CAS# 78-87-5 (PCE) (trans-1,2-DCE) Tetrachloride Chloroform Dilution Sample Date of CAS# 75-35-4 CAS# 75-34-3 CAS# 71-55-6 CAS# 79-01-0 CAS# 127-18-CASV 156-59-2 CASN 156-60-5 CASN 56-23-5 CAS# 67-66-3 Factor ppb ppb ppb ppb ppb ppb ppb ppb ppb NC Groundwater Quality Standard 200 0,6 0.7 70 100 0.3 70 8/1/1997 <1 < <u>1</u> < 1 <1 <1 < 1 <1 <1 < 1 8/13/1997 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <] <1 3/25/1998 <<u>1</u> <1 <1 <1 <1 <1 <1 5/28/1998 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/9/1998 <I <1 <1 <1 <<u>1</u> <1 <1 <1 <1 <u><1</u> 12/30/1998 <1 <1 <1 <1 <1 <1 3/24/1999 <1 <1 <1 <1 <1 <1 <i <1 <1 1.6 5/20/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/15/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 12/20/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 3/24/2000 <1 <1 <1 <1 <1 <1 <1 <1 6/15/2000 <1 <1 <1 <1 <1 <1 <! <1 <1 <1 9/8/2000 <1 <1 <1 <1 <1 <1 <I <1 <1 <1 12/28/2000 <! <1 <1 <1 <1 <1 <1 <1 <1 <1 2/28/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 6/7/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 8/27/2001 <1 <1 <1 <1 <1 <1 <1 <1 1.1 3/28/2002 <1 <1 <1 <1 <1 <1 <1 <1 <1 1.3 10/1/2002 <1 <1 <1 <1 <! <1 <1 1.6 3/25/2003 <1 <1 <1 <1 <1 <1 2.1 <1 <1 <1 9/25/2003 <1 <1 <1 <1 <1 <1 <1 <1 <1 3.9 9/29/2004 <1 <1 <1 <1 <1 <1 < 1 1/18/2005 <1 <1 <1 <1 <1 <1 <1 <1 <1 3.1 5/27/2005 <1 <1 <1 <1 <1 <1 <1 <1 ≤1 2,8 9/27/2005 <1 <1 <u>۔</u> 1> <1 <1 <1 <1 <] <1 3/10/2006 <1 <1 <1 <1 <1 <1 <1 <1 <1 2.7 10/19/2006 68 <<u>l</u> <1 <1 <1 <1 <1 <1 <1 4.1 3/7/2007 5.4 < 1 < 1 < 1 < 1 < 1 5.6 < 1 9/20/2007 5.5 <1 <1 <1 <1 <1 <l <1 <1 6.9 3/11/2008 30 <1 <1 <1 <1 <1 <1 <1 3.7 9/22/2008 10 <1 <1 <1 <1 <1 <1 <1 <1 2.8 4/29/2009 51 <1 <1 <1 <1 <1 <1 <1 <1 2.9 9/15/2009 <1 52 <1 <1 <1 <1 <1 <1 <i 3.2 3/23/2010 <1 21 <1 <1 <1 <1 <1 <1 <1 2.8 15 <1 <1 <1 <1 <1 <1 <1 <1 2.1 MW-40 <1 8/1/1997 < 1 < 1 < 1 < 1 <1 <1 < i < 1 <1 8/12/1997 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 <1 3/27/1998 3.4 <1 <1 <1 <1 <1 1.7 <1 5/28/1998 <1 <1 <1 <1 <1 <1 <l <! <1 <1 9/9/1998 <1 <1 <1 <1 <1 <1 <1 <1 <<u>l</u> <1 12/29/1998 <1 <1 <1 <1 <1 <1 <1 <1 3/22/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 5/20/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/16/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 12/16/1999 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 3/24/2000 4.6 <1 <1 <1 <1 <<u>l</u> <1 <1 <1 <1 6/15/2000 <1 <1 <I <1 <1 <1 <1 <1 <1 <1 9/11/2000 <1 <1 <1 <1 <1 <1 <1 <1 1.5 12/28/2000 <l <1 <1 <I <1 1.3 <1 <1 <1 <1 3/1/2001 3,4 <1 <1 <1 <1 <1 <1 <1 <1 <1 6/7/2001 1.1 <1 <1 <1 <1 <1 <1 <1 <1 8/30/2001 <1 <1 <1 <1 <1 <1 <1 <1 <1 1.7 3/28/2002 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/30/2002 1.9 <1 <1 <1 <1 <! <1 <1 <1 3/25/2003 1.1 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/25/2003 <1 <1 <1 <1 <u><1</u> <1 <1 <1 <1 <1 9/30/2004 17 <1 <1 <1 <1 <1 <1 <1 < 1 1/17/2005 <1 <1 <1 <1 <1 0.35 <1 <1 <1 0.5 6/23/2005 <1 <1 <1 <1 <1 <1 <1 <! <1 <1 9/27/2005 <1 <1 < 1 < 1 <1 < 1 <1 < 1 < 1 < 1 3/10/2006 2,4 <1 <I <1 <1 <1 <1 <1 <1 10/16/2006 110 <u><1</u> <1 <1 <1 <1 <1 <1 <1 <1 3/7/2007 1/4 140 < 1 < 1 < 1 < 1 < 1 < 1 <1 < 1 < 1 9/20/2007 7.3 <1 <1 <1 <1 <1 <1 <1 <1 3/11/2008 19 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/22/2008 17 <1 <1 <1 <1 <1 <1 <1 <1 0.94 J 4/29/2009 1/10 640 <1 <1 <1 <1 <1 <1 <1 <1 <1 9/14/2009 91 <1 <1 <1 <1 <1 <1 <1 <1 <1 30 <1 <1 <1 <1 <1 <1 <1 <1 <1 geden (i.e. <1

Table 3

Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells Summary of Detected Purgeable Halocarbons (Chlorinated Compounds) Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Sammary of Detected Parageable Halocarbona (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

	Drive, Gastonia, N	1	9.4	1		1	1		1	· ·	T.	
			1,1-Dichlora-	1,1-Dichloro-	1,1,1-Trichloro-		Trichloro-	Tetrachloro-	cis-1,2-Dichloro-	trans-1,2-Dichloro-		
			ethene	ethane	ethane	1,2-Dichloro-	ethene	ethene	ethene	ethene	Carbon	
			(1,1-DCE)	(1,1-DCA)	(1,1,1-TCA)	propane	(TCE)	(PCE)	(cis-1,2-DCE) CAS# 156-59-2	(trans-1,2-DCE) CAS# 156-60-5	Tetrachloride CAS# 56-23-5	Chlorofo
Sample Location	Date of Sampling	Dilution Factor	CAS# 75-35-4 ppb	CAS# 75-34-3 ppb	CAS# 71-55-6 ppb	CAS# 78-87-5 ppb	CAS# 79-01-6 ppb	CAS# 127-18-4 ppb	ppb	Dpb	ppb	ppb
	er Quality Standard	1	7	6	200	0.6	3	0.7	70	100	0.3	70
DM-7	2/8/1994	1	<1	<1	<1	<1	< 1	< 1	< 1	<1	<1	< 20
2	5/26/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< 25
CP-I	2/10/1994	20	1,000	< 20	< 20	< 20	< 20	< 20	< 20	< 20	< 20	< 10
Cr•1	5/26/1994	25	2,000	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 25	< 50
	5/26/1994	20	610	< 10	9.6 J	< 20	8.8 J	6.4 J	< 10	< 10	< 10	<1
	7/28/1994	50	1,300	< 50	< 50	< 50	< 50	< 50	< 50	< 50	< 50	< 630
		1000			<1		<1	<1	<1	<1	<1	< 1,00
	1/30/1997		30,000	<1		<1		< 630	<630	< 630	< 630	< 1,00
	3/18/1997	625	27,000	< 630	< 630	< 630	< 630		< 1,000	< 1,000	< 1,000	< 1,00
	3/21/1997	1,000	34,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000			< 1,000	< 1,00
	3/26/1997	1,000	31,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,00
	3/28/1997	1,000	32,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000		
	4/2/1997	1,000	34,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,00
	4/4/1997	1,000	34,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	<1,00
	4/7/1997	1,000	31,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	< 1,000	<10
	3/18/1998	1,000	22,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1,000	<1
	6/3/1998	10	30,000	<10	<10	<10	<10	<10	<10	<10	<10	_ <1
	8/27/1998	1	20,000	<1	<1	4	4	<1	1	< 1	<1	<1
	12/30/1998	1	9,500	<1	<1	2	3	<1	<1	<1	<1	2
	3/19/1999	1	5,500	<1	<1	<1	1	<1	<1	2	<1	<40
	5/19/1999	1	15,000	11		3	8	<1	7	<1	<1	<40
	9/20/1999	40	13,000	<40	<40	<40	<40	<40	<40	<40	<40	<400
	12/20/1999	40	11,000	<40	<40	<40	<40	<40	<40	<40	<40	<400
	3/22/2000	400	10,000	<400	<400	<400	<400	<400	<400	<400	<400	<1
	6/14/2000	400	19,000	<400	<400	<400	<400	<400	<400	<400	<400	<100
	9/8/2000	1	8,100	1.9	<1	2,5	15	<1	15	<1	<1	<400
	12/1/2000	1000	9,300	<1000	<1000	<1000	<1000	<1000	<1000	<1000	<1000	<400
	2/21/2001	400	11,000	<400	<400	<400	<400	<400	<400	<400	<400	<400
	6/5/2001	400	13,000	<400	<400	<400	<400	<400	<400	<400	<400	<400
	8/23/2001	400	11,000	<400	<400	<400	<400	<400	<400	<400	<400	<40
	3/14/2002	400	4,900	<400	<400	<400	<400	<400	<400	<400	<400	<40
	10/2/2002	40	3,800	<40	<40	<40	<40	<40	<40	<40	<40	<400
	3/25/2003	40	4,400	<40	<40	<40	<40	<40	<40	<40	<40	<10
	9/23/2003	400	3,300	<400	<400	<400	<400	<400	<400	<400	<400	< 10
	10/1/2004	10	4,100	<10	<10	<10	36	<10	46	<10	<10	<1
	1/17/2005	10	4,300	<10	<10	<10	48	<10	56	<10	<10	<10
	5/27/2005	10	4,600	<10	<10	<10	30	<10	39	<10	<10	<10
	9/27/2005	10/100	1,900	< 10	< 10	< 10	40	< 10	51	< 10	< 10	<1
	3/8/2006	10/100	1,900	< 10	< 10	< 10	51	< 10	73	< 10	< 10	< 10
	3/6/2007	10/100	3,100	< 10	< 10	< 10	37	< 10	47	< 10	< 10	< 10
	9/24/2007	10/100	5,500	< 10	< 10	< 10	28	< 10	45	< 10	< 10	< 10
	3/11/2008	10/100	3,200	< 10	< 10	< 10	38	< 10	57	< 10	< 10	< 10
	9/22/2008	10/100	2,200	< 10	< 10	< 10	49	< 10	63	< 10	< 10	< 10
	4/30/2009	10	440	< 10	< 10	< 10	36	< 10	78	< 10	< 10	< 10
	9/17/2009	10	1,100	< 10	< 10	< 10	41	< 10	62	< 10	< 10	< 10
	3/11/2003	10/100	1,400	< 10	< 10	< 10	25	< 10	31	< 10	< 10	< 10
	8 (70) (70)	10/100	1,500	< 10	< 10	< 10	71	< 10	33	< 10	< 10	< 10
CP-2	2/10/1994	1	4	<1	< i	<1	<1	<1	< 1	<1	<1	<1
WS-3	2/16/1994	i	_5	2	2	3	5	<1	2	<1	<1	<1
,, 3-3	5/25/1994	1	4	<1	5	4	2	<1	<1	<1	<1	<1
	7/26/1994	1	3	<1	2	2	3	3	<1	<1	<1	<1
WS-6	5/26/1994	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	< 10
		1		<1	<1	<1	<1	<1	<1	<1	<1	< 10
WS-7	8/25/1994		<1					< 10	< 10	< 10	< 10	<1
	3/15/1996	10	480	< 10	< 10	< 10	< 10					
	3/15/1996	10	590	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	<1
	12/20/1996	5	110	<1	<1	<1	<1	<1	<1	<1	<1	<1
	1/16/1997	1	24	<1	<1	<1	<1	<1	<1	<1	<1	<1
	4/18/1997	1	5.6	<1	<1	<1	<1	<u> </u>	<1	<1	<1	<1
	1/30/1997	1	15	<1	< i	<1	1.2	<1	<1	<1	<1	< 4
	3/3/1997	1	7.4	<1	< 1	< 1	1.2	< 1	<1	<1	< 1	<1
	7/31/1997	4	18	< 4	< 4	<4	<4	< 4	<4	<4	< 4	<500

Table 3
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Pergeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

Sample Location IC Groundwate	Date of Sampling er Quality Standard	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CASN 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CASE 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CAS# 71-55-6 ppb 200	1,2-Dichloro- propane CAS# 78-87-5 ppb	Trichloro- ethene (TCE) CAS# 79-01-6 ppb	Tetrachloro- cthene (PCE) CAS# 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASN 156-59-2 ppb	trans-1,2-Dichloro- ethene (trans-1,2-DCE) CAS# 156-60-5 ppb	Carbon Tetrachloride CASN 56-23-5 ppb 0,3	Chloroform CAS# 67-66-3 ppb 70
EW-2	1/30/1997	250	4,900	<1	<1	<1	<1	<1	<1	<1	<1	<1
	3/18/1998	500	6,200	<500	<500	<500	<500	<500	<500	<500	<500	<1
	5/27/1998	1	9,500	<1	7	<1	<1	<1	<1	<1	<1	<1
	8/27/1998	1	<1	<1	<1	<1	11	<1	< 1	<1	<1	<1
	12/30/1998	1	9,800	<1	<1	<1	<1	<1	<1	<1	<1	< <u>l</u>
	3/19/1999	1	2,500	<1	3	<1	<1	<1	<1	<1	<1	1.6
	5/19/1999	- 1	9,700	<1	3	1	<1	<1	<1	1	<1	<40
	9/20/1999	40	10,000 6,400	<1 <40	5,2 <40	<1 <40	<1	<1	<1	1.4	<1	1.6
	3/21/2000	1	7,500	<1	2	<1	<40 2.1	<40 <1	<40 <1	<40 2,1	<40 <1	<100 <100
	6/14/2000	100	20,000	<100	<100	<100	<100	<100	<100	<100	<100	<100
	9/8/2000	100	5,300	<100	<100	<100	<100	<100	<100	<100	<100	<40
	12/1/2000	100	2,800	<100	<100	<100	<100	<100	<100	<100	<100	<100
	2/21/2001	40	3,100	<40	<40	<40	<40	<40	<40	<40	<40	<100_
i	6/5/2001	100	2,900	<100	<100	<100	<100	<100	<100	<100	<100	<40
	8/23/2001	100	2,800	<100	<100	<100	<100	<100	<100	<100	<100	<100
	3/14/2002	40	4,200	<40	<40	<40	<40	<40	<40	<40	<40	<40
	10/2/2002	100	5,100	<100	<100	<100	<100	<100	<100	<100	<100	<400
	3/25/2003	40	4,800	<40	<40	<40	<40	<40	<40	<40	<40	<10
	9/23/2003	400	3,800	<400	<400	<400	<400	<400	<400	<400	<400	<100
	10/1/2004	100	4,400	<10	<10	<10	<10	<10	<10	<10	<10	<1
	5/27/2005	100	4,600 3,500	<10 <100	<10 <100	<10 <100	<10 <100	<10 <100	<10 <100	<100	<100 <100	<100
	9/27/2005	100	9,600	<100	<100	<100	<100	<100	<100	<100	<100	<100
	3/10/2006	100	3,100	<100	<100	<100	<100	<u><100</u>	<100	<100	<100	<100
l	10/13/2006	100	12,000	<100	<100	<100	<100	<100	<100	<100	<100	<100
Į	3/6/2007	100	3,900	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 100	< 100
[9/24/2007	100	4,200	<100	<100	<100	<100	<100	<100	<100	<100	<100
	3/11/2008	10/100	2,800	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10	< 10
,	9/22/2008	10/100	2,800	<10	<10	<10	<10	<10	<10	<10	<10	<10
	4/30/2009	10/100	4,900	<10	<10	<10	<10	<10	<10	<10	<10	<10
ļ	9/17/2009	10/100	1,900	<10	<10	<10	<10	<10	<10	<10	<10	<10
ŀ	19/23/2010	10/100	1,400	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/30/2010	10	960	<10	<10	<10	<10	<10	<10	<10	<10	<10
EW-3	1/30/1997	25	300	<1	<1	<1	<1	<1	<1	<1	<1	2
ŀ	3/18/1998 5/27/1998	10	100 33	<10 < 1	3	<10 < 1	<10 < 1	<10 <1	<10 < 1	<10 <1	<10	2 2
	8/27/1998		53	<1	6	<1	<1	<1	<1	<1	<1	<1
Ì	12/30/1998	1	35	<1	5	<1	<1	<1	<1	<1	<1	2
Ī	3/19/1999	1	11	<1	4	<1	<1	<1	<1	<1	<1	<1
	5/19/1999	1	36	<1	5	<l< td=""><td><1</td><td><i< td=""><td><1</td><td><1</td><td><1</td><td>1.1</td></i<></td></l<>	<1	<i< td=""><td><1</td><td><1</td><td><1</td><td>1.1</td></i<>	<1	<1	<1	1.1
	9/20/1999	1	30	<l< td=""><td>4.5</td><td><1</td><td><l< td=""><td><<u>l</u></td><td>2.6</td><td><1</td><td><1</td><td><i< td=""></i<></td></l<></td></l<>	4.5	<1	<l< td=""><td><<u>l</u></td><td>2.6</td><td><1</td><td><1</td><td><i< td=""></i<></td></l<>	< <u>l</u>	2.6	<1	<1	<i< td=""></i<>
ļ	12/20/1999	1	23	<1	5.1	<1	<1	<1	<1	<1	<1	<1
- [3/21/2000	1	25	<1	2.6	<1	2.4	<1	<1	<1	<1	<1
-	6/15/2000	1	16	<1	<1	<1	<1	<1	<1	<1	<1	<1
	6/5/2001	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
L	8/23/2001	1	4.8	<1	<1	</td <td><1</td> <td><1</td> <td><1</td> <td><1</td> <td><<u>l</u></td> <td><u> </u></td>	<1	<1	<1	<1	< <u>l</u>	<u> </u>
-	3/14/2002	1	9	<1	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1	<1	<1
-	10/2/2002 3/25/2003	1 1	18 91	<1 <1	1.2	<1 <1	<1 <1	<1	<1	<1 <1	<1 <1	1.4 <1
F	9/23/2003	1	89	<1		<1	<1	<1	<1	<1	<1	<1 <1
ľ	10/1/2004	1	510	<1	<1	<1	<1	<1	<1	<1	<1	<1
Ī	1/17/2005	10	180	<10	<10	<10	<10	<10	<10	<10	<10	<10
	6/23/2005	1	86	<1	<1	<1	<1	<1	<1	<1	<1	< 2.5
	9/27/2005	1	150	<1	<1	<1	<1	<1	<1	<1	<1	<1
ļ	3/10/2006	1	58	<1	<1	<l< td=""><td><l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<></td></l<>	<l< td=""><td><1</td><td><1</td><td><1</td><td><1</td><td><1</td></l<>	<1	<1	<1	<1	<1
L	10/13/2006	1/40	610	<1	<1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td></l<>	<1	<1
-	3/6/2007	1/10	350	<1	<1	<1	0.86 J	<1	0, 65	<1	<1	< 1
-	9/24/2007	1/10	270	<1	<1	<1	<1	<1	<1	<1	<1	<1
-	3/11/2008	1/10	180	<1	<1	<1	<1	<u><1</u>	</td <td><u> </u></td> <td><1</td> <td>. <1</td>	<u> </u>	<1	. <1
}	9/22/2008	1/10	100	<1	<1	<1	<1	<1	<1	<1	<1	<1
H	4/30/2009	1/10	540	<1	<1	<1	<1	<1	<1	<1	<1	<1
3	9/17/2009	1/100	1,500	<1	<1	<1	<1 <1	< <u>l</u>	<1	<1	<1	<1
3	97:020 (0	1/40	140	<1	<1	<1	<u></u>	<1	<1	<1	<1	<1

A312 Table 3

1 abic 0
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Halocarbons (Chlorinated Compounds)
207 Telegraph Drive, Gastonia, North Carolina

Sample Location	Date of Sampling	Dilution Factor	1,1-Dichloro- ethene (1,1-DCE) CASS 75-35-4 ppb	1,1-Dichloro- ethane (1,1-DCA) CAS# 75-34-3 ppb	1,1,1-Trichloro- ethane (1,1,1-TCA) CAS# 71-55-6 ppb	1,2-Dichloro- propane CASN 78-87-5 ppb	Trichloro- ethene (TCE) CAS# 79-01-6 ppb	Tetrachloro- ethene (PCE) CASN 127-18-4 ppb	cis-1,2-Dichloro- ethene (cis-1,2-DCE) CASN 156-59-2 ppb	trans-1,2-Dichloro- ethens (trans-1,2-DCE) CASS 156-60-5 ppb	Carbon Tetrachloride CAS# 56-23-5 ppb 0.3	Chloroforr CAS# 67-66- ppb
	er Quality Standard		7	6	200	0,6	3	0.7				
EW-4	7/31/1997	2,5	15	13	6.6	< 2.5	72	4.8	32 16	< 2.5 < 1	< 2.5 < 1	<u><1</u> <1
	8/22/1997 12/30/1998	1	6	11 <1	2.6 <1	<1	31 17	1.8 <1	16	<1	<1	<u> </u>
	3/19/1999	1	<1	<1	<1	<1	4	<1	3	<1	<1	<u> </u>
	5/19/1999	1	9	1	<1	<1	17	<1	15	<1	<1	<1
	9/20/1999	1	9.7	<1	<1	<1	26	<1	3,3	<1	<1	<1
	12/20/1999	1	6.4	<1	<1	<i< td=""><td>20</td><td><1</td><td>28</td><td>1,7</td><td><1</td><td><1</td></i<>	20	<1	28	1,7	<1	<1
	3/21/2000	1	13	1.7	<1	<1	15	<1	16	<1	<1	<l< td=""></l<>
	6/14/2000	1	17	2.9	<1	<1	16	<1	20	<1	<1	<1
	9/8/2000	1	14	1.3	<1	<1	14	<1	19	<1	<1	<1_
	9/8/2000	1	14	1.3	<1	<1	14	<1	19	<1	<1	<1
	12/1/2000	1	18	3,3	<1	<1	11	<1	12	<1	<1	<1
	2/21/2001	1	32	<1	<1	<1	16	<1	25	<1	<1	<1
	6/5/2001	1	38	1.4	<1	<1	15	<1	18	<1	<1	<1
	8/23/2001	1	38	<1	<1	<1	12	<1	25	<1	<1	<u><1</u>
	3/14/2002		25	<1	<1	<1	6.6	<1	11	<1	<1	<1
	10/2/2002	1	77	1.3	<1	<1	16	<1	28	<1	<1 <1	< <u> </u> <
	3/25/2003	1	49	<1	<1	<1	12 11	<1 <1	17 17	<1	<1	<1 <1
	9/23/2003	1	58 49	1.1 <1	<1	<1 <1	9.2	<1	14	<1	<1	<1
	10/1/2004	1	64	<1 <1	<1	<1	9.2	<1	20	<1	<1	1.6
	6/23/2005	1	37	<1	<1	<1	9.7	<1	14	<1	<1	< 200
	9/28/2005	1	48	<1	<1	<1	9,7	<1	16	<1	<1	<100
	3/10/2006	1	54	<1	<1	<1	11	<i< td=""><td>23</td><td><<u>l</u></td><td><1</td><td><l< td=""></l<></td></i<>	23	< <u>l</u>	<1	<l< td=""></l<>
	10/16/2006	1	15	<1	<1	<l< td=""><td>4</td><td><l< td=""><td>4.9</td><td><1</td><td><1</td><td><1</td></l<></td></l<>	4	<l< td=""><td>4.9</td><td><1</td><td><1</td><td><1</td></l<>	4.9	<1	<1	<1
	3/11/2008	1	23	<1	<1	<1	7.2	<1	11	<1	<1	<1
	9/23/2008	1	23	<1	<1	<1	8,8	<1	14	<1	<1	<1
	4/30/2009	1	8	<1	<1	<1	1.3	<1	1.6	<1	<1	<1
	9/17/2009	11	17	<l< td=""><td><1</td><td><1</td><td>6.2</td><td><1</td><td>9.7</td><td><<u>1</u></td><td><1</td><td><1</td></l<>	<1	<1	6.2	<1	9.7	< <u>1</u>	<1	<1
	374/2010	11	13	<l< td=""><td><1</td><td><1</td><td>5.1</td><td><1</td><td>7.3</td><td><1</td><td><<u>l</u></td><td><1</td></l<>	<1	<1	5.1	<1	7.3	<1	< <u>l</u>	<1
	1000000	1	12	<1	<1	<1	3.8	<1	6.5	<i< td=""><td><1</td><td><1</td></i<>	<1	<1
EW-5	8/1/1997	200	< 200	< 200	< 200	< 200	< 200	< 200	< 200	< 200	< 200	<100
	12/30/1998	100	<100	<100	<100	<100	<100	<100	<100	<100	<100	<10
	3/19/1999	100	<100	<100	<100	<100	<100	<100	<100	<100	<100	<40 <40
	5/19/1999	10	<10	<10	<10	<10	<10	<10 <40	<10 <40	<10 <40	<10 <40	<10
	9/20/1999	40	<40	<40	<40 <40	<40 <40	<40 <40	<40	<40	<40	<40	<10
	12/20/1999	40	<40 <10	<40 <10	<10	<10	<10	<10	<10	<10	<10	<10
	3/22/2000 6/14/2000	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/8/2000	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/8/2000	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	12/1/2000	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	2/21/2000	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	6/5/2001	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	8/30/2001	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	3/14/2002	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	10/2/2002	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	3/25/2003	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	9/23/2003	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	10/1/2004	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	< 10
	1/17/2005	10	<10	<10	<10	<10	<10	<10	<10	<10	<10	<10
	6/23/2005	10	<10	<10	<10	<10	<10 < 10	<10 <10	<10 < 10	<10 <10	<10 < 10	< 10 < 10
	9/28/2005	10	< 10	< 10	< 10 <10	< 10 <10	<10	<10	<10	<10	<10	<10
	3/10/2006	10	<10	<10 <10	<10	<10	<10	<10	<10	<10	<10	<10
	10/13/2006 3/9/2007	10	<10 < 1	<10 <1	<10	<10	2.8	<10	3.3	<1	<10	<1
	3/9/2007	1/10	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	9/23/2008	1	<1	<1	<1	<1	<1	<1	<1	<1	<1	<1
	4/30/2009	1/10	<1	<1	<1	<1	1.2	<1	<1	<1	<1	<1
	9/18/2009	1/10	<1	<1	<1	<1	1,3	<1	<1	<1	< 1	< 1
	7,74200	1/10	<1	<1	<1	< 1	<1	<1	<1	<1	<1	< 1

NOTES:

Sample results are expressed as a concentration or as less than the quantitation limit (e.g. <1).

The quantitation limit is equal to the dilution factor multiplied by the undiluted quantitation limit.

J = Result is detected below the reporting limit or is an estimated concentration because the result fell outside of the calibration range of the analytical instrument.

CAS# = Chemical Abstracts Service registry number assigned by the American Chemical Society to compounds included in the CAS Registry System.

Bold text indicates an exceedance of a NC Groundwater Quality Standard

NS = Well not sampled due to insufficient water column

NA = Sample not analyzed for this compound.

Table 4

Groundwater Analytical Data Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds) 207 Telegraph Drive, Gastonia, North Carolina

207 Telegraph l	Drive, Gastonia, l	North Car	olina	T				1	
Sample Location	Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropyl- ether (IPE) CAS# 108-20-3 ppb
NC Groundwa	ter Quality Stand	lard	1	50	600	600	500	20	70
MW-1A	2/8/1994	1	<1	< 1	<1	< 1	<1	<1	<1
	2/8/1994	1	<1	<1	< 1	<1	<1	<1	<1
	9/19/1996	1	<1	<1	<1	< 1	< 1	<1	<1
MW-1B	1/18/1995	1	< 1	<1	<1	< 1	< 1	<1	< 1
	9/18/1996	1	< 1	< 1	<1	< 1	< 1	<1	< 1
MW-3	2/8/1994	1	< 1	< 1	< 1	< 1	<1	<1	< 1
	9/18/1996	1	< 1	< 1	< 1	< 1	<1	< 1	< 1
MW-4A	2/7/1994	1	<1	< 1	<1	<1	< 1	<1	<1
	7/25/1994	1	< 0.5	< 0.5	< 0.5	< 0.5	3.2	NA NA	NA
	9/18/1996	1	< 1	<1	<1	< 1	<1	<1	< 1
MW-4B	1/17/1995	1	<1	<1	< 1	<1	<1	<1	< 1
	9/18/1996	1	< 1	<1	<1	<1	<1	<1	< 1
MW-5	2/8/1994	1	< 1	<1	<1	< 1	< 1	<1	< 1
MW-6	2/8/1994	1	<1	<1	<1	<1	<1	6	< 1
	9/19/1996	_1	<1	<1	<1	<1	< 1	<1	< 1
	8/1/1997	1	<1	<1	< 1	<1	< 2	4.4	<1
MW-7A	2/8/1994	1	<1	<1	<1	<1	< 1	<1	< 1
	9/17/1996	., 1	<1	<1	< 1	< 1	<1	< 1	<1
MW-7B	9/17/1996	1	<1	< 1	< 1	1.3	6.6	< 1	<1
	8/12/1997	25	< 25	NA	< 25	< 25	< 25	< 25	NA
MW-8	2/8/1994	1	<1	<1	< 1	< 1	< 1	<1	< 1
	9/17/1996	1	<1	<1	<1	< 1	<1	< 1	< 1
	9/29/2003	1	<1	<1	<1	< 1	<3	<5	<5
MW-10A	2/9/1994	50	3,400	< 50	540	2,000	3,900	2,500	520
	7/27/1994	100	2,700	< 100	770	1,600	4,300	3,300	1,300
	11/3/1994	25	1,800	< 25	440	1,500	3,100	2,100	580
	1/17/1995	50	3,000	< 50	560	1,600	3,800	5,000	510
	4/17/1995	100	2,100	< 100	280	820	2,200	1,800	250
	7/20/1995	50	4,400	< 50	610	1,700	5,100	4,200	570
	10/18/1995	100	2,300	< 100	1,100	1,200	6,400	680	990
	1/25/1996	100	7,500	< 100	3,000	10,000	32,000	1,400	4,200
	9/18/1996	10	2,440	< 10	904	137	7,360	2,710	437
	3/24/1997	100	3,000	<1	550	620	4,000	2,300	<1
	6/26/1997	50	4,200	<1	670	710	5,400	3,400	550
	9/30/1997	50	2,900	<1	110	270	1,800	1,800	94
	11/7/1997	50	1,300	<1	<50	<50	560	1,900	<50
MW-10B	9/9/1994	1	45	<1	<1	3	2	77	2
	11/4/1994	5	360	<5	<5	47	40	370	20
	1/18/1995	10	330	< 10	< 10	16	< 10	540	18
MW-11	2/9/1994	1	2	<1	<1	<1	<1	<1	<1
	2/9/1994	1	<1	<1	<1	<1	<1	<1	<1
	7/27/1994	1	<1	<1	<1	<1	<1	<1	<1
	11/3/1994	1	<1	<1	<1	<1	<1	<1	<1
	1/18/1995	1	<1	<1	<1	<1	<1	1	<1
	11/27/1995	1 1	<1	<1	<1	<1	<1	<1	<1 <1
	1/25/1996	1	<1	<1	<1	<1	<1	<1	<1
	9/18/1996	1	<1	<1	<1	<1	<1	<1 <1	<1
	3/24/1997	1	<1	<1	<1	<1	<1	<1	<1 <1
	6/26/1997	1,	<1	<1	<1 <1	<1 <1	<1 <1	<1	<1
	9/30/1997	1	<1	<1 <1	<1	<1	<1	<1	<1
	1/7/1997	1	<1	<u></u>	<u></u>	~1	~1	71	-1

Table 4 Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds) 207 Telegraph Drive, Gastonia, North Carolina

o relegraph	Drive, Gastonia,	Horia Car	oma	Т	T		1	tort Dutul	1
Sample	Date of	Dilution	Benzene CAS# 71-43-2	Chloro- benzene CAS# 108-90-7	Ethyl- benzene CAS# 100-41-4	Toluene CAS# 108-88-3	Xylenes (total) CAS#a	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4	Diisopropyl- ether (IPE) CAS# 108-20-3
Location VC Groundwa	Sampling ater Quality Stand	Factor	ppb 1	ppb 50	ppb 600	ppb 600	ppb 500	ppb 20	ppb
MW-12	2/10/1994	1	8	<1	1	15	4		
MW-12	7/28/1994	2	46	<2	<2	40	<2	11	25 80
	11/4/1994	5	16	<5	<5	10	< 5	97	38
	1/18/1995	1	11	<1	<1	6	5	32	28
	4/17/1995	1	18	<1	1.4	16	17	80	44
	7/20/1995	1	19	<1	<1	9.3	8.5	16	35
	10/18/1995	25	190	<25	63	130	< 25	940	130
	1/25/1996	1	19	<1	<1	6.1	15	8.8	16
	9/18/1996	1	1.3	<1	<1	<1	<1	<1	<1
	3/24/1997	1	<1	<1	<1	1.4	2.6	<1	<1
	6/26/1997	1	<1	<1	<1	<1	<1	<1	<l< td=""></l<>
	9/30/1997	1	1.1	<1	<1	<1	<l< td=""><td><1</td><td>1.7</td></l<>	<1	1.7
	11/7/1997	1	4.1	<1	<1	1,5	4	<1	6.4
MW-13	2/9/1994	1	1	<1	<1	<1	<1	5	<1
	7/27/1994	1	<1	<1	<1	<1	<1	13	<1
	11/1/1994	1	<1	<1	<1	<1	<1	<1	<1
	1/18/1995	1	<1	<1	<1	<1	<1	<1	<1
	1/18/1995	1	<1	<1	<1	<1	< 1	<1	<1
	4/17/1995	1	<1	<1	<1	<1	< 1	4.2	<1
	7/20/1995	1	<1	<1	<1	<1	<1	<1	<1
	11/27/1995	1	1.7	<1	<1	< 1	< 1	<1	<1
	1/25/1996	250	< 250	< 250	< 250	< 250	< 250	< 250	< 250
	9/18/1996	1	<1	<1	<1	<1	<1	<1	<1
	3/24/1997	1	< 1	<1	<1	<1	<1	<1	<1
	6/26/1997	1	< 1	<1	< 1	<1	<1	< 1	<1
	9/30/1997	11	<1	<1	<1	<1	<1	3	<1
	11/7/1997	1	<1	<1	<1	<1	<1	1.6	<1
	9/2/1998	1	3	<1	3	7	19	<5	<5
	9/20/1999	1	2.4	<1	<1	<1	<3	<5	<5
	9/8/2000	1	<1	<1	<1	<1	<3	<5	<5
	8/28/2001	1	<1	<1	<1	<1	<3 →	5.6	<5
•	10/1/2002	1	58	<1	<1	<1	<3	89	<5
	9/29/2003	10	1200 870	<10 <10	<10	<10	38	1800	<50
	10/1/2004	10		İ	<10	<10	92 190	480	<50
	10/3/2003	10/100	640 1400	< 10 < 10	< 10 < 10	< 10 < 10	99	720	< 50 < 50
	9/21/2007	10/100	2700	<10	<10	<10	61	1200	43 J
	9/22/2008	10/100	2100	<10	<10	11	70	490	< 50
	9/16/2009	1	85	<1	<1	2.1	4.7	49	< 5.0
	Section to the	1/10	130	<1	9.3	<1	9.2	48	< 5.0
MW-14	2/9/1994	1	84	<1	<1	2	14	78	<1
	7/27/1994	10	280	< 10	< 10	< 10	77	230	< 10
	11/3/1994	25	120	< 25	< 25	< 25	220	<25	< 25
	1/17/1995	2	170	< 2	< 2	3	50	17	130
	4/17/1995	1	86	<1	<1	1.3	27	78	<1
	7/20/1995	5	180	< 5	< 5	5,5	42	240	< 5
	11/27/1995	5	190	< 5	< 5	< 5	56	260	< 5
	3/7/1996	5	230	< 5	< 5	< 5	68	170	< 5
	9/18/1996	1	194	< 1	<1	<1	67.3	95	<1
	6/26/1997	1	59	<1	<1	<1	14	8.5	<1
	9/30/1997	1	22	<1	<1	<1	4.3	4	<1
	11/7/1997	1	<1	<1	<1	<1	<i< td=""><td>74</td><td><1</td></i<>	74	<1

Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells

Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)

207 Telegraph Drive, Gastonia, North Carolina

Sample Location NC Groundwa	Date of Sampling ter Quality Stand	Dilution Factor	Benzene <i>CAS# 71-43-2</i> ppb 1	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb 600	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb 20	Diisopropylether (IPE) CAS# 108-20-3 ppb 70
MW-15A	2/7/1994	1	<1	<1	< 1	<1	<1	<1	<1
	7/27/1994	1	2	<1	2	15	16	2	< 1
	1/16/1995	1	<1	<1	<1	<1	<1	<1	<1
	9/18/1996	10	< 10	< 10	< 10	< 10	< 10	< 10	< 10
	9/18/1998	1	< 1	<1	<1	<1	< 3	< 5	< 5
	9/24/1999	1	<1	<1	<1	<1	∢	<5	<5
	9/11/2000	1	<1	<1	<l< td=""><td><1</td><td>⊲</td><td><5</td><td><5</td></l<>	<1	⊲	<5	<5
	8/28/2001	1	<1	<1	<1	<1	⊲	<	<5
	10/1/2002	1	<1	<1	<1	<1	∢	<5	<5
	9/29/2003	1	<1	<1	<1	<1	⋖₃	<5	<5
	10/1/2004	1	<1	<1	<1	<1	⋖₃	<5	<5
	9/30/2005	1	0.7	< 1	0.81 J	2.4	4	< 5	< 5
	10/20/2006	1	<0.5	<1	<1	<1	⊲	<5	<5
	9/21/2007	1	9.7	<1	<1	<1	<1	19	<5
	9/22/2008	1	< 0.5	<1	<1	< <u>i</u>	♡	<5	<5
	9/15/2009	1	< 0.5	<1	<1	<1	⋖₃	<5	<5
	5/10/000	1	< 0.5	<1	<1	<1	⋖	<5	<5
MW-15B	2/7/1994	200	< 200	< 200	< 200	< 200	< 200	< 200	< 200
	2/7/1994	25	< 25	< 25	< 25	< 25	<25	< 25	< 25
	7/27/1994	25	< 25	< 25	< 25	< 25	< 25	< 25	< 25
	7/27/1994	25	170	< 25	< 25	< 25	52	590	< 25
	1/16/1995	5	< 5	< 5	< 5	< 5	< 5	< 5	< 5
	9/18/1996	20	< 20	< 20	< 20	< 20	< 20	< 20	< 20
	9/18/1998	1	2	<1	< 1	<1	< 3	< 5	< 5
	9/24/1999	1	<1	<1	<1	<1	<3	<5	<5
	9/18/2000	1	<1	<1	<1	<1	<3	্ত	<5
	8/28/2001	1	<1	<1	<1	<1	⊲3	<5	<5
	10/1/2002	1	<1	<1	<1	<1	<3	<5	<5
	9/29/2003	1	<1	<1	<1	<1	<3	<5	<5
	9/30/2004	1	<1	<1	<1	<1	⊲3	<5	<5
	10/18/2005	1	< 0.5	<1	<1	<1	<3	< 5	< 5
	10/20/2006	1	< 0.5	<1	<1	<1	<3	< 5	< 5
	9/21/2007	1	0.91 J	<1	<1	<1	<1	9,5	< 5
		1	< 0.5	<1	<1	<1	<1	<5	< 5
	9/23/2008	1	< 0.5	<1	<1	<1	- 3	< 5	< 5
	9/15/2009	1	< 0.5	<1	<1	<1	3	< 5	< 5
102 150				<1	<1	<1	<1	<1	<1
MW-15C	1/2/1994	1	<1	<1	<1	<1	<1	2	1
	1/17/1995		<u> </u>	<1	<1	<1	<1	<1	<1
	9/18/1996	1	<1	NA NA	<1	<1	4.2	<1	NA NA
	8/12/1997	1		NA <1	<1	<1	< 3	<5	<5
	9/18/1998	1	<1	<1	<1	<1	<3	<s< td=""><td><5</td></s<>	<5
	9/24/1999	1 1	<1				3	<5	<5
	9/18/2000	10	<1	<10	<10	<1	<30	<50	<50
	8/28/2001	10	<10	<10	<10	<10		<5	<5
	10/3/2002	1	4.7	<1	<1	<1	♂	<5	<5
	9/26/2003	1	<1	<1	<1	<1		Î	<5
	9/30/2004	1	3.3	<1	<1	<1	<3	<5	
	10/3/2005	1	4.5	<1	<1	<1	<3	< 5	< 5
	10/18/2006	1	4.1	< 1	<1	<1	<3	< 5	< 5
	9/24/2007	1	3,6	<1	<1	<1	<3	37	< 5
	9/25/2008	1/10	2.3	<1	<1	<1	< 3	< 5	< 5
	9/15/2009	1	<0.5	<1	<1	<1	< 3	< 5	< 5
	表的意思的意思	1/10	<0.5	<1	<1	<1	< 3	< 5	< 5

A3|16

Table 4
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)
207 Telegraph Drive, Gastonia, North Carolina

Sample Location NC Groundwa	Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb 600	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropylether (IPE) CAS# 108-20-ppb 70
MW-15D	11/4/1994	1	1	< 1	<1	1	< 1	4	<1
	11/4/1994	1	<1	<1	<1	<1	<1	3	<1
	1/17/1995	i	<1	<1	<1	<1	<1	11	<1
	9/18/1996	1	<1	<1	<1	<1	<1	4.2	<1
	8/12/1997	1	<1	NA NA	<1	<1	<1	7	NA NA
	9/22/1998	1	<1	<1	<1	<1	< 3	< 5	
	9/24/1999	1	<1	<1	<1	<1 <1	<3	<5	< 5 < 5
	9/18/2000	1	<1	<1	<1	<1	<3	5.4	<5
	8/28/2001	1	<1	<1	<1	<1	<3	<5	- √ -<5
	10/3/2002	1	<1	<1	<1	<1	<3	<5	
	9/26/2003	1	<1	<1	<1				
			<1			<1	3	7.1	<5
	9/30/2004	1		<1	<1	<1	<3	3.0 J	<5
	10/3/2005	1	< 0.5	<1	<1	1.2	< 3	< 5	< 5
	10/20/2006	1	< 0.5	<1	<1	1.2	< 3	2.2	< 5
	9/25/2007	1	<1	<1	</td <td><1</td> <td>< 3</td> <td><5</td> <td><5</td>	<1	< 3	<5	<5
	9/26/2008	1	<1	<1	<1	<1	< 3	3.1 J	<5
	9/18/2009	1	<0.5	<1	<1	<1	< 3	3.1 J	<5
	9/29/2010	1	<0.5	<1	<1	<1	< 3	3.2 J	<5
MW-16A	1/18/1995	1	< 1	< 1	< 1	<1	<1	<1	<1
	9/19/1996	100	< 100	< 100	< 100	< 100	< 100	< 100	< 100
MW-16B	1/18/1995	1	<1	< 1	<1	<1	< 1	<1	<1
	9/19/1996	1	<1	< 1	< 1	< 1	<1	< 1	<1
MW-17A	2/7/1994	1	<1	<1	<1	< 1	< 1	< i	<1
	1/18/1995	1	<1	< 1	<1	< 1	<1	<1	<1
	9/19/1996	1	<1	<1	< 1	< 1	<1	<1	< 1
MW-17B	2/7/1994	20	< 20	< 20	< 20	< 20	< 20	< 20	1,700
	1/16/1995	1	<1	< 1	< 1	< 1	<1	< 1	<1
	9/19/1996	500	< 500	< 500	< 500	< 500	< 500	< 500	< 500
·	9/19/1996	500	< 500	< 500	< 500	< 500	< 500	< 500	< 500
MW-17C	1/17/1995	1	<1	<1	<1	< 1	<1	< 1	<1
	9/19/1996	1	< 1	< 1	<1	< 1	<1	1.4	< 1
	8/12/1997	1	< 1	NA	<1	< 1	<1	10	NA
	3/27/1998	1	<1	<1	<1	<1	<1	<1	<1
	9/14/1998	1	<1	<1	<1	<1	<3	<5	<5
	9/16/1999	1	2,6	<1	<1	<1	<3	5,6	<5
	3/23/2000	40	<40	<40	<40	<40	<100	<200	<200
	2/28/2001	1	1.1	<1	<1	<1	⊲3	7.7	<5
	8/30/2001	1	<1	<1	<1	<1	<3	<5	<5
	3/28/2002	1	<1	<1	<1	<1	3	<5	<u>√</u> <5
	9/30/2002	1	1.4	<1	<1	<1	3	6.2	<5
ì	3/25/2003	1	1.4	<1	<1	<1	3	6.0	্
ŀ	9/25/2003	1	<1	<1	<1	<1	3	5.9	<u> </u>
	9/29/2004	1	<1	<1	<1	<1			
Ì							3	5.7	<5
ŀ	1/14/2005	1	<1	<1	<1	<1	<3	7.0	<5
	6/22/2005	10	<10	<10	<10	<10	<10	<10	<10
}	9/28/2005	1	0.63	<1	<1	<1	< 3	5.8	< 5
}	3/9/2006	. 1	< 0.5	<1	<1	<1	< 3	6.6	< 5
}	10/18/2006	1	< 0.5	<1	<1	<1	<3	4.2	< 5
ŀ	3/8/2007	1	< 0.5	<1	<1	<1	< 3	3.9	< 5
}	9/21/2007	1	<1	<1	<1	<1	⊲	7.4	< 5
-	3/12/2008	1	< 0.5	<1	<1	<1	⋖	5.4	< 5
1	9/22/2008	1	<0.5	<1	<1	<1	_ ⊲	6.7	< 5
ļ	4/30/2009	1	<0.5	<1	<1	<1	<3	4.9 J	< 5
ļ	9/16/2009	1	<0.5	<1	< 1	<1	⋖₃	4.3 J	< 5
	September 1	1	<0.5	<1	<1	<1	⊲	4.2 J	< 5
	1997 (1093) (10 9 8)	1	<0.5	< 1	< 1	<1	<3	5.3	< 5

Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)
207 Telegraph Drive, Gastonia, North Carolina

	Drive, Gastonia, I	l car		Chloro-	Ethyl-		Xylenes	tert-Butyl methyl ether	Diisopropyl-
Sample	Date of	Dilution	Benzene CAS# 71-43-2	benzene CAS# 108-90-7	benzene CAS# 100-41-4	Toluene CAS# 108-88-3	(total) CAS#a	(MTBE) CAS# 1634-04-4	ether (IPE) CAS# 108-20-3
Location NC Crowndays	Sampling ter Quality Stand	Factor	ppb 1	ppb 50	ppb 600	ppb 600	ppb 500	ppb 20	ppb 70
		1					< 1	<1	<1
MW-18A	2/9/1994	1	<1	<1	<1	<1	<1	<1	<1
	1/19/1995	1	<1	<1	<1	<1	<1	<1	<1
	9/17/1996	1	<1	<1	<1	<1	<3	<5	<5
	9/5/1998	1	<1 <1	<1	<1	<1	7 7	<5	<5
	9/28/1999	1	<1	<1	<1	<1	⟨₹	<5	<5
	9/12/2000 8/29/2001	1	<1	<1	<1	<1	<3	<5	<5
		1	<1	<1	<1	<1	<3	<5	<5
	9/30/2002	1	<1	<1	<1	<1	⊲3	<5	<
	9/29/2003		<1	<1	<1	<1	⊲3	্	<5
	9/28/2004	1		<1	<1	<1	<3	< 5	< 5
	9/27/2005	1 .	< 0.5 < 0.5	<1	<1	<1	<3	< 5	<5
	9/26/2007	1	< 0.5 <1	<1	<1	<1	<3	< 5	<5
	9/25/2008	1	<0.5	<1	<1	<1	<3	< 5	<5
	9/17/2009	1	<0.5	<1	<1	<1	<3	<5	< 5
	9/17/2009	1	<0.5	<1	<1	<1	<3	< 5	<5
MW-18B	7/28/1994	1	<1	<1	<1	< 1	<1	<1	<1
MW-10D	1/20/1995	1	<1	<1	<1	<1	<1	3	<1
	9/17/1996	1	<1	<1	<1	<1	<1	<1	<1
	9/5/1998	1	<1	<1	<1	<1	<3	<5	<5
	9/28/1999	1	<1	<1	<1	<1	<3 :	<5	<5
	9/12/2000	1	<1	<1	<1	<1	<3	<5	<5
	8/29/2001	1	<1	<1	<1	<1	⊲3	<5	<5
	10/4/2002	1	<1	<1	<1	<1	<3	<5	<5
	9/25/2003	1	<1	<1	<1	<1	∢	<5	<5
	9/28/2004	1	<1	<1	<1	<1	⋖₃	<5	<5
	9/26/2005	1	<0.5	<1	<1	<1	<3	<5	<5
	10/19/2006	1	<0.5	<1	<1	<1	<3	<5	<5
	9/26/2007	1	<1	<1	<1	<1	∢	<5	<5
	9/25/2008	1	<0.5	<1	<1	<1	⊲	<5	<5
	5,7,00,700	1	<0.5	<1	<1	<1	<3	<5	<5
MW-19A	2/7/1994	2	100	100	100	100	2	19	11
	7/26/1994	2	200	< 2	<2	< 2	28	65	<2
	9/14/1998	10	720	<10	<10	<10	100	410	<50
	9/20/1999	10	2900	<10	<10	29	580	2400	<50
	9/8/2000	10	910	<10	<10	<10	49	600	<50
	8/28/2001	10	300	<10	<10	<10	<30	160	<50
	10/1/2002	1	<1	<1	<1	<1	∢	<5	ব
	9/29/2003	1	5.9	<l< td=""><td><1</td><td><1</td><td>⊲</td><td>15</td><td><5</td></l<>	<1	<1	⊲	15	<5
	10/1/2004	1	9.2	<1	<1	<1	∢	8.4	<5
	10/3/2005	1	69	<1	<1	<1	<1	27	< 5
	10/17/2006	1	5.7	<1	<1	<1	< 3	3	< 5
	9/26/2007	1/10	600	<1	<1	4.1	15.5	370	15
	9/22/2008	1	1.4	<1	<1	<1	< 3	4.0 J	< 5
	9/17/2009	1	1.1	<1	<1	<1	< 3	< 5	< 5
	Secretaria de	1	1.7	<1	< 1	< 1	< 3	< 5	< 5

Semple	μ						•	nactive Water Supp s (Petroleum-Relat olina	Aromatic		fonitoring We ummary of De
New New	Diisopropyl- ether (IPE) CAS# 108-20- ppb	ethyl ether (MTBE) # 1634-04-4	m	(total) CAS#a	CAS# 108-88-3	benzene CAS# 100-41-4	benzene CAS# 108-90-7	Benzene CAS# 71-43-2	Dilution	Date of	Sample
10	70	20						1	ard	ter Quality Stand	NC Groundwa
326/1998	<1	65	<u> </u>	3	<1	19	<1	< 1	1	2/8/1994	MW-19B
\$2,000	< 10	120		< 10	< 10	< 10	< 10	13	10	7/27/1994	
920/1999 1	<1	31	_	3.3	<1	<1	<1		11	3/26/1998	
MW-20A 1 2.1	<5		+			-					
98,7000	<5		+-	· · · · · · · · · · · · · · · · · · ·							
31/2001	<5	1	+								
891/2001 1 2.4 < < < < < < < < <	<5		+								
3/28/2002	<5		1	T				•			
101/12002	< <u></u> <5 <5		1		·						
306/2003 1	<5		+								
P352003	<u> </u>										
Page Page	<5		+								
	<5										
MW-20A 9777996 1	<5		1								
Page 2005	<1		1								
MW-20A MW-21A MW-21A MW-22A M	< 5		1						1		
3/8/2007 1 <0.5 <1 <1 <1 <3 5.6 9/21/2007 1 <1 <1 <1 <1 <1 <1 <3 7.4 3/12/2008 1 1.7 <1 <1 <1 <1 <1 <3 7.4 3/12/2008 1 1.7 <1 <1 <1 <1 <1 <3 12 4/30/2009 1 4.9 <1 <1 <1 <1 <1 <3 11 9/15/2009 1 2 <1 <1 <1 <1 <3 9.7 3/30/308 1 1.5 <1 <1 <1 <1 <3 9.7 3/30/308 1 1.5 <1 <1 <1 <1 <3 9.8 3/30/309 1 1.5 <1 <1 <1 <1 <3 9.8 3/30/309 1 1.5 <1 <1 <1 <1 <3 9.8 3/30/309 1 1.5 <1 <1 <1 <1 <1 <3 9.8 3/30/309 1 1.5 <1 <1 <1 <1 <1 <1 <3 9.8 3/30/309 1 1.5 <1 <1 <1 <1 <1 <1 <1 <	< 5	9.2	1	< 3	<1	<1	<1	< 0.5	1	3/9/2006	
MW-20A MW-20A MW-21A MW-22A MW-22A MW-22A MW-22A MW-22B MW-23A M	< 5			< 3	<1	< 1	<1	1.4	1	10/17/2006	
MW-20A MW-21A MW-21A MW-22A MW-22A MW-22A MW-23A M	< 5	5.6		< 3	<1	<1	<1	< 0.5	1	3/8/2007	
9/23/2008	< 5	7.4		< 3	<1	< 1	<1	<1	1	9/21/2007	
AF90/2009	< 5	12		< 3	<1	< 1	< 1	1.7	ı	3/12/2008	
MW-20A MW-20B M	< 5	12		< 3	<1	<1	<1	4.1	1	9/23/2008	i
MW-20A 9/17/1996 1	< 5	11		< 3	<1	<1	< 1	4.9	1	4/30/2009	
MW-20A 9/17/1996 1	< 5	9.7	<u> </u>	< 3	<1	< 1	< 1	2	1	9/15/2009	
MW-20A 9/17/1996 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1 < 1	< 5	9.8	ļ	< 3	<1	< 1	<1	0.8	1	No. 2019/2010 (1970) A 1980 (1970) (1	
MW-20B	< 5	9.6	-	< 3	<1	< 1	<1	1.5	1	9/30/2010	
MW-20B 1/20/1995 1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	< 1	<1	ļ	2.1	2.1	<1	< 1	< 1			MW-20A
9/17/1996 1 <1 <1 <1 <1 <1 <1 <1	NA	<1	├	< 1	<1	< 1	NA	< 1	1	8/12/1997	
8/12/1997 1 <1 NA <1 <1 <1 <1 <1 <1 <1 <	< 1	<1	-			<1	< 1	< 1			MW-20B
MW-21 2/9/1994 1 <1	< 1									-7117777	
1/20/1995	NA		-			Î	i				
9/16/1996	<1		ļ. —						- 1		MW-21
8/13/1997 1 <1 NA <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <	<1		-								
MW-22A 1/19/1995 1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	<1										
9/16/1996 1 <1 <1 <1 <1 <1 <1 <2 <1 <2 <1 <1	NA										MW 22 *
MW-22B	<1		 								MW-22A
MW-22B 1/19/1995 1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	< 1 NA										ł
9/16/1996 1	<1										MW-22P
MW-23A 7/28/1994 250 9,400 < 250 1,400 18,000 9,400 < 250 11/4/1994 250 4,800 < 250	<1										141 44 -22D
11/4/1994 250 4,800 <250 800 9,300 5,100 3,100 9/18/1998 1 2,600 <1	< 250										MW-23A
9/18/1998 1 2,600 <1 <1 6,300 5,500 100 9/28/1999 1 <1	370							1			
9/28/1999 1 <1	56						Î				
9/13/2000 1 9,6 <1	<5										
8/28/2001 1 <1	<5										1
9/29/2003 100 1,000 <100 2,300 10,000 9,300 <500 9/29/2004 1 80 <1	<5						1				Ì
9/29/2003 100 1,000 <100 2,300 10,000 9,300 <500 9/29/2004 1 80 <1	<5								1		Ī
9/30/2005 1 7.3 <1	<500	<500		9,300	10,000	2,300	<100		100	9/29/2003	[
10/16/2006 1 39 <1	<5	<5		34	31		<1		1	9/29/2004	[
9/26/2007 1 110 <1 800 50 81 <5 9/25/2008 1 25 <1	< 5	< 5		4.4	3,4	2.4	<1	7.3	1	9/30/2005	
9/25/2008 1 25 <1 83 12 10 <5	< 5	< 5		65	49	190	<1	39	1	10/16/2006	[
	< 5	< 5		81	50	800	<1	110	1	9/26/2007	
	< 5	< 5		10	12	83	<1	25	1	9/25/2008	
9/18/2009 1/10 44 <1 750 230 620 <5	< 5	< 5		620	230	750	<1	44	1/10	9/18/2009	

Groundwater Analytical Data
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)
207 Telegraph Drive, Gastonia, North Carolina

207 Telegraph	Drive, Gastonia, I	TOTAL CALL	onna			T		I and Dute I	
Sample Location	Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb 600	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropylether (IPE) CAS# 108-20-3 ppb 70
· · · · · · · · · · · · · · · · · · ·	iter Quality Stand		ı ı	50	600				
MW-23B	7/28/1994	25	1,100	< 25	75	520	1,300	160	240
	11/4/1994	10	940	< 10	52	180	630	480	210
	9/21/1998	1	110	<1	5	18	9	200	17
	9/28/1999	1	<1	<1	<1	<1	<3	<5	<5
	9/12/2000	1	<1	<1	<1	<1	<3	<5	<5
	8/28/2001	1	<1	<1	<1	<1	<3	<5	<5
	9/30/2002	1	<1	<1	<1	<1	<3	<5	<5
	9/29/2003	1	29	<1	<1	9.1	32	<5	25
	9/29/2004	1	<1	<1	<1	<1	<3	<5	<5
	9/27/2005	1	< 0.5	<1	<1	<1	< 3	< 5	< 5
	10/17/2006	1	< 0.5	<1	< 1	<1	< 3	< 5	< 5
	9/26/2007	1	<1	<1	<1	<1	< 3	< 5	< 5
	9/25/2008	1	< 0.5	<1	<1	<1	< 3	< 5	< 5
	9/17/2009	1	< 0.5	<1	< 1	<1	< 3	< 5	< 5
	9/28/2010	1	1.4	< 1	< 1	<1	< 3	< 5	< 5
MW-24	9/17/1996	1	<1	<1	<1	<1	<1	< 1	<1
	9/17/1996	1	< 1	<1	< 1	<1	<1	<1	<1
MW-25	7/28/1994	500	8,000	< 500	1,600	22,000	11,000	2,500	< 500
	11/1/1994	250	6,000	<250	1,400	18,000	9,400	2,400	620
	9/2/1998	400	610	<400	2,000	7,800	10,000	<2,000	<2,000
	9/30/1999	10	160	<10	430	190	200	<50	77
	9/19/2000	11	190	<1	430	100	210	28	95
	8/29/2001	40	270	<40	2,900	2,400	8,500	<200	490
	10/1/2002	100	930	<100	3,100	5,900	15,000	<500	2,500
	9/29/2003	40	290	<40	2,700	2,900	10,000	<200	<200
	10/1/2004	40	65	<40	1,700	470	3,600	<200	<200
	9/30/2005	40	28	< 40	890	130	1,460	< 200	< 200
	10/18/2006	40	<20	< 40	1500	170	1990	< 200	< 200
	9/26/2007	40	31 J	< 40	1700	220	2530	< 200	< 200
	9/26/2008	. 10	59	< 10	780	67	790	< 50	<50
	9/17/2009	10	32	< 10	1200	81	1450	< 50	<50
	\$2,000,000	10	19	< 10	660	46	405	< 50	<50
MW-26	7/28/1994	500	2,400	< 500	2,000	27,000	13,000	< 500	< 500
	11/4/1994	250	1,100	< 250	1,500	15,000	10,000	1,300	890
	9/19/1998	100	< 100	< 100	1,900	2,300	10,000	< 500	< 500
	9/30/1999	10	<10	<10	1,200	1,600	6,500	77	450
	9/13/2000	10	16	<10	1,400	1,000	5,900	<50	540
	8/28/2001	10	16	<10	750	410	2,000	<50	370
	9/30/2002	10	20	<10	440	160	910	<50	83
	9/29/2003	10	<10	<10	860	160	2,400	<50	<50
	10/4/2004	1	<1	<1	<1	<1	5.4	<5	<5
	9/30/2005	1/10	< 0.5	<1	610	62	1,170	< 5	< 5
	10/17/2006	1/10	< 0.5	<1	700	42	1340	< 5	< 5
	9/25/2007	1/10	<1	<1	760	21	1220	< 5	< 5
	9/25/2008	1	9.9	<1	780	11	690	< 5	< 5
	9/17/2009	1/10	< 0.5	<1	390	8.6	600	< 5	< 5
	Sec. 1001/1010(2)	1/10	5.1	<1	170	5.1	74	21	< 5

Table 4
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)
207 Telegraph Drive, Castonia. Narth Carolina

Sample Location	Drive, Gastonia, 1 Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropyl- ether (IPE) CAS# 108-20-3 ppb
NC Groundwa	ter Quality Stand	ard	1	50	600	600	500	20	70
MW-27	7/28/1994	25	63	< 25	130	450	1,800	55	120
	7/28/1994	25	68	< 25	160	520	2,000	65	140
	11/3/1994	5	46	< 5	55	150	490	73	48
	9/9/1998	1	2.0	<1	20	17	150	<5	7
	9/27/1999	1	1.0	<1	19	100	180	<5	24
	9/12/2000	1	2.7	<1	21	66	120	<5	13
	8/28/2001	1	<1	<1	7.9	16	37	<5	13
	9/30/2002	1	1.3	<1	<1	2.9	10	<5	<5
	9/29/2003	1	<1	<1	1.7	1.3	7.6	<5	<5
	9/29/2004	1	<1	<1	<1	<1	⋖₃	<5	<5
	9/28/2005	1	< 0.5	<1	<1	<1	4.8	< 5	< 5
	10/16/2006	1	< 0.5	<1	31	20	157	< 5	< 5
	9/25/2007	1	<1	<1	22	11	75	< 5	< 5
	9/26/2008	1	3.3	<1	31	25	131	< 5	< 5
	9/17/2009	1	< 0.5	< 1	<1	<1	<3	< 5	< 5
	10/1/2010	1	< 0.5	<1	< 1	<1	⋖₃	< 5	< 5
MW-29	7/28/1994	1	<1	<1	<1	<1	< 1	<1	<1
	11/3/1994	1	3	<1	<1	<1	7	11	5
	1/19/1995	1	3	<1	< 1	1	4	<1	<1
	9/9/1998	1	<1	<1	<1	<1	Ø	<5	<5
	9/27/1999	1	<1	<1	<1	<]	⋖₃	<5	<5
	9/13/2000	1	<1	<1	<1	<1	⋖	<5	<5
	8/28/2001	1	<1	<1	<1	<1	⋖₃	<5	<5
	9/30/2002	1	<1	<1	<1	<1	⋖	<5	<5
	9/29/2003	1	<1	<1	<1	<1	⊲	<5	<5
	9/29/2004	1	<1	<1	<1	<1	⊲	<5	<5
	9/28/2005	1	< 0.5	<1	< 1	<1	< 3	< 5	< 5
	10/16/2006	1	< 0.5	< 1	< 1	<1	< 3	< 5	< 5
	9/25/2007	1	<1	<1	<1	<1	< 3	< 5	< 5
	9/26/2008	1	<0.5	<1	< 1	<1	< 3	< 5	< 5
	9/17/2009	1	<0.5	< 1	< 1	< 1	< 3	< 5	< 5
	10/17/610	1	<0.5	<1	< 1	<1	<3	< 5	< 5
MW-30	9/9/1994	10	320	<1	< 1	<1	< 1	410	18
	11/2/1994	5	350	< 5	< 5	16	16	170	23
	1/19/1995	5	310	< 5	< 5	8	7	330	16
	1/19/1995	5	320	<5	< 5	7	5	320	8

Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)

Sample Location	Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb 600	Toluene CAS# 108-88-3 ppb 600	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropyl- ether (IPE) CAS# 108-20-3 ppb
NC Groundwat	er Quality Stand	lard	1						
MW-31A	11/4/1994	25	1,100	< 25	78	980	1,100	520	110
	11/4/1994	25	1,300	< 25	92	1,100	1,200	470	160
	1/19/1995	25	1,100	< 25	34	830	1,100	< 25	96
	3/18/1998	11	73	<1	<1	1.3	<1	4.7	1.8
	5/27/1998	1	63	<1	<1	5	⊲	<5	7
	9/5/1998	1	42	<1	<1	2	<1	8	<1
	12/31/1998	1	5	<1	<1	<1	<3	15	<5
	9/20/1999	1	1	<1	<1	<1	_ ⊲	<5	<5
	12/16/1999	1	<1	<1	<1	<1	⊲	<5	<5
	3/23/2000	1	<1	<1	<1	<1	<3	<5	<5
	6/16/2000	1	<1	<1	<1	<1	<3	<5	<5
	9/7/2000	1	<1	<1	<1	<1	∢	<5	<5
	12/27/2000	1	1.1	<1	<1	<1	⊲	<5	<5
	2/28/2001	1	<1	<1	<1	<1	⋖	<5	<5
	6/7/2001	1	<1	<1	<1	<1	⊲	<5	<5
	8/31/2001	1	<1	<1	<1	<l< td=""><td>⊲</td><td><5</td><td><5</td></l<>	⊲	<5	<5
	3/15/2002	1	<1	<1	<1	<1	<3	<5	<5
	10/1/2002	1	<1	<1	<1	<1	<3	<5	<5
	3/26/2003	1	<1	<1	<1	<1	<3	<5	<5
	9/24/2003	1	<1	<1	<1	<1	⋖₃	<5	<5
	9/29/2004	1	<1	<1	<1	<1	<3	<5	<5
	1/17/2005	1	<0.5	<1	<1	<1	⋖₃	<5	<5
	6/23/2005	1	<1	<1	<1	<1	<1	<1	<1
	9/26/2005	1	<1	<1	<1	<1	⊲	<5	<5
	3/10/2006	1	<0.5	<1	<1	<1	⊲	<5	<5
	10/17/2006	1	<0.5	<1	<1	<1	<3	<5	<5
	3/8/2007	i	<0.5	<1	<1	<1	<3	<5	<5
	9/24/2007	1	<1	<1	<1	<1	<3	<5	<5
	3/10/2008	1	<1	<1	<1	<1	<3	<5	<5
	9/23/2008	1	<0.5	<1	<1	<1	<3	<5	<5
	4/30/2009	1	<0.5	<1	<1	<1	<3	<5	<5
	9/16/2009	1	<0.5	<1	<1	<1	<3	<5	<5
		1	<0.5	<1	<1	<1	<3	<5	<5
	9/28/2010		<0.5	<1	<1	<1	⊲	<5	<5
	isa mananan Mananan	1	<0.5	<1	<1	<1	3	<5	<5

Table 4

Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells

Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)

207 Telegraph Drive, Gastonia, North Carolina

207 Telegraph l	Drive, Gastonia, I	North Car	olina		1		,		
Sample Location	Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropyl- ether (IPE) CAS# 108-20-3 ppb
NC Groundwat	ter Quality Stand	lard	1	50	600	600	500	20	70
MW-31B	11/3/1994	1	< 1	<1	<1	<1	< 1	<1	< 1
	1/19/1995	1	2	<1	<1	<1	2	2	< 1
	3/26/1998	1	<1	<1	<1	<1	<1	<1	</td
	5/29/1998	1	<1	<1	<1	<1	- ⊲	<5	<5
	9/5/1998	1	<1	<1	<1	<1	∢	<5	<5
	12/31/1998	1	<1	<1	<1	<1	ಶ	<5	<5
	9/28/1999	1	<1	<1	<1	<1	⋖	<5	<5
	12/16/1999	11	<1	<1	<1	<1	⊲	<5	<5
	3/23/2000	1	<1	<1	<1	<1	<3	<5	<5
	6/16/2000	1	<1	<1	<1	<1	⋖	<5	<5
	9/7/2000	1	< <u>1</u>	<1	<1	<1	<3	<5	<5
	12/28/2000	1	<1	<1	<1	<1	<3	<5	<5
	2/28/2001	1	<1	<1	<1	<1	⋖₃	<5	<5
	6/7/2001	1	<1	<1	<1	<1	⋖₃	<5	<5
	8/31/2001	1	<1	<1	<1	<1	⋖₃	<5	<5
	3/15/2002	1	<1	<1	<1	<1	∢	<5	<5
	10/1/2002	1	<1	</td <td><1</td> <td><1</td> <td>⋖₃</td> <td><5</td> <td><5</td>	<1	<1	⋖₃	<5	<5
	3/26/2003	1	<1	<1	<1	<1	⋖	<5	<5 .
	9/24/2003	1	<1	<1	<1	<1	⋖₃	<5	<5
	9/28/2004	1	<1	<1	<1	<1	⊲	<5	<5
	1/17/2005	<0.5	<0.5	<1	<1	<1	⊲	<5	<5
	6/23/2005	1	<1	<1	<1	<1	<1	<1	<1
ļ	9/26/2005	1	<0.5	<1	<1	<1	∢	<5	<5
	3/10/2006	1	<0.5	<1	<1	<1	⊲	<5	<5
	10/16/2006	1	<0.5	<1	<1	<1	⋖	<5	<5
	3/8/2007	1	<0.5	<1	<1	<1	<3	<5	<5
	9/24/2007	1	<1	<1	<1	<1	⋖	<5	<5
	3/10/2008	1	<0.5	<1	<1	<1	<3	<5	<5
ļ	9/23/2008	1	<0.5	<1	<1	<1	<3	<5	<5
]	4/30/2009	1	<0.5	<1	<1	<1	<3	<5	<5
ļ	9/16/2009	1	<0.5	<1	<1	<1	⋖	<5	<5
ļ	9/28/2010	1	<0.5	<1	<1	<1	⋖	<5	<5
	3/24/2010	1	<0.5	<1	<1	<1	⋖	<5	<5
	9/28/2010	1	<0.5	<1	<1	<1	⋖₃	<5	<5

Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)
207 Telegraph Drive, Gastonia, North Carolina

Sample Location NC Groundwa	Date of Sampling ter Quality Stand	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb 50	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb 600	Xylenes (total) CAS#a ppb 500	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb 20	Diisopropyl- ether (IPE) CAS# 108-20-3 ppb
MW-32	11/4/1994	1	12	<1	<1	4	11	7	3
	1/19/1995	1	< 1	<1	<1	<1	<1	<1	<1
	3/18/1998	1	<1	<1	<1	<1	<1	<1	<1
	9/5/1998	1	<1	<1	<1	<1	<3	<5	<5
	9/29/1999	1	<1	<1	<1	<1	∢	<5	<5
	3/27/2000	1	<1	<1	<1	<1	♡	<5	<5
	9/13/2000	1	<1	<1	<1	<1	<3	<5	<5
	2/28/2001	1	<1	<1	<1	<1	∢	<5	<5
	8/31/2001	1	<1	<1	<1	<1	<3	<5	<5
	3/15/2002	1	<1	<1	<1	<1	<3	<5	<5
	10/1/2002	1	<1	<1	<1	<1	⊲	<5	<5
	3/26/2003	1	<1	<1	<1	<1	<3	<5	<5
	9/24/2003	1	<1	<1	<1	<1	<3	<5	<5
	9/28/2004	1	<1	<1	<1	<1	<3	<5	<5
	1/17/2005	1	<0,5	<1	<1	<1	<3	<5	<5
	6/23/2005	1	<1	<1	<1	<1	<1	<1	<1
	9/26/2005	1	<0.5	<1	<1	<1	⊲	<5	<5
	3/7/2006	1	<0.5	<1	<1	<1	<3	<5	<5
	10/17/2006	1	<0.5	<1	<1	<1	∢	<5	<5
	3/8/2007	1	<1	<1	<1	<1	<3	<5	<5
	9/24/2007	1	<1	<1	<1	<1	<3	<5	<5
	3/11/2008	1	< 0.5	<1	<1	<1	<3	<5	<5
	9/23/2008	1	<0.5	<1	<1	<1	♡	<5	<5
	4/30/2009	1	<0.5	<1	<1	<1	<3	<5	<5
	9/16/2009	1	<0.5	<1	<1	<1	⊲	<5	<5
	9/28/2010	1	<0.5	<1	<1	<1	⋖	<5	<5
		1	<0.5	<1	<1	<1	⋖	<5	<5
		1	<0.5	<1	<1	<1	⋖₃	<5	<5
MW-33	9/19/1996	1	<1	<1	<1	<1	< 1	<1	<1
MW-34A	9/16/1996	1	<1	<1	<1	< 1	<1	<1	<1
MW-34B	9/16/1996	1	<1	<1	<1	< 1	< 1	<1	<1
	9/18/2009	1	<0.5	<1	<1	< 1	< 3	< 5	< 5
MW-35	1/19/1995	1	<1	<1	<1	< 1	<1	<1	< 1
	9/19/1998	1	<1	<1	<1	< 1	<1	<1	<1
	9/28/1999	1	<1	<1	<1	<1	⋖₃	<5	<5
	9/13/2000	1	<1	<1	<1	<1	⋖₃	<5	<5
	8/29/2001	1	<1	<1	<1	<1	⋖₃	<5	<5
	10/1/2002	1	<1	<1	<1	<i< td=""><td>∢</td><td><5</td><td><5</td></i<>	∢	<5	<5
	9/29/2003	1	<1	<1	<1	<1	∢	<5	<5
	9/29/2004	1	<1	<1	<1	<1	⊲	<5	<5
	9/27/2005	1	< 0.5	<1	<1	<1	< 3	< 5	< 5
	10/16/2006	1	< 0.5	<1	<1	<1	< 3	< 5	< 5
	9/26/2007	1	<1	<1	<1	<1	< 3	< 5	<5
	9/26/2008	1	<0.5	<1	< 1	<1	< 3	< 5	<5
	9/18/2009	1	<0.5	<1	<1	<1	< 3	< 5	<5
	Aldino, des	1	<0.5	<1	<1	<1	< 3	< 5	<5

Table 4

Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)
207 Telegraph Drive, Gastonia, North Carolina

Sample Location NC Groundwa	Date of Sampling ter Quality Stand	Dilution Factor	Benzene CAS# 71-43-2 ppb 1	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb 600	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropylether (IPE) CAS# 108-20-3 ppb 70
MW-36	12/12/1995	1	<1	<1	<1			<1	<1
MW-30	3/7/1996	1	<1	<1	<1	<1 <1	<1 <1	<1	<1
	3/7/1996	1	<1	<1	<1	<1	<1	<1	<1
MW-37A	3/7/1996	1	<1	<1	<1	<1	<1	<1	<1
MW-37R	3/7/1996	i	4.1	<1	<1	<1	1.7	<1	<1
MW-38	8/13/1997	1	<1	NA NA	<1	<1	< 1	<1	NA NA
MW-39	8/1/1997	1	<1	NA NA	<1	<1	<1	<1	NA NA
IVI VV -39	8/12/1997	1	<1	NA NA	<1	<1	<1	<1	NA NA
	8/12/1997	1	<1	NA NA	<1	<1	< 1	1	NA.
MW-40	8/1/1997	1	<1	NA NA	<1	<1	<1	<1	NA NA
14114-40	8/12/1997	1	<1	NA NA	<1	<1	1.0	<1	NA NA
DM-7	2/8/1994	1	<1	<1	<1	<1	< 1	<1	<1
CP-1	2/10/1994	1	<1	<1	<1	<1	<1	<1	<1
C1-1	7/28/1994	1	<1	<1	<1	<1	<1	<1	<1
	3/18/1998	1	<1	<1	<1	<1	<1	<1	<1
	6/3/1998	1	<10	<10	<10	<10	<30	<50	<50
	8/27/1998	1	<1	<1	<10	<1	<3	<5	<5
	12/30/1998	1	<1	<1	<1	<1	∢ 3	<5	<u>√</u> <5
	3/19/1999	1	<1	<1	<1	<1	3	<5	<5
	5/19/1999	1	<1	5.0	<i< td=""><td><1</td><td>⟨3</td><td><5</td><td><u></u></td></i<>	<1	⟨3	<5	<u></u>
	9/20/1999	40	<40	<40	<40	<40	<100	<200	<200
	12/20/1999	40	<40	<40	<40	<40	<100	<200	<200
	3/22/2000	400	<400	<400	<400	<400	<1000	<2000	<2000
	6/14/2000	400	<400	<400	<400	<400	<1000	<2000	<2000
	9/8/2000	1	<1	<1	<1	<1	<3	<5	<5
	12/1/2000	1000	<1000	<1000	<1000	<1000	√3000	<5000	<5000
	2/21/2001	400	<400	<400	<400	<400	<1000	<2000	<2000
	6/5/2001	400	<400	<400	<400	<400	<1000	<2000	<2000
	8/23/2001	400	<400	<400	<400	<400	<1000	<2000	<2000
	3/14/2002	400	<400	<400	<400	<400	<1000	<2000	<2000
	10/2/2002	40	<40	<40	<40	<40	<100	<200	<200
	3/25/2003	40	<40	<40	<40	<40	<100	<200	<200
	9/23/2003	400	<400	<400	<400	<400	<1000	<2000	<2000
	10/1/2004	10	<10	<10	<10	<10	<30	<50	<50
	1/18/2005	10	<5	<10	<10	<10	<30	<50	<50
	5/27/2005	10	< 5	< 10	< 10	< 10	<30	< 50	< 50
	9/28/2005	10/100	< 5	< 10	< 10	< 10	< 30	< 50	< 50
	3/8/2006	10	< 5	< 10	< 10	< 10	<30	< 50	< 50
	3/6/2007	10	< 5	< 10	< 10	< 10	< 30	< 50	< 50
	9/24/2007	10	<10	< 10	< 10	< 10	< 30	< 50	< 50
	3/11/2008	10/100	<10	< 10	< 10	< 10	<30	< 50	< 50
	9/22/2008	10/100	<5	<10	<10	<10	<30	<50	<50
	4/30/2009	10	<5	<10	<10	<10	<30	<50	<50
	9/17/2009	10	<5	<10	<10	<10	<30	<50	<50
	Section 1	10/100	<5	<10	<10	<10	<30	<50	<50
	\$1/610/01106 B	10/100	<5	<10	<10	<10	<30	<50	<50

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Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells

Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)

207 Telegraph Drive, Gastonia, North Carolina

Sample Location	Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropyl- ether (IPE) CAS# 108-20-3 ppb
NC Groundwa	ter Quality Stanc	lard	1	50	600	600	500	20	70
CP-2	2/10/1994	1	< 1	< 1	< 1	< 1	2	<1	<1
WS-3	2/14/1992	1	< 5	NA	< 5	< 5	<1	NA	NA
	11/1/1993	1	. 86	NA	< 10	5.6	33	460	NA
	2/16/1994	5	190	< 5	< 5	< 5	6	290	< 5
	5/25/1994	2	11	< 2	< 2	< 2	< 2	110	2
	7/26/1994	1	2	< 1	<1	< 1	<1	48	<1
WS-6	5/26/1994	1	2	<1	<1	5	6	6	<1
WS-7	8/25/1994	5	240	< 5	15	230	300	400	50
	3/15/1996	1	<1	<1	<1	<1	<1	<1	<1
	3/15/1996	1	<1	<1	<1	< 1	<1	<1	<1
	1/16/1997	2	<1	< 1	<1	< 1	<1	180	<1
	3/3/1997	2	84	<1	< 1	3.3	3.2	210	<1
	4/18/1997	2.5	110	< 1	<1	< 1	<1	230	<1
	7/31/1997	4	140	NA	<4	< 4	< 4	190	NA
EW-2	3/18/1998	1	<1	<1	<1	<1	<l< td=""><td><1</td><td><1</td></l<>	<1	<1
	5/27/1998	1	<1	<1	<1	<1	ℴ	<5	<5
	8/27/1998	1	<1	<1	<1	<1	∢	<5	<5
	12/30/1998	1	<1	<1	<1	<1	<3	<5	<5
	3/19/1999	1	<1	<1	<1	<1	<3	<5	<5
	5/19/1999	1	<1	9	<1	<1	<3	<5	<5
	9/20/1999	1	<1	<1	<1	<1	∢	<5	<5
	12/20/1999	40	<40	<40	<40	<40	<100	<200	<200
	3/21/2000	1	<1	<1	<1	<1	<3	<5	<5
	6/14/2000	100	<100	<100	<100	<100	<300	<500	<500
	9/8/2000	100	<100	<100	<100	<100	<300	<500	<500
	12/1/2000	100	<100	<100	<100	<100	<300	<500	<500
	2/21/2001	40	<40	<40	<40	<40	<100	<200	<200
	6/5/2001	100	<100	<100	<100	<100	<300	<500	<500
	8/23/2001	100	<100	<100	<100	<100	<300	<500	<500
	3/14/2002	40	<40	<40	<40	<40	<100	<200	<200
	10/2/2002	100	<100	<100	<100	<100	<300	<500	<500
	3/25/2003	40	<40	<40	<40	<40	<100	<200	<200
	9/23/2003	400	<400	<400	<400	<400	<1000	<2000	<2000
	10/1/2004	10	<10	<10	<10	<10	<30	<50	<50
	1/17/2005	100	<50	<100	<100	<100	<300	<500	<500
	5/27/2005	100	< 50	< 100	< 100	< 100	< 300	< 500	< 500
	9/27/2005	100	< 50	< 100	< 100	< 100	< 300	< 500	< 500
	3/8/2006	100	< 50	< 100	< 100	< 100	< 300	< 500	< 500
	10/13/2006	100	< 50	< 100	< 100	< 100	<300	< 500	< 500
	3/6/2007	100	< 50	<100	< 100	< 100	<300	< 500	< 500
	9/24/2007	100	<100	< 100	< 100	< 100	<300	< 500	< 500
	3/11/2008	10/100	<5	<100	<100	<10	<30	<50	<10
			<5	<10	<10	<10	<30	<50	<50
	9/22/2008	10/100	<5	<10	<10	<10	<30	<50	<50
	4/30/2009	10/100		<10	<10	<10	<30	<50	<50
	9/17/2009	10	<5		<10	<10	<30	<50	<50
	No receive a	10 10	< <u></u>	<10 <10	<10	<10	<30	<50	<50

Table 4
Groundwater Analytical Data
Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells
Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)
207 Telegraph Drive, Gastonia, North Carolina

Sample Location NC Groundwa	Date of Sampling	Dilution Factor	Benzene CAS# 71-43-2 ppb 1	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb 600	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb 20	Diisopropylether (IPE) CAS# 108-20-3 ppb 70
EW-3	3/18/1998	1	<1	<1	<1	<1	<1	<1	<1
2.,, 3	5/27/1998	1	<1	<1	<1	<1	<3	<5	<5
	8/27/1998	1	<1	<1	<1	<1	3	<5	<5
	12/30/1998	1	<1	<1	<1	<1	<3	<5	<5
	3/19/1999	1	<1	<1	<1	<1	⟨3	<5	<5
	5/19/1999	1	<1	<1	<1	<1	⟨3	<5	<5
	9/20/1999	1	<1	<1	<1	<1	3	<5	
	12/20/1999	1	<1	<1	<1	<1	3	<5	<5
	3/21/2000	1	<1	<1	<1	<1	<3	<5	<5
	6/15/2000	1	<1	<1	<1	<1	⊲3	<5	<5
	6/5/2001	1	<1	<1	<1	<1	<3	<5	<5
	8/23/2001	1	<l< td=""><td><1</td><td><1</td><td><1</td><td><3</td><td><5</td><td><5</td></l<>	<1	<1	<1	<3	<5	<5
	3/14/2002	1	<1	<1	<1	<1	<3	<5	<5
	10/2/2002	1	<1	<1	<1	<1	<3	<5	<5
	3/25/2003	1	<1	<1	<1	<1	<3	<5	<5
	9/23/2003	1	<1	<1	<1	<1	⊲	<5	<5
	10/1/2004	1	<1	<1	<1	<1	<3	<5	<5
	1/17/2005	10	<5	<10	<10	<10	<30	<50	<50
	6/23/2005	1	<1	<1	<1	<1	<1	<1	<1
	9/27/2005	1	< 0.5	<1	< 1	<1	<3	< 5	< 5
	3/9/2006	1	< 0.5	<1	<1	<1	< 3	< 5	< 5
	10/13/2006	1	< 0.5	<1	< 1	<1	< 3	< 5	< 5
	3/6/2007	1	< 0.5	<1	< 1	<1	< 3	< 5	< 5
	9/24/2007	1	<1	< 1	< 1	<1	< 3	< 5	< 5
	3/11/2008	1/10	<0.5	<1	< 1	< 1	< 3	< 5	< 5
	9/22/2008	1	<0.5	<1	< 1	<1	< 3	< 5	< 5
	4/30/2009	1	<0.5	< 1	< 1	<1	< 3	< 5	< 5
	9/17/2009	1	<0.5	<1	<1	<1	< 3	< 5	< 5
	3/23/2000	1	<0.5	<1	< 1	<1	< 3	< 5	< 5
	9/10/2010	1	<0.5	< 1	<1	<1	< 3	< 5	< 5

Table 4

Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells

Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds)

Sample Location	Date of Sampling	Dilution Factor	Benzene <i>CAS# 71-43-2</i> ppb	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropyl- ether (IPE) CAS# 108-20-3 ppb
	ter Quality Stand	1	1	50	600	600	500	20	
EW-4	7/31/1997	2.5	8.1	< 2.5	< 2.5	< 2.5	< 2.5	6.4	< 2.5
	8/22/1997	1	8.7	NA	<1	<1	<1	6.1	NA
	12/30/1998	1	6	<1	<1	<1	<3	12	<5
	3/19/1999	1	6	< <u>l</u>	<1	<1	<3	<5	<5
	5/19/1999	1	8	<1	2	<1	<3	<5	<5
	9/20/1999	1	4.1	<1	<1	<1	<3	24	<5
	12/20/1999	1	5.5	<1	<1	<1	<3	32	<5
	3/21/2000	1	6	<1	<1	<1	<3	24	<5
	6/14/2000	11	9.5	<1	<1	<1	_ ⊲	25	<5
	9/8/2000	1	12	<1	<	<1	<3	27	<5
	12/1/2000	1	28	<1	<1	<1	⊲	46	<5
	2/21/2001	1	47	<1	<1	<1	⊲	63	<5
	6/5/2001	1	40	<1	<1	<1	<3	<5	<5
	8/23/2001	1	29	<1	<1	<1	<3	41	<5
	3/14/2002	1	18	<1	<1	<1	<3	26	<5
	10/2/2002	1	13	<1	<1	<1	⊲	19	<5
	3/25/2003	1	5.3	<1	<1	<1	<3	6.6	<5
	9/23/2003	1	3.4	<1	<1	<1	<3	7.1	<5
	10/1/2004	1	1.1	<1	<1	<1	<3	2.8J	<5
	1/17/2005	1	1.2	<1	<1	<1	<3	4.8	<5
	6/23/2005	1	<1	<1	<1	<1	<1	5.4	<1
	9/28/2005	1	3.1	<1	<1	< 1	< 3	7.5	< 5
	3/9/2006	1	3.1	<1	<1	< 1	< 3	12	< 5
	10/16/2006	1	7.3	<1	<1	<1	< 3	13	< 5
	3/11/2008	1	4.2	<1	< 1	<1	< 3	17	< 5
	9/23/2008	1	5	<1	<1	<1	< 3	20	< 5
	4/30/2009	1	3.1	<1	<1	<1	< 3	5.7	< 5
	9/17/2009	1	8.5	<1	< 1	<1	< 3	14	< 5
		1	6.7	<1	<1	<1	< 3	11	< 5
	Establish (Per	1	5.6	<1	<1	< 1	<3	9	< 5

Table 4

Groundwater Analytical Data

Monitoring Wells, Extraction Wells, and Inactive Water Supply Wells

Summary of Detected Purgeable Aromatics (Petroleum-Related Compounds) 207 Telegraph Drive, Gastonia, North Carolina

Sample Location IC Groundwa	Date of Sampling ter Quality Stand	Dilution Factor	Benzene CAS# 71-43-2 ppb 1	Chloro- benzene CAS# 108-90-7 ppb	Ethyl- benzene CAS# 100-41-4 ppb	Toluene CAS# 108-88-3 ppb	Xylenes (total) CAS#a ppb	tert-Butyl methyl ether (MTBE) CAS# 1634-04-4 ppb	Diisopropyl- ether (IPE) CAS# 108-20- ppb
EW-5	8/1/1997	200	1,300	NA	820	5,500	4,100	240	NA
211 3	12/30/1998	100	2,000	<100	460	3,100	2,800	<500	<500
	3/19/1999	100	1,700	<100	550	2,100	2,200	<500	<500
	5/19/1999	10	1,500	<100	460	1,400	1,700	460	290
	9/20/1999	40	800	<40	180	690	890	400	<200
	12/20/1999	40	890	<40	180	680	380	280	<200
	3/22/2000	10	1,000	<10	240	1,000	1,800	340	280
	6/14/2000	10	820	<10	410	830	1,900	99	380
	9/8/2000	10	580	<10	32	280	560	150	70
	12/1/2000	10	590	<10	100	320	640	210	130
	2/21/2001	10	<10	<10	47	240	520	220	79
	6/5/2001	10	490	<10	28	320	430	<50	<50
	8/30/2001	10	540	<10	24	200	330	110	<50
	3/14/2002	10	430	<10	47	240	470	120	<50
	10/2/2002	10	420	<10	100	160	290	98	<50
	3/25/2003	10	580	<10	840	1,100	2,300	54	51
	9/23/2003	10	580	<10	1,700	2,100	4,600	34 J	27 J
	10/1/2004	10	1,100	<10	640	950	950	340	33 J
	1/17/2005	10	1,300	<10	570	900	860	420	<50
	6/23/2005	10	790	<10	470	740	790	260	73
	9/28/2005	10	700	< 10	570	640	900	240	< 50
	3/9/2006	10	32	< 10	320	41	223	< 50	< 50
	10/13/2006	10	37	< 10	630	89	450	< 50	< 50
	3/9/2007	10	17	<1	180	89	137	< 5	< 5
	3/12/2008	10	44	<1	400	82	440	< 5	< 5
	9/23/2008	10	43	<1	830	38	438	< 5	< 5
	4/30/2009	1/10	12	< 1	170	9.8	187	< 5	< 5
	9/18/2009	1/10	18	<1	180	15	118	< 5	< 5
	1999	1/10	13	<1	240	16	108	< 5	< 5
	1710/1/2010	1/4	7.6	<1	150	8.2	67.8	< 5	< 5

Sample results are expressed as a concentration or as less than the quantitation limit (e.g. <1). The quantitation limit is equal to the dilution factor multiplied by the undiluted quantitation limit.

NA = Sample not analyzed for this compound.

CAS# = Chemical Abstracts Service registry number assigned by the American Chemical Society to compounds included in the CAS Paristre Service. included in the CAS Registry System.

aCAS#s: o-xylene = 95-47-6; m-xylene = 108-38-3; p-xylene = 106-42-3.

Bold text indicates an exceedance of a NC Groundwater Quality Standard

2010 data

Table 5
Summary of Parameters Measured During Groundwater Sampling Activities
207 Telegraph Drive, Gastonia, North Carolina

			Approximate	Field Parameters Measured at End of Well Purge Specific					
			Volume		7 70				
Sample Location	Sam Date	pling Time	of Purge (gallons)	pH (pH units)	Conductance (mS/cm)	Temperature (°C)			
MW-1A	09/30/10	15:10	6.50	5.80	0.110	19.90			
MW-1B	09/27/10	15:15	27.00	6.00	0.150	19.10			
MW-3	09/30/10	16:45	0.50	5.86	0.540	19.40			
MW-4A	09/27/10	17:10	3.62	5.88	0.290	19.40			
MW-4B	09/27/10	17:25	22.00	6.00	0.080	18.80			
MW-7A	09/30/10	9:45	2.25	6.20	0.330	18.90			
MW-7B	09/27/10	11:35	22.20	5.94	0.190	18.90			
MW-8	09/30/10	10:45	0.75	5.90	0.070	19.90			
MW-13	09/30/10	17:40	6.00	5.80	0.300	19.90			
MW-15A	09/30/10	13:15	4.50	5.60	0.230	20.40			
MW-15B	09/28/10	17:30	13.75	5.40	0.300	19.50			
MW-15C	09/28/10	15:40	15.00	5.70	0.150	19.00			
MW-15D	09/29/10	16:10	62.00	8.46	0.240	21.10			
MW-16A	09/27/10	10:25	22.50	5.80	0.130	20.90			
MW-16B	09/27/10	12:10	17.50	6.50	0.200	19.90			
MW-17A	09/29/10	11:20	11.75	4.90	0.680	17.60			
MW-17B	09/30/10	14:55	19.00	5.49	0.350	17.80			
MW-17C	09/30/10	16:00	8.50	7.68	0.310	18.10			
MW-18A	09/29/10	17:40	10.25	5.40	0.120	16.90			
MW-18B	09/29/10	15:45	25.00	7.30	0.200	17.90			
MW-19A	09/30/10	16:35	9.00	5.30	0.340	19.40			
MW-19B	09/30/10	14:00	11.25	7.54	0.370	18.90			
MW-20A	09/27/10	15:55	14.00	5.72	0.100	18.40			
MW-20B	09/28/10	13:00	14.00	6.40	0.180	18.70			
MW-21	09/28/10	16:20	9.00	6.03	0.120	18.70			
MW-22A	09/28/10	13:25	6.25	5.85	0. <u>140</u>	21.60			
MW-22B	09/28/10	11:15	9.25	6.16	0.150	20.80			
MW-23A	10/01/10	10:20	5.00	6.26	0.250	17.20			
MW-23B	09/28/10	18:25	7.00	5.67	0.110	18.10			
MW-24	09/30/10	12:15	9.50	7.06	0.180	18.70			
MW-25	10/01/10	11:35	6.75	6.09	0.260	18.60			
MW-26	10/01/10	10:30	4.50	5.90	0.150	17.00			
MW-27	10/01/10	9:15	8.00	5.64	0.090	18.60			
MW-29	10/01/10	9:30	7.50	5.80	0.090	19.00			
MW-31A	09/28/10	15:30	11.00	6.06	0.100	17.10			
MW-31B	09/28/10	16:30	8.75	6.07	0.110	16.60			
MW-32	09/28/10	17:35	6.25	5.88	0.090	17.70			
MW-33	09/28/10	10:35	17.00	6.30	0.140	16.80			
MW-34A	09/29/10	17:30	9.00	5.51	0.110	17.70			
MW-34B	09/29/10	11:10	45.00	8.03	0.240	20.40			
MW-35	10/01/10	12:05	9.50	5.10	0.080	19.00			
MW-38	09/30/10	10:00	11.45	6.37	0.170	16.20			
MW-39	09/30/10	11:15	11.25	6.12	0.150	17.00			
MW-40	09/27/10	13:50	11.35	6.27	0.130	18.10			

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urface Water Analytical Data Summary of Detected Purgeable Halocarbons and Aromatics (Chlorinated and Petroleum-Related Compounds) 207 Telegraph Drive, Gastonia, North Carolina 1,1-Dichloro-Chloro Xvlenes methyl ether Disopropyl-(1,1-DCE) (total) (MTBE) Toluene Benzene CASI 108-20-3 Sample Date of Dilution CAST 75-35-4 CAS# 71-43-2 CASH 108-90-7 CAS# 100-41-4 CASI 108-88-3 CASE CAST 1634-04-4 ppb ppb ppb ppb ampling ppb ppb 58 20 1/13/1997 <1 22 < 1 <1 32 59 7.2 130 90 4/8/1997 <1 97 <1 18 7/2/1997 <1 69 <1 10 96 76 85 10 11/10/1997 <1 29 <1 2.6 31 79 62 10 92 59 40 11 3/25/1998 <1 79 <1 20 <1 <1 25 <1 5/27/1998 <1 <1 <1 55 68 6 8/27/1998 <1 41 <1 10 32 <5 12/30/1998 <] <1 <1 <3 17 <1 <3 <5 <5 3/19/1999 <1 <1 <1 <3 12 <5 5/10/1999 <1 <3 9/14/1999 <1 <} <1 <1 <1 12/16/1999 <l <1 <1 <1 <1 <3 <5 <5 <1 <1 <I <1 <3 <5 <5 3/22/2000 <<u>l</u> <1 <3 <5 <5 6/13/2000 <1 <3 <1 9/11/2000 <1 <1 <1 <1 <5 <3 11/30/2000 <1 <1 <1 <1 <1 <5 2/20/2001 <1 <1 <1 <I <3 <5 <5 <1 <1 <3 <5 <5 6/5/2001 <1 <1 <1 3/14/2002 <1 <5 <5 <3 3/25/2003 <1 <1 <1 <1 <1 9/29/2003 <] <1 <1 <1 <1 <3 <5 <5 10/1/2004 <! <1 <1 <1 <3 <5 <5 <1 <1 <5 < 0.5 6/23/2005 <1 < 3 9/29/2005 < 1 < 0.5 < 1 < 1 3/7/2006 < 1 < 0.5 < 1 < 1 < 1 < 3 < 5 < 5 10/16/2006 < 0.5 < 1 < 1 < 3 < 5 < 5 < 0.5 < 1 < 3 < 5 3/6/2007 < 1 < 3 < 5 < 1 < 1 1.7 9/25/2007 < 1 1.8 < 5 < 3 < 5 3/11/2008 < 1 < 0.5 < 1 < 1 9/22/2008 DRY < 1 < 0.5 < 1 < 1 < 3 < 5 < 5 4/30/2009 DRY 9/18/2009 < 1 < 3 < 5 3/24/2010 < 1 < 0.5 < 1 1.9 < 3 < 1 < 0.5 < 1 <1 <! <1 <1 <1 4.3 <1 1/13/1997 <1 <1 <1 <<u>1</u> <1 <1 4/8/1997 <1 <1 <1 <1 <1 7/2/1997 <I <1 <1 11/10/1997 <1 <1 <1 <1 <1 <1 <1 3/25/1998 <1 <1 <1 <1 <1 6.1 <1 <1 <1 <1 <1 <1 <1 5/27/1998 <1 <1 <1 <1 <1 <1 <1 8/27/1998 <1 <1 <5 12/30/1998 <1 <1 <1 <1 <1 ⋖3 <5 <1 <1 <1 <1 <1 <3 <5 <5 3/19/1999 ⋖ <5 <5 <1 <1 <1 5/10/1999 <1 <5 <3 <1 <1 <1 9/14/1999 <1 <1 <5 12/16/1999 <1 <1 <1 <1 <1 <3 <5 <1 <1 <1 <1 <1 <3 <5 <5 3/22/2000 <3 <5 <5 <1 <1 6/13/2000 <1 <1 <1 <5 <5 <1 9/11/2000 <1 <l 11/30/2000 <1 <1 <1 <1 <1 ⋖3 <5 <1 <1 <1 <1 <1 <₃ <5 <5 2/20/2001 <1 <1 <3 <5 <5 <1 <1 6/5/2001 <1 <1 <3 <5 <5 <1 8/23/2001 <1 <5 <5 3/14/2002 <1 <1 <1 <1 <1 <3 <1 <1 <1 <1 <1 <3 <5 <5 3/25/2003 <1 <1 <3 <5 <5 9/29/2003 <3 <5 <5 <1 <1 <1 10/1/2004 <1 <3 6/23/2005 <1 < 0.5 <1 <1 <1 9/29/2005 <1 < 0.5 < 1 < 1 < 1 < 3 < 5 < 5 <1 <1 <1 <1 <1 <1 <1 <1 3/8/2006 <1 <1 <1 <1 <1 <1 3.9 10/16/2006 < 3 < 5 < 5 3/6/2007 2.3 < 0.5 <1 <1 <1 9/25/2007 <1 <1 <1 <1 <1 <1 <1 <1 3/11/2008 < 1 < 0.5 < 1 <1 DRY 9/22/2008 < 0.5 < 1 < 3 < 5 < 5 4/30/2009 < 0.5 < 1 < 1 < 3 < 5 < 5 9/18/2009 < 5 < 0.5 < 1 < 1 < 1 < 3 < 5

NOTES:

Sample results are expressed as a concentration or as less than the quantitation limit (e.g. <1).

The quantitation limit is equal to the dilution factor multiplied by the undiluted quantitation limit.

CAS# = Chemical Abstracts Service registry number assigned by the American Chemical Society to compounds included in the CAS Registry System.

*CAS#s: o-xylene = 95-47-6; m-xylene = 108-38-3; p-xylene = 106-42-3.

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Table 7

Estimated Mass of VOCs Removed From Groundwater

September 18, 2009 to September 7, 2010

207 Telegraph Drive, Gastonia, North Carolina

	ľ	stem Influent	Treatment Sy				_ 2	_	3		Cumulative Contaminant	
	Concen	tration	Concen	tration		Percent	Removeď	Remov	al Rate	Rem	oved'	VOC
Date of	1,1-DCE	Toluene	1,1-DCE	Toluene	Flowrate	1,1-DCE	Toluene	1,1-DCE	Toluene	1,1-DCE	Toluene	Removed ⁴
Sampling	(μg/L)	(μg/L)	(μg/L)	(μg/L)	(gpm)	(%)	(%)	(lbs/day)	(lbs/day)	(lbs)	(lbs)	(lbs)
09/18/09	1,100	<5	< 0.5	<0.5	55.0	99.95	90.00	0.73	0.00	0.00	0.00	0.00
10/23/09	1,000	<5	< 0.5	<0.5	54.5	99.95	90.00	0.65	0.00	24.17	0.10	24.27
11/17/09	910	<5	< 0.5	< 0.5	54.5	99.95	90.00	0.60	0.00	39.79	0.18	39.97
12/11/09	770	<5	< 0.5	<0.5	54.5	99.94	90.00	0.50	0.00	52.98	0.25	53.23
01/04/10	770	<5	< 0.5	< 0.5	53.5	99.94	90.00	0.49	0.00	64.96	0.32	65.28
02/05/10	750	<0.5	<0.5	< 0.5	54.0	99.93	0.00	0.49	0.00	80.66	0.36	81.02
03/05/10	610	<5	<0.5	< 0.5	54.6	99.92	90.00	0.40	0.00	93.06	0.41	93.47
04/01/10	630	<5	<0.5	< 0.5	55.0	99.92	90.00	0.42	0.00	104.08	0.49	104.56
05/07/10	550	<5	<0.5	< 0.5	55.1	99.91	90.00	0.36	0.00	118.11	0.59	118.70
06/10/10	610	<5	<0.5	< 0.5	55.5	99.92	90.00	0.41	0.00	131.20	0.69	131.90
07/12/10	580	<5	<0.5	<0.5	55.7	99.91	90.00	0.39	0.00	143.91	0.79	144.70
08/05/10	640	<5	<0.5	< 0.5	55.4	99.92	90.00	0.43	0.00	153.67	0.86	154.53
09/07/10	660	<5	<0.5	< 0.5	55.0	99.92	90.00	0.44	0.00	167.88	0.96	168.85
				ESTIMATED T	TOTAL MA	SS OF VOC	REMOVEI	9/18/09 thr	ough 9/07/10	: 167.88	0.96	168.85

NOTES:

gpm = gallons per minute

BP/DB

¹ When lab data was reported as below the reporting limit (BRL), half of the reporting limit value was used in this table, and the value was preceded by "<".

² Percent Contaminant Removed calculated as follows: [(influent conc. - effluent conc.)/influent conc.] x 100

³ Contaminant Removal Rate calculated as follows: [(influent conc. - effluent conc.) x flowrate x (1.2014E-5)], where 1.2014E-5 is the unit conversion factor.

⁴ Cumulative Contaminant Removed calculated as follows: [((Ave. contaminant removal rate between 2 consecutive sampling events) x (# days between 2 consecutive sampling events)) + previous cumulative contaminant removed]

Table 8							
Summary of Remedial Technologies							
207 Telegraph Drive, Gastonia, North Ca	rolina						
Containment							
Hydraulic	Five groundwater extraction wells currently operating; providing plume containment and mass reduction.						
Physical Barriers (Slurry wall, grouting)	Not practical to install slurry walls as deep as needed. Grouting is possible but not likely to be completely effective. Would address migration through bedrock fractures.						
Permeable Reactive Barriers (Zero Valent Iron, bio, e-barriers)	Depth is too great and would be difficult to install into the bedrock.						
Phytoremediation	Depth is too great.						
	one Restoration/Partial Mass Removal						
Electrical Resistive Heating (three-phase or six-phase heating)	Installation of electrodes into the bedrock paired with a vapor extraction system. Resistive heating would "boil off" the VOCs, which would then be captured and treated. High potential for removing a significant mass of DNAPL. Installation into the bedrock would be technically challenging and very expensive. Potential concern with enhancing migration of possible DNAPL as the matrix is heated. May have to heat first from below the possible DNAPL zone ("hot floor" method - not readily applicable to bedrock).						
Steam Flushing	Contact with contaminants not likely to be complete in fractured bedrock.						
In-Situ Oxidation	Injection of oxidizing chemicals such as hydrogen peroxide or potassium permanganate. Difficult to inject the chemicals into the bedrock fractures where possible DNAPL exists.						
In-Situ Reduction	Injection of reducing chemicals such as a slurry of zero-valent iron. Difficult to locate and inject the chemicals into the bedrock fractures where possible DNAPL exists.						
Dual-Phase Extraction	Application of a high vacuum to deep wells to extract mobile DNAPL, contaminated groundwater, and vapors. If it is possible to dewater the formation, soil vapor extraction can be performed. Even if possible to implement this technology, it will still leave a mass of possible DNAPL.						
Enhanced Reductive Dechlorination	Injection of vegetable oil or other electron donor to stimulate reductive dechlorination. Difficult to locate and inject the chemicals into the bedrock fractures where possible DNAPL exists.						

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Appendices

Appendix A

Monitoring Well Construction Data

Table A-1 Summary of Monitoring Well Construction Data 207 Telegraph Drive, Gastonia, NC

Well Location	Hydrogeologic Unit	Ground Surface	NAD 83	Coordinates	Well	Surface Casing	Top of PVC Well	Screen Interval	Screen Interval
	Screened	Elevation			Diameter	Depth	Casing		Elevation
		(Feet Above MSL)	(N)	(E)	(Inches)	(Feet)	(Feet Above MSL)	(Depth - Feet)	(Feet Above MSL)
MW-1A	Saprolite	735.16	535,096.851	1,340,540.328	2	N/A	735.17	23.5 - 33.5*	711.66 - 701.66
MW-1B	Bedrock	735.62	535,105.883	1,340,538.490	2	43.5	735.49	49 - 74	686.62 - 661.62
MW-3	Saprolite	720.06	535,139.745	1,340,746.244	2	N/A	722.56	10 - 20*	710.06 - 700.06
MW-4A	Saprolite	712.77	535,011.289	1,340,788.137	2	N/A	714.32	8 - 18*	704 .77 - 694.77
MW-4B	Bedrock	713.57	535,022.065	1,340,779.695	2	45	716.38	48 - 58	665.57 - 655.57
MW-5	Saprolite	762.64	535,292.352	1,340,058.189	2	N/A	762.55	40.5 - 50.5*	722.14 - 712.14
MW-7A	Saprolite	710.50	534,939.366	1,340,937.908	2	N/A	712.80	13 - 23*	697.50 - 687.50
MW-7B	Bedrock	710.54	534,946.228	1,340,949,126	2	35.5	713.23	49 - 64	661.54 - 646.54
MW-8	Saprolite	722.59	534,988.244	1,341,086.482	2	N/A	724.52	21.5 - 31.5*	701.09 - 691.09
MW-10A	Saprolite	759.77	535,540.476	1,340,354.549	2	N/A	759.52	36.5 - 46.5*	723.27 - 713.27
MW-10B	Bedrock	759.45	535,539.060	1,340,344.569	2	73	759.15	203.5 - 243.5	555.95 - 515.95
MW-11	Saprolite	758.10	535,466.350	1,340,345.661	2	N/A	757.73	34.5 - 44.5*	723.60 - 713.60
MW-12	Bedrock	760.63	535,633,107	1,340,238,560	4	25	760.36	25 - 46**	735.63 - 714.63**
MW-13	Saprolite	743.16	535,494.44	1,340,484.48	2	N/A	743.36	19.5 - 34.5*	723.66 - 708.66
MW-15A	Saprolite	725.40	535,444.884	1,340,693,162	2	N/A	725.23	10.5 - 20.5	714.90 - 704.90
MW-15B	Bedrock	725.80	535,438.132	1,340,687.553	2	22.5	725.71	23.5 - 33.5	702.30 - 692.30
MW-15C	Bedrock	724.61	535,436,793	1,340,695.053	2	36	724.25	60.5 - 75.5	664.11 - 649.11
MW-15D	Bedrock	725.67	535,460.183	1,340,690.515	2	123	725.37	203 - 243	522.67 - 482.67
MW-16A	Saprolite	735.72	535,272.738	1,340,530.030	2	N/A	735.42	30 - 55	705.72 - 680.72
MW-16B	Bedrock	735.69	535,268.467	1,340,532.660	2	76	735.25	90 - 115	645.69 - 620.69
MW-17A	Saprolite	724.28	535,277.519	1,340,728.255	2	N/A	723.96	21 - 36	703.28 - 688.28
MW-17B	Bedrock	724.40	535,282,376	1,340,726,755	2	42	724.11	44 - 54	680.40 - 670.40
MW-17C	Bedrock	724.20	535,266.867	1,340,725.223	2	53.5	723.70	105 - 120	619.20 - 604.20
MW-18A	Saprolite	729.40	535,411,452	1,340,925,009	2	N/A	729.13	20 - 35	709.40 - 694.40
MW-18B	Bedrock	728.75	535,416.217	1,340,918.850	2	75	728.45	80 - 100	648.75 - 628.75
MW-19A	Saprolite	742.80	535,526.046	1,340,563.919	2	N/A	742.49	28 - 38	714.80 - 704.80
MW-19B	Bedrock	742.81	535,520.709	1,340,561.257	2	43	742.54	52 - 77	690.81 - 665.81
MW-20A	Saprolite	728.17	535,127.194	1,341,005,012	2	N/A	730.92	33 - 63	695.17 - 665.17
MW-20B	Bedrock	728.55	535,119.860	1,341,009.204	2	79	730.96	83 - 98	645.55 - 630.55
MW-21	Bedrock	741.00	535,614,481	1,340,862,614	2	36	740.79	44 - 69	697.00 - 672.00
MW-22A	Saprolite	732.50	535,307.242	1,341,003.874	2	N/A	734.88	34.5 - 54.5	698.00 - 678.00
MW-22B	Bedrock	732.34	535,302.221	1,341,002.917	2	56/633	734.77	68 - 93	664.34 - 639.34
MW-23A	Saprolite	729.21	535,648.730	1,340,573.615	2	N/A	731.27	4 - 19	725.21 - 710.21
MW-23B	Bedrock	729.80	535,651.712	1,340,567.858	2	26	732.03	35 - 45	694.80 - 684.80
MW-24	Bedrock	729.97	534,993.783	1,341,188.236	2	77	732.54	95 - 110	634.97 - 619.97
MW-25	Saprolite	744.48	535,615.857	1,340,469.870	2	N/A	744.12	18 - 38	726.48 - 706.48
MW-26	Saprolite	742.19	535,737.426	1,340,470.780	2	N/A	745.13	16 - 31	726.19 - 711.19

Table A-1 Summary of Monitoring Well Construction Data 207 Telegraph Drive, Gastonia, NC

Well Location	Hydrogeologic Unit Screened	Ground Surface Elevation	NAD 83 (Coordinates	Well Diameter	Surface Casing Depth	Top of PVC Well Casing	Screen Interval	Screen Interval Elevation
		(Feet Above MSL)	(N)	(E)	(Inches)	(Feet)	(Feet Above MSL)	(Depth - Feet)	(Feet Above MSL)
MW-27	Bedrock	755.92	535,779.245	1,340,420.406	2	N/A	¹ 754.56	28 - 48	727.92 - 707.92
MW-28	Bedrock	767.88	535,722.796	1,340,146.005	2	N/A	771.00	38 - 68	729.88 - 699.88
MW-29	Bedrock	756.55	535,857.023	1,340,360.088	2	N/A	759.44	25 - 50	731.55 - 706.55
MW-30	Bedrock	767.36	535,460.533	1,339,974.503	2	50	767.16	174 - 194	593.36 - 573.36
MW-31A	Saprolite	727.45	535,763.174	1,340,681.243	2	N/A	729.88	15 - 30	712.45 - 697.45
MW-31B	Bedrock	728.20	535,763.788	1,340,691.307	2	42	730.68	55 - 70	673.20 - 658.20
MW-32	Saprolite	725.60	535,629.453	1,340,704.119	2	N/A	727.86	10 - 20	715.60 - 705.60
MW-33	Bedrock	710.90	534,806.923	1,340,888.215	2	73	713.73	80 - 100	630.90 - 610.90
MW-34A	Bedrock	740.27	535,277.975	1,341,199.988	2	N/A	741.86	88 - 103	652.27 - 637.27
MW-34B	Bedrock	739.88	535,252.080	1,341,209.006	2	122	741.29	172 - 202	567.88 - 537.88
MW-35	Saprolite	729.11	535,903.670	1,340,615.540	2	N/A	730.84	11 - 26	718.11 - 703.11
MW-36	Saprolite	743.63	535,920.370	1,340,956.650	2	N/A	743.41	34.5 - 54.5	709.13 - 689.13
MW-37A	Saprolite	753.56	535,254.070	1,340,199.730	2	N/A	753.41	45 - 65	708.56 - 688.56
MW-37B	Bedrock	756.12	535,275.750	1,340,195.500	6	90	755.89	90 - 220**	666.12 - 536.12**
MW-38	Bedrock	707.81	534,640.040	1,340,959.670	2	75	709.72	82 - 102	625.81 - 605.81
MW-39	Bedrock	711.86	534,838.609	1,341,076.498	2	77	714.15	82 - 102	629.86 - 609.86
MW-40	Bedrock	728.56	535,074.984	1,341,077.425	2 .	N/A	731.16	82 - 102	646.56 - 626.56
DM-7	Bedrock	768.44	535,598.796	1,340,008.372	2	N/A	768.12	31 - 81	737.41 - 687.41
CP-1	Bedrock	716	535,294.250	1,340,895.126	6	0 - 40	716.87	40 - 48 ²	679 - 671
EW-2	Saprolite/Bedrock ³	722	535,093.092	1,340,965.352	6	NA	723.09	58 - 103	667 - 622
EW-3	Saprolite/Bedrock ³	708	534,933.559	1,340,920.837	6	NA	709.26	30 - 80	681 - 631
EW-4	Saprolite/Bedrock ³	726	535,475.731	1,340,683.001	6	NA	724.6	17 - 67	709 - 659
EW-5	Saprolite/Bedrock ³	745	535,609.711	1,340,475.372	6	NA	742.1	7 - 77	738 - 668

NOTES:

N/A = Not applicable.

^{*}Screen interval based on the measured depth to the bottom of the well. This interval differs from the screen interval shown in the well construction diagram for this well. **Open borehole interval in bedrock.

MW-27 was converted from a stick-up well to a flush mount, the Top of Casing Elevation for MW-27 was surveyed by CH2M HILL on 1-12-07

CP-1 is an open bedrock borehole from 40 to 48 feet below grade. EW-2, EW-3, EW-4, and EW-5 are screened across the saprolite/bedrock interface.

Not included in Fiscal Analysis

Appendix B

Laboratory Analytical Report