

**Annual Monitoring Report**

**April 2004**

**Carroll and Dubies Superfund Site  
Town of Deerpark, Orange County, New York**

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**July 2004**

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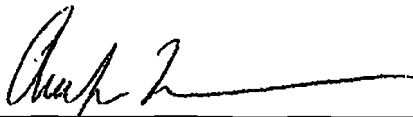
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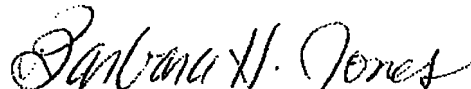
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July 2004

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## 1.0 Introduction

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This *Annual Monitoring Report* has been prepared for the Carroll and Dubies (C&D) Superfund Site, on behalf of the Carroll and Dubies Superfund Site Potentially Responsible Parties (PRP) Group, by Shield Environmental Associates, Inc. (Shield Environmental). In a letter dated January 30, 2003, the United States Environmental Protection Agency (U.S. EPA) Region 2 approved a request to change the monitoring frequency for this site from semiannually to annually beginning in 2003. This report documents field activities and presents analytical data from field work completed between April 19 and April 26, 2004. The documents used as guidance for the field activities were:

- *Field Sampling and Analysis Plan Addendum* (Shield Environmental, 1998)
- *Final Remedial Design Work Plan Addendum* (Shield Environmental, 1998a)
- *Quality Assurance Project Plan Addendum* (Shield Environmental, 1998b)
- *Remedial Action Work Plan, Appendix B, Health and Safety Plan* (Shield Environmental, 1998c)

Historical data from the *Addendum to Supplemental Hydrogeologic Remedial Investigation* (Remediation Technologies Inc., 1995) and *Preliminary Remedial Investigation Results* (Blasland, Bouck, & Lee, Inc., 1992) have been incorporated into this report as well as data collected during previous Shield Environmental sampling events (Appendices A and B). These reports and earlier monitoring reports provide descriptions of the hydrogeology of the site and baseline water quality.

The remedial activities for this site are separated into two operable units. Operable Unit 1 (OU-1) addressed source control through remediation of the lagoons, which was completed in 1999. This report covers the Operable Unit 2 (OU-2) activities only, which address groundwater impacts and remediation.

### 1.1 Site Location and History

The three-acre C&D Superfund Site is located in the Town of Deerpark in Orange County, New York, which is approximately 3,000 feet northeast of the City of Port Jervis, New York (Figure 1). In 1970, the site began operating as a disposal facility. Liquid industrial wastes were received from approximately 1971 to 1979. The facility also accepted septic waste and municipal solid waste until 1989, which were the

majority of wastes disposed of at the site. The wastes disposed at the site were removed during remediation activities conducted in 1999. These activities are documented in the OU-1 Remedial Action Report (Shield Environmental, 2000).

The site is situated in the Neversink Valley. Gold Creek lies approximately 1,500 feet to the east, and the Neversink River is located approximately 2,000 feet beyond Gold Creek. The immediate surrounding area includes undeveloped woodlands to the north; undeveloped woodlands, a sand and gravel quarry pit, and the closed City of Port Jervis landfill to the east; the Orange County Transfer Station and a concrete products fabrication company to the south; and a sparsely vegetated, shale bedrock, hillside to the west.

## **1.2 Work Plan Variations**

During this sampling event, there were two variations from the work plans:

- The sediment sample collection method was modified to reduce the moisture content of the samples, which has been a problem in the past. First, a piece of polyvinyl chloride (PVC) pipe was driven into the sediments with a hammer to a depth of 1 foot below the surface. A decontaminated hand pump was then used to draw off water from the surface of the sediments, and the sediment samples were collected using a stainless-steel spoon.
- Semivolatile organic compounds (SVOCs) remain on the analyte list for monitoring well OW-10R only per the January 30, 2003 correspondence from the U.S. EPA.

## **2.0 Annual Monitoring Activities**

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This section outlines the procedures used for field sampling and monitoring activities, including measurement of groundwater elevations, collection of groundwater samples, measurement of field parameters in groundwater, surface water sampling, and sediment sampling. The results and interpretation of the monitoring program are provided in Section 3.0.

### **2.1 Groundwater Elevations**

Groundwater elevations were determined based on measured depths to water from the reference point elevations before sampling began. The depth to groundwater was measured using a Solinst® electronic water-level meter and recorded in the field logbook to the 1/100<sup>th</sup> of a foot.

In addition to gauging groundwater levels, three surface water elevations were also measured. Three permanent staff gauges were installed, two along Gold Creek and one adjacent to the quarry pond east of monitoring well OW-8 during the June 2000 sampling event. These locations are shown in Figure 2 and are identified as SW-1, SW-2, and SW-3 (quarry pond). During the April 2004 sampling event, the surface water elevations at SW-1, SW-2, and SW-3 were directly measured using a survey transit and graduated staff. This method of obtaining water levels at the surface water locations is recommended for future sampling events.

### **2.2 Monitoring Well Sample Collection**

Dedicated low-flow purging and sampling pumps are utilized to sample the monitoring wells. After stabilization procedures were completed, groundwater samples were taken directly from the Tygon® tubing dedicated to each respective well pump.

Groundwater purged from the monitoring wells was generally clear and contained little suspended sediment. All groundwater samples were collected at a flow rate of between 175 and 250 milliliters per minute (mL/min). All monitoring wells maintained 0.33 foot of drawdown or less during purging and stabilization.

During purging of each monitoring well, temperature, dissolved oxygen (DO), redox potential, specific conductance, pH, and turbidity were monitored and recorded on field forms (Appendix C) in average intervals of 5 minutes. The wells were pumped for

periods ranging from 30 to 65 minutes. The goal was to obtain three consecutive readings of the field parameters within the following ranges:

- $\pm 1.0$  degrees centigrade ( $^{\circ}\text{C}$ ) for temperature
- $\pm 10$  percent (%) or  $\pm 0.3$  mg/L for DO (whichever is greater)
- $\pm 10$  millivolts (mV) for redox potential (redox)
- $\pm 3\%$  for specific conductance (conductivity)
- $\pm 0.1$  for pH
- $\pm 10\%$  or  $\pm 2$  nephelometric turbidity units (NTUs) for turbidity (whichever is greater)

## 2.3 Groundwater Field Parameters

Groundwater field parameters were measured with a YSI Model 556 MPS-10 multi-parameter unit equipped with a flow-through cell and a Lamotte Turbidity Meter Model 2020, which were calibrated prior to sampling activities. Field parameters measured and recorded included:

- Temperature ( $^{\circ}\text{C}$ )
- DO (mg/L)
- Redox potential (mV)
- Specific conductance (micromhos per centimeter [ $\mu\text{mhos/cm}$ ])
- pH (standard units)
- Turbidity (NTUs)

The final stabilized field parameter values are presented in Table 1.

### Temperature

Although a range for temperature fluctuations was not established in the *Field Sampling and Analysis Plan Addendum* (Shield Environmental, 1998), fluctuations were less than  $1.0^{\circ}\text{C}$  for a minimum of three consecutive readings prior to sampling all of the monitoring wells. Groundwater temperatures ranged from  $9.61^{\circ}$  to  $12.65^{\circ}\text{C}$ .

### Dissolved Oxygen

DO concentrations stabilized for a minimum of three consecutive measurements within the recommended  $\pm 10\%$  or 0.3 mg/L range in all monitoring wells. DO is an important parameter in the interpretation of natural attenuation trends. DO readings across the site ranged from 0.11 to 4.39 mg/L.

### Redox

Redox stabilized within the recommended  $\pm 10$  mV range for three consecutive measurements in all monitoring wells. Redox is also an important parameter in the interpretation of natural attenuation trends. A wide range of redox readings were found in the wells, varying from -49.3 to 203.3 mV.

### Specific Conductance

A minimum of three consecutive readings for specific conductance was achieved within the recommended  $\pm 3\%$  range for all monitoring wells prior to sampling. Specific conductance varied from 85 to 867 umhos/cm across the site.

### pH

A minimum of three consecutive readings for pH within the recommended  $\pm 0.1$  range was achieved for all monitoring wells prior to sampling. Groundwater samples were within the pH range of 5.67 to 6.51.

### Turbidity

A minimum of three consecutive readings for turbidity were achieved within the recommended  $\pm 10\%$  or  $\pm 2$  NTUs range for monitoring wells prior to sampling. Turbidity readings varied from 0.00 to 22.00 NTUs across the site.

## **2.4 Surface Water Sampling**

Two surface water samples were collected from Gold Creek and are identified as SW-1, the downstream sample, and SW-2, the upstream sample.

Samples were collected for volatile organic compounds (VOCs) in accordance with the *Field Sampling and Analysis Plan* (Shield Environmental, 1998), using a disposable container provided by the laboratory to collect and transfer the sample water at each location to the VOC sample vials. The sample bottles were labeled appropriately, placed in a cooler with ice, and sent to a laboratory for analysis.

## **2.5 Sediment Sampling**

Two sediment samples were collected from Gold Creek. The sediment sampling locations, labeled as SED-1 and SED-2, are identified in Figure 2, coinciding with

surface water sample locations SW-1 and SW-2. The samples were collected at the sediment/water interface.

The sampling approach was designed to collect samples with a relatively lower moisture content. First, a decontaminated 8-inch diameter PVC pipe segment, approximately 2 feet in length, was driven into the sediments to about 1 foot below the sediment surface. A decontaminated hand pump was then used to draw off water from the surface of the sediments, and the upper layer of muck and debris was scraped from the surface. The samples were collected using a stainless-steel spoon by the field technician wearing clean disposable nitrile gloves. Between sediment sample locations, the stainless-steel spoon, pump, and PVC pipe were decontaminated with a non-phosphate detergent and rinsed with distilled water. This technique was effective in improving the solids content of the sediment samples, compared to April 2003. The average solids content was about 41% using this technique, compared to about 16% in 2003. Higher solids concentrations improve the analytical reporting limits.

## **3.0 Sampling Results**

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### **3.1 Groundwater Elevations**

The groundwater elevations for this sampling round are presented in Table 2. Associated groundwater elevation contours are shown in Figure 3.

The elevations of adjacent surface water bodies have been included in the groundwater contours map shown in Figure 3. Measurements of surface water elevations were collected on the same day that the monitoring wells were gauged.

The groundwater flow direction on site is toward the southeast in the direction of Gold Creek. The groundwater gradient across the former lagoon site is approximately 0.080. This gradient transitions to a lower gradient, at about the location of the towpath. From the towpath to Gold Creek, the gradient is very shallow, approximately 0.007. The steeper gradient on the western side of the site is due to the depth to bedrock along the valley wall. As the depth to bedrock increases towards the valley floor, the thickness of the alluvial fill increases and the groundwater gradient flattens.

### **3.2 Groundwater Quality**

Detected VOC analytes from the April 2004 sampling event are presented in Table 3 and are included in Figure 4. No SVOCs were detected in duplicate samples from OW-10R, the one monitoring well that continues to be monitored for SVOCs. Historical data of detected organic compounds have been combined with the most recent data and are presented in Appendices A (VOCs) and B (SVOCs). Complete laboratory analytical packages for this sampling episode are provided as electronic files on disk in Appendix D.

#### **3.2.1 Volatile Organic Compounds**

VOCs are the primary contaminants of concern. The concentrations of VOCs for this annual monitoring event have been compared to results from previous sampling events.

Eleven VOCs were detected in various wells during this sampling event, of which six exceeded regulatory limits (Table 3). The VOCs that exceeded regulatory limits are benzene, chlorobenzene, 1,2-dichloroethene (1,2-DCE) (total), tetrachloroethene (PCE), trichloroethene (TCE), and vinyl chloride. The detected compounds and

concentrations are shown beside each well in Figure 4. The principal VOCs were benzene, 1,2-DCE (total), and PCE, which is consistent with previous sampling episodes.

Figure 5 presents a graphical representation of concentration trends since January 2000 for benzene, 1,2-DCE (total), and PCE in wells where regulatory criteria have been exceeded. Although there have been variations in the detected concentrations over time, the overall trends are downward for benzene. For OW-5 (1,2-DCE and PCE) and OW-13 (1,2-DCE), the overall trends are also downward. In OW-2, downgradient of former Lagoon 2, concentrations of 1,2-DCE and PCE appear to be cyclic, and influenced by groundwater elevations. For example, as shown in Figure 6, the pattern of fluctuations in groundwater elevations in OW-2 mirrors the pattern seen for 1,2-DCE concentrations. Even though groundwater elevations fluctuate throughout the site, as shown in a comparison of hydrographs for four representative monitoring wells (Figure 7), contaminant concentrations appear to be most affected in OW-2, based on its location close to the original source of impact. It is anticipated that over time fluctuations will decrease as concentrations decline through natural attenuation.

Changes in the VOC groundwater concentrations over time are depicted graphically in Figures 8, 9, 10, and 11. Figures 8 and 9 represent the concentrations of benzene and 1,2-DCE (total) in three wells over time, OW-10/OW-10R, OW-13, and OW-18. The graphs are shown chronologically from February 1999 (pre-remediation sampling) to the most recent monitoring event in April 2004. Figures 10 and 11 indicate the relationship between benzene and 1,2-DCE (total) in these wells, with respect to distance from the source area over time.

The wells used in the graphical assessments (Figures 8, 9, 10, and 11) were chosen based on their detected VOC concentrations and position along the gradient of the contaminant plume. Monitoring well OW-10/OW-10R is at the edge of the lagoon area along the towpath. OW-13 is located approximately 150 feet downgradient from OW-10R in the direction of groundwater flow, and OW-18 is located near Gold Creek, approximately 700 feet downgradient of OW-13.

The graphs (Figures 8, 9, 10, and 11) indicate that there has been a decrease in the concentrations of both benzene and 1,2-DCE (total) at most monitoring wells over time since the remediation activities were completed in 1999.

### **3.2.2 Semivolatile Organic Compounds**

In a letter dated January 30, 2003, U.S. EPA Region 2 approved elimination of SVOCs from the analyte list for monitoring wells OW-13 and OW-18. U.S. EPA requested that OW-10R continue to be monitored for SVOCs. No SVOCs were detected in the sample or duplicate sample from monitoring well OW-10R. A historical summary of detected SVOC constituents in groundwater is provided in Appendix B. This well was also nondetectable for SVOCs when sampled in April 2003.

### **3.2.3 Monitored Natural Attenuation Parameters**

Field measurements and laboratory analyses were performed on all groundwater samples to evaluate natural attenuation trends. The physical and chemical parameters measured as part of this monitoring program show that natural attenuation is occurring at the site. Since wastes and contaminated soils that acted as source material have been removed, concentrations of organic constituents in the groundwater continue to decrease over time due to natural attenuation processes.

Table 4 summarizes the natural attenuation parameter results, which include:

- Alkalinity
- Chloride
- DO
- Ethane/ethene
- Ferrous iron
- Methane
- Nitrate
- Redox
- Sulfate
- Sulfide
- Total organic carbon (TOC)

The field parameters (Appendix C; Appendix E) were measured during sample collection using a multiparameter instrument. Natural attenuation analytical parameters for samples collected since February 1999 are provided in Appendix F.

#### **General Observations**

The contaminant plume at the C&D site is relatively complex, in that it shows impacts from chlorinated VOCs, nonchlorinated VOCs, and organic carbon as lagoon leachate. The general observations regarding natural attenuation data are:

- Several parameters indicate that reductive dechlorination of chlorinated VOCs is occurring (presence of chlorinated daughter products, historical presence of ethane/ethene and methane within the plume, elevated chloride concentrations downgradient of the lagoons, redox readings less than 50 mV within the plume).
- Indicators of reductive dechlorination are most consistently found in association with the former Lagoon 8 portion of the contaminant plume, which contains both benzene and chlorinated VOCs. The former Lagoons 1 and 2 portion of the plume contains mainly chlorinated VOCs.
- DO readings were less than 0.5 mg/L in most of the wells, indicating low-oxygen conditions, under which reductive dechlorination can occur.

### Alkalinity

Alkalinity greater than two times the background concentration is indicative of the ultimate oxidation of chlorinated VOCs to carbon dioxide (CO<sub>2</sub>). Alkalinity greater than or equal to 210 mg/L (twice the value found in MW-1) was measured in OW-10R, OW-13, OW-18, OW-19, OW-21, and OW-22.

### Chloride

Elevated chloride concentrations (equal to or greater than two times the background concentration of 4.6 mg/L in MW-1) are indicative of reductive dechlorination. Chloride concentrations greater than or equal to 9.2 mg/L were found in MW-4, OW-5, OW-16, OW-18, OW-19, and OW-22, where chlorinated VOCs and degradation products have been detected.

### Dissolved Oxygen

Groundwater with DO concentrations less than 0.5 mg/L, indicating low oxygen conditions that favor degradation of chlorinated compounds, were present in 11 of the 14 wells sampled during this monitoring period. DO readings were less than 0.5 mg/L in wells MW-4, OW-5, and OW-8, downgradient from former Lagoons 1 and 2. DO readings of less than 0.5 mg/L were found in wells OW-10R, OW-13, OW-18, OW-19, OW-21, and OW-22, south of the former lagoon area.

### Ethane/Ethene

These compounds are end degradation products of chlorinated VOCs. Ethane was detected in OW-13 at 0.19 micrograms per liter (ug/L). This value is J-qualified as an estimated result below the reporting limit. Ethene was detected at 0.385 ug/L in OW-13.

### Ferrous Iron

Ferrous iron is generated when ferric iron is used as an electron acceptor under anaerobic conditions. Ferrous iron concentrations greater than 1 mg/L may indicate reductive dechlorination. Ferrous iron concentrations greater than 1 mg/L were measured in all monitoring wells except MW-1, OW-2, OW-5, OW-6, OW-10R, and OW-16. The elevated ferrous iron concentrations were found downgradient of all of the former lagoons.

### Methane

Methane is an ultimate reductive dechlorination daughter product, and concentrations greater than 0.1 mg/L (100 ug/L) may indicate this process. However, the presence of methane at this site could also be an indicator of impacts from septic waste disposal or the closed landfill. Methane was measured at concentrations of about 0.1 mg/L (100 ug/L) or greater in MW-4, OW-10R, OW-13, OW-15, OW-18, OW-19, and OW-22.

### Nitrate

Low nitrate concentrations (less than 1 mg/L) potentially indicate the occurrence of anaerobic degradation processes and reductive dechlorination in groundwater. Nitrate concentrations of 1 mg/L or less were found in all monitoring wells, except MW-1 (the background well), OW-2 and OW-5 downgradient of the Lagoons 1 and 2 contaminant plume, and OW-16, a monitoring well downgradient of Lagoons 3 and 4. The relatively high nitrate concentrations in OW-2 and OW-5 are most likely attributable to septage wastes that were once disposed in former Lagoons 1 and 2.

### Redox

Redox readings below approximately +750 mV indicate that anaerobic processes can occur. A reductive dechlorination pathway is possible when redox readings are +50 mV or less. Redox values for most of the wells located near and downgradient of the former lagoons exhibited redox readings less than +50 mV, indicative of conditions conducive to dechlorination. Background well MW-1 and downgradient wells OW-2, OW-5, OW-6, and OW-16, located further from the source areas, had redox values of greater than +50 mV.

### Sulfate/Sulfide

Sulfate is an electron acceptor under anaerobic, strongly reducing conditions, and sulfide is produced under these conditions. Relatively low sulfate (less than 20 mg/L),

combined with detectable sulfide (greater than 1 mg/L) may be indicative of this reductive dechlorination pathway. Sulfate concentrations less than 20 mg/L were measured in MW-1, OW-8, OW-13, OW-15, OW-18, OW-19, and OW-22, and detectable sulfide (greater than 1 mg/L) was detected in OW-18 and OW-22.

### TOC

Organic carbon, as measured by TOC, provides a carbon and energy source for biodegradation, and can drive dechlorination. TOC was not measured in any well at the rule of thumb concentration of 20 mg/L or greater. Concentrations greater than the background concentration of 1 mg/L (MW-1) were found in wells MW-4, OW-10R, OW-13, OW-15, OW-18, OW-19, OW-21, and OW-22.

## **3.3 Surface Water Quality**

Two surface water samples, SW-1 (downstream) and SW-2 (upstream), were collected during this sampling event and analyzed for VOCs. VOCs were not detected in samples from either surface water monitoring point. SVOC analysis in surface water is no longer required (see letter from the U.S. EPA Region 2 dated September 3, 2002). VOCs and SVOCs that have been detected in surface water during previous episodes are shown in Table 5.

Historically, VOCs and SVOCs have not been detected above a regulatory drinking water standard in surface water, with the exception of a detection of 1,2-DCE above the NYSDEC SGV of 0.6 ug/L at 0.61 ug/L in March 2001 at SW-1, at a J-qualified value, meaning an estimated concentration below the reporting limit. Based on these findings, there is no evidence that surface water has been impacted by the site.

## **3.4 Sediment Quality**

Acetone and 2-butanone were detected in both the downstream (SED-1) and upstream (SED-2) samples at low concentrations below the reporting limit (J values) during this sampling period (Table 6). Acetone was also detected in the associated trip and field blanks. SVOC analysis is no longer required for sediments.

## 3.5 Data Quality Review

A Tier II data quality review of sample data packages was completed using U.S. EPA guidelines. The Tier II data evaluation consisted of a review of data package completeness and a quality control (QC) review, as summarized in the QC forms provided by the laboratory. The completeness review covered:

- Signed transmittal page
- Data package narrative
- Sample transmittal documentation
- Standard QC forms for:
  - Surrogate recovery
  - Matrix spike/matrix spike duplicate (MS/MSD) recovery
  - Laboratory control samples
  - Method blank summary
  - Instrument performance check
  - Internal standard summary and retention time (RT) summary
  - Form Is and raw data for field samples
  - Initial calibration data
  - Continuing calibration data
  - Form Is and raw data for blanks, laboratory control samples, MS/MSDs
  - Copies of logbook pages documenting sample preparation, moisture determination, extract transfer, instruments, and sample tracking

The QC review covered the same items as in the completeness review, and:

- Holding times
- Form Is and raw data for field and QC samples
- Other data quality considerations (field procedures and sediment handling, moisture content of sediment samples, field duplicates and field blanks)

### 3.5.1 Results of Data Review

Refer to Appendix G for Tier II data review summary tables (VOC, SVOC, dissolved gases, and general chemistry) and marked Form Is for two sample lots, 4D24136 and A4D270137. For both sample lots, the data packages were complete and appropriately organized, and all relevant supporting information was provided.

Holding times were met, and field duplicates and MS/MSD duplicate samples were analyzed where required.

Although minor deviations from QC parameters were noted in some instances, the only results that required qualification through the data validation process were low concentrations of acetone or methylene chloride (J-qualified, below the reporting limit) in the following samples:

- MW-4
- OW-2
- OW-10R
- OW-10R Dup
- OW-18
- OW-19
- OW-22
- SW-1

The associated method, field, or trip blanks for field samples also contained low concentrations of these common laboratory contaminants. Using U.S. EPA data validation methods, the field results were reported as nondetectable (U) at the reporting limit, as shown in Tables 3 and 5. Acetone in the sediment samples was flagged as FB for field blank or TB for trip blank because acetone was detected in the associated aqueous trip and field blanks.

### **3.5.2 Data Quality and Usability**

The data review process indicates that the analytical results are of acceptable quality, and no results have been rejected. Acceptable field techniques are indicated by the comparable field duplicate results and the improved solids concentrations in the sediment samples.

## **4.0 Summary and Recommendations**

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The 14 monitoring network wells were sampled between April 23 and April 26, 2004 for analysis of VOCs and natural attenuation parameters. One monitoring well (OW-10R) in the network was sampled for analysis of SVOCs. Two surface water samples and two sediment samples were collected and analyzed for VOCs. All laboratory analytical samples were analyzed by STL, North Canton, Ohio.

### **4.1 Summary**

In summary, the sampling data show:

- In general, the concentrations of VOCs in groundwater continue to show a downward trend following the OU-1 remedial action.
- Concentrations of all VOCs in two downgradient wells (OW-8 and OW-16) have consistently been below maximum contaminant levels and the NYSDEC Standards and Guidance Levels, for at least five sampling rounds, and have been detectable for VOCs for three or more sampling rounds.
- Fluctuations in concentrations in OW-2 appear to be related to cyclic fluctuations in groundwater elevations.
- Conditions remain favorable for natural attenuation, and the results indicate that the process is ongoing.
- No VOCs were detected in surface water samples.
- Acetone and 2-butanone were detected in the downstream (SED-1) and upstream (SED-2) sediment locations in low concentrations typical of what has been previously reported.

### **4.2 Recommendation**

It is recommended that the groundwater monitoring program continue on an annual basis for the parameters analyzed in April 2004, with the following exceptions:

- Elimination of SVOCs as an analytical requirement for OW-10R. SVOCs were not detected in samples from this monitoring well in 2003 or 2004. There has been no detection of SVOCs in this well in concentrations exceeding criteria since 2001.
- Elimination of OW-8 and OW-16 from the monitoring program. There is sufficient coverage by other monitoring wells, detected VOCs in these wells have always been below criteria, and results have been nondetectable for three or more sampling rounds.

## 5.0 References

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Blasland, Bouck, and Lee, Inc., 1992, *Preliminary Remedial Investigation Results, Carroll & Dubies Site*, Port Jervis, New York.

Remediation Technologies, Inc., 1995, *Addendum to Supplemental Hydrogeologic Remedial Investigation: Results of Field Investigation at the Carroll and Dubies Site During April, 1995*.

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## Tables

**Table 1**  
**Groundwater Field Parameters**  
**April 2004**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**

Well ID	Date	Temperature (°C)	Dissolved Oxygen (mg/L)	Redox (mV)	Specific Conductance (uS/cm)	pH (standard units)	Turbidity <sup>(1)</sup> (NTUs)	Ferrous Iron (mg/L)	Nitrate Nitrogen (mg/L)
MW-1	04/25/04	9.94	0.13	149.7	237	6.04	0.00	0.0	0.0
MW-4	04/26/04	11.03	0.27	38.7	867	6.51	2.90	0.1	0.0
OW-2	04/26/04	10.90	1.61	203.3	153	5.67	0.00	0.0	7.47
OW-5	04/25/04	11.11	0.40	151.3	360	5.98	0.00	0.0	0.7
OW-6	04/25/04	10.42	4.39	181.5	85	5.59	1.60	0.0	0.4
OW-8	04/25/04	9.81	0.11	-27.7	142	6.29	3.73	3.4	0.0
OW-10R	04/26/04	11.23	0.13	-14.0	476	6.39	0.70	3.4	0.4
OW-13	04/25/04	11.35	0.15	-11.0	394	6.19	0.35	2.8	0.0
OW-15	04/23/04	10.83	0.33	-49.3	293	6.26	3.30	3.2	0.0
OW-16	04/25/04	9.61	0.83	162.1	224	5.58	0.05	0.0	0.0
OW-18	04/23/04	9.95	0.08	-42.0	605	6.41	4.30	2.5	0.0
OW-19	04/24/04	11.06	0.40	-29.0	455	6.36	22.00	3.6	0.0
OW-21	04/24/04	12.51	0.33	-8.6	444	6.33	0.30	3.4	0.0
OW-22	04/24/04	12.65	0.37	-39.5	505	6.32	0.15	3.2	0.0

**Notes:**

<sup>(1)</sup>Negative turbidity readings due to calibration value of zero being greater than true zero; negative values are considered zero.

mg/L = milligrams per liter

mV = millivolts

uS/cm = microsiemens per centimeter

NTU = nephelometric turbidity units

**Table 2**  
**Groundwater and Surface Water Elevation Data<sup>(1)</sup>**  
**April 20, 2004**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**

Well No.	Top of Casing Elevation or Staff Gauge <sup>(2)</sup>	Screened Interval	Depth to Groundwater or Surface Water	Groundwater or Surface Water Elevation
MW-1	469.39	28.5 - 43.5	31.76	437.63
MW-4	470.13	35.3 - 50.3	38.20	431.93
OW-2	472.33	30.0 - 47.0	40.35	431.98
OW-3	472.70	30.0 - 46.5	41.11	431.59
OW-4	473.33	26.5 - 27.5	34.92	438.41
OW-5	459.85	25.5 - 45.5	28.02	431.83
OW-6	464.40	31.4 - 51.4	32.52	431.88
OW-7	459.31	24.5 - 34.5	27.69	431.62
OW-8	464.63	34.6 - 54.6	32.60	432.03
OW-9	472.91	25.3 - 35.3	28.96	443.95
OW-10R	469.27	29.0 - 39.0	28.95	440.32
OW-13	458.00	24.8 - 34.8	26.42	431.58
OW-15	472.05	22.0 - 32.0	11.85	460.20
OW-16	453.90	18.0 - 28.0	22.67	431.23
OW-17	447.18	11.0 - 21.0	16.03	431.15
OW-18	444.57	11.0 - 21.0	13.57	431.00
OW-19	438.69	5.0 - 15.0	8.14	430.55
OW-21	467.46	37.1 - 47.1	36.32	431.14
OW-22	467.10	38.0 - 48.0	35.97	431.13
OW-23	444.73	29.0 - 39.0	13.74	430.99
SW-1 <sup>(3)</sup>	432.06	-	-	429.26
SW-2 <sup>(3)</sup>	432.03	-	-	430.94
SW-3 <sup>(3)</sup>	440.10	-	-	436.80

**Notes:**

<sup>(1)</sup>Data reported in feet; elevations relative - mean sea level; 1988 National Geodetic Vertical Datum.

<sup>(2)</sup>Top of casing and gauge staff elevations surveyed by Maser Consulting P.A.

<sup>(3)</sup>Water elevation measured using surveying transit.

**Table 3**  
**Summary of Detected TCL Volatile Organic Compounds in Groundwater (ug/L)**  
**April 2004**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**

Compound	NYSDEC SGV	U.S. EPA MCL	MW-1 04/25/04	MW-4 04/26/04	OW-2 04/26/04	OW-5 04/25/04	OW-6 04/25/04	OW-8 04/25/04	OW-19R 04/26/04	OW-10R DUP 04/26/04	OW-13 04/25/04	OW-15 04/23/04	OW-16 04/25/04	OW-18 04/23/04	OW-19 04/24/04	OW-21 04/24/04	OW-22 04/24/04
Acetone	50 (G)	NE	<10	10 U	67 U	<10	<10	<10	<10	10 U	<17	<10	<10	10 U	10 U	<10	10 U
Benzene	1 (S)	5	<1.0	2.5	1.5 J	0.38 J	<1.0	<1.0	8.1	7.9	45	1.2	<1.0	3.3	2.1	2.9	4.2
Chlorobenzene	5 (S)*	100	<1.0	<1.0	<6.7	<1.0	<1.0	<1.0	2.2	2.1	0.38 J	0.67 J	<1.0	7.3	6.0	<1.0	5.9
Chloroethane	5 (S)*	NE	<2.0	<2.0	<13	<2.0	<2.0	<2.0	0.80 J	0.93 J	<3.3	<2.0	<2.0	0.74 J	1.3 J	<2.0	0.24 J
1,1-Dichloroethane	5 (S)*	NE	<1.0	<1.0	<6.7	<1.0	<1.0	<1.0	<1.0	<1.0	<1.7	0.29 J	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-Dichloroethene (total)	5 (S)*	70	<1.0	0.50 J	170	8.6	21	<1.0	0.57 J	0.61 J	4.9	<1.0	<1.0	<1.0	0.58 J	<1.0	<1.0
Ethylbenzene	5 (S)*	700	0.24 J	<1.0	<6.7	<1.0	<1.0	<1.0	0.22 J	0.26 J	1.0 J	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
Methylene Chloride	5 (S)*	5	<1.0	<1.0	<6.7	<1.0	<1.0	<1.0	1.0 U	1.0 U	<1.7	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
Tetrachloroethene	5 (S)*	5	<1.0	0.72 J	110	5.2	32	<1.0	<1.0	<1.0	<1.7	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
Toluene	5 (S)*	1,000	<1.0	<1.0	1.3 J	<1.0	<1.0	<1.0	0.27 J	0.28 J	0.31 J	<1.0	<1.0	<1.0	<1.0	<1.0	0.23 J
Trichloroethene	5 (S)*	5	<1.0	<1.0	22	2.5	6.3	<1.0	<1.0	<1.0	0.67 J	<1.0	<1.0	<1.0	<1.0	<1.0	<1.0
Vinyl Chloride	2 (S)	2	<2.0	<2.0	<13	<2.0	<2.0	<2.0	0.99 J	0.98 J	3.2 J	0.90 J	<2.0	<2.0	0.56 J	0.56 J	0.32 J
Xylenes (total)	5 (S)*	10,000	1.0	<1.0	<6.7	<1.0	<1.0	<1.0	<1.0	<1.0	<1.7	<1.0	<1.0	1.2	<1.0	<1.0	1.1

**Notes:**

TCL = Target Compound List

NYSDEC SGV = New York State Department of Environmental Conservation Standards (S) and Guidance (G) Values for groundwater

U.S. EPA MCL = United States Environmental Protection Agency Maximum Contaminant Level for drinking/groundwater

NE = Not established; no criteria specified

< = Analyte not detected at reporting limit

U (DATA VALIDATION QUALIFIER) = The analyte was analyzed for, but was not detected above the reported quantitation limit. In this table, "U" instead of "<" indicates a result modified through the data validation process.

J = Estimated result; result is less than reporting limit

\* = The principal organic contaminant (POC) standard for groundwater of 5 ug/L applies to this substance.

Red = Concentrations detected at or above regulatory limit

Blue = Analyte detected at less than regulatory limit, or analyte detected but no regulatory criteria specified.

**Table 4**  
**Natural Attenuation Parameters**  
**April 2004**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**

Well ID	Date	Alkalinity (mg/L)	Chloride (mg/L)	Dissolved Oxygen (mg/L)	Ethane (ug/L)	Ethene (ug/L)	Ferrous Iron (mg/L)*	Methane (ug/L)	Laboratory Nitrate (mg/L)	Redox (mV)	Sulfate (mg/L)	Sulfide (mg/L)	TOC (mg/L)
MW-1	04/25/04	120	4.6	0.13	<0.5	<0.5	0.0	0.26 J	5.4	149.7	12.1	<1.0	1
MW-4	04/26/04	200	149	0.27	<0.5	<0.5	3.4	340	1.0	38.9	128	<1.0	2
OW-2	04/26/04	45	5.2	-1.61	<0.5	<0.5	0.0	0.93 J	3.7	203.3	27.5	<1.0	<1
OW-5	04/25/04	110	20.5	0.40	<0.50	<0.50	0.0	0.21 J	3.4	151.3	73.6	<1.0	1
OW-6	04/25/04	27	1.2	-4.39	<0.5	<0.5	0.0	0.16 J	0.37	181.5	21.4	<1.0	<1
OW-8	04/25/04	64	2.4	0.11	<0.5	<0.5	3.4	1.0	<0.1	-27.7	16.1	<1.0	<1
OW-10R	04/26/04	280	2.4	0.13	<2.5	<2.5	0.1	500	<0.1	-14.0	35.3	<1.0	2
OW-13	04/25/04	250	2.7	0.15	0.19 J	0.38 J	2.8	140	<0.10	-11.0	14.4	<1.0	3
OW-15	04/23/04	150	3.3	0.33	<2.5	<2.5	3.2	440	<0.10	-49.3	5.9	<1.0	4
OW-16	04/25/04	82	10.8	-0.83	<0.5	<0.5	0.0	0.25 J	1.6	162.1	38.6	<1.0	<1
OW-18	04/23/04	320	10.2	0.08	<2.5	<2.5	2.5	470	<0.10	-42.0	15.2	2.5	11
OW-19	04/24/04	220	11.6	0.40	<2.5	<2.5	3.6	400	<0.10	-29.0	12.1	<1.0	7
OW-21	04/24/04	220	8.3	0.33	<0.50	<0.50	3.4	56	<0.10	-8.6	53.2	<1.0	2
OW-22	04/24/04	260	9.3	0.37	<2.5	<2.5	3.2	540	<0.10	-39.5	10	1.6	6

**Notes:**

mg/L = milligrams per liter

ug/L = micrograms per liter

mV = milliVolts

TOC = total organic carbon

" < " = Analyte not detected at method reporting limit.

B = Method blank contamination. The associated method blank contains the target analyte at a reportable level.

D = Result was obtained from the analysis of a dilution.

J = Estimated result; result is less than the reporting limit.

\*Ferrous iron was measured in the field (Hach kit).

NA = Not available

Detected TCL Volatile and Semivolatile Organic Compounds in Surface Water (ug/L)

1 of 5

Carroll and Dubies Superfund Site  
Town of Deerpark, Orange County, New York

Analyte	Acetone	Benzene	2-Butanone	Chloroethane	1,2-Dichloroethane	1,2-Dichloroethene (total)	Methylene Chloride	Toluene	Vinyl Chloride	Di-n-butyl phthalate
NYSDEC SGV	50 (G)	1 (S)	NE	5 (S)*	0.6 (S)	5 (S)*	5 (S)*	5 (S)*	2 (S)	50 (S)
U.S. EPA MCL	NE	5	NE	NE	5	70	5	1,000	2	NE

Sample ID	Date	VOCs									SVOC
SW-1	02/18/99	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	<10
	01/04/00	<10	0.19 J,B	<10	0.85 J	<1.0	0.15 J,B	<1.0	<1.0	0.99 J	<10
	06/08/00	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	<10
	03/15/01	<10	<1.0	<10	0.97 J	0.61 J	0.39 J	<1.0	<1.0	0.52 J	<10
	08/28/01	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	<10
	04/23/02	<10	<1.0	0.60 J	<2.0	<1.0	1.5 U	1.0 U	<1.0	<2.0	NA
	09/17/02	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	NA
	04/08/03	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	NA
SW-2	04/23/04	10 U	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	NA
	02/18/99	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	<10
	01/04/00	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	<10
	06/08/00	<10	<1.0	<10	<2.0	<1.0	<1.0	<1.0	<1.0	<2.0	<10
SW-2 / Duplicate	08/28/01	<10 / <1.0	<1.0 / <1.0	<10 / <1.0	<2.0 / <2.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<2.0 / <2.0	<10
	04/23/02	10 U / 10 U	<1.0 / <1.0	<10 / <1.0	<2.0 / <2.0	<1.0 / <1.0	1.2 U / 1.0 U	<1.0 / <1.0	<1.0 / <1.0	<2.0 / <2.0	0.77 J / <10
	09/17/02	<10 / <1.0	<1.0 / <1.0	<10 / <1.0	<2.0 / <2.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<2.0 / <2.0	NA
	04/08/03	1.3 J / 2.0 J	<1.0 / <1.0	<10 / <1.0	<2.0 / <2.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<2.0 / <2.0	NA
Duplicate	04/23/04	<10 / <1.0	<1.0 / <1.0	<10 / <1.0	<2.0 / <2.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<1.0 / <1.0	<2.0 / <2.0	NA

Notes:

TCL = Target Compound List

NYSDEC SGV = New York State Department of Environmental Conservation Standards (S) and Guidance (G) values for groundwater.

\* = The principal organic contaminant (POC) standard for groundwater of 5 ug/L applies to this substance.

U.S. EPA MCL = United States Environmental Protection Agency Maximum Contaminant Level for drinking/groundwater.

NE = Not established; no criteria specified.

> = Analyte not detected at reporting limit.

J = Estimated result; result is less than reporting limit.

B = Method blank contamination. The associated method blank contains the target analyte at a reportable level.

U (DATA VALIDATION QUALIFIER) = The analyte was analyzed for, but was not detected above the reported sample quantitation limit.

LA = Lab accident; during the concentration process, sample was inadvertently concentrated to dryness due to analyst error. Insufficient sample volume was received to re-extract; sample could not be analyzed.

NA = Not analyzed

Blue = Analyte detected at less than regulatory limit, or analyte detected but no regulatory criteria specified.

Red = Analyte detected at or above SGV or MCL.

**Table 6**  
**Detected TCL Volatile and Semivolatile Organic Compounds in Sediment (ug/kg)**  
**April 2004**  
**Carroll and Dubles Superfund Site**  
**Town of Deepark, Orange County, New York**

Analyte	Date	VOCs					SVOCs		
		Acetone	Benzene	2-Butanone	Carbon Disulfide	1,2-Dichloro-ethene (total)	Toluene	Vinyl Chloride	bis(2-Ethylhexyl) phthalate
	09/27/94	58	ND	<20	NR	ND	ND	<20	ND
	02/18/99	<28	<6.9	<28	<6.9	<6.9	<14	<450	77 J
	01/04/00	370	<31	82 J	<31	<31	6.9 J	<2,000	<2,000
	06/08/00	60 J,B	<13	17 J	<13	<13	<27	590 J	<880
	03/15/01	55 J	<16	<62	<16	<16	<31	<1,000	<1,000
	08/28/01	27 J	2.1 J	9.4 J	<12	4.6 J	<24	<790	<790
	04/23/02	1,100 R	<57 R	260 R	<57 R	<57 R	<110 R	<3,800 R	<3,800 R
	09/17/02	180 B,J	2.7 J	58 J	3.5 J	2.1 J	54 J	34 UJ	NA
	04/08/03	110 J	3.4 J	34 J	<21	5.7 J	<21	3.0 J	NA
	04/23/04	28 J,FB,TB	<10	7.2 J	<10	<10	<20	NA	NA
SED 1	09/27/97	76	ND	<23	NR	ND	ND	<23	ND
	02/18/99	140 J,B	<44	50 J	<44	<44	<88	<2,900	<2,900
	01/04/00	180 J / <190	<55 / <47	<220 / <190	<55 / <47	<55 / <47	<110 / <94	<3,600 / <3,100	<3,600 / <3,100
	06/08/00	150 J / <160	<46 / <41	49 J / <160	<46 / <41	<46 / <41	<91 / <81	2,900 J / 1,500 J	<3,000 / <2,700
	03/15/01	36 UJ / 69 UJ	17 UJ / 17 UJ	70 UJ / 69 UJ	<17 / <17	17 UJ / 17 UJ	35 UJ / 35 UJ	<1,200 / <1,100	<1,200 / <1,100
	08/28/01	44 J / 22 J	<16 / <13	14 J / 7.6 J	<16 / <13	<16 / <13	<32 / <25	<1,100 / <830	<1,100 / <830
	04/23/02	63 J / 85 UJ	30 UJ / 21 UJ	21 J / 85 UJ	30 UJ / 21 UJ	30 UJ / 21 UJ	59 UJ / 42 UJ	2,000 UJ / 1,400 UJ	2,000 UJ / 1,400 UJ
	09/17/02	40 B / 29 J,B	<9.6 / <9.3	17 J / 9.3 J	<9.6 / <9.3	<9.6 / <9.3	1.0 J / 0.91 J	<19 / <19	NA
SED-2 / Duplicate	04/08/03	79 J / 27 J	<41 / <44	21 J / <180	<41 / <44	<41 / <44	<41 / <44	<83 / <88	NA
	04/23/04	38 J,FB / <53	<14 / <13	12 J / <53	<14 / <13	<14 / <13	<14 / <13	<28 / <27	NA

**Notes:**  
TCL = Target Compound List  
ND = Not detected at reporting limit prior to 06/08/00.  
< = Not detected at the method detection limit.  
NR = Analyte not reported  
B = Method blank contamination. The associated method blank contains the analyte at a reportable level.  
R (DATA VALIDATION QUALIFIER) = The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet QC criteria. The presence or absence of the analyte cannot be verified.  
# = This value represents a probable combination of 3-methylphenol (m-creso) and 6-methylphenol (p-creso).  
UJ (DATA VALIDATION QUALIFIER) = Analyte not detected above the reporting limit; however, the reporting limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.  
Methylene Chloride (2.6 ug/kg J, FB, TB) was detected in SED-1 during 4/04 sampling round.

## Figures





1. THE MONITORING/RECOVERY WELLS AND LIMITED SITE LOCATION OF PHYSICAL FEATURES SHOWN HEREON ARE BASED ON A FIELD SURVEY MADE FOR PROPER ORIENTATION WITH A FULL SITE PLAN BEING PREPARED BY SHIELD ENVIRONMENTAL ASSOC., INC.

2. THE HORIZONTAL POSITION OF THE HEREIN SURVEYED "CARROLL AND DUBES" SUPERFUND SITE IS BASED ON NEW YORK GEODETIC CONTROL MONUMENTS PID # LY2611 (LAUREL) AND PID # AB3870 (DIANNE).

ADJUSTED TO NAD 1983 WITHIN THE NEW YORK STATE PLANE COORDINATE SYSTEM

NOTE: THE OUTLINES OF THE FORMER LAGOONS ARE APPROXIMATE BASED ON THE ACTUAL EXCAVATION

SOURCE: MASER CONSULTING P.A., MONITORING WELL LOCATION PLAN.  
INDEX N). SU0009, MARCH 3, 1999.

SED-2/SW-2 O

O SED-1/SW-1



**SHIELD**  
ENVIRONMENTAL ASSOCIATES, INC.  
MONROEVILLE, PENNSYLVANIA

CARROLL AND DUBIES SUPERFUND SITE  
TOWN OF DEERPARK, ORANGE COUNTY, NEW YORK

698-0110

FIGURE 2  
SITE PLAN/MONITORING  
WELL LOCATIONS  
APRIL 2004

							CADD FILE	80110-22	CURRENT DATE:	05/04/04
							DRAWING NO.	FIGURE 2	REVISION	

OW-15 MONITORING WELL ID  
(436.81) GROUNDWATER/SURFACE WATER ELEVATION  
● MONITORING WELL LOCATION  
○ SEDIMENT/SURFACE WATER  
SAMPLE LOCATION  
CONTOUR INTERVAL AS INDICATED  
CONTOUR LINE DASHED WHERE INFERRED.



1. THE MONITORING/RECOVERY WELLS AND LIMITED SITE LOCATION OF PHYSICAL FEATURES SHOWN HEREON ARE BASED ON A FIELD SURVEY MADE FOR PROPER ORIENTATION WITH A FULL SITE PLAN BEING PREPARED BY SHIELD ENVIRONMENTAL ASSOC., INC.
2. THE HORIZONTAL POSITION OF THE HEREIN SURVEYED "CARROLL AND DUBES" SUPERFUND SITE IS BASED ON NEW YORK GEODETIC CONTROL MONUMENTS PID # LY2611 (LAUREL) AND PID # AB3870 (DIANNE).  
ADJUSTED TO NAD 1983 WITHIN THE NEW YORK STATE PLANE COORDINATE SYSTEM

**SOURCE:** MASER CONSULTING P.A., MONITORING WELL LOCATION PLAN.  
INDEX N). SU0009, MARCH 3, 1999.



**698-0110**

**FIGURE 3  
GROUNDWATER CONTOUR MAP  
APRIL 2004**

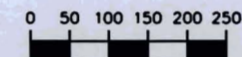
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0								
NO	DRWN	DATE	REVISION		CHKD	DATE	APPRD	DATE

CADD FILE	80110-22	CURRENT DATE:	05/03/04
DRAWING NO.	FIGURE 3	REVISION	

## OW-15 MONITORING WELL ID

- ALL CONCENTRATIONS IN  $\mu\text{g/L}$ .

RED VALUES EXCEED SGV OR MCL



### INFORMATION OF FACT

1. THE MONITORING/RECOVERY WELLS AND LIMITED SITE LOCATION OF PHYSICAL FEATURES SHOWN HEREON ARE BASED ON A FIELD SURVEY MADE FOR PROPER ORIENTATION WITH A FULL SITE PLAN BEING PREPARED BY SHIELD ENVIRONMENTAL ASSOC., INC.
2. THE HORIZONTAL POSITION OF THE HEREIN SURVEYED "CARROLL AND DUBES" SUPERFUND SITE IS BASED ON NEW YORK GEODETIC CONTROL MONUMENTS PID # LY2611 (LAUREL) AND PID # AB3870 (DIANNE).

ADJUSTED TO NAD 1983 WITHIN THE NEW YORK STATE PLANE COORDINATE SYSTEM

NOTE: THE OUTLINES OF THE FORMER LAGOONS ARE APPROXIMATE BASED ON THE ACTUAL EXCAVATION.

**SOURCE:** MASER CONSULTING P.A., MONITORING WELL LOCATION PLAN.  
INDEX N). SU0009, MARCH 3, 1999.



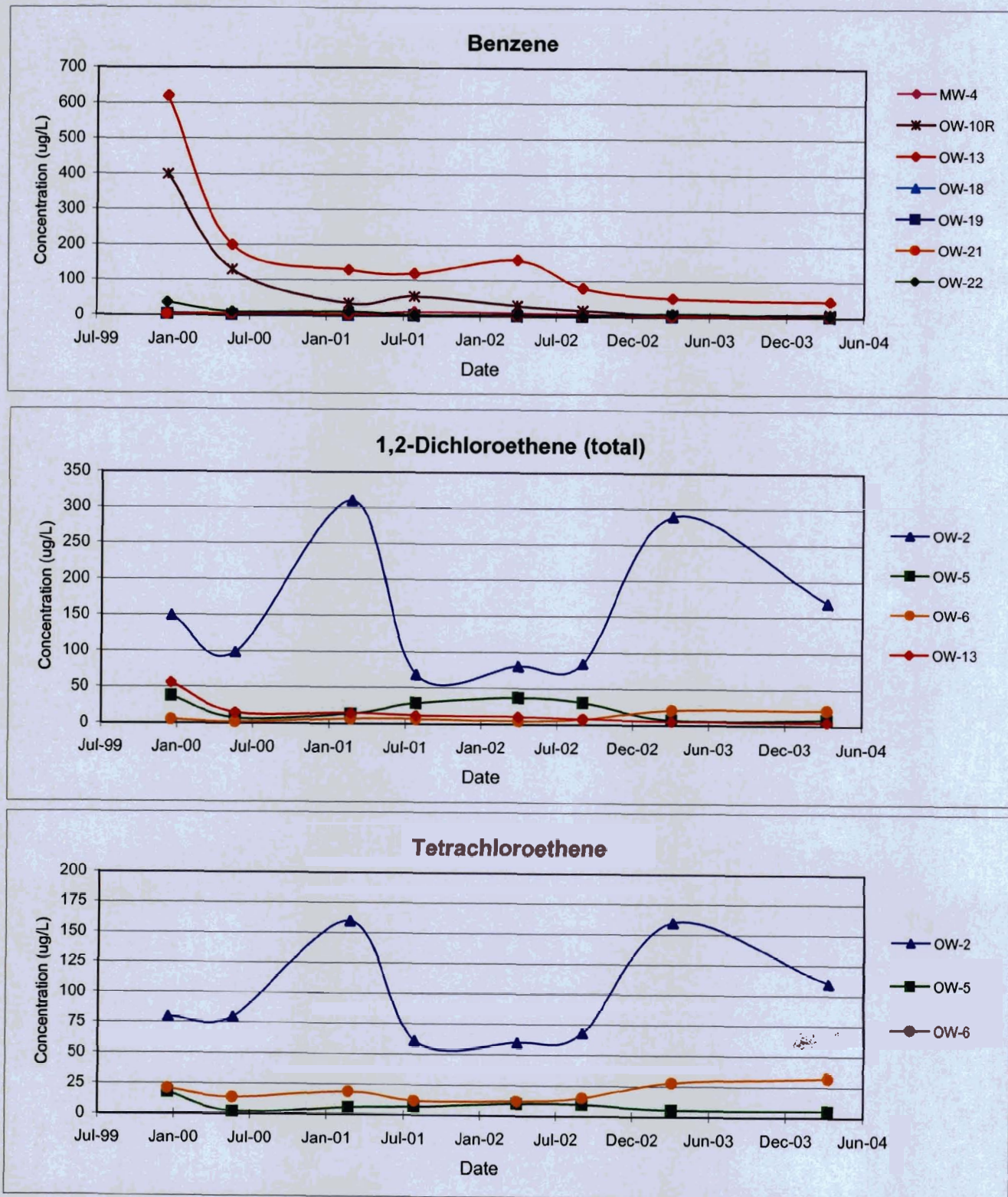
**CARROLL AND DUBIES SUPERFUND SITE**  
**TOWN OF DEER PARK, ORANGE COUNTY, NEW YORK**  
 698-0110

1							
0							
NO	DRWN	DATE	REVISION		CHKD	DATE	APPRD DATE

**FIGURE 4  
DETECTED VOLATILE ORGANIC  
COMPOUNDS IN GROUNDWATER  
APRIL 2004**

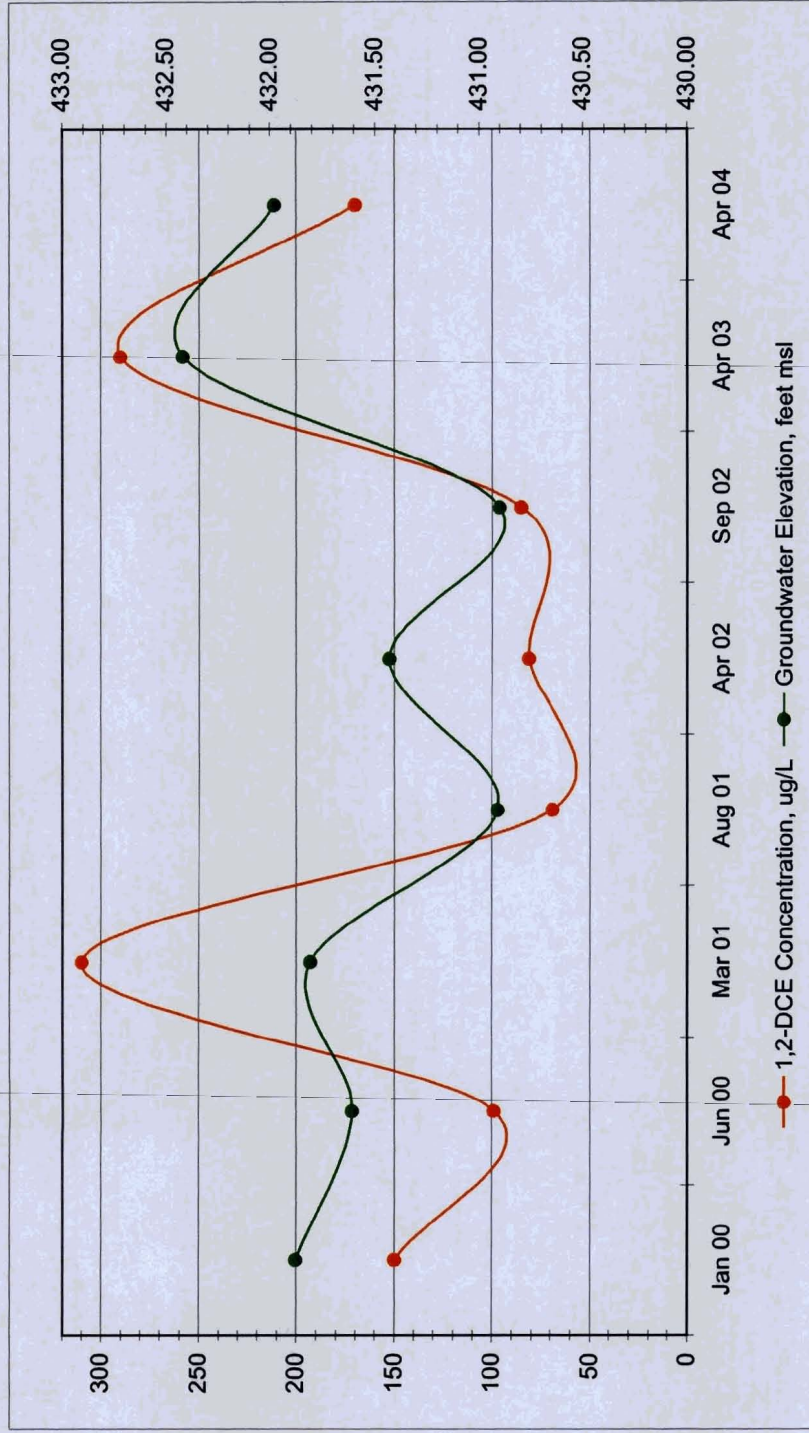
CADD FILE	80110-22	CURRENT DATE:	05/17/04
DRAWING NO.	FIGURE 4	REVISION	

**Figure 5**  
**Concentration Trends for Principal VOCs in Groundwater<sup>(1)</sup>**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**

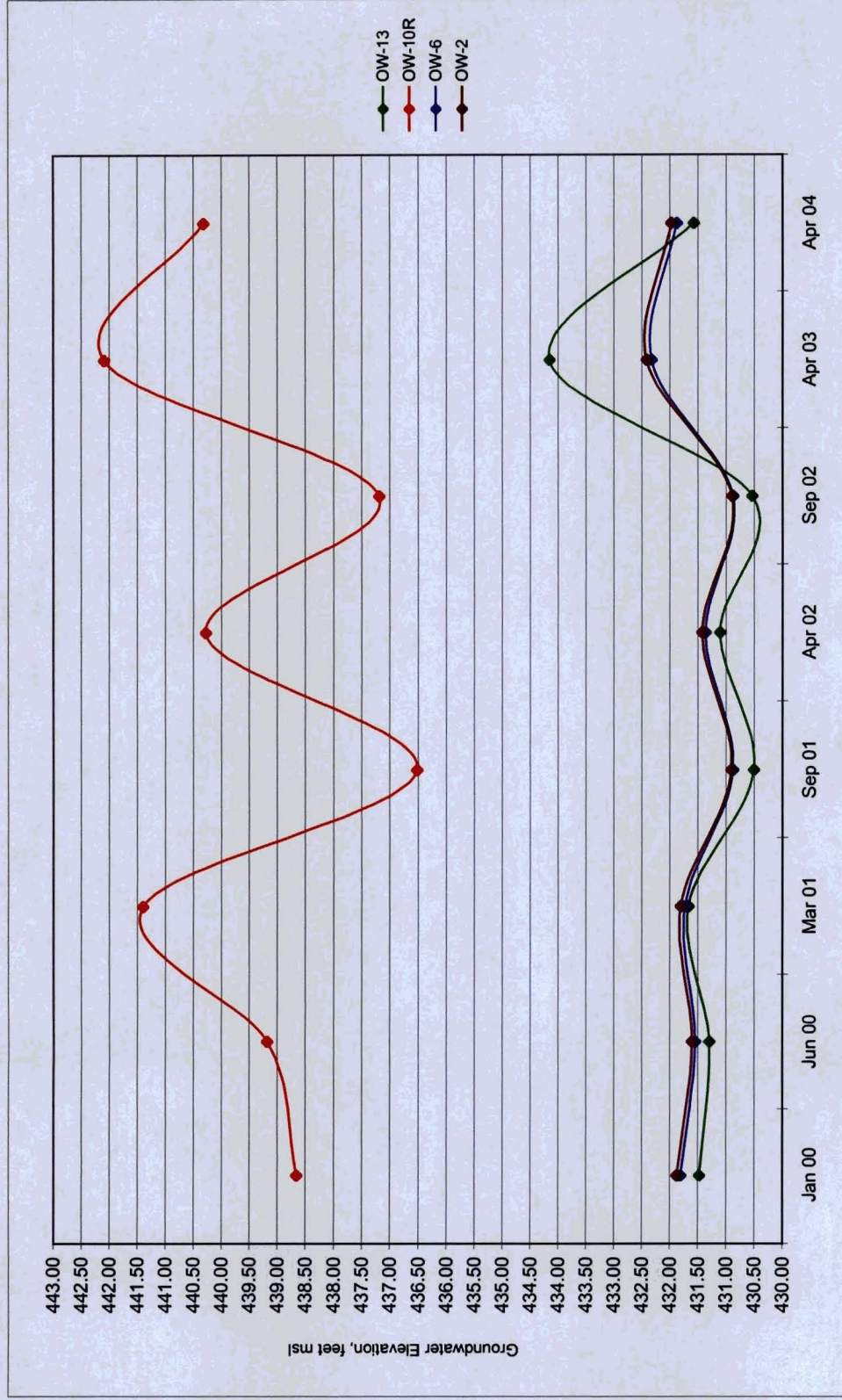


<sup>(1)</sup>In monitoring wells where criteria have been exceeded.

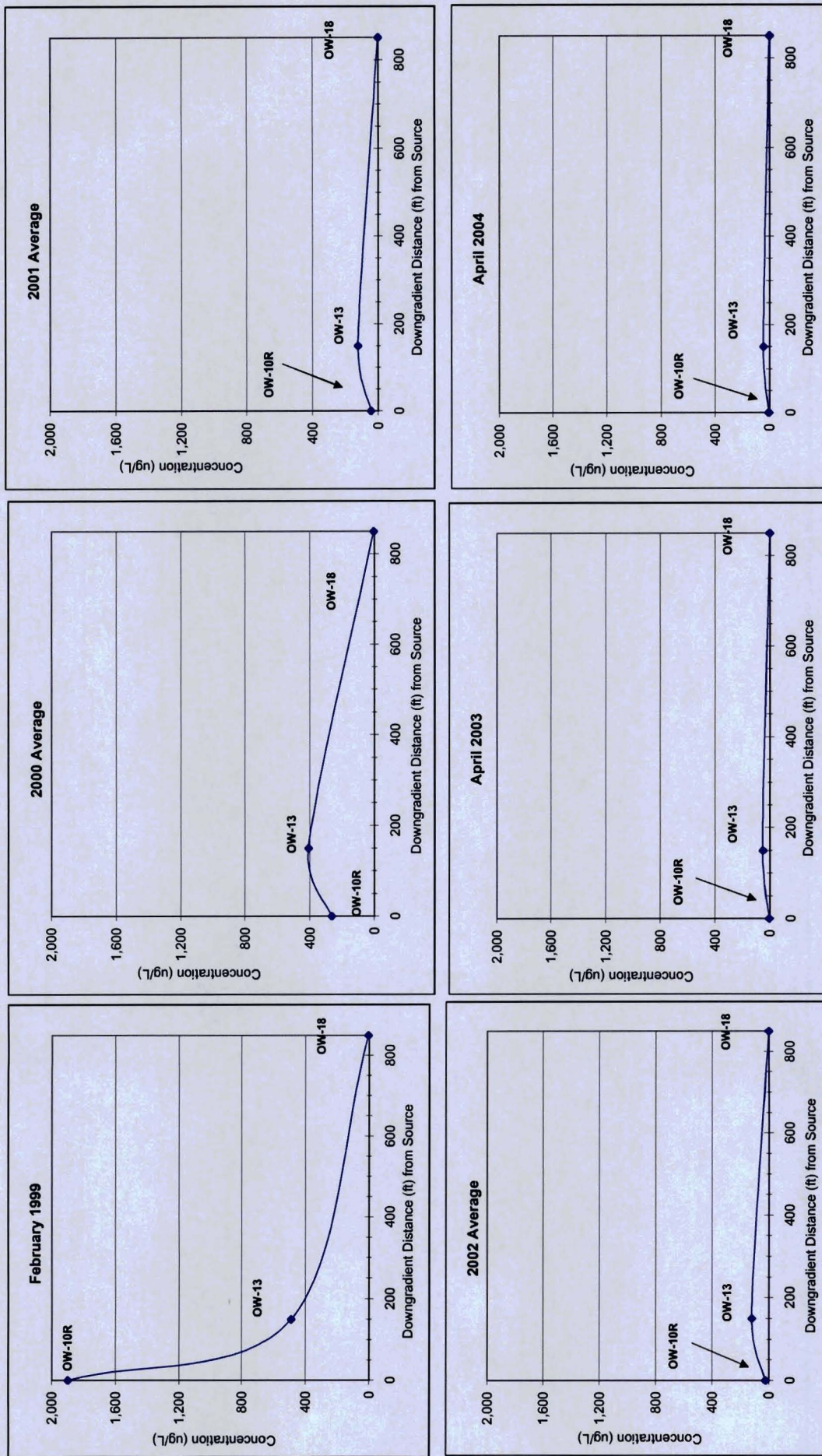
**Figure 6**  
**Relationship Between Groundwater Elevations**  
**and 1,2-Dichloroethene Concentrations in OW-2**



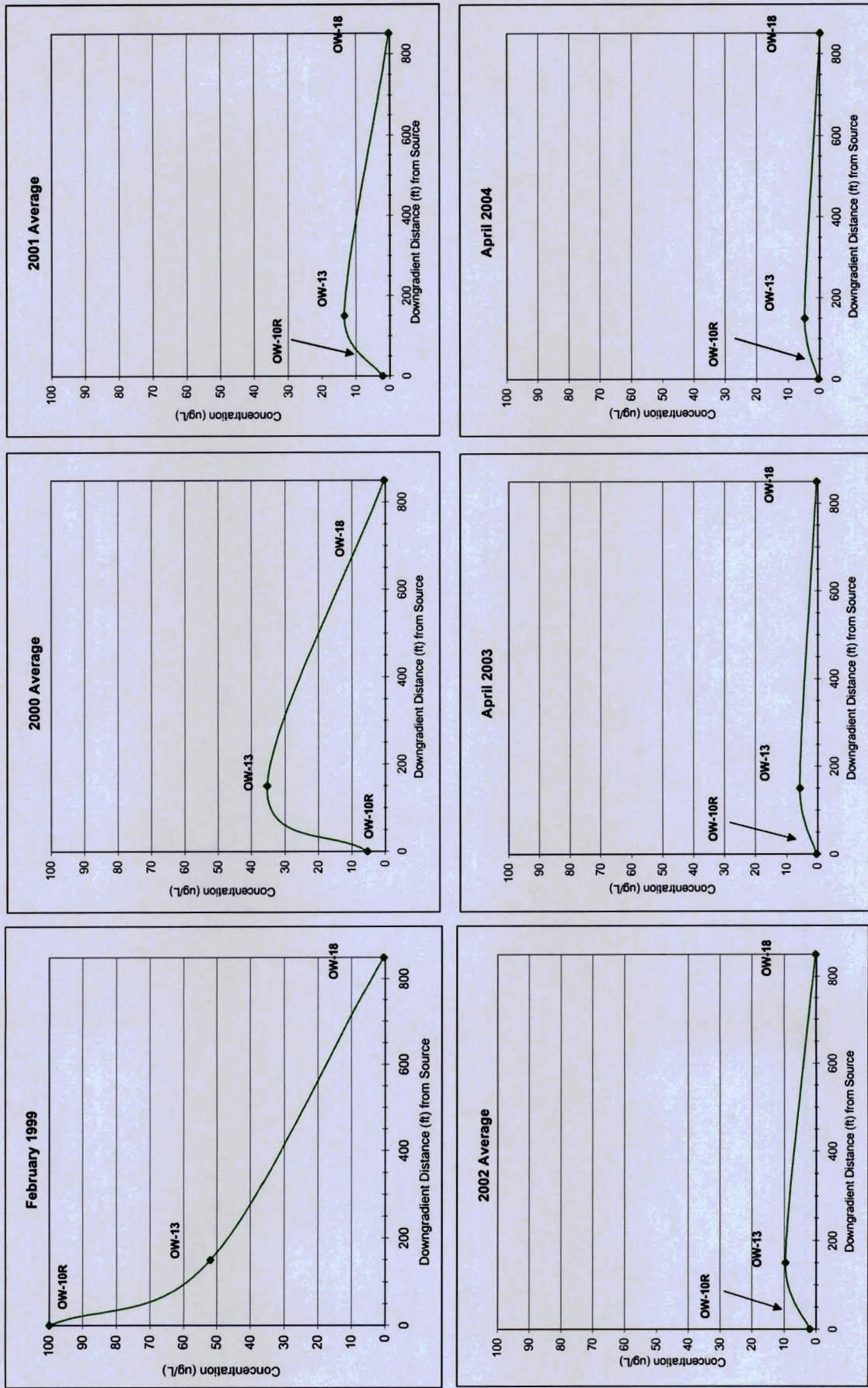
**Figure 7**  
**Hydrographs for Representative Monitoring Wells**



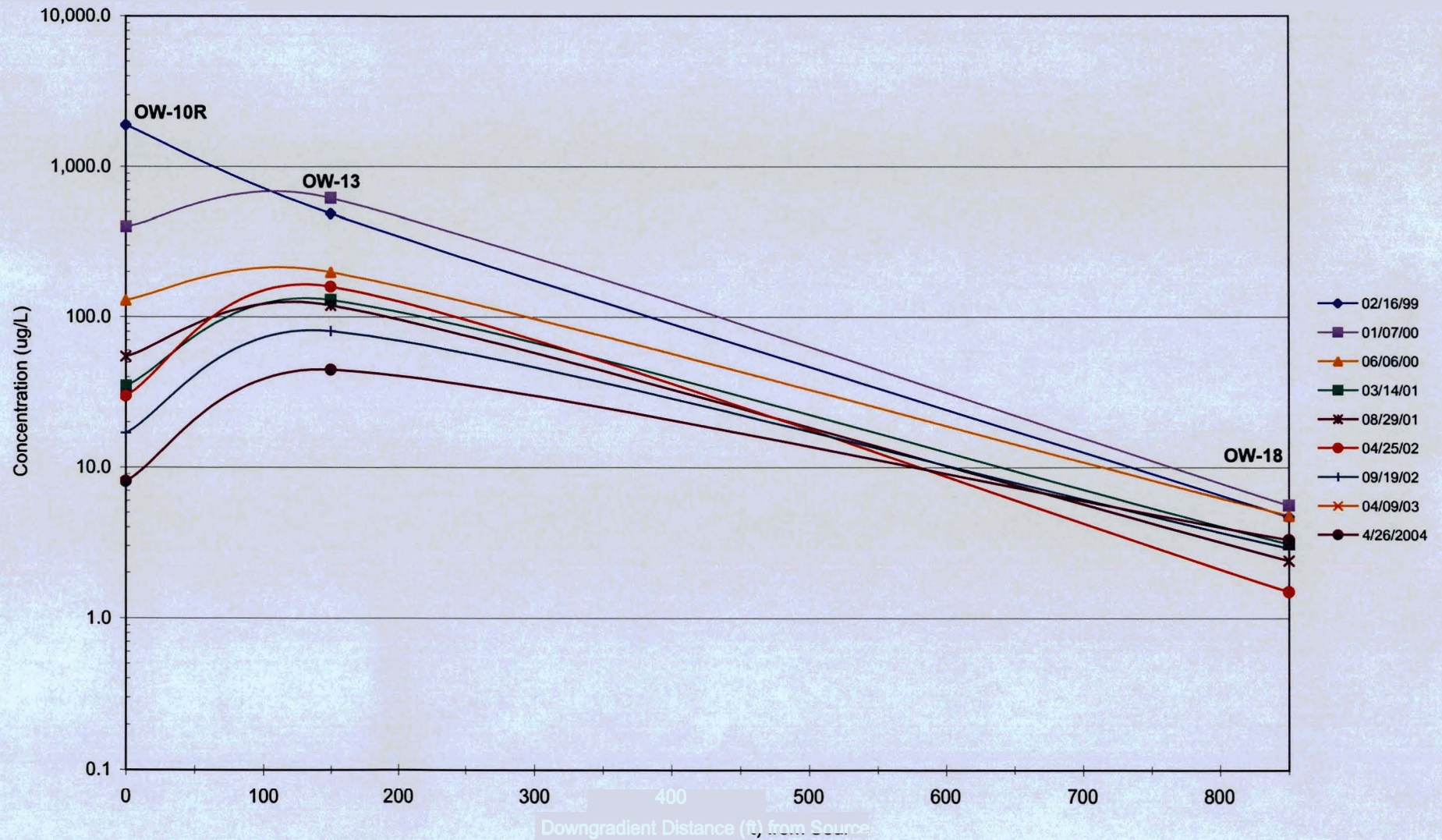
**Figure 8**  
**Concentration Trends Across Plume for Benzene in Groundwater**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**



**Figure 9**  
**Concentration Trends Across Plume for 1,2-DCE (total) in Groundwater**  
**Carroll and Dubies Superfund Site**  
**Town of Deepark, Orange County, New York**



**Figure 10**  
**Benzene Concentration in Groundwater Over Duration of the Monitoring Period**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**



**Figure 11**  
**1,2-DCE (total) Concentration in Groundwater Over Duration of the Monitoring Period**  
**Carroll and Dubies Superfund Site**  
**Town of Deerpark, Orange County, New York**

