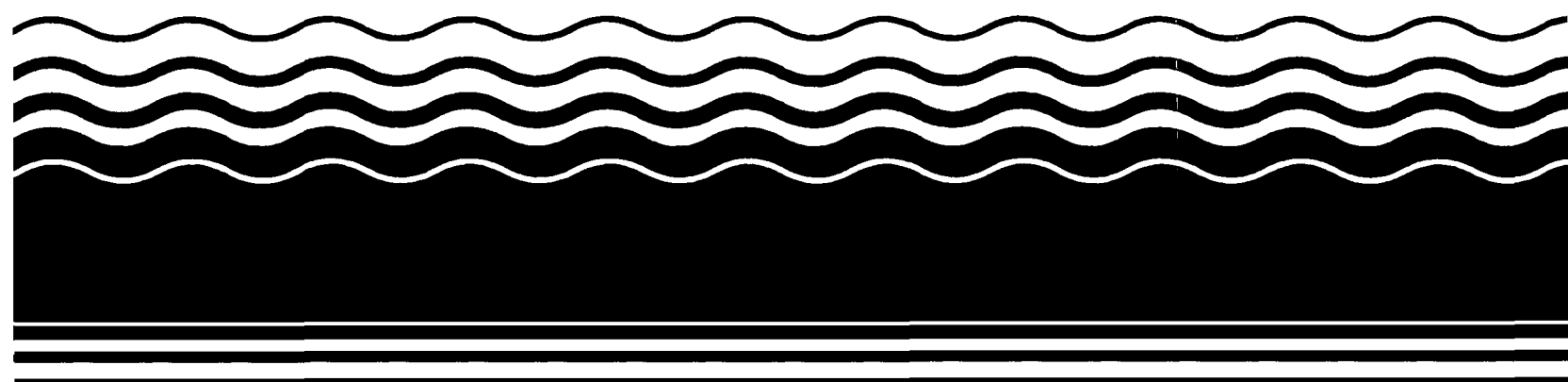


EPA/ROD/R02-95/250
August 1995

EPA Superfund
Record of Decision:

Chemical Insecticide Corp., OU2
Edison Township, NJ
3/28/95



DECLARATION FOR THE RECORD OF DECISION

SITE NAME AND LOCATION

Chemical Insecticide Corporation

Edison Township, Middlesex County, New Jersey

STATEMENT OF BASIS AND PURPOSE

This decision document presents the selected remedial action for addressing contaminated soil and sediment in off-site residential areas and areas in and immediately adjacent to the unnamed tributary and Mill Brook associated with the Chemical Insecticide Corporation site. The remedial action was chosen in accordance with the requirements of the Comprehensive Environmental Response, Compensation, and Liability Act of 1980, as amended, and to the extent practicable, the National Oil and Hazardous Substances Pollution Contingency Plan. This decision document explains the factual and legal basis for selecting the remedy for the site.

The New Jersey Department of Environmental Protection concurs with the selected remedy. The information supporting this remedial action decision is contained in the administrative record for the site.

ASSESSMENT OF THE SITE

Actual or threatened releases of hazardous substances from the Chemical Insecticide Corporation site, if not addressed by implementing the response action selected in this Record of Decision, may present an imminent and substantial endangerment to public health, welfare, or the environment.

DESCRIPTION OF THE SELECTED REMEDY

The response action described in this document represents the second of three planned remedial phases for the Chemical Insecticide Corporation site. It addresses contaminated soil and sediment in residential areas and areas in and adjacent to the unnamed tributary and Mill Brook.

A previous Record of Decision, signed on September 29, 1989, selected an interim remedial action to control contaminated runoff from the site. Construction of this interim action, representing the first remedial phase for the site, was completed in September 1994.

The final remedy for the site will be selected after the completion of additional soil washing treatability studies. It will address contaminated soil on the site and in off-site industrial areas as well as any associated groundwater contamination.

The major components of the selected remedy for this second remedial phase include the following:

- excavation of approximately 10,000 cubic yards of soil and sediment containing arsenic at levels greater than 20 parts per million;
- appropriate off-site disposal of contaminated soil and sediment; and
- restoration of the excavated areas to the extent practicable.


The selected remedy provides protection of human health and the environment by removing contaminated soil and sediment from the above-described areas and restoring such areas.

Because this remedy does not include on-site containment of contaminated material, long-term management and controls will not be necessary.

DECLARATION OF STATUTORY DETERMINATIONS

The selected remedy is protective of human health and the environment, complies with Federal and State requirements that are legally applicable or relevant and appropriate, and is cost-effective. It utilizes permanent solutions and alternative treatment (or resource recovery) to the maximum extent practicable for the areas to be remediated. However, treatment of the principal threats associated with the areas to be remediated was not found to be practicable due to the unavailability of an effective treatment technology at the present time. Therefore, this remedy does not satisfy the statutory preference for treatment as a principal element.

A five-year review of the remedy will not be necessary, because it will not result in hazardous substances remaining in the remediated areas above health-based levels.



William J. Muszyński, P.E.
Deputy Regional Administrator

3-28-95
Date

RECORD OF DECISION DECISION SUMMARY

**Chemical Insecticide Corporation Site
Edison Township, Middlesex County, New Jersey**

**United States Environmental Protection Agency
Region II
New York, New York**

TABLE OF CONTENTS

	<u>PAGE</u>
SITE LOCATION AND DESCRIPTION	1
SITE HISTORY AND ENFORCEMENT ACTIVITIES	1
HIGHLIGHTS OF COMMUNITY PARTICIPATION	3
SCOPE AND ROLE OF ACTION	4
SUMMARY OF SITE CHARACTERISTICS	5
SUMMARY OF SITE RISKS	7
Human Health Assessment	7
Ecological Risk Assessment	9
REMEDIAL ACTION OBJECTIVES	12
DESCRIPTION OF REMEDIAL ALTERNATIVES	13
SUMMARY OF COMPARATIVE ANALYSIS OF ALTERNATIVES	15
SELECTED REMEDY	20
STATUTORY DETERMINATIONS	22

ATTACHMENTS

APPENDIX I.	FIGURE 1
APPENDIX II.	DATA
APPENDIX III.	ADMINISTRATIVE RECORD INDEX
APPENDIX IV.	STATE LETTER OF CONCURRENCE
APPENDIX V.	RISK ASSESSMENT DOCUMENTS
APPENDIX VI.	RESPONSIVENESS SUMMARY

SITE LOCATION AND DESCRIPTION

The Chemical Insecticide Corporation (CIC) site is located at 30 Whitman Avenue in Edison Township, Middlesex County, New Jersey. The site occupies approximately 6 acres, bordered on the north by Route 287 and on the east, west and south by industrial properties. There are no permanent surface water bodies on the CIC site. After heavy precipitation, the surface water runoff drains toward the northeast corner of the site where it discharges into an underground conduit which flows into an unnamed tributary of Mill Brook. Mill Brook, in turn, flows into the Raritan River approximately four miles downstream of the site. Both the unnamed tributary and Mill Brook run through residential areas, including two apartment complexes (Edison Glen and Edison Woods). Figure 1 of Appendix I indicates the site location and the surface water flow pathway. The residents of these complexes obtain potable water from a public water supply system and none of these water bodies are used as a drinking water source downstream of the site.

SITE HISTORY AND ENFORCEMENT ACTIVITIES

CIC owned and operated the site from 1954 to 1970. During this time, the site was used for manufacturing insecticides, fungicides, rodenticides and herbicides. These manufacturing activities combined with poor housekeeping led to widespread chemical contamination at the site, as well as migration of contaminants to off-site areas.

Over the period of CIC operations, the Edison Department of Health and Human Resources (EDHHR) became concerned about activities on the site due to numerous neighborhood complaints of odors, documented off-site discharges and releases, and the frequency of on-site fires. In the late 1960s, EDHHR ordered the cessation of on-site discharges of wastewater, required the disposal of leaking drums to eliminate the odor problem, and the closing of on-site lagoons. CIC declared bankruptcy in 1970. The property was purchased in anticipation of future development by Piscataway Associates, which demolished the production facilities on the site in 1975.

Triggered primarily by the potential for the presence of dioxin (a contaminant generated in the production of 2,4,5-trichlorophenoxyacetic acid, a herbicide which was handled on the premises), both the U.S. Environmental Protection Agency (EPA) and the New Jersey Department of Environmental Protection (NJDEP) performed on-site and off-site field investigations at CIC, testing soils and surface water for dioxin in 1983. In 1984, NJDEP further sampled on-site soils for the presence of other pollutants.

Based upon the analytical results from these investigations, EPA Region II authorized a remedial investigation and feasibility

study (RI/FS) for the site. On-site and off-site field investigations were performed over the period from July 1987 through March 1988. In August 1990, EPA included the CIC site on its National Priorities List of Superfund sites.

Concurrent with the RI/FS work, EPA conducted several immediate actions to mitigate risks associated with the continuing problem of contaminated surface water runoff from the site. These included installation of a fence around site drainage areas, improvements to site drainage controls and cleanup of contaminated runoff from the adjacent Metroplex parking lot. However, these limited response actions only partially addressed the surface water runoff problem, in that the contaminated runoff would continue to migrate to downstream waterways (i.e., the unnamed tributary and Mill Brook), creating the potential for adverse impacts to human health and the environment.

Interim Remedy

On September 29, 1989, EPA issued a Record of Decision (ROD) selecting an interim remedial action to control contaminated runoff from the CIC site. The major features of the selected interim remedial action include: clearing and grading of the site; covering the site with a surficial cap; construction of a surface water runoff diversion system; and controlled release of collected, uncontaminated surface water runoff from the site. This action represents the first cleanup phase for the entire site. Construction of this interim remedy was completed in September 1994. Although it was anticipated that this interim action would be completed in 1992, remedial construction was delayed due to contractual problems as well as to address concerns regarding the potential for buried explosives at the site. Based on a subsequent explosives investigation, no buried explosives were identified within the top two feet of surface soil.

Final Remedy

While proceeding with the interim remedial action, EPA continued with the RI/FS, collecting additional samples and evaluating final solutions for site-wide contamination. The combination of chemical contaminants at the CIC site (herbicides, pesticides and metals) requires the performance of treatability tests prior to the identification of appropriate alternatives to remedy the site. EPA has performed bench-scale incineration, solidification and soil washing treatability testing since the issuance of the interim remedy ROD in September 1989. Soil washing was determined to be ineffective in remediating site soils using the soil washing procedure applied during the treatability test. Therefore, the draft final feasibility study, which EPA issued in March 1994, dismissed soil washing from further evaluation as a potential remedial alternative for the CIC site.

EPA has been working closely with the CIC Technical Assistance Grant (TAG) Committee and its technical advisor regarding the selection of a final remedy for the site. Based on its review of the draft feasibility study, the TAG Committee suggested further study of soil washing as a potential viable remedial alternative for the CIC site. EPA has also further explored the viability of soil washing with experts at its Office of Research and Development which recommended pursuit of further optimization tests based on new information. In addition, on July 27, 1994, the Municipal Council of the Township of Edison adopted a resolution requesting that EPA further evaluate the feasibility of soil washing at CIC. Consequently, EPA has decided to perform a follow-up soil washing treatability study involving the soils on the CIC site. EPA expects that the additional treatability study will be completed in 1996.

If, upon further study, soil washing appears effective as a potential remedial alternative, it will be evaluated in a revised draft feasibility study report. Based on the results of the feasibility study, EPA will release another proposed plan for public comment which will identify the Agency's preferred final remedy to address contaminated soil and groundwater. EPA expects to propose this final remedy during 1997.

Remedy for Off-site Areas

Concurrent with the RI/FS and construction of the interim remedy, EPA performed additional investigations in off-site areas to determine the nature and extent of off-site contamination. Based on the data generated during these investigations, EPA is documenting its selected remedy for addressing contaminated soil and sediment in residential areas and areas in and immediately adjacent to the unnamed tributary and Mill Brook.

Enforcement Activities

Since the Chemical Insecticide Corporation is no longer in existence, EPA has not been able to take enforcement action against CIC for the problems caused by the company. However, EPA has notified Mr. Arnold M. Livingston, who was the president of CIC, of his potential liability with respect to the site.

HIGHLIGHTS OF COMMUNITY PARTICIPATION

EPA has been working closely with the Edison Wetlands Association, CIC Technical Assistance Grant (TAG) Committee, CIC Citizen's Advisory Committee, public officials and all other interested and concerned members of the community. Their participation and contributions to the site investigation/remediation process have been and continue to be beneficial towards achieving the Agency's goal of effectively protecting

human health and the environment.

The Proposed Plan and supporting documentation for the second cleanup phase associated with the CIC site were released to the public for comment on November 7, 1994. These documents were made available to the public in the administrative record repositories maintained at the EPA Region II office (formerly, 26 Federal Plaza and currently, 290 Broadway, New York, New York), the Edison Township Municipal Complex (100 Municipal Boulevard, Edison, New Jersey), the Edison Library (340 Plainfield Avenue, Edison, New Jersey), and the Metuchen Library (480 Middlesex Avenue, Metuchen, New Jersey). A notice of availability for these documents was published in The Star-Ledger on November 7, 1994. A public comment period on the documents was held from November 7, 1994 to December 7, 1994.

On November 28, 1994, EPA and the New Jersey Department of Environmental Protection conducted a public meeting at the Edison Township Municipal Building, to inform local officials and interested citizens about the Superfund process, to review the planned remedial activities at the site, and to respond to any questions from area residents and other attendees.

Responses to the comments recieved at the public meeting and in writing during the public comment period are included in the Responsiveness Summary (see Appendix V).

SCOPE AND ROLE OF ACTION

The CIC site, as characterized by the field investigations, is extremely complex due to the number and variety of contaminants present, the concentrations of contaminants, the physical and geological characteristics of the site, and the many potential migration routes for these contaminants. Consequently, EPA has divided the response actions for the site into several remedial phases as follows:

Interim Remedy: this remedial phase addressed the contaminated surface water runoff conditions at the CIC site.

Final Remedy: this remedial phase will address contaminated soil on the site and in off-site industrial areas, as well as groundwater contamination associated with the CIC site. A final remedy will be selected after the completion of additional soil washing treatability studies, as described above.

Off-Site Remedy: this will address the remediation of contaminated soil and sediment in off-site areas associated with the CIC site, as outlined in this ROD.

Since the purpose of this ROD is to document EPA's selected

remedy for addressing soil and sediment contamination in off-site areas, the subsequent summary of site characteristics focuses only on the off-site investigation efforts.

SUMMARY OF SITE CHARACTERISTICS

Prior to construction of the above-described interim remedy, contaminated surface water runoff drained toward the northeast corner of the site where it discharged into an underground conduit which flows into an unnamed tributary of Mill Brook. As a result, off-site areas in and around the unnamed tributary and Mill Brook are contaminated with contaminants from the CIC site.

Arsenic was used in pesticides produced by CIC during the 1950s and 1960s and is the primary contaminant of concern involving the off-site areas. Arsenic also occurs naturally in soil. NJDEP has determined the upper limit of naturally occurring arsenic for New Jersey soils to be 20 parts per million (ppm).

In October 1992, EPA collected five soil samples in residential areas near the unnamed tributary and Mill Brook and found elevated levels of arsenic. Although the levels detected (up to 79.7 ppm) did not indicate an immediate risk to human health, additional soil sampling to evaluate the long-term risk due to exposure to arsenic contaminated soils was considered appropriate.

As a result, in July 1993, the EPA Environmental Response Team (ERT) conducted additional soil sampling in areas near the aforementioned streams. These areas included the Edison Glen and Edison Woods developments, areas in Metuchen near Route 287, and properties on Prince Street, Patrick Street and Cortlandt Street in Edison, New Jersey. Four samples of dirt and/or dust from building interiors and one well water sample were also collected.

The July 1993 sampling results indicated arsenic concentrations were below 20 ppm in all but one off-site residential surface soil sample (located within the Edison Glen condominium complex), while higher levels (up to 720 ppm) were found in and immediately adjacent to the unnamed tributary and Mill Brook. The analytical results of the dirt and/or dust samples indicated arsenic levels consistent with background, or naturally occurring concentrations. These results are summarized in EPA's October 20, 1993 correspondence in Appendix II.

In an attempt to define the extent of off-site contamination associated with the CIC site, additional soil, sediment, surface water, groundwater and interior dust sampling was performed by EPA during March and April 1994. EPA conducted extensive sampling at a number of locations including: 1) certain residential properties located along Wilshire Road and Rodak Circle, 2)

the Edison Glen and Edison Woods residential complexes in Edison, 3) areas in and immediately adjacent to the unnamed tributary and Mill Brook, and 4) areas in a Mill Brook tributary located along the southern edge of the Edison Woods residential complex.

During this investigation, soil and sediment samples were obtained in and around the unnamed tributary and Mill Brook at thirty-one specific transect locations. Each transect location is shown in Figure 1 (see Appendix I) as a solid line, designated with an alphabetical letter or letters (A to Z and AA, BB, CC, DD and EE), running perpendicularly through the unnamed tributary and Mill Brook.

The following data resulting from the March/April 1994 sampling effort indicate a similar distribution of arsenic as the July 1993 data:

- o All but one of the residential surface soil samples (those samples taken from a residential yard or common ground within the Edison Glen and Edison Woods residential complexes) indicated arsenic concentrations below 20 ppm.
- o The analyses of soil samples taken from areas immediately adjacent to the unnamed tributary and Mill Brook revealed arsenic concentrations ranging from 0.2 to 1110 ppm. The majority of the detected arsenic concentrations were below 20 ppm.
- o The analyses of sediment samples taken from areas in the unnamed tributary and Mill Brook generated arsenic concentrations ranging from 1.1 to 366 ppm. The majority of the detected arsenic concentrations were below 20 ppm.
- o The analyses of interior dirt and/or dust samples taken from several apartments within the Edison Glen residential complex revealed arsenic concentrations ranging from 0.8 ppm up to 11.3 ppm, indicating levels consistent with back-ground, or naturally occurring concentrations.

EPA analyzed approximately ten percent of the samples for other contaminants, in addition to arsenic. None of the residential surface soil samples indicated the presence of any other contaminants at levels of concern. The soil and sediment samples taken in and immediately adjacent to the unnamed tributary and Mill Brook revealed elevated levels of several compounds, including semi-volatile organic compounds [benzo(b)flouranthene-1.2 ppm, benzo(a)pyrene-0.77-1.4 ppm, benzo(g,h,i)perylene-0.68 ppm, benzo(k)flouranthene-1.7 ppm, benzo(a)anthracene-1.3 ppm], one pesticide (dieldrin-0.005-0.099 ppm) and several inorganic compounds (antimony-25 ppm, beryllium- 1.1-1.9 ppm, lead-103-244 ppm).

The three surface water samples which were collected from specific locations along the unnamed tributary and Mill brook revealed the presence of several volatile organic compounds, herbicides, pesticides, and inorganic compounds, some of which exceeded applicable federal or state surface water quality criteria.

The results of EPA's 1994 sampling effort are summarized in EPA's July 11, 1994 correspondence and data tables in Appendix II.

SUMMARY OF SITE RISKS

Based upon the results of the off-site investigations and assuming a reasonable maximum exposure scenario (as described below), a baseline risk assessment (see Appendix IV) was conducted to estimate the risks associated with current and future exposure to soils and sediments in and around the unnamed tributary and Mill Brook. The baseline risk assessment estimates the human health and ecological risk which could result from the contamination in these off-site areas if no remedial action were taken. Based on a residential exposure scenario, which assumes daily direct contact with contaminants, EPA also evaluated the risks associated with the one residential area where elevated levels of arsenic were detected (see Appendix IV).

Human Health Risk Assessment

As part of the Human Health Risk Assessment, the reasonable maximum human exposure is evaluated by utilizing a four step process for assessing site-related human health risks. The four steps consist of: Hazard Identification--identifies the contaminants of concern at the site based on several factors such as toxicity, frequency of occurrence, and concentration. Exposure Assessment--estimates the magnitude of actual and/or potential human exposures, the frequency and duration of these exposures, and the pathways (e.g., ingesting contaminated wellwater) by which humans are potentially exposed. Toxicity Assessment--determines the types of adverse health effects associated with chemical exposures, and the relationship between magnitude of exposure (dose) and severity of adverse effects (response). Risk Characterization--summarizes and combines outputs of the exposure and toxicity assessments to provide a quantitative assessment of site-related risks.

Unnamed Tributary and Mill Brook Areas

The baseline risk assessment began with selecting contaminants of concern which would be representative of off-site risks. In this case, arsenic was determined to be the primary contaminant of concern. Risk posed by other contaminants (excluding arsenic) which were detected in soils and sediment in off-site areas was

insignificant in comparison to arsenic and would not change the conclusions of EPA's risk assessment.

The baseline risk assessment then identified the reasonable maximum exposure scenario; it was presumed that adolescents playing at the tributary or the Brook would be the most sensitive potential receptors to the contamination. Adolescents (ages 7-18) were assumed to play at the unnamed tributary or Mill Brook once a week throughout the year, for 12 years. Exposure is primarily related to the incidental ingestion of soils and sediments contaminated with arsenic. Under current EPA guidelines, the likelihood of carcinogenic (cancer-causing) and non-carcinogenic effects due to exposure to site chemicals are considered separately. It is assumed that the toxic effects of site-related chemicals are additive. Thus, carcinogenic and non-carcinogenic risks associated with exposures to individual compounds of concern are respectively summed, to indicate the potential risks associated with mixtures of potential carcinogens and non-carcinogens. A summary of the potential carcinogenic and non-carcinogenic health effects is provided below.

o Carcinogenic Risk

Potential carcinogenic risks were evaluated using the cancer slope factors developed by EPA for the contaminants of concern. Cancer slope factors (SFs) have been developed by EPA's Carcinogenic Risk Assessment Verification Endeavor for estimating excess lifetime cancer risks associated with exposure to potentially carcinogenic chemicals. SFs, which are expressed in units of kilogram-day/milligrams (kg-day/mg), are multiplied by the estimated intake of a potential carcinogen, in mg/kg-day, to generate an upper-bound estimate of the excess lifetime cancer risk associated with exposure to the compound at that intake level. The term "upper bound" reflects the conservative estimate of the risks calculated from the SF. Use of this approach makes underestimation of the risk highly unlikely. The oral SF for arsenic is presented in Appendix V.

For known or suspected carcinogens, EPA considers excess upper-bound individual lifetime cancer risks of between 10^{-4} to 10^{-6} to be acceptable. This level indicates that an individual has not greater than approximately one in ten thousand to one in a million chance of developing cancer as a result of site-related exposure to a carcinogen over a 70-year lifetime under the specific exposure conditions at the site.

The excess cancer risk for an adolescent exposed to arsenic in the soils and sediment of the unnamed tributary and Mill Brook (using the reasonable maximum scenario described above) is 5×10^{-6} , which is within EPA's acceptable risk range. This can be interpreted to mean that an individual would have a five in a million excess risk of developing cancer if exposed to arsenic

under the above-described scenario.

o Non-Carcinogenic Risk

Non-carcinogenic risks were assessed using a hazard index (HI) approach, based on a comparison of expected contaminant intakes and safe levels of intake (Reference Doses). Reference doses (RfDs) have been developed by EPA for indicating the potential for adverse health effects. RfDs, which are expressed in units of milligrams/kilogram-day (mg/kg-day), are estimates of daily exposure levels for humans which are thought to be safe over a lifetime (including sensitive individuals). Estimated intakes of chemicals from environmental media (e.g., the amount of a chemical ingested from contaminated drinking water) are compared to the RfD to derive the hazard quotient for the contaminant in the particular medium. The HI is obtained by adding the hazard quotients for all compounds across all media that impact a particular receptor population.

An HI greater than 1.0 indicates that the potential exists for non-carcinogenic health effects to occur as a result of site-related exposures. The HI for non-carcinogenic effects from ingestion of arsenic in soils and sediments of the unnamed tributary and Mill Brook (using the reasonable maximum exposure scenario) was calculated to be 0.05 (see Appendix V). Therefore, non-carcinogenic effects are highly unlikely to occur from the exposure scenario evaluated in the risk assessment.

Residential Areas

In addition to the reasonable maximum exposure scenario evaluated for the areas in and around the unnamed tributary and Mill Brook, EPA also evaluated the risks associated with the one residential area where elevated levels of arsenic were detected. Based on a purely residential exposure scenario, which assumes daily direct contact with contaminants, an area with arsenic contamination significantly above 20 ppm (the upper concentration limit of naturally occurring arsenic for New Jersey soils) in a residential area may pose a human health threat on a long-term exposure basis. For the area behind Building 14 of the Edison Glen Condominium complex, based on the reasonable maximum exposure scenario, the carcinogenic risk was estimated to be 2×10^{-4} , which is at the high-risk end of EPA's acceptable risk range (see Appendix V). In addition, the non-carcinogenic risk to a child receptor was estimated, and an HI of 3.2 was calculated. This indicates the potential for adverse non-carcinogenic effects.

Ecological Risk Assessment

A four-step process is utilized for assessing site-related ecological risks for a reasonable maximum exposure scenario: Problem Formulation--a qualitative evaluation of contaminant

release, migration, and fate; identification of contaminants of concern, receptors, exposure pathways, and known ecological effects of the contaminants; and selection of endpoints for further study. Exposure Assessment--a quantitative evaluation of contaminant release, migration, and fate; characterization of exposure pathways and receptors; and measurement or estimation of exposure point concentrations. Ecological Effects Assessment--literature reviews, field studies, and toxicity tests, linking contaminant concentrations to effects on ecological receptors. Risk Characterization--measurement or estimation of both current and future adverse effects.

As with the Human Health Risk Assessment, arsenic was identified as the contaminant of concern in the ecological risk assessment (see Appendix IV). The assessment quantitatively evaluated the exposure pathways through which ecological receptors could be exposed to arsenic. The most probable exposure pathways for species inhabiting the areas in and immediately adjacent to the unnamed tributary and Mill Brook include ingestion of contaminated biota in the food chain and contact with or ingestion of contaminants present in surface soils, surface water and sediments. Receptor species, such as the deer mouse and eastern cottontail rabbit, could be directly exposed to arsenic through burrowing and grooming activities.

Potential risks to ecological receptors from arsenic present in surface soil and sediments were assessed by calculating the ratio of the medium-specific average and maximum contaminant concentrations to the criteria. Criteria utilized include a benchmark arsenic concentration at which the potential for chronic risk to small mammals exists. If the resulting ratio or hazard index is greater than 1.0, the biota may be at risk of an adverse effect from the arsenic. When there is more than one contaminant of concern in a media, a total hazard index is calculated by summing all the chemical-specific hazard indices for each media. In this case, since arsenic is the only contaminant of concern, there is only one hazard index per media of concern. It follows that a total hazard index greater than 1.0 indicates that exposure to all contaminants (in this case, only arsenic) of ecological concern within that medium may pose a risk to organisms. A literature search was also performed to gather and use all the valuable existing information regarding arsenic effects on the environment.

Results of the ecological risk assessment indicate that the average and maximum total chronic hazard indices for the surface soils are 2.0 and 27.4, respectively (calculated for small mammals as represented by the deer mouse). This indicates that adverse chronic ecological effects may occur in small mammals such as the deer mouse. This risk is driven by arsenic, which is the primary contaminant of concern. However, acute effects to small mammals are unlikely as the receptor would not likely

forage exclusively at locations where the potential exists for an acute effect to begin in the small mammal population (only 6 in over 115 locations with arsenic concentrations in surface soil exceeded 245 milligrams per kilogram (mg/kg)-- the potential threshold for acute effects).

Based on a chronic exposure scenario, it was determined that, at approximately 25 mg/kg of arsenic in surface soil, the potential for risk to the deer mouse population exists. As a result of a review of the existing information regarding arsenic effects in the environment, it was also determined that chronic microbiota effects begin at 375 mg/kg, acute earthworm effects begin at 150 to 165 mg/kg, and reduced plant productivity is noted at 25 to 85 mg/kg.

Risk to organisms at higher trophic levels (for example, at levels above the deer mouse) via exposure through the food chain was also assessed. It was determined that no risk to such organisms via this pathway of exposure is anticipated.

Uncertainties

The procedures and inputs used to assess risks in this evaluation, as in all such assessments, are subject to a wide variety of uncertainties. In general, the main sources of uncertainty include:

- environmental chemistry sampling and analysis
- environmental parameter measurement
- fate and transport modeling
- exposure parameter estimation
- toxicological data.

Uncertainty in environmental sampling arises in part from the potentially uneven distribution of chemicals in the media sampled. Consequently, there is significant uncertainty as to the actual levels present. Environmental chemistry-analysis error can stem from several sources including the errors inherent in the analytical methods and characteristics of the matrix being sampled.

Uncertainties in the exposure assessment are related to estimates of how often an individual would actually come in contact with the chemicals of concern, the period of time over which such exposure would occur, and in the models used to estimate the concentrations of the chemicals of concern at the point of exposure.

Uncertainties in toxicological data occur in extrapolating both from animals to humans and from high to low doses of exposure, as well as from the difficulties in assessing the toxicity of a mixture of chemicals. These uncertainties are addressed by

making conservative assumptions concerning risk and exposure parameters throughout the assessment. As a result, the Risk Assessment provides upper-bound estimates of the risks to populations near the site, and is highly unlikely to underestimate actual risks related to the site.

Summary of Human Health and Ecological Risk

The results of the human health risk assessment indicate that, based on the reasonable maximum exposure scenario evaluated for the soils and sediments in and around the unnamed tributary and Mill Brook, the excess carcinogenic risk is within EPA's acceptable risk range, and adverse non-carcinogenic effects are not likely to occur. However, under a residential exposure scenario, an area with arsenic contamination significantly above 20 ppm may pose a human health threat on a long-term exposure basis.

The ecological risk assessment concluded that the analytical results of surface soil samples collected in the vicinity of the unnamed tributary and Mill Brook indicate the presence of arsenic at levels sufficient to generate acute risk to soil invertebrates and chronic risk to soil microbiota and small mammals. Contaminants other than arsenic which were found in soil during the investigation do not pose any significant risk to the terrestrial community. In addition, surficial stream sediments in some areas of the unnamed tributary and Mill Brook contain sufficient arsenic to pose a risk to the benthic community (organisms living on the tributary and brook bottoms). Isolated sediment sampling locations also indicated sufficient levels of semi-volatile organic compounds, pesticides and inorganic compounds to generate risk to the benthic community.

Actual or threatened releases of hazardous substances from the CIC site, if not remediated, may present a current or potential threat to public health and the environment.

REMEDIAL ACTION OBJECTIVES

Remedial action objectives are specific goals to protect human health and the environment. These objectives are based on available information and standards such as applicable or relevant and appropriate requirements (ARARs) and risk-based levels established in the risk assessment. In order to address potential human health and ecological risks, the remedial action objective for the off-site remedy is to eliminate the potential for exposure to contaminated soils and sediment in residential areas and areas in and immediately adjacent to the unnamed tributary and Mill Brook.

The human health risk assessment indicated that, based on the reasonable maximum exposure scenario evaluated for the soils and sediments in and around the unnamed tributary and Mill Brook, the excess carcinogenic risk (5×10^{-6}) is within EPA's acceptable risk range, and adverse non-carcinogenic effects are not likely to occur. However, under a residential exposure scenario, an area with arsenic contamination significantly above 20 ppm may pose a human health threat on a long-term exposure basis.

The ecological risk assessment indicated the potential for adverse ecological effects if no remedial action is taken. Because the unnamed tributary and Mill Brook currently exist as a mature habitat corridor in an otherwise heavily developed area, the habitat value of this area is increased and destruction of the area in connection with the removal of contaminated soils and sediment should be mitigated. However, its high ecological value also increases the attraction of receptors into this potentially hazardous habitat. Therefore, achieving a balance between mitigating the potential risk to ecological receptors and preserving the value of the habitat (or minimizing destruction of the habitat through remedial activity) is preferred.

EPA has determined that the highest levels of arsenic are located in the soil and/or sediment in and around the area of transects A, B, I, J, K, L, M, N, P, Q, S, V, W, X, Y, Z, and AA along the unnamed tributary and Mill Brook, as shown in Figure 1 of Appendix I. These transect locations include all the areas identified by the NJDEP as containing an arsenic concentration in soil and/or sediment at or above 100 ppm. EPA believes that by focusing remediation of soil and/or sediment in these areas, the majority of the contamination would be removed while achieving a reasonable and acceptable balance in preserving the ecology of the area.

EPA plans to rely on NJDEP's determination of the upper limit of naturally occurring arsenic for New Jersey soils (20 ppm) as a criteria in remediating the above-described areas targeted for cleanup. This same approach will be utilized to restore contaminated residential areas (a backyard or common ground within an apartment complex) such that any long-term risks associated with these areas are removed. Although an arsenic cleanup goal of 25 ppm would address the risks identified in the ecological risk assessment, applying a soil/sediment cleanup criteria of 20 ppm (or achieving an average residual arsenic level of 20 ppm) also enables EPA to further mitigate the human health risks.

DESCRIPTION OF REMEDIAL ALTERNATIVES

CERCLA requires that each selected remedy be protective of human

health and the environment, be cost effective, comply with other statutory laws, and utilize permanent solutions and alternative treatment technologies and resource recovery alternatives to the maximum extent practicable. In addition, the statute includes a preference for the use of treatment as a principal element for the reduction of toxicity, mobility, or volume of the hazardous substances.

EPA's ability to exercise a preference for alternative treatment technologies in identifying potential remedial alternatives for the contaminated soils and sediment was limited for the following reasons:

- insufficient information is available to evaluate treatment technologies and their effectiveness in treating the contaminated soils and sediment without performing additional treatability studies;
- the expeditious schedule required for remediation; and
- logistical problems associated with storage (until sufficient additional treatability study work can be performed) of the off-site contaminated soil and sediment at the CIC site due to the recent completion of the above-described interim remedy.

In addition, containment of the contaminated material was also considered but screened out prior to EPA's final evaluation of alternatives. Because of the free-flowing and naturally unpredictable conditions present in the areas of the unnamed tributary and Mill Brook, in-situ containment was deemed impracticable and excluded from the subsequent alternatives evaluation.

Therefore, EPA has evaluated two remedial alternatives for addressing off-site contaminated soil and sediment.

These alternatives are:

Alternative 1: No Action

Estimated Capital Cost: \$0
Estimated Operation & Maintenance Cost: \$0
Estimated Present Worth Cost: \$0
Estimated Implementation Time: 0

The Superfund program requires that the "no-action" alternative be considered as a baseline for comparison with other alternatives. Under this alternative, EPA would not take any action to remediate contaminated off-site soil and sediment. Because this alternative would result in CIC contaminants remaining in off-site areas, CERCLA requires that these areas be reviewed every five years. If justified by the review, remedial

actions may be implemented to remove or treat the contaminants.

Because the average arsenic level throughout the area would remain above 20 ppm, NJDEP would recommend land use restrictions (based on the New Jersey Industrial Site Recovery Act) to eliminate the potential for direct contact.

Alternative 2: Excavation/Off-Site Disposal

Estimated Capital Cost: \$8,583,000

Estimated Operation & Maintenance Cost: \$0

Estimated Present Worth Cost: \$0

Estimated Time to Design and Implement: 24 months

Alternative 2 includes excavation and off-site disposal of contaminated soil and/or sediment within the areas of transects A, B, I, J, K, L, M, N, P, Q, S, V, W, X, Y, Z, and AA along the unnamed tributary and Mill Brook (as shown in Figure 1 of Appendix I). In addition, contaminated soil in a grassy area behind Building 14 of the Edison Glen Condominium Complex would be excavated to 20 ppm arsenic for off-site disposal. This is the only residential area warranting remediation.

The volume of soil and sediment to be excavated is estimated to be 10,000 cubic yards, which is approximately equivalent to 14,100 tons of soil and sediment.

The estimated capital cost of \$8,583,000 required to implement Alternative 2 is a conservative estimate based on disposal of the contaminated soils and sediment as hazardous waste at an EPA-approved off-site hazardous waste landfill. Since it is possible that this material may not be classified as a hazardous waste, it may be disposed of at a non-hazardous waste facility, if such authorized facility is willing to accept it. Disposal of the material as non-hazardous waste at a non-hazardous waste facility would reduce the estimated capital cost required to implement Alternative 2 to \$2,431,000.

The cost of this alternative also includes the cost of restoring the excavated areas to the extent practicable. This would include backfilling and revegetation to stabilize the excavated areas.

Because average residual levels of arsenic will be below 20 ppm, these areas would not be reviewed every five years and NJDEP would not recommend any land use restrictions. Furthermore, NJDEP's Stream Encroachment Regulations would serve to control any future intrusive activities in and around the unnamed tributary and Mill Brook.

SUMMARY OF COMPARATIVE ANALYSIS OF ALTERNATIVES

During the detailed evaluation of remedial alternatives, each alternative is assessed utilizing nine evaluation criteria as set forth in the National Oil and Hazardous Substances Pollution Contingency Plan (NCP) and Office of Solid Waste and Emergency Response (OSWER) Directive 9355.3-01. These criteria were developed to address the requirements of Section 121 of CERCLA to ensure all important considerations are factored into remedy selection decisions.

The following "threshold" criteria are the most important, and must be satisfied by any alternative in order to be eligible for selection:

- o Overall protection of human health and the environment addresses whether or not a remedy provides adequate protection and describes how risks posed through each pathway are eliminated, reduced, or controlled through treatment, engineering controls, or institutional controls.
- o Compliance with applicable or relevant and appropriate requirements addresses whether or not a remedy will meet all of the applicable or relevant and appropriate requirements of other federal and state environmental statutes and requirements or provide grounds for invoking a waiver.

The following "primary balancing" criteria are used to make comparisons and to identify the major trade-offs between alternatives:

- o Long-term effectiveness and permanence refers to the ability of a remedy to maintain reliable protection of human health and the environment over time, once cleanup goals have been achieved.
- o Reduction of toxicity, mobility, or volume through treatment is the anticipated performance of the treatment technologies a remedy may employ.
- o Short-term effectiveness addresses the period of time needed to achieve protection and any adverse impacts on human health and the environment that may be posed during the construction and implementation period until cleanup goals are achieved.
- o Implementability is the technical and administrative feasibility of a remedy, including the availability of materials and services needed to implement a particular option.
- o Cost includes estimated capital and operation and

maintenance costs, and net present worth costs.

The following "modifying" criteria are considered fully after the formal public comment period on the Proposed Plan is complete:

- o State acceptance indicates whether, based on its review of the Proposed Plan and supporting documentation, the state concurs, opposes, and/or has identified any reservations with the preferred alternative.
- o Community acceptance refers to the public's general response to the alternatives described in the Proposed Plan and supporting documentation. Factors of community acceptance to be discussed include support, reservation, and opposition by the community.

A comparative analysis of these alternatives based upon the evaluation criteria noted above follows.

- o Overall Protection of Human Health and the Environment

Alternative 1, no action, would not be protective of human health and the environment as there would not be any action taken to eliminate potential for direct contact with or ingestion of contaminated soils and sediment. Since no remedial activities would be implemented under this alternative, the risks posed to human health and the environment would be the same as those identified in the risk assessment. Alternative 1 is not consistent with remedial action objectives.

Alternative 2 provides protection of human health and the environment by removing the soils and sediment in the above-described contaminated areas, and restoring such areas. By eliminating the potential human health and ecological risks, Alternative 2 would satisfy the remedial action objectives. Furthermore, it balances the preservation of a valuable ecological habitat, while providing for removal of the majority of the arsenic contamination. Removal of contaminated soils and sediment also contributes to the improvement of surface water quality in the unnamed tributary and Mill Brook.

- o Compliance with ARARs

ARARs are those federal or state environmental and public health regulations that apply to remedial activities at the site [or area(s) to be remediated]. There are three classifications of ARARs: chemical-specific, which are health- or risk-based concentration limits; location-specific, which are based on the geographical location of the site and its surroundings; and action-specific, which are controls on particular types of remedial activities.

EPA plans to use 20 ppm as a criteria in remediating the contaminated soils and sediment in residential areas and areas in and immediately adjacent to the unnamed tributary and Mill Brook. Although this is not a promulgated chemical-specific standard, and therefore not an ARAR, it does represent the upper limit of naturally occurring arsenic concentrations in New Jersey soils. Use of this criteria allows EPA to remove a large majority of the arsenic contamination while achieving a reasonable and acceptable balance in preserving a majority of the valuable ecology existing in these areas.

Alternative 1 would not attain the chemical-specific criteria in soils and sediment as it does not involve active remediation. Action- and location-specific ARARs are not applicable as Alternative 1 does not involve implementation of remedial activities.

Alternative 2 is expected to attain all chemical-, location- and action-specific ARARs. The chemical-specific cleanup criteria for soils and sediment would be achieved, since the residual average levels of arsenic will be below 20 ppm. All action- and location-specific ARARs for remedial activities in wetlands and floodplains would be achieved, including Section 404 of the Clean Water Act, Executive Order 11990 (Protection of Wetlands), the Flood Hazard Area Control Act (N.J.S.A. 16:50A) and the Freshwater Wetlands Protection Act (N.J.S.A. 13:9B). The Occupational Safety and Health Act (OSHA) would apply to all workers conducting the remedial activities specified under Alternative 2.

In addition and as discussed above, the excavated material may be classified as a hazardous waste requiring adherence to regulations involving the transport and off-site disposal of hazardous waste. If necessary, the requirements of the Resource Conservation and Recovery Act (RCRA), the Department of Transportation (DOT) and the New Jersey Solid and Hazardous Waste Regulations would be achieved through proper handling and shipment of the contaminated material to an EPA-approved disposal facility. A determination of the material classification would be made during design/implementation of Alternative 2 in order to select an appropriate off-site disposal facility.

o Long-Term Effectiveness and Permanence

Alternative 1 is not considered to be effective over the long term as it does not include remediation of any off-site contamination. Therefore, this alternative would not achieve the remedial action objectives, since it would not reduce exposure to contaminated soils and sediment in the subject areas. As required by CERCLA, areas where contamination is left in place must be reviewed every five years. If justified by the review, remedial actions may be implemented to address the contaminated areas.

Alternative 2 includes removal of the majority of the arsenic contamination, thereby eliminating the potential for human exposure to and adverse ecological effects from contaminants at levels of concern. As a result, a review of these areas every five years is not required. This alternative is considered effective over the long term and represents a permanent remedy to address these areas, and does not require future monitoring, operation or maintenance.

o Reduction in Toxicity, Mobility, or Volume

Alternative 1 does not include treatment of contaminated soils and sediment, and therefore, does not reduce the toxicity, mobility, or volume of contamination in the areas subject to remediation. The contaminated material would remain in these areas.

As with Alternative 1, Alternative 2 is not expected to include treatment. However, if so determined, the material excavated under Alternative 2 may require treatment before it can be disposed of in an off-site landfill. Such treatment may result in a reduction of the toxicity and mobility of contamination.

o Short-Term Effectiveness

Since there are no remedial activities being implemented under Alternative 1, there would be no additional short-term risks posed to human health or the environment.

The time required to implement Alternative 2 is estimated to be 24 months. No additional risks to human health or the environment are expected as a result of the implementation of these activities. Due to the intrusive nature of soil and sediment removal, there may be potential risks posed to workers during the remedial activities. Worker protection would be required to prevent direct contact with contaminated material during the excavation effort. In addition, workers would be trained in health and safety, and protective equipment would be provided during construction activities.

The excavation effort would cause significant disruption to areas in and around the unnamed tributary and Mill Brook. While EPA plans to minimize the destruction of trees in these areas, it is expected that a number of trees would be removed. The related activities would also require stream encroachment and disturbance of sediments. However, EPA would take the appropriate steps to adequately control water flow and sediment runoff. Upon completion of the excavation activities, the disturbed areas would be restored and stabilized by backfilling and revegetating.

Since the remedial activities would occur in and around residential areas, EPA anticipates that these activities may be

considered disruptive to some or all residents. For example, a high volume of traffic is expected to occur during the soil and sediment removal activities. EPA would take appropriate and reasonable measures to ensure that proper traffic controls are implemented and that impacts to affected residents are minimized. EPA would work closely with the community during the remedial design period to coordinate these construction-related activities.

o Implementability

There are no implementability concerns posed by Alternative 1 since no remedial action would be taken under this alternative.

Alternative 2 would be difficult to implement due to the large volume of material to be excavated and the limited accessibility to the areas to be remediated. Temporary access roadways would be constructed to provide necessary vehicular equipment access to areas to be remediated in and immediately adjacent to the unnamed tributary and Mill Brook.

In addition, attempts would be made during implementation of the remedial action to minimize tree removal, which would likely require additional time for the excavation work.

o Cost

Since Alternative 1 does not include any remedial activities, there are no costs associated with this alternative.

Since Alternative 2 does not require operation and maintenance, present worth and operation and maintenance costs were not estimated. The estimated capital cost associated with Alternative 2 is \$8,583,000 and \$2,431,000 assuming off-site disposal of contaminated material as hazardous and non-hazardous waste, respectively.

o State Acceptance

The State of New Jersey concurs with EPA's preference of Alternative 2.

o Community Acceptance

In general, both public officials and community residents expressed support for Alternative 2. A more detailed discussion of community concerns is presented in the Responsiveness Summary.

SELECTED REMEDY

Based upon consideration of the requirements of CERCLA, the detailed analysis of the alternatives, and public comments, both

EPA and the New Jersey Department of Environmental Protection have determined that Alternative 2 is the most appropriate remedy to address soil and sediment contamination related to the CIC site in and around off-site residential areas.

Alternative 2 includes excavation and off-site disposal of contaminated soil and/or sediment within the areas of transects A, B, I, J, K, L, M, N, P, Q, S, V, W, X, Y, Z, and AA along the unnamed tributary and Mill Brook (as shown in Figure 1 of Appendix I). In addition, contaminated soil in a grassy area behind Building 14 (also shown in Figure 1 of Appendix I) of the Edison Glen Condominium Complex will also be excavated for off-site disposal.

Approximately 10,000 cubic yards of soil and sediment will be removed and the remediated areas will be appropriately restored. This approach enables EPA to restore contaminated residential areas such that any long-term risk associated with these areas is removed and no property use restrictions will be required. By also targeting specific contaminated areas in and near the unnamed tributary and Mill Brook, EPA is able to remove a significant portion of the CIC contamination while achieving a reasonable and acceptable balance in preserving the ecology of the area. Furthermore, because the average residual levels of arsenic will be to be below 20 ppm, it will not be necessary to perform five-year review in connection with the off-site areas or require the imposition of land use restrictions.

The selected remedy is the most protective of human health and the environment because it eliminates the risk associated with exposure to contaminated soil and sediment by both human and ecological receptors. Alternative 1 is not protective of human health and the environment.

Due to the large volume of material to be excavated and the limited accessibility to the areas to be remediated, Alternative 2 will be difficult to implement. However, the benefits of the selected remedy outweigh any associated implementability issues. EPA will work closely with the community during the remedial design period and to coordinate construction activities.

Future use of the off-site areas being targeted for remediation is also an important consideration. These areas would likely be restricted from future use under Alternative 1 in order to control exposure to contaminated soils and sediment. However, under the selected remedy, a greater degree of flexibility exists regarding future use of the areas targeted for remediation, as land use restrictions will not be required.

The selected remedy is much higher in cost than Alternative 1. However, when evaluating the cost effectiveness of the alternatives, which is determined by weighing the cost against

the alternative's ability to achieve ARARs and remedial action objectives, the selected remedy is cost effective. Unlike Alternative 1, the selected remedy achieves a balance in preservation of the ecology while eliminating the potential for exposure to contaminated soils and sediments in the areas targeted for remediation.

The selected remedy provides the best balance of trade-offs among alternatives with respect to the evaluation criteria. EPA and NJDEP believe that the selected remedy will be protective of human health and the environment, will comply with ARARs, will be cost effective, and will utilize permanent solutions and alternative treatment technologies or resource recovery technologies to the maximum extent practicable.

STATUTORY DETERMINATIONS

Under its legal authorities, EPA's primary responsibility at Superfund sites is to undertake remedial actions that are protective of human health and the environment. In addition, Section 121 of CERCLA establishes several other statutory requirements and preferences. These specify that when complete, the selected remedial action must comply with applicable or relevant and appropriate environmental standards established under federal and state environmental laws unless a statutory waiver is justified. The selected remedy also must be cost effective and utilize permanent solutions and alternative treatment technologies or resource recovery technologies to the maximum extent practicable. Finally, the statute includes a preference for remedies that employ treatment that permanently and significantly reduce the volume, toxicity, or mobility of hazardous wastes, as available. The following sections discuss how the selected remedy meets these statutory requirements.

Protection of Human Health and the Environment

The selected alternative provides protection of human health and the environment by removing the soils and sediment in the above-described contaminated areas, and restoring such areas. The excavated contaminated material will be disposed at a secure landfill, thereby preventing any future negative impacts to the surrounding environment. Alternative 2 balances the preservation of an ecological habitat, while providing for removal of the majority of the arsenic contamination. Because this remedy involves removal of a limited portion of the areas in and around the unnamed tributary and Mill Brook, short-term and long-term impacts to the ecosystem are minimized. Removal of contaminated soils and sediment also contributes to the improvement of surface water quality in the unnamed tributary and Mill Brook.

Compliance with ARARs

The selected remedy is expected to comply with all ARARs necessary to achieve the above-described remedial action objectives. The remedy is designed to comply with all action- and location-specific ARARs that pertain to remedial activities in wetlands and floodplains. The chemical-specific cleanup criteria for soils and sediment will be achieved since the residual average levels of arsenic will be below 20 ppm. All action-specific ARARs identified under OSHA will apply to all workers conducting the remedial activities specified under the selected remedy. In addition and as discussed above, the selected remedy may require classification of the excavated material as a hazardous waste and, therefore, be required to adhere to regulations involving the transport and off-site disposal of hazardous waste. If the soils and sediment are classified as hazardous waste, the remedy will comply with the requirements of the Resource Conservation and Recovery Act, the Department of Transportation and the New Jersey Solid and Hazardous Waste Regulations will be achieved through proper handling and shipment of the contaminated material to an EPA-approved disposal facility. A determination of the material classification will be made during design/implementation of the selected remedy in order to select an appropriate off-site disposal facility.

Cost Effectiveness

When evaluating the cost effectiveness of the alternatives, which is determined by weighing the cost against the alternative's ability to achieve ARARs and remedial action objectives, the selected remedy has been determined to be the most cost effective.

Utilization of Permanent Solutions and Alternative Treatment Technologies to the Maximum Extent Practicable

The selected remedy represents the maximum extent to which permanent solutions and treatment technologies can be utilized in an expeditious and cost effective manner to remediate CIC contamination in and around off-site residential areas. Although the selected remedy does not involve the reduction of toxicity, mobility, or volume through treatment, it provides long- and short-term effectiveness, and is cost effective.

Preference for Treatment as a Principal Element

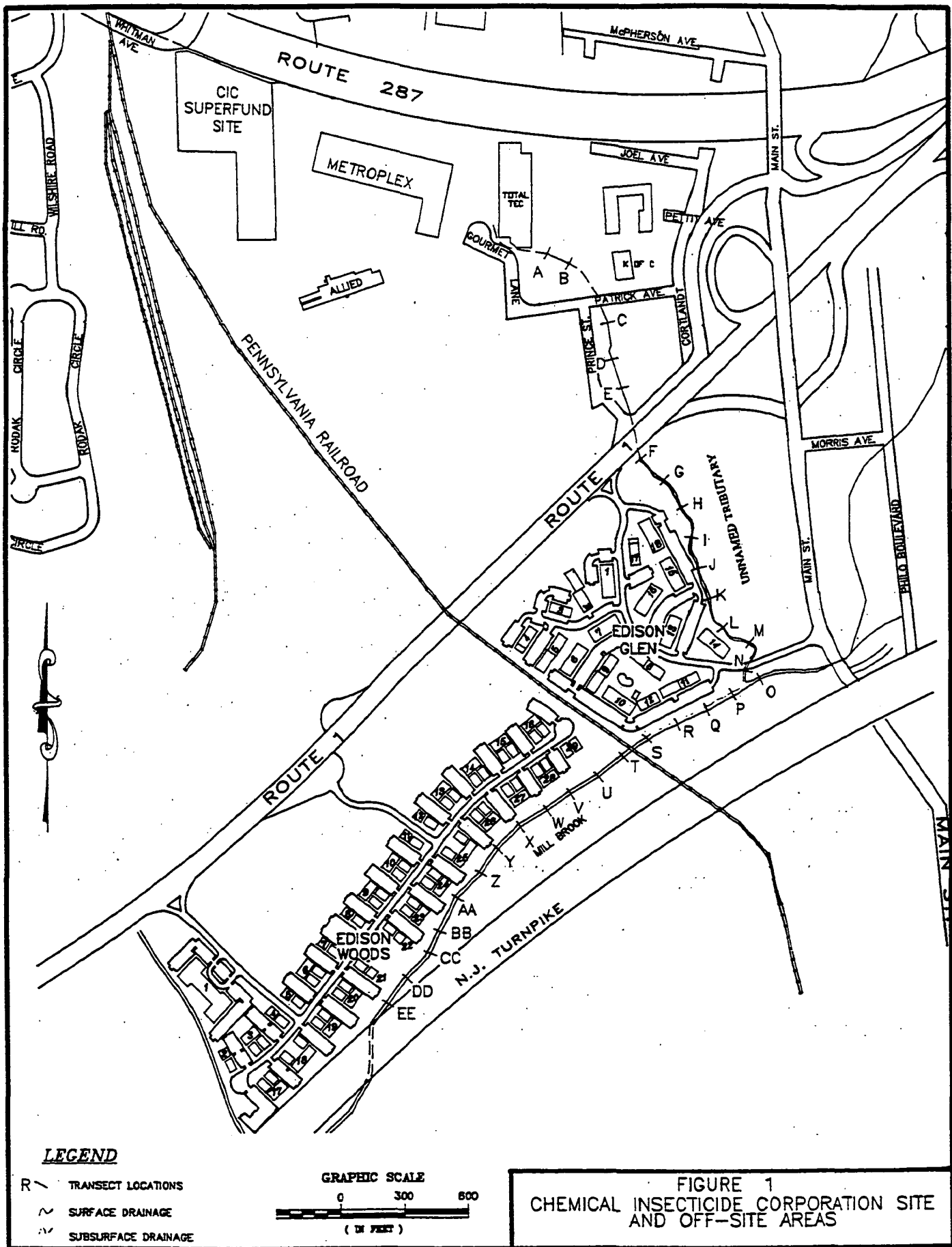
The selected remedy does not satisfy the preference for treatment as a principal element. Treatment of the principal threats of the areas to be remediated was not found to be practicable because no effective treatment technology is readily available at this time.

DOCUMENTATION OF SIGNIFICANT CHANGES

There have been no significant changes in the selected remedy from the preferred remedy described in the Proposed Plan.

APPENDIX I

Figure 1



APPENDIX II

Data



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

REGION II

JACOB K. JAVITS FEDERAL BUILDING

NEW YORK, NEW YORK 10278-0012

October 20, 1993

Dear Resident:

The purpose of this letter is to inform you of the test results for the soil samples collected by the U. S. Environmental Protection Agency (EPA) from July 6 through 8, 1993. The testing was performed to better define the extent of contamination associated with the Chemical Insecticide Corporation (CIC) Superfund site at 30 Whitman Avenue, Edison, New Jersey. The samples were collected at a number of locations including: 1) properties on Prince Street, Patrick Avenue and Cortland Street in Edison, 2) the Edison Glen and Edison Woods residential complexes in Edison, and 3) areas in Metuchen near Route 287. Four samples of dirt and/or dust from building interiors and one well water sample were also collected.

This sampling effort was a follow-up to soil sampling performed by EPA in October of 1992 near Mill Brook and an unnamed creek, two streams that receive drainage from the CIC site. Some levels of arsenic found in the 1992 samples were higher than levels typically found in New Jersey soils. However, because those samples were taken mainly on the banks of the two streams, it was not known whether the elevated concentrations of arsenic extended from the streams to areas closer to the nearby buildings. The July 1993 sampling effort included additional sampling on the banks of the streams as well as soil samples taken closer to the nearby buildings. The July 1993 project involved the collection of sixty-seven samples, as compared to only nine samples collected in October of 1992.

Arsenic was used in pesticides produced by the Chemical Insecticide Corporation during the 1950's and 1960's. Arsenic also occurs naturally in soils and is typically found at concentrations between 5 to 30 parts per million (ppm) in New Jersey. Higher or lower concentrations are found in different geographic areas. An extensive study was conducted to establish naturally occurring background concentrations of arsenic for a Superfund site in Middlesex County. Background concentrations for arsenic were found to be 28 ppm.

The July 1993 sampling results are presented in an EPA document entitled: "Final Report, Off-Site Characterization, Chemical Insecticide Site, Edison, NJ" dated September, 1993. The results are summarized on the attached map. Arsenic concentrations were below 28 ppm in all but one off-site soil samples except those in the immediate vicinity of the unnamed creek and Mill Brook.

EPA's preliminary evaluation the data indicates that the potential exposure to arsenic contaminated soil does not present an immediate health risk. Although the highest concentrations of arsenic for the July 1993 samples are greater than the highest concentrations found in samples collected in October 1992, the recent data indicates that soil arsenic levels decrease to normal levels with increasing distance from the streams. The new sampling results indicate that the arsenic contamination is limited chiefly to the outskirts of properties along the two streams. This limited spacial distribution of the arsenic contamination reduces the potential for human exposure to the arsenic. However, there is concern that exposure to arsenic levels found near the creek and brook may be of concern over the long-term.

EPA has made the July 1993 sampling results available to the Federal Agency for Toxic Substances and Disease Registry (ATSDR) and to the New Jersey Department of Health (NJDOH) for review and comment. The initial review of the sampling results by ATSDR and NJDOH confirmed EPA's preliminary evaluation. ATSDR and NJDOH concurred that the arsenic levels documented in the July 1993 sampling event do not pose an immediate public health threat. However, ATSDR and NJDOH did express the concern that the arsenic levels may pose a public health threat on a long-term exposure basis. EPA, ATSDR and NJDOH plan to continue to evaluate the potential health risks associated with the off-site arsenic contamination.

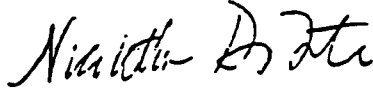
Since there is no concern about immediate health effects due to potential short-term exposure to arsenic contamination, EPA plans to address the concerns related to long-term exposure as part of the complete remedy for the site. EPA anticipates selecting a cleanup plan for the site in the Spring of 1994. Excavation of off-site soils would be expedited, following the selection of a remedy.

Anticipating that community residents, property owners and others may have questions about this matter, EPA has scheduled a public meeting to provide additional information and answer questions about the CIC site. ATSDR and NJDOH also plan to participate in this meeting. The time and place are:

Monday, October 25, 1993 from 7 PM to 9:30 PM
Metuchen Borough Hall
Main Street and Middlesex Avenue (Route 27)
Metuchen, New Jersey 08840

You will be kept informed regarding future EPA activities for the CIC site. In addition, feel free to contact Mr. Jonathan Josephs, Project Manager at 212-264-8098, or Ms. Cecilia Echols, Community Relations Coordinator at 212-264-0949 if you require further information.

Sincerely,

A handwritten signature in cursive script, appearing to read "Nicoletta Di Forte".

Nicoletta Di Forte, Chief
Northern New Jersey Section
New Jersey Superfund Branch II

Attachment

**QUESTIONS AND ANSWERS
CONCERNING OFF-SITE SAMPLING RESULTS
FOR THE CHEMICAL INSECTICIDE SITE, EDISON, NEW JERSEY**

Is there an immediate problem that needs to be addressed on an emergency basis?

The concentrations of arsenic found off-site do not pose a short-term health threat; therefore, an immediate action is not required. EPA uses a range of 1000 to 2000 ppm of arsenic to determine when an immediate action is warranted.

Is there a concern about effects from long-term exposure?

In limited areas around the unnamed creek and Mill Brook, long-term exposure to the concentrations of arsenic found may be of concern. This concern is based on repeated ingestion of the most contaminated soils over many years. However, the location of contaminated soils reduces the likelihood for exposure to arsenic.

Have the appropriate health agencies reviewed the data?

Both the Agency for Toxic Substances and Disease Registry and the New Jersey Department of Health have reviewed the results. These Agencies have independently concluded that while there may be a concern about the effects of long-term exposures, there is no immediate public health threat associated with the concentrations of arsenic found off-site.

Will the contamination be cleaned up?

Since there is no immediate public health threat associated with off-site contaminated soils, EPA plans to clean up the areas around the unnamed creek and Mill Brook as part of the complete remedy for the CIC site. This remedy should be selected by the Spring of 1994. Excavation of off-site soils will be expedited, once the remedy is selected.

Will additional off-site soil sampling be performed?

There is sufficient sampling to determine that a remedy is warranted for off-site soils. Additional samples will be taken as part of the complete remedy to define the boundaries of off-site areas to be excavated.

What levels of arsenic in off-site soils will require long-term cleanup?

EPA is currently working with the New Jersey Department of Environmental Protection and Energy to establish a cleanup level for the CIC site. An extensive study was conducted to determine background concentrations of arsenic for another Superfund site in Middlesex County. The study indicated that 28 ppm was background for the area. EPA anticipates that the cleanup level will be in the 20 ppm range.

What off-site areas require cleanup?

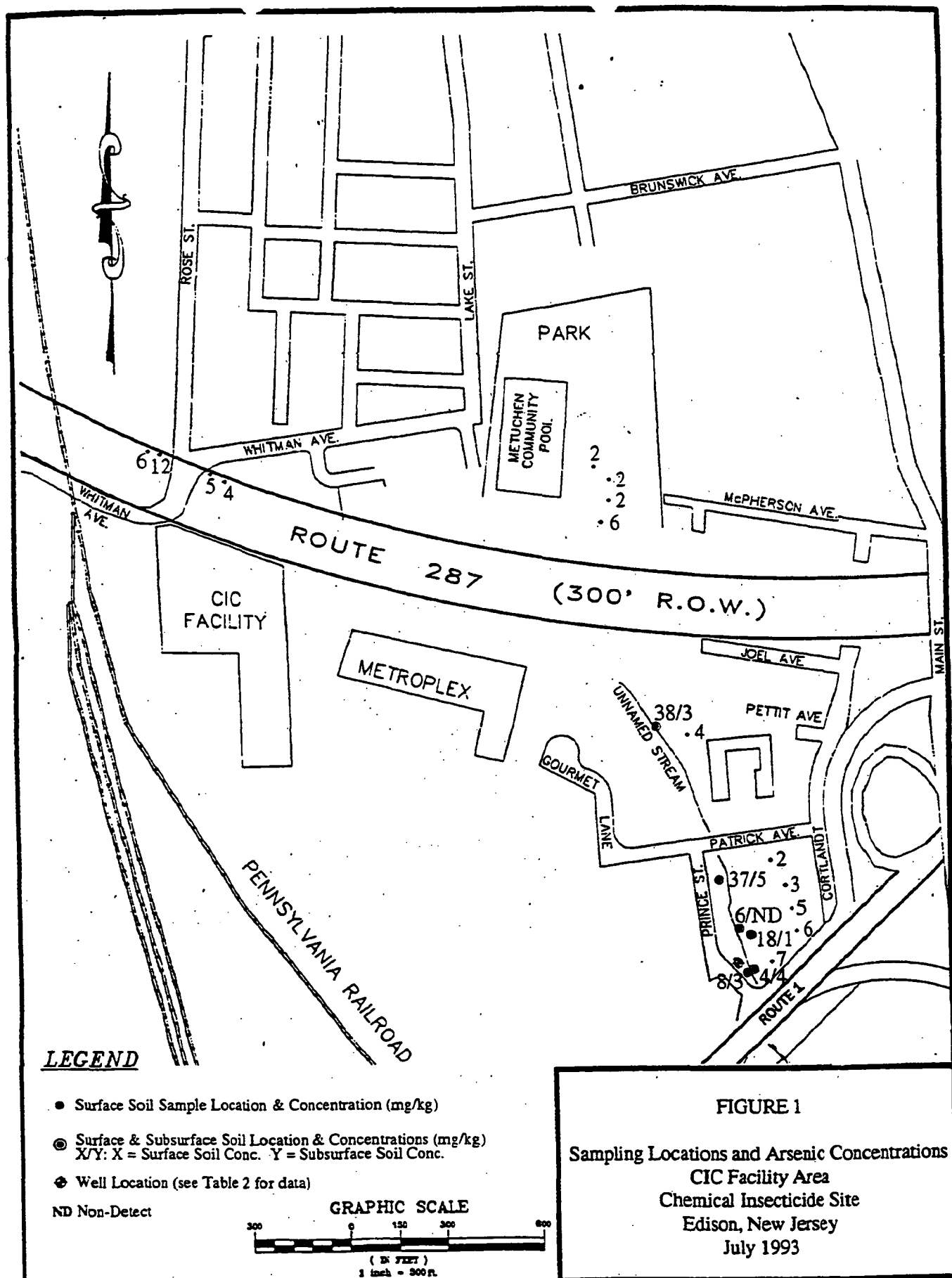
Of the 62 soil samples collected, 20 exceed 20 ppm. All of the samples that exceeded 20 ppm are adjacent to the unnamed creek or Mill Brook.

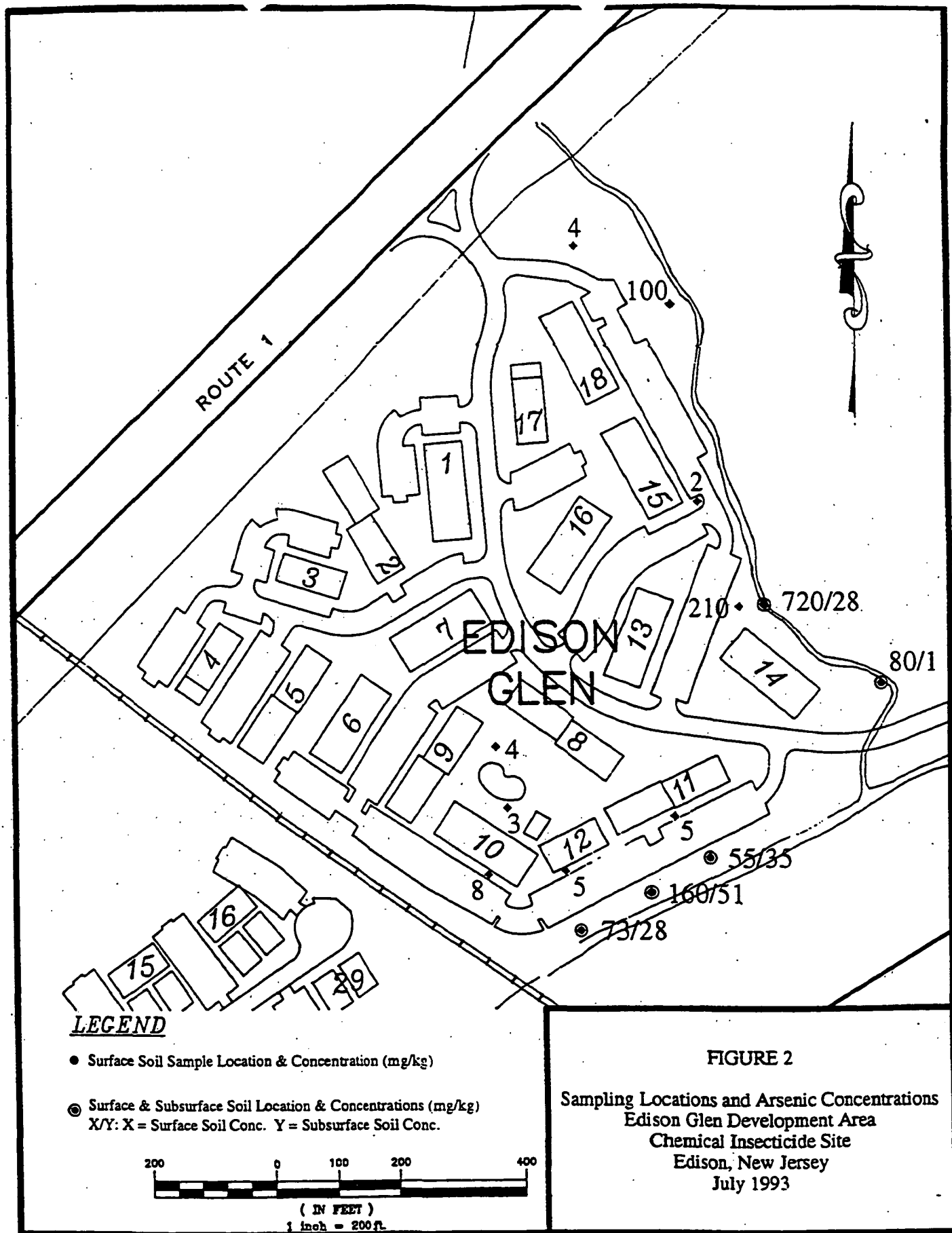
Will the interim cap address off-site contamination?

The installation of a cap would prevent future migration of contaminated stormwater runoff from the CIC site, but would not address the current off-site conditions.

When will the interim cap be installed on the CIC site?

The contractor which will be conducting an investigation for explosives waste has begun mobilizing equipment to the site. The explosives waste investigation is scheduled to begin shortly. If the explosives investigation shows that there is not a significant problem, then mobilization for cap construction should begin in December.





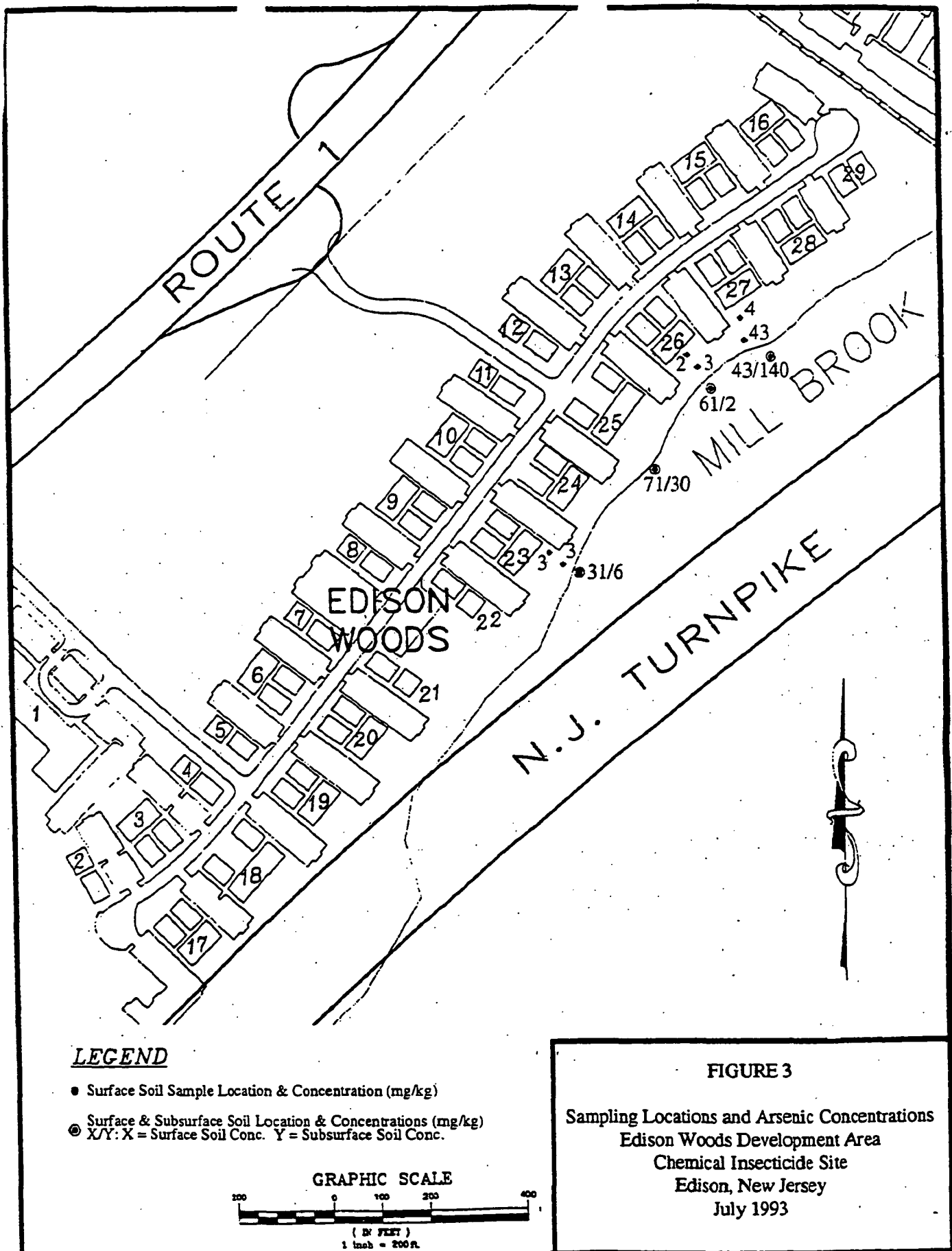


Table 1

Results of Arsenic Analysis of Dust Samples
Chemical Insecticide Site
Edison, N.J.
July 1993

Sample Location	Arsenic Concentration (mg/kg)
14 Prince Street	3
1 Cortlandt Street	9
1 Cortlandt Street (crawlspace)	2
Knights of Columbus Hall	1

Table 2

Organics/Inorganics Concentrations Detected in Groundwater Sample
Chemical Insecticide Site
Edison, N.J.
July 1993

Parameter	Concentration (ug/L)
Aluminum	290
Barium	110
Beryllium	0.6
Cadmium	3.4
Calcium	46000
Cobalt	30
Copper	3100
Iron	190
Lead	8
Magnesium	11000
Manganese	380
Nickel	57
Potassium	3400
Sodium	49000
Zinc	570
Aroclor 1254 (a polychlorinated biphenyl)	0.7



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY

REGION II

JACOB K. JAVITS FEDERAL BUILDING

NEW YORK, NEW YORK 10278-0012

July 11, 1994

Re: Chemical Insecticide Corporation Superfund Site

Dear Resident and/or Property Owner:

This letter is to inform you of the results of soil, sediment, surface water and groundwater sampling performed by the U.S. Environmental Protection Agency (EPA) from March 14 through April 1, 1994. EPA is pleased to report that the recent sampling results are consistent with previous testing, and indicate that there is no immediate health threat posed by off-site arsenic contamination.

The sampling was intended to further measure the extent of potential off-site contamination associated with the Chemical Insecticide Corporation (CIC) Superfund site at 30 Whitman Avenue, Edison, New Jersey. The samples were collected at a number of locations including: 1) certain residential properties located along Wilshire Road and Rodak Circle, 2) the Edison Glen and Edison Woods residential complexes in Edison, 3) areas in and immediately adjacent to the unnamed stream and Mill Brook, and 4) areas in a Mill Brook tributary located along the Southern edge of the Edison Woods residential complex. Three samples of dirt and/or dust from certain apartments and one well water sample were also collected. The results are summarized in the attached figures.

As you may be aware, arsenic was used in pesticides produced by the Chemical Insecticide Corporation during the 1950's and 1960's and is the primary contaminant of concern for the site. Arsenic also occurs naturally in soils and is typically found at concentrations between 5 to 30 parts per million (ppm) in New Jersey. Higher or lower concentrations are found in different geographic areas. The New Jersey Department of Environmental Protection and Energy has determined the upper limit of naturally occurring arsenic concentration for New Jersey (including Edison) to be 20 ppm.

This recent sampling effort was a follow-up to soil sampling performed by EPA in July of 1993 at a number of locations including properties on Prince Street, Patrick Avenue and Cortland Street in Edison; the Edison Glen and Edison Woods residential complexes in Edison; and areas in Metuchen near Route 287. The July 1993 sampling results indicated arsenic concentrations were below 20 ppm in all but one off-site residential soil sample, while higher levels were found in and immediately adjacent to the unnamed stream and Mill Brook. The attached arsenic data resulting from the March/April 1994 sampling effort indicate a similar distribution of arsenic as follows:

- All but one of the residential soil sample (those samples taken from a residential yard or common ground within Edison Glen and Edison Woods residential complexes) analyses indicated arsenic concentrations below 20 ppm. Figures 1, 2, 3, and 5 provide the data associated with each of such sampling locations. The dumpster area shown in Figure 3 indicates the location of the one residential soil sample which generated an arsenic concentration above 20 ppm.
- The analyses of soil and sediment samples taken from areas in and immediately adjacent to the the unnamed stream and Mill Brook generated arsenic concentrations ranging from less than 1 up to 1100 ppm. The majority of the detected concentrations were below 20 ppm. Figure 6 provides the data associated with each of these samples.
- The three dirt and/or dust samples collected from specific apartments within the Edison Glen condominium complex indicated arsenic concentrations which ranged from approximately 0.8 ppm up to 11.3 ppm. Figure 4 provides the data associated with each of these apartments.

Soil in Residential Yards/Common Areas

EPA's evaluation of the off-site sampling data indicates that the potential exposure to arsenic does not present an immediate health risk. As stated above, with the exception of one sample, all soil in residential areas contained arsenic at concentrations of less than 20 ppm, which is consistent with naturally occurring levels in New Jersey. However, since arsenic contamination significantly above 20 ppm in a residential yard or common grounds within Edison Glen or Edison Woods residential complexes may pose a long-term risk, this contamination will be remediated by EPA.

Soil/Sediment in Unnamed Stream/Mill Brook

A Risk Assessment was conducted to estimate the human health risks associated with potential exposures to arsenic detected in the soils and sediments in and immediately adjacent to the Unnamed Stream and Mill Brook. The risk assessment was conducted using a highly conservative estimate of exposure, which is likely to overestimate the health risks related to the unnamed stream and Mill Brook. This risk assessment identified adolescents playing at the stream or the Brook as the most sensitive potential receptors to the contamination. Adolescents (ages 7-18) were assumed to play at the unnamed stream or Mill Brook once a week throughout the year, for 12 years. Exposures were assumed to occur primarily through incidental ingestion of soils and sediments contaminated with arsenic.

Based on the Risk Assessment for soils and sediments of the stream and brook, there is no unacceptable risk to human health posed by exposure to these areas. For known or suspected carcinogens such as arsenic, EPA considers excess upper-limit individual lifetime cancer

risks of between 10^{-4} and 10^{-6} to be acceptable. This range indicates that an individual has not greater than approximately a one in ten thousand to one in a million chance of developing cancer as a result of site-related exposure to a carcinogen under the specific exposure conditions at the site. The excess cancer risk for an adolescent exposed to arsenic in the soils and sediments in and immediately adjacent to the unnamed stream and Mill Brook (using the conservative assumptions outlined above) is 5×10^{-4} , which is well within EPA's acceptable risk range. In addition, non-carcinogenic effects are also highly unlikely to occur from the exposure scenario evaluated in the Risk Assessment.

Interior Dust

The level of arsenic found in the interior dust samples at the Edison Glen complex is generally consistent with the concentrations found in the surficial soils at Edison Glen, thus, suggesting that the major contributing factor to the interior arsenic levels is the soil derived dust (with naturally occurring levels of arsenic below 20 ppm) from outside the dwellings. The risk associated with the arsenic levels detailed above would also fall within EPA's acceptable excess lifetime cancer risk range of between 10^{-4} and 10^{-6} and would not pose a non-carcinogenic hazard for residents of the Edison Glen.

Public Availability Session and Meeting

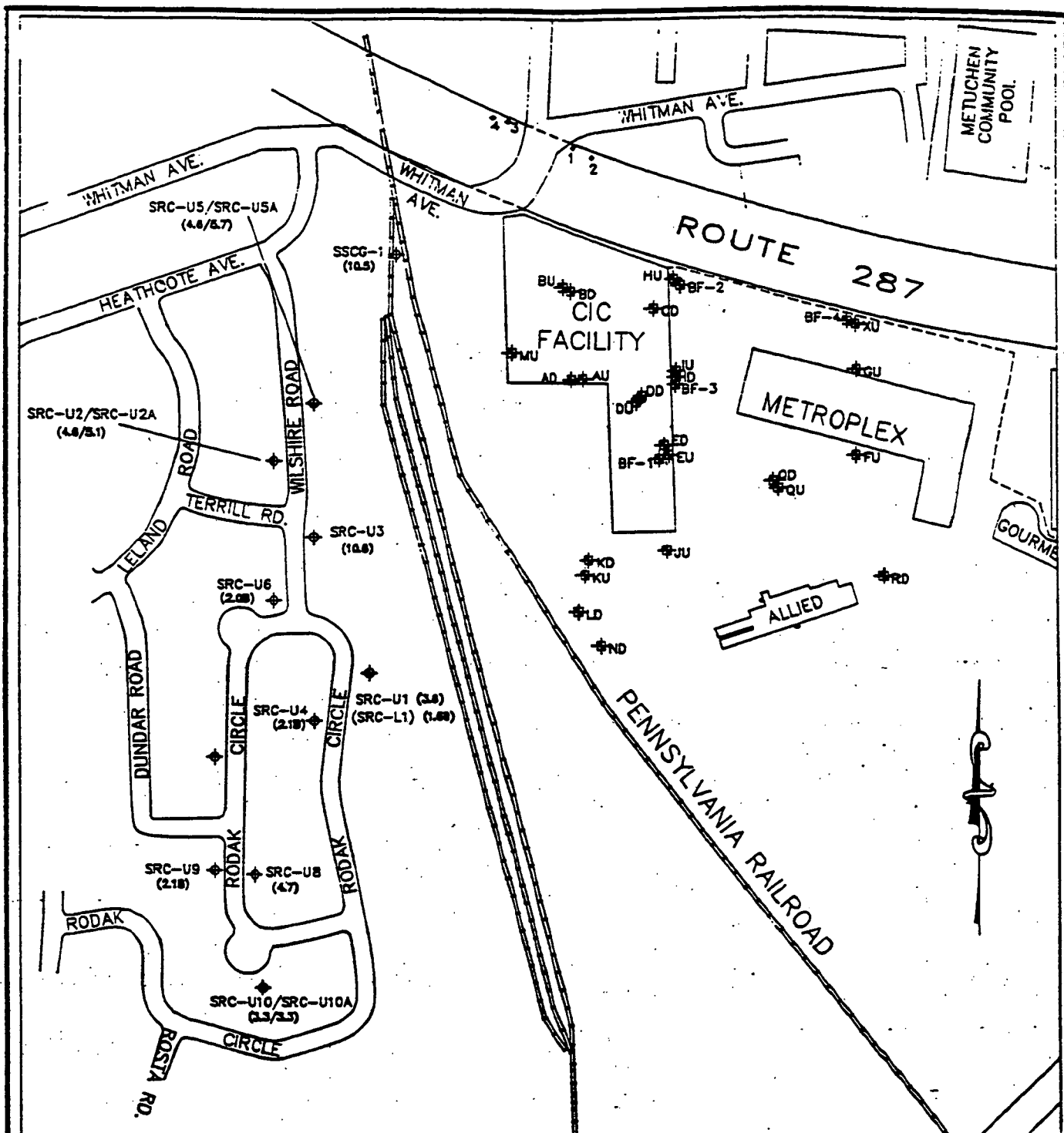
EPA has scheduled an availability session and public meeting to provide additional information and answer questions that community residents, property owners and others may have regarding the attached results and the CIC site. The availability session will be held from 1:00-4:00 PM and will be followed by a public meeting in the evening from 7:00-10:00 PM on Thursday, July 14, 1994 at the Stelton Community Center (Auditorium) located at 328 Plainfield Avenue in Edison, New Jersey (908-248-7309).

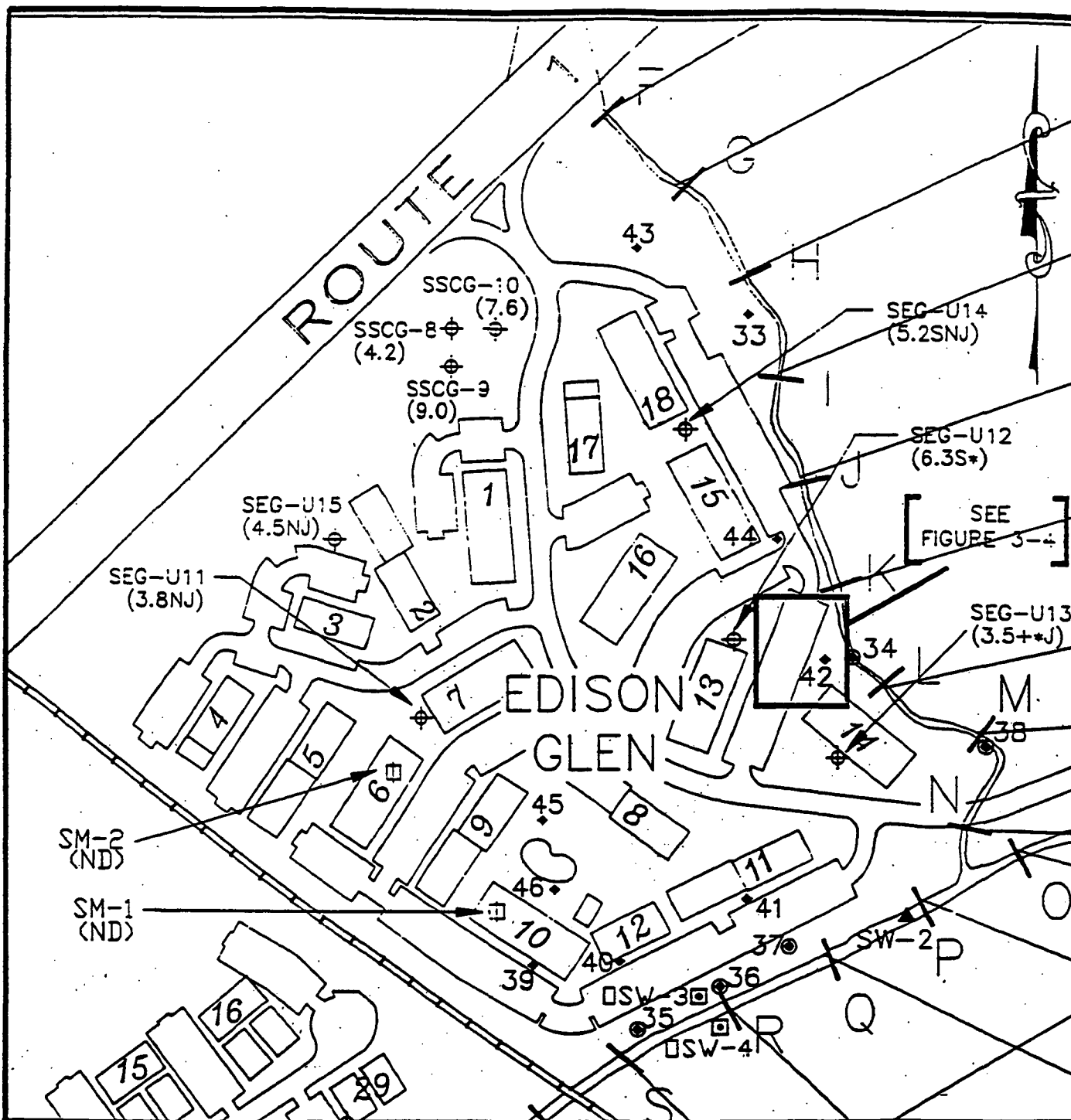
Please feel free to contact me at (212) 264-6311, or Ms. Cecilia Echols, Community Relations Coordinator, at (212) 264-0949.

Sincerely,



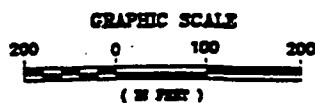
Pat Evangelista, Project Manager
Central New Jersey Section II
Emergency & Remedial Response Division





LEGEND

- ◆ REAC SURFACE SOIL SAMPLE AND DESIGNATION
- REAC SURFACE & SUBSURFACE SOIL SAMPLE AND DESIGNATION
- SEG-U11
- ⊕ SOIL SAMPLE LOCATION AND DESIGNATION (1994 INVESTIGATION) (3.8NJ) (ARSENIC CONCENTRATIONS ARE PRESENTED IN MG/KG)
- R/ TRANSECT LOCATIONS
- ~ SURFACE DRAINAGE

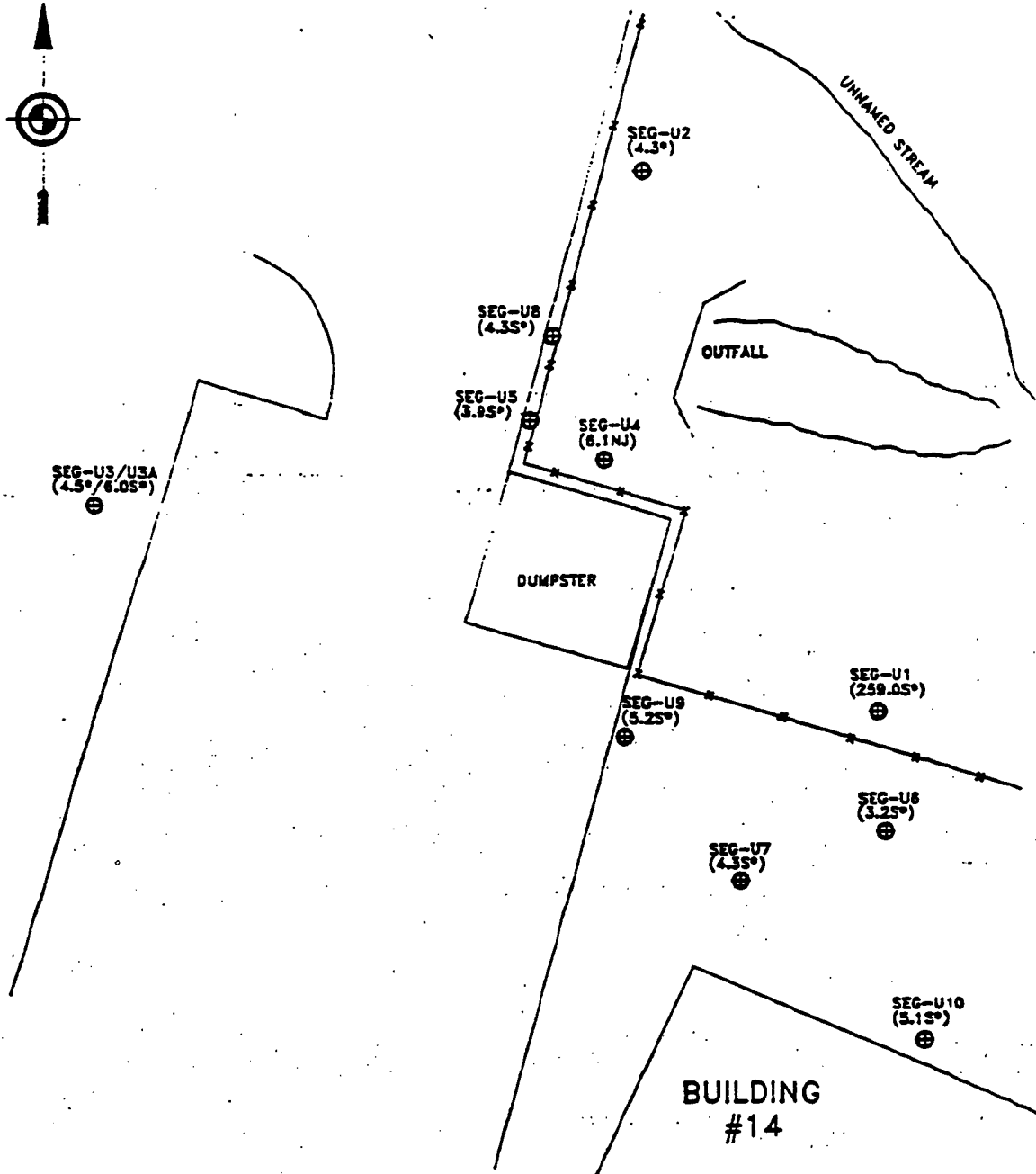


UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
EDISON CHEMICAL INSECTICIDE CORPORATION SITE NEW JERSEY

WESTON

FIGURE 2
SOIL SAMPLING LOCATIONS
INDICATING ARSENIC RESULTS
EDISON GLEN DEVELOPMENT

DATE	BY	DATE	BY
B. MAC		TRY-1	1
1" = 200'	04200-023-021	1	1



LEGEND

FIGURE IS NOT TO SCALE. DIMENSIONS ARE APPROXIMATE.

SEG-U4 SOIL SAMPLING LOCATION AND DESIGNATION

(6.1NJ) ARSENIC CONCENTRATIONS ARE PRESENTED IN MG/KG

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
EDISON CHEMICAL INSECTICIDE CORPORATION SITE NEW JERSEY

WESTON

FIGURE 3
SOIL SAMPLING LOCATIONS INDICATING ARSENIC
RESULTS IN PROXIMITY TO THE DUMPSTER
(USEPA-ERT SAMPLING LOC. #42)
EDISON GLEN DEVELOPMENT

DATE	BY	DATE	BY
8. MAC		DUMPSTER	0
1" = NTS	04200-023-021	1	1

REVISION 2 0/1/84 FILE INFO DUMPSTER

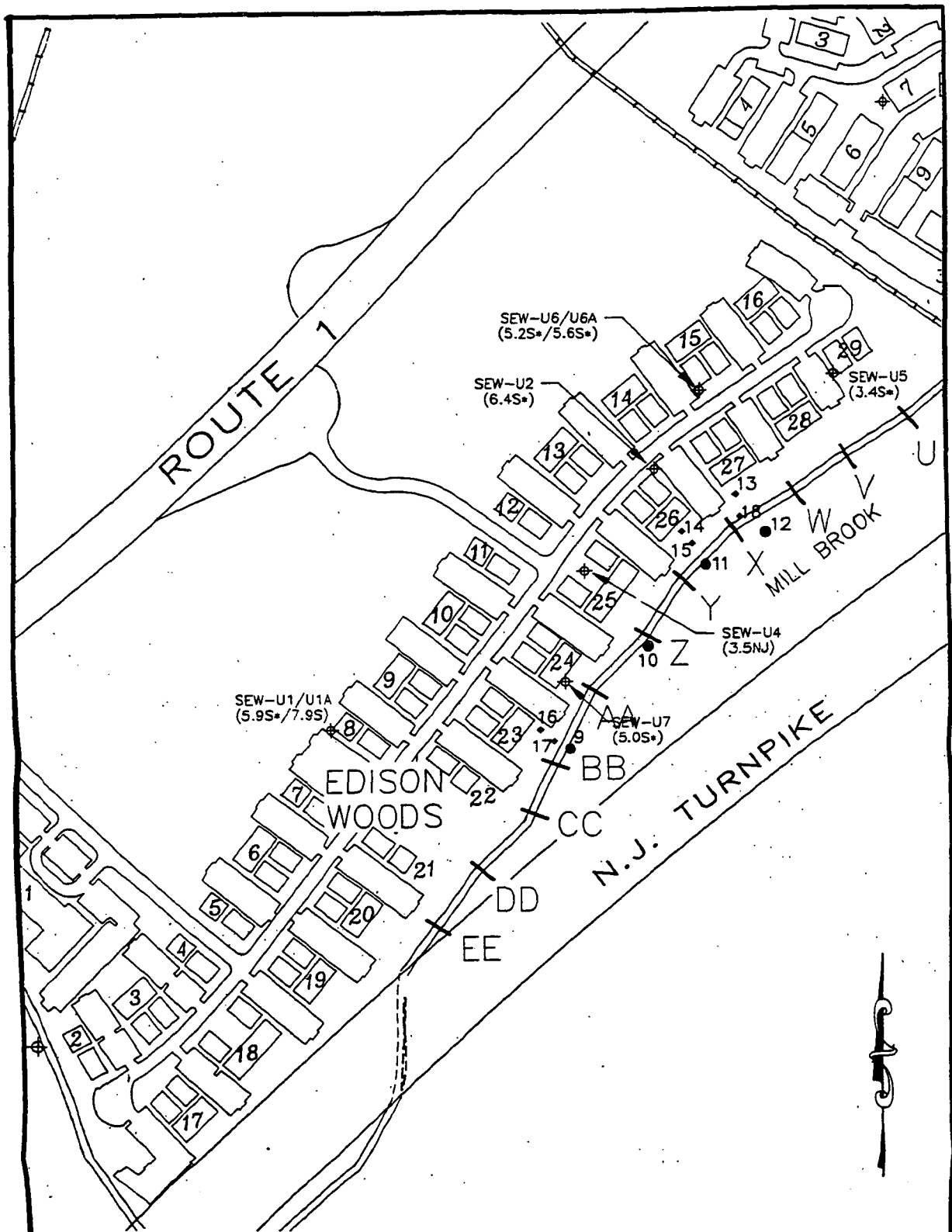
FIGURE 4
ARSENIC RESULTS - DUST/DIRT
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	SAMPLE DATE/TIME	SAMPLING LOCATION	RESULTS .ppm
DD-1	SB5060	3-30-94/1110	Apt. #1202	5.4
DD-2	SB5061	3-30-94/1110	Apt. #1202	3.92
DD-3	SB5062	3-30-94/1150	Apt. #1109 (vacant)	11.3
DD-4	SB5063	3-30-94/1215	Apt. #1409 (vacant)	0.77

NOTE: All samples were analyzed specifically for arsenic.

DD-2 is a duplicate of DD-1

DD-1 is a Matrix Spike



LEGEND

AA/ TRANSECT SAMPLING LOCATIONS

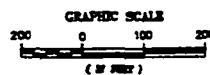
● REAC SURFACE & SUBSURFACE SOIL SAMPLE

• REAC SURFACE SOIL SAMPLE

SEW-U7 (3.5IU) SOIL SAMPLE LOCATION AND DESIGNATION (1994 INVESTIGATION) (ARSENIC CONCENTRATIONS ARE PRESENTED IN MG/KG)

— SURFACE DRAINAGE

— SUBSURFACE DRAINAGE



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
CHEMICAL INSECTICIDE CORPORATION SITE NEW JERSEY

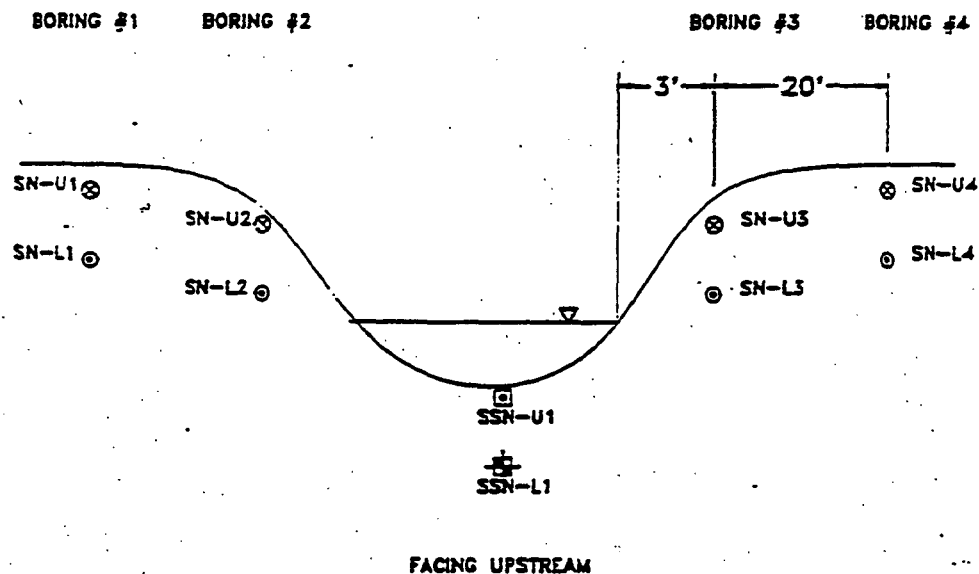
WISIN

FIGURE 5
SOIL SAMPLING LOCATIONS
INDICATING ARSENIC RESULTS
EDISON WOODS DEVELOPMENT

DATE	BY	REV	DATE	BY	REV
04/200	MAC	1	04/200	TRY-1	0
1" = 200'	04/200-023-021	1	1		

DUE TO THEIR LARGE SIZE,
FIGURES 6 AND 7 COULD NOT BE INCLUDED
WITH THIS DOCUMENT
AND ARE AVAILABLE FROM THE EPA REGION.

Enlargement of transect N from Figure 6. Other transects are the same except for their letter designation.



LEGEND

- SSN-U1 SEDIMENT SAMPLE (0.5-1.0 FOOT BELOW BOTTOM OF STREAM)
- ⊕ SSN-L1 SUBSURFACE SEDIMENT SAMPLE (2.5-3.0 FEET BELOW BOTTOM OF STREAM)
- ⊗ SN-U3 SURFACE SOIL SAMPLE (0.5-1.0 FOOT BELOW GRADE)
- ⊙ SN-L3 SUBSURFACE SOIL SAMPLE (2.5-3.0 FEET BELOW GRADE)

- NOT TO SCALE -

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY			
EDISON		CHEMICAL INSECTICIDE CORPORATION SITE	
NEW JERSEY			
WESTON			
FIGURE 8			
IDEALIZED TRANSECT SAMPLING LOCATIONS			
DATE	BY	NO. OF	NO. OF
8. MAC		4-8	1
DATE	BY	NO. OF	NO. OF
1" = NTS	04200-023-021	1	1

TABLE 3-2
VOLATILE ORGANIC RESULTS - SURFACE WATER/SUMPS/GROUNDWATER
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER SAMPLE LOCATION NO. CLP SAMPLE CODE DEPTH INTERVAL UNITS DATE SAMPLE COLLECTED	SUMP 1 SM-1 885040	TRIP 885042	SUMP 2 SM-2 885043	MW-1 885045	DUP MW-1 MW-2 885047	TRIP 885049	SW-1 885050	SW-2 885052	SW-3 885054	DUP SW-3 SW-4 885056	Field Bk 885058
	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l
Chloromethane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Bromomethane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Vinyl Chloride	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Chloroethane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Methylene Chloride	2U	2U	2U	2U	2U	2U	2U	2U	2U	2U	4
Acetone	R	R	R	R	R	R	R	R	R	R	R
Carbon Disulfide	1U	1U	1U	1U	1U	0.7J	1U	1U	1U	15U	0.4J
1,1-Dichloroethane	1U	1U	1U	1U	0.5U	1U	1U	1U	1U	1U	1U
1,1-Dichloroethane	1U	1U	1U	1U	1J	1U	1U	1U	1U	1U	1U
1,2-Dichloroethane (Total)	1U	1U	1U	1U	22	1U	1U	1U	1U	1U	1U
Chloroform	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
1,2-Dichloroethane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	0.2J
2-Butanone	1U	1U	1U	1U	3	1U	1U	1U	1U	1U	1U
1,1,1-Trichloroethane	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U
Carbon Tetrachloride	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Bromodichloromethane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
1,2-Dichloropropane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
cis-1,3-Dichloropropane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Trichloroethane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Dibromochloromethane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
1,1,2-Trichloroethane	1U	1U	1U	1U	10	1U	0.3U	0.6U	1	1U	1U
Benzene	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
trans-1,3-Dichloropropane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Bromocloro	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
4-Methyl-2-Pentanone	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
2-Hexanone	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Tetrachloroethene	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U
1,1,2,2-Tetrachloroethane	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U	5U
Toluene	1U	1U	1U	1U	1U	1U	0.4U	0.3U	1U	0.9U	1U
Chlorobenzene	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Ethylbenzene	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Styrene	1U	1U	1U	1U	1U	1U	0.2U	1U	1U	1U	1U
Xylene (Total)	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
1,3-Dichlorobenzene	1U	1U	1U	1U	1U	1U	1U	1U	0.3U	0.3U	1U
1,4-Dichlorobenzene	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
1,2-Dichlorobenzene	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
1,2-Dibrom-3-chloropropane	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
TOTAL TIC	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U	1U
Total TIC Concentration											

U = Analyte was not detected at the instrument detection limit given
J = Estimated Value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence of analyte; estimated quantity
R = Rejected during data validation

TABLE 3-3
SEMI-VOLATILE ORGANIC RESULTS - SURFACE WATER/SUMPS/GROUNDWATER
OPP-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	SM-1	SM-2	MW-1	MW-1 Dup	SW-1	SW-2	SW-3	SW-3 Dup	Field Bk
SAMPLE LOCATION NO.	SB5040	SB5043	SB5045	SB5047	SB5050	SB5052	SB5054	SB5056	SB5058
CLP SAMPLE CODE									
DEPTH INTERVAL									
UNITS	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l
DATE SAMPLE COLLECTED									
Phenol									
bb(2-Chloroethyl) ether									
2-Chlorophenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
1,3-Dichlorobenzene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
1,4-Dichlorobenzene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
1,2-Dichlorobenzene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2-Methylphenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2,2'-oxybis(1-Chloropropane)	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
4-Methylphenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
N-Nitroso-d-n-propylamine	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Hexachlorocyclopentadiene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Nitrobenzene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Isophorone	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2-Nitrophenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2,4-Dimethylphenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
bb(2-Chloroethoxy)methane	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2,4-Dichlorophenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
1,2,4-Trichlorobenzene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Naphthalene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
4-Chloroaniline	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Hexachlorobutadiene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
4-Chloro-3-methylphenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2-Methylnaphthalene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Hexachlorocyclopentadiene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2,4,6-Trichlorophenol	20 U	20 U	210 U	20 U	20 U	20 U	20 U	20 U	20 U
2,4,5-Trichlorophenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2-Chloronaphthalene	20 U	20 U	21 U	20 U	20 U	20 U	20 U	20 U	20 U
2-Nitroaniline	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Dimethylnaphthalene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Acenaphthylene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2,6-Dinitrotoluene	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U
3-Nitroaniline	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Acenaphthene									
2,4-Dinitrophenol	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U
4-Nitrophenol	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U
Dibenzofuran	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
2,4-Dinitrotoluene									
Diethylnthalate	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
4-Chlorophenyl-phenylether	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Fluorene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U

TABLE 3-3 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - SURFACE WATER/SUMPS/GROUNDWATER
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	SM-1	SM-2	MW-1	MW-1 Dup	SW-1	SW-2	SW-3	SW-3 Dup	Field Bk.
SAMPLE LOCATION NO.	SB5040	SB5043	SB5045	SB5047	SB5050	SB5052	SB5054	SB5056	SB5058
CLP SAMPLE CODE									
DEPTH INTERVAL									
UNITS	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l
DATE SAMPLE COLLECTED									
4-Nitroaniline	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
4,6-Dinitro-2-methylphenol	20 U	20 U	21 U	20 U	20 U	20 U	20 U	20 U	20 U
N-Nitrosodiphenylamine (1)	20 U	20 U	21 U	20 U	20 U	20 U	20 U	20 U	20 U
4-Bromophenyl-phenylether	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Hexachlorobenzene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Pentachlorophenol	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Phenanthrene	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U	20 U
Anthracene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Carbazole	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Di-n-butylphthalate	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Fluoranthene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Pyrene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Butylbenzylphthalate	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
3,3'-Dichlorobenzidine	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Benzo(a)anthracene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Chrysene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
bis(2-Ethylhexyl)phthalate	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Di-n-octylphthalate	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Benzo(b)fluoranthene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Benzo(k)fluoranthene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Benzo(a)pyrene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Indeno(1,2,3-cd)pyrene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Dibenz(a,h)anthracene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
Benzo(ghi)perylene	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U	5 U
TIC Count									
Total TIC Concentration									

U = Analyte was not detected at the instrument detection limit given
B = Analyte was detected in blank
NJ = Presumptive evidence for presence of analyte; estimated quantity
J = Estimated value
R = Rejected during data validation
(1) = Cannot be separated from Diphenylamine

TABLE 3-4
HERBICIDE/PESTICIDE/PCB RESULTS - SURFACE WATER/SUMPS/GROUNDWATER
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER SAMPLE LOCATION NO. CLP SAMPLE CODE DEPTH INTERVAL UNITS DATE SAMPLE COLLECTED	SM-1 SB5040	SM-2 SB5043	NW-1 SB5045	NW-1 Dup SB5047	SW-1 SB5050	SW-2 SB5052	SW-3 SB5054	SW-3 Dup SB5056	Field Bk SB5058
	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l
alpha-BHC	0.010 U	0.010 U	0.010 U	0.010 U	0.022	0.20 D	0.0082 JN	0.0070	0.010
beta-BHC	0.010 U	0.010 U	0.010 U	0.010 U	0.010 U	0.057	0.010 U	0.010	0.010
delta-BHC	0.010 U	0.010 U	0.010 U	0.010 U	0.010 U	0.023 P	0.010 U	0.010	0.010
gamma-BHC (Lindane)	0.010 U	0.010 U	0.010 U	0.010 U	0.010 U	R	0.051 NJ	0.048	0.010
Heptachlor	0.010 U	0.010 U	0.010 U	0.010 U	0.010 U	0.041	0.012 NJ	0.014	0.010
Aldrin	0.010 U	0.010 U	0.010 U	0.01 U	0.010 U	0.010 U	0.010 U	0.010	0.010
Heptachlor epoxide	0.010 U	0.010 U	0.010 U	0.010 U	0.0049 J	0.01 U	0.010 U	0.010	0.010
Endosulfan I	0.010 U	0.010 U	0.010 U	0.010 U	0.010 U	0.010	0.010 U	0.010	0.010
Dieldrin	0.020 U	0.020 U	0.02 U	0.02 U	0.020 U	0.075	0.020 U	0.020	0.020
4,4'-DDE	0.020 U	0.020 U	0.020 U	0.020 U	0.018 J	0.58 D	0.020 U	0.020	0.020
Endrin	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	R	0.020 U	0.02	0.020
Endosulfan II	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020	0.020
4,4'-DDD	0.020 U	0.020 U	0.020 U	0.020 U	0.02 U	R	0.020 U	0.020	0.020
Endosulfan Sulfate	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020	0.020
4,4'-DDT	0.020 U	0.020 U	0.020 U	0.020 U	0.002	3.3 D	0.020 U	0.020	0.020
Methoxychlor	0.10 U	0.10 U	0.10 U	0.10 U	0.1 U	0.10 U	0.10 U	0.10	0.10
Endrin ketone	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.012 J	0.020 U	0.020	0.020
Endrin Aldehyde	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020 U	0.020	0.020
alpha-Chlordane	0.010 U	0.010 U	0.010 U	0.010 U	0.01 U	0.12 PJ	0.010 U	0.01	0.010
gamma-Chlordane	0.010 U	0.010 U	0.030 U	0.030 U	0.0080 J	0.14	0.010 U	0.010	0.010
Toxaphene	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U
Aroclor-1016	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U
Aroclor-1221	0.40 U	0.40 U	0.40 U	0.40 U	0.40 U	0.40 U	0.40 U	0.40 U	0.40 U
Aroclor-1232	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U
Aroclor-1242	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U
Aroclor-1248	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U
Aroclor-1254	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U
Aroclor-1260	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U

U = Analyte was not detected at the instrument detection limit given
J = Estimated value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence of analyte; estimated quantity
P = There is a greater than 25% difference for detected concentrations
between the two GC columns; the lower of the two values is reported.
R = Rejected during data validation

TABLE 3-5
INORGANIC RESULTS - SURFACE WATER/SUMP/SURROUNDWATER
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER SAMPLE LOCATION NO. CLP SAMPLE CODE DEPTH INTERVAL UNITS DATE SAMPLE COLLECTED	← SUMP 1 →		← SUMP 2 →		← HW-1 →		← HW-1 DUP →		SW-1	SW-2	SW-3	SW-3 DUP	Field Blank (sum)	Field Blank (Extraction)
	(total) MBNP01	(dissolved) MBNP02	(total) MBNP03	(dissolved) MBNP04	(total) MBNP05	(dissolved) MBNP06	(total) MBNP07	(dissolved) MBNP08	(total) MBNP09	(total) MBNP11	(total) MBNP13	(total) MBNP15	MBNP17	MBNP18
	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l	ug/l
Aluminum	484	424 B	608	607 B	127 B	204 B	142 B	208	1030	2200	1370	1300	206 U	206 U
Antimony	283 U	283 U	283 U	283 U	283 U	283 U	283 U	374	285 B	283 U	285 B	285 B	283 U	286 U
Arsenic	14 B	17 B	52 B	84 B	13 U	13 U	13 U	13	128	112	31 B	34 B	13 U	13 U
Barium	206 B	203 B	205 BE	201 B	621 B	642 B	664 B	661	785 B	716 B	300 B	31 B	645 B	640 U
Beryllium	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U
Cadmium	27 U	27 U	27 U	27 U	3 U	3.0 B	2.7 U	2.7	44 B	2.7 U	3 U	27 U	27 U	27 U
Calcium	48500 E	81700	27800 E	28000	30400	42500	38900	40300	14800	13900	9050	9100	383 B	231 U
Chromium	28 U	28 U	28 U	28 U	28 U	28 U	28 U	28	81 B	7.7 B	46 B	28 B	28 U	28 U
Cobalt	28 U	28 U	28 U	28 U	32 B	32 B	38 U	29	28 U	42 B	46 B	28 U	28 U	28 U
Copper	71 B	44 B	27 B	138 B	24 U	24 U	24 U	24	412	188 B	121 B	118 B	168 B	24 U
Iron	803 E	444 B	804 E	588 B	3450	3080	3400	2620	3100	3410	1910	1850	216 B	81 U
Lead	43 WH	10 B	33 WH	690 B	24 B	690 B	45	41	282	286	75	73	616 B	1140 U
Magnesium	7030 E	7890	3740 BE	3040 B	15300	16800	14800	15900	3270 B	3680 B	2320 B	2320 B	218 U	268 U
Manganese	98 BE	18 B	80 BE	17 U	604	635 00	587	600	108	111	88 B	90 B	17 U	17 U
Mercury	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U
Nickel	108 U	108	108 U	108 U	108 U	138 B	108 U	108	108 U	108 U	108 U	108 U	108 U	108 U
Potassium	13800	14800	7116	7580	6870	7280	8400	7080	2000 B	2110 B	1780 B	1560 B	348 U	348 U
Selenium	12 U	12 U	12 U	12 U	12 U	12 U	12 U	12	12 U	12 U	12 U	12 U	12 U	12 U
Silver	28 U	24 U	28 U	28 U	28 U	28 U	28 U	28	28 U	28 U	28 U	28 U	28 U	28 U
Sodium	84000 E	25800	12800 E	28900	28900	31100	27100	28200	47700	34500	13900	13300	158 B	108 B
Thallium	10 U	10 U	10 U	10 U	10 U	10 U	10 U	10 U	10 U	10 U	10 U	10 U	10 U	10 U
Vanadium	28 B	23 U	23 U	23 U	23 U	23 U	23 U	23 U	44 U	62 B	38 B	32 B	23 U	23 U
Zinc	101	52 B	29 B	244	154 B	128 B	164	121	121	109	650	650	88 B	31 U
Cyanide	100 U		100 U		100 U		100		100 U	100 U	100 U	100 U	100 U	

U = Analyte was not detected at the instrument detection limit given
B = Reported value is between the instrument detection limit and the contract required detection limit
E = Value is estimated due to interference
N = Spiked sample recovery was not within control limits
* = Duplicate analysis was not within control limits
+ = Correlation coefficient for the MSA is less than 0.999
J = Estimated value
S = Determined by method of standard addition (MSA)
W = Post-digestion spikes for Furnace AA analysis out of control limits, while sample absorbance is less than 80% of spike absorbance
R = Rejected during data validation
M = Duplicate injection precision criteria was not met
NA = Not Analyzed

TABLE 3-6
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SA-L1	SB5844	Soil	3-22-94/1110	2.5 - 3.0	As	Hand Auger	8.4 +NJ
SA-U1	BMN01 MBNQ01	Soil	3-22-94/1050	0.5 - 1.0	TCL TAL/CN	Split Spoon Hand Auger	21.3 J
SA-U2	SB5843	Soil	3-22-94/1040	0.5 - 1.0	As	Hand Auger	32.1 NJ
SA-U3	SB5847	Soil	3-22-94/1015	0.5 - 1.0	As	Hand Auger	26.2 SNJ
SA-L3	SB5848	Soil	3-22-94/1015	2.0 - 2.5	As	Hand Auger	21.3 SNJ
SA-U4	SB5849	Soil	3-22-94/1025	0.5 - 1.0	As	Hand Auger	14.9 SNJ
SA-L4	SB5850	Soil	3-22-94/1025	2.5 - 3.0	As	Hand Auger	55.8
SSA-U1	BMN02 MBNQ02	Sediment	3-22-94/1030	0.2 - 0.8	TCL TAL/CN	Split Spoon	106.0 + (R)
SSA-L1	SB5846	Sediment	3-22-94/1056	1.5 - 2.0	As	Split Spoon	30.7 SNJ
SB-U1	SB5833	Soil	3-22-94/0935	0.5 - 1.0	As	Hand Auger	5.7 SNJ
SB-L1	SB5834	Soil	3-22-94/0940	1.5 - 2.0	As	Hand Auger	7.6 SNJ
SB-U2	BMN03 MBNQ03	Soil	3-22-94/0905	0.5 - 1.0	TCL TAL/CN	Hand Auger	11.0 J
SB-U3	SB5838	Soil	3-22-94/0845	0.5 - 1.0	As	Hand Auger	13.4 SNJ
SB-L3	SB5842	Soil	3-22-94/0905	2.5 - 3.0	As	Hand Auger	5.0 SNJ
SB-U4	SB5841	Soil	3-22-94/0905	0.5 - 1.0	As	Hand Auger	16.5 SNJ
SB-L4	SB5840	Soil	3-22-94/0925	2.0 - 2.5	As	Hand Auger	7.7 SNJ
SSB-U1	SB5836	Sediment	3-22-94/0920	0.5 - 1.0	As	Split Spoon	366.0
SSB-L1	SB5837	Sediment	3-22-94/0935	1.5 - 2.0	As	Split Spoon	89.9
SC-U1	SB5826	Soil	3-22-94/0800	0.5 - 1.0	As	Hand Auger	6.7 SNJ
SC-U2	SB5827	Soil	3-22-94/0838	0.5 - 1.0	As	Hand Auger	36.3 SNJ
SC-U3	SB5832	Soil	3-22-94/0810	0.5 - 1.0	As	Hand Auger	61.4
SC-L3	BMN04 MBNQ04	Soil	3-22-94/0824	1.0 - 1.5	TCL TAL/CN	Hand Auger	6.1 NJ
SC-U4	SB5831	Soil	3-22-94/0800	0.5 - 1.0	As	Hand Auger	5.0 SNJ
SSC-U1	SB5828	Sediment	3-22-94/0802	0.5 - 1.0	As	Split Spoon	20.1 SNJ
SSC-L1	SB5829	Sediment	3-22-94/0810	1.5 - 2.0	As	Split Spoon	4.4 SNJ
SD-U1	SB5812	Soil	3-21-94/1550	0.5 - 1.0	As	Hand Auger	1.7B

PAGES 3-11 TO 3-16 PURPOSELY OMITTED.

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SD-L1	SB5813	Soil	3-21-94/1600	2.0 - 2.5	As	Hand Auger	1.4B
SD-U2	SB5814	Soil	3-21-94/1550	0.5 - 1.0	As	Hand Auger	11.9
SD-U3	SB5819	Soil	3-21-94/1540	0.5 - 1.0	As	Hand Auger	9.8
SSD-U1	SB5816	Sediment	3-21-94/1600	0.2 - 1.0	As	Split Spoon	37.4
SSD-U1A	SB5817	Sediment	3-21-94/1600	0.2 - 1.0	As	Split Spoon	27.2
SSD-L1	SB5818	Sediment	3-21-94/1605	1.5 - 2.0	As	Hand Auger	38.6
SE-U1	SB5803	Soil	3-21-94/1440	0.5 - 1.0	As	Hand Auger	9.2
SE-L1	SB5804	Soil	3-21-94/1500	2.5 - 3.0	As	Hand Auger	2.8
SE-U2	SB5805	Soil	3-21-94/1441	0.5 - 1.0	As	Hand Auger	5.2
SE-L2	BMN05 MBNQ05	Soil	3-21-94/1500	2.5 - 3.0	TCL TAL/CN	Hand Auger	15.4 S
SE-U3	SB5808	Soil	3-21-94/1435	0.5 - 1.0	As	Hand Auger	6.4
SE-L3	SB5809	Soil	3-21-94/1500	2.5 - 3.0	As	Hand Auger	3.3
SE-U4	SB5810	Soil	3-21-94/1430	0.5 - 1.0	As	Hand Auger	7.2
SE-L4	SB5811	Soil	3-21-94/1450	2.5 - 3.0	As	Hand Auger	6.2
SSE-U1	SB5806	Sediment	3-21-94/1451	0.3 - 1.0	As	Split Spoon	5.8
SSE-U1A	SB5839	Sediment	3-21-94/1451	0.3 - 1.0	As	Split Spoon	24.6
SSE-L1	SB5807	Sediment	3-21-94/1510	1.5 - 2.0	As	Split Spoon	4.3
SF-U1	SB5797	Soil	3-21-94/1235	0.5 - 1.0	As	Hand Auger	3.2J
SF-L1	SB5798	Soil	3-21-94/1245	2.5 - 3.0	As	Hand Auger	3.8J
SF-U4	SB5799	Soil	3-21-94/1230	0.5 - 1.0	As	Hand Auger	11.8J
SF-U4A	SB5800	Soil	3-21-94/1230	0.5 - 1.0	As	Hand Auger	6.2J
SF-L4	SB5801	Soil	3-21-94/1235	2.5 - 3.0	As	Hand Auger	3.1
SG-U1	SB5788	Soil	3-21-94/1130	0.5 - 1.0	As	Hand Auger	1.7B
SG-L1	SB5789	Soil	3-21-94/1135	2.5 - 3.0	As	Hand Auger	1.6B
SG-U2	SB5790	Soil	3-21-94/1156	0.6 - 1.0	As	Hand Auger	37.5
SG-L2	BMN07 MBNQ07	Soil	3-21-94/1200	2.0 - 2.5	TCL TAL/CN	Hand Auger	0.23 BWJ
SG-U3	SB5793	Soil	3-21-94/1140	0.5 - 1.0	As	Hand Auger	12.6
SG-L3	SB5794	Soil	3-21-94/1155	2.5 - 3.0	As	Hand Auger	2.6

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SG-U4	SB5795	Soil	3-21-94/1235	0.5 - 1.0	As	Hand Auger	12.3
SG-L4	SB5796	Soil	3-21-94/1250	2.0 - 2.5	As	Hand Auger	3.7J
SSG-U1	SB5791	Sediment	3-21-94/1125	0.5 - 1.0	As	Split Spoon	4.5
SSG-L1	SB5792	Sediment	3-21-94/1141	1.5 - 2.0	As	Split Spoon	1.7B
SH-U1	BMN35 MBNQ35	Soil	3-21-94/1025	0.5 - 1.0	TCL TAL/CN	Hand Auger	12.0J
SH-U1A	BMN36 MBNQ36	Soil	3-21-94/1025	0.5 - 1.0	TCL TAL/CN	Hand Auger	6.9 SJ
SH-L1	SB5780	Soil	3-21-94/1025	2.5 - 3.0	As	Hand Auger	2.9
SH-U2	SB5781	Soil	3-21-94/1105	0.5 - 1.0	As	Hand Auger	9.0
SH-L2	SB5782	Soil	3-21-94/1110	0.5 - 1.0	As	Hand Auger	6.4
SH-U3	SB5785	Soil	3-21-94/1025	0.5 - 1.0	As	Hand Auger	8.6
SH-L3	SB5786	Soil	3-21-94/1035	2.5 - 3.0	As	Hand Auger	8.0
SH-U4	BMN37 MBNQ37	Soil	3-21-94/1055	0.5 - 1.0	TCL TAL/CN	Hand Auger	17.9 S
SH-L4	SB5787	Soil	3-21-94/1105	2.5 - 3.0	As	Hand Auger	7.3
SSH-U1	SB5783	Sediment	3-21-94/1030	0.5 - 1.0	As	Split Spoon	1.5B
SSH-L1	SB5784	Sediment	3-21-94/1038	1.5 - 2.0	As	Split Spoon	3.9
SI-U1	BMN13 MBNQ13	Soil	3-18-94/0950	0.5 - 1.0	TCL TAL/CN	Hand Auger	8.1 S*
SI-L1	SB5772	Soil	3-18-94/1000	2.0 - 2.5	As	Hand Auger	5.8 SNJ
SI-U2	SB5773	Soil	3-18-94/1025	0.5 - 1.0	As	Hand Auger	131.0
SI-L2	SB5774	Soil	3-18-94/1030	2.5 - 3.0	As	Hand Auger	30.5 NSJ
SI-U3	SB5777	Soil	3-18-94/1010	0.5 - 1.0	As	Split Spoon	50.7
SI-L3	SB5778	Soil	3-18-94/1030	2.5 - 3.0	As	Hand Auger	56.5
SSI-U1	SB5775	Sediment	3-18-94/1128	0.5 - 1.0	As	Hand Auger	42.5 NSJ
SSI-U1A	SB5779	Sediment	3-18-94/1128	0.5 - 1.0	As	Hand Auger	83.3
SSI-L1	SB5776	Sediment	3-18-94/1130	1.5 - 2.0	As	Split Spoon	97.9
SJ-U1	SB5759	Soil	3-18-94/0835	0.5 - 1.0	As	Hand Auger	2.6 SN* (R)
SJ-L1	SB5760	Soil	3-18-94/0850	2.5 - 3.0	As	Hand Auger	7.6 SN* (R)

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SJ-U2	SB5761	Soil	3-18-94/0840	0.5 - 1.0	As	Hand Auger	39.2 SN* (R)
SJ-L2	BMN08 MBNQ08	Soil	3-18-94/0850	2.0 - 2.5	TCL TAL/CN	Refusal at 2.5	85.6 NJ
SJ-U3	SB5766	Soil	3-18-94/0930	0.5 - 1.0	As	Hand Auger	150.0*
SJ-U4	SB5768	Soil	3-18-94/0900	0.5 - 1.0	As	Hand Auger	8.2 SN* (R)
SJ-U4A	SB5769	Soil	3-18-94/0900	0.5 - 1.0	As	Hand Auger	8.2 SN* (R)
SJ-L4	SB5770	Soil	3-18-94/0915	2.0 - 2.5	As	Hand Auger	212.0*
SSJ-U1	SB5762	Sediment	3-18-94/1217	0.3 - 1.2	As	Split Spoon	7.5 SN* (R)
SSJ-U1A	SB5771	Sediment	3-18-94/1230	0.3 - 1.2	As	Split Spoon	2.0 BSN* (R)
SSJ-L1	SB5763	Sediment	3-18-94/1230	1.5 - 2.0	As	Split Spoon	3.4 SN* (R)
SK-U1	SB5754	Soil	3-18-94/0759	0.5 - 1.0	As	Hand Auger	5.7 SN* (R)
SK-L1	SB5755	Soil	3-18-94/0817	2.5 - 3.0	As	Hand Auger	42.5 SN* (R)
SK-U2	SB5756	Soil	3-18-94/0750	0.5 - 1.0	As	Hand Auger	336.0*
SK-L2	SB5757	Soil	3-18-94/0805	2.0 - 2.5	As	Hand Auger	1100.0*
SK-U3	SB5764	Soil	3-18-94/0800	0.5 - 1.0	As	Hand Auger	13.2 SN* (R)
SK-L3	SB5765	Soil	3-18-94/0820	2.5 - 3.0	As	Hand Auger	66.5*
SSK-U1	BMN09 MBNQ09	Sediment	3-18-94/1250	0.3 - 1.2	TCL TAL/CN	Split Spoon	17.6S*
SSK-L1	SB5758	Sediment	3-18-94/1310	1.5 - 2.0	As	Split Spoon	58.5*
SL-U1	SB5747	Soil	3-17-94/1435	0.5 - 1.0	As	Hand Auger	4.1
SL-L1	SB5748	Soil	3-17-94/1440	2.0 - 2.5	As	Hand Auger	327.0
SL-U2	SB5749	Soil	3-17-94/1445	0.5 - 1.0	As	Hand Auger	71.0
SL-L2	SB5750	Soil	3-17-94/1450	2.5 - 3.0	As	Hand Auger	11.4
SL-U3	BMN11 MBNQ11	Soil	3-17-94/1510	0.5 - 1.0	TCL TAL/CN	Hand Auger	130.0 NJ
SL-L3	SB5702	Soil	3-17-94/1515	2.5 - 2.9	As	Hand Auger	27.5 J
SSL-U1	SB5751	Sediment	3-18-94/1305	0.3 - 1.2	As	Split Spoon	1.3 BWN* (R)
SSL-L1	SB5752	Sediment	3-18-94/1350	1.5 - 2.0	As	Split Spoon	0.89 BWN* (R)
SM-U1	SB5740	Soil	3-17-94/1350	0.5 - 1.0	As	Hand Auger	680.0
SM-L1	BMN10	Soil	3-17-94/1405	2.5 - 3.0	TCL	Hand Auger	96.3 NJ

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
	MBNQ10				TAL/CN		
SM-U2	SB5741	Soil	3-17-94/1350	0.5 - 1.0	As	Hand Auger	380.0
SM-L2	SB5742	Soil	3-17-94/1355	2.5 - 3.0	As	Hand Auger	14.0
SM-U3	SB5745	Soil	3-17-94/1410	0.5 - 1.0	As	Hand Auger	4.1
SM-L3	SB5746	Soil	3-17-94/1415	2.5 - 3.0	As	Hand Auger	114.0
SM-U4	SB5724	Soil	3-17-94/1420	0.5 - 1.0	As	Hand Auger	3.0
SM-L4	SB5727	Soil	3-17-94/1430	1.5 - 2.0	As	Hand Auger	3.4
SSM-U1	SB5743	Sediment	3-18-94/1410	0.5 - 1.0	As	Split Spoon	5.3 SN* (R)
SSM-L1	SB5744	Sediment	3-18-94/1450	1.5 - 2.0	As	Split Spoon	13.8 NSJ
SN-U1	SB5729	Soil	3-17-94/1055	0.5 - 1.0	As	Hand Auger	5.0
SN-L1	SB5730	Soil	3-17-94/1120	2.5 - 3.0	As	Hand Auger	10.2
SN-U2	SB5731	Soil	3-17-94/1105	0.5 - 1.0	As	Hand Auger	58.1
SN-U2A	SB5732	Soil	3-17-94/1105	0.5 - 1.0	As	Hand Auger	52.9
SN-L2	SB5733	Soil	3-17-94/1115	2.5 - 3.0	As	Hand Auger	10.0
SN-U3	SB5736	Soil	3-17-94/1157	0.5 - 1.0	As	Hand Auger	32.4
SN-U4	SB5737	Soil	3-17-94/1150	0.5 - 1.0	As	Hand Auger	86.3
SSN-U1	SB5734	Sediment	3-17-94/1130	0.5 - 1.0	As	Hand Auger	41.6
SSN-L1	SB5735	Sediment	3-21-94/0840	1.5 - 2.0	As	Split Spoon	7.8
SO-U1	SB5721	Soil	3-17-94/1233	0.5 - 1.0	As	Hand Auger	35.1
SO-L1	SB5722	Soil	3-17-94/1240	2.0 - 2.5	As	Hand Auger	9.8
SO-U2	SB5723	Soil	3-17-94/1254	0.5 - 1.0	As	Hand Auger	8.4
SO-U3	SB5726	Soil	3-17-94/1300	0.5 - 1.0	As	Hand Auger	3.9
SO-U4	SB5728	Soil	3-17-94/1315	0.5 - 1.0	As	Hand Auger	4.8
SO-L4	BMN14 MBNQ14	Soil	3-17-94/1327	2.5 - 3.0	TCL TAL/CN	Hand Auger	5.6 NSJ
SSO-U1	BMN12 MBNQ12	Sediment	3-21-94/0920	0.3 - 1.2	TCL TAL/CN	Split Spoon	4.4
SSO-L1	SB5725	Sediment	3-21-94/0925	1.5 - 2.0	As	Split Spoon	5.2
SP-U1	SB5712	Soil	3-17-94/0950	0.5 - 1.0	As	Hand Auger	17.6
SP-L1	SB5713	Soil	3-17-94/1000	2.5 - 3.0	As	Hand Auger	111.0

TABLE 3-6 CONTINUED
ARSENIC RESULTS – SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SP-U2	SB5709	Soil	3-17-94/0940	0.5 - 1.0	As	Hand Auger	224.0
SP-U2A	SB5710	Soil	3-17-94/0940	0.5 - 1.0	As	Hand Auger	190.0
SP-L2	SB5711	Soil	3-17-94/0947	1.5 - 2.0	As	Hand Auger	78.0
SP-U3	SB5716	Soil	3-17-94/1010	0.5 - 1.0	As	Hand Auger	14.3
SP-U4	SB5718	Soil	3-17-94/1030	0.5 - 0.8	As	Hand Auger	2.9
SSP-U1	SB5714	Sediment	3-17-94/1010	0.5 - 1.0	As	Hand Auger	46.1
SSP-L1	SB5715	Sediment	3-21-94/0855	1.5 - 2.0	As	Split Spoon	24.6
SQ-U1	SB5700	Soil	3-17-94/0753	0.5 - 1.0	As	Hand Auger	51.3
SQ-L1	BMN15 MBNQ15	Soil	3-17-94/0806	2.0 - 2.5	TCL TAL/CN	Hand Auger	25.4 NJ
SQ-U2	SB5701	Soil	3-17-94/0811	0.5 - 1.0	As	Hand Auger	20.8
SQ-U3	SB5705	Soil	3-17-94/0845	0.5 - 1.0	As	Hand Auger	13.6
SQ-L3	SB5706	Soil	3-17-94/0910	2.0 - 2.5	As	Hand Auger	47.7
SQ-U4	SB5707	Soil	3-17-94/0922	0.5 - 1.0	As	Split Spoon	5.1
SQ-L4	SB5708	Soil	3-17-94/0935	1.5 - 2.0	As	Hand Auger	14.2
SSQ-U1	SB5703	Sediment	3-17-94/0821	0.5 - 1.1	As	Split Spoon	2.6
SSQ-L1	SB5704	Sediment	3-17-94/0855	1.5 - 2.0	As	Split Spoon	5.2
SR-U1	SB5690	Soil	3-16-94/1520	0.5 - 1.0	As	Hand Auger	23.7
SR-L1	SB5691	Soil	3-16-94/1530	2.0 - 2.5	As	Hand Auger	82.2
SR-U2	SB5692	Soil	3-16-94/1520	0.5 - 1.0	As	Hand Auger	30.7
SR-L2	BMN16 MBNQ16	Soil	3-16-94/1530	2.5 - 3.0	TCL TAL/CN	Hand Auger	7.0 NSJ
SR-U3	SB5696	Soil	3-16-94/1550	0.5 - 1.0	As	Hand Auger	15.9
SSR-U1	SB5693	Sediment	3-16-94/1550	0.5 - 1.0	As	Split Spoon	4.2
SSR-U1A	SB5694	Sediment	3-16-94/1550	0.5 - 1.0	As	Split Spoon	2.8
SSR-L1	SB5695	Sediment	3-16-94/1600	2.0 - 2.6	As	Split Spoon	1.1B
SS-U1	SB5681	Soil	3-16-94/1325	0.5 - 1.0	As	Hand Auger	5.7
SS-L1	SB5682	Soil	3-16-94/1335	2.0 - 2.5	As	Hand Auger	46.7
SS-U2	SB5683	Soil	3-16-94/1325	0.5 - 1.0	As	Hand Auger	88.4
SS-L2	SB5684	Soil	3-16-94/1340	2.5 - 3.0	As	Hand Auger	69.7

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SS-U3	SB5687	Soil	3-16-94/1410	0.5 - 1.0	As	Hand Auger	18.3
SS-U4	SB5688	Soil	3-16-94/1420	0.5 - 1.0	As	Hand Auger	4.2
SSS-U1	SB5685	Sediment	3-16-94/1350	0.5 - 1.2	As	Split Spoon	3.2
SSS-L1	SB5686	Sediment	3-16-94/1410	1.5 - 2.0	As	Split Spoon	294.0
ST-U1	SB5671	Soil	3-16-94/0916	0.5 - 1.0	As	Hand Auger	25.0
ST-L1	SB5672	Soil	3-16-94/0927	1.0 - 1.5	As	Hand Auger	27.6
ST-U2	SB5673	Soil	3-16-94/0912	0.5 - 1.0	As	Hand Auger	13.2
ST-L2	SB5674	Soil	3-16-94/0920	1.2 - 1.5	As	Hand Auger	21.0
ST-U3	SB5678	Soil	3-16-94/0945	0.5 - 1.0	As	Hand Auger	4.3
ST-L3	SB5679	Soil	3-16-94/1045	2.0 - 2.5	As	Split Spoon	26.5
ST-U4	SB5680	Soil	3-16-94/1120	0.5 - 1.0	As	Split Spoon	2.8
ST-L4	BMN18 MBNQ18	Soil	3-16-94/1126	2.5 - 3.0	TCL TAL/CN	Split Spoon	11.5 NSJ
SST-U1	SB5675	Sediment	3-16-94/1205	0.5 - 1.0	As	Split Spoon	8.5
SST-U1A	SB5676	Sediment	3-16-94/1205	0.5 - 1.0	As	Split Spoon	10.3
SST-L1	SB5677	Sediment	3-16-94/1226	1.5 - 2.0	As	Split Spoon	15.0
SU-U1	SB5662	Soil	3-16-94/0800	0.5 - 1.0	As	Hand Auger	12.9
SU-U2	SB5664	Soil	3-16-94/0815	0.5 - 1.0	As	Hand Auger	14.7
SU-L2	SB5665	Soil	3-16-94/0818	2.5 - 3.0	As	Hand Auger	23.7
SU-U3	SB5667	Soil	3-16-94/0820	0.5 - 1.0	As	Hand Auger	6.5
SU-L3	SB5668	Soil	3-16-94/0830	2.0 - 2.5	As	Hand Auger	35.5
SU-U4	SB5669	Soil	3-16-94/0820	0.5 - 1.0	As	Hand Auger	3.6
SU-L4	SB5670	Soil	3-16-94/0830	2.5 - 3.0	As	Hand Auger	3.3
SSU-U1	BMN19 MBNQ19	Sediment	3-16-94/1000	0.4 - 0.8	TCL TAL/CN	Split Spoon	5.2 NJ
SSU-U1A	BMN34 MBNQ34	Sediment	3-16-94/1000	0.4 - 0.8	TCL TAL/CN	Split Spoon	9.8 NSJ
SSU-L1	SB5666	Sediment	3-16-94/1030	1.5 - 2.0	As	Split Spoon	4.2
SV-U1	SB5652	Soil	3-15-94/1420	0.5 - 1.0	As	Hand Auger	22.1 SNJ
SV-L1	BMN 20	Soil	3-15-94/1545	2.5 - 3.0	TCL	Hand Auger	4.1 NSJ

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
	MBNQ 20	Soil		2.5 - 3.0	TAL/CN		
SV-U2	SB5653	Soil	3-15-94/1420	0.5 - 1.0	As	Hand Auger	124.0
SV-L2	SB5661	Soil	3-15-94/1425	1.5 - 2.0	As	Hand Auger	44.9 NSJ
SV-U3	SB5656	Soil	3-15-94/1450	0.5 - 1.0	As	Hand Auger	9.2 NSJ
SV-L3	SB5657	Soil	3-15-94/1625	1.5 - 2.0	As	Hand Auger	15.6 NSJ
SV-U4	SB5658	Soil	3-15-94/1620	0.5 - 1.0	As	Hand Auger	2.5 NSJ
SV-L4	SB5659	Soil	3-15-94/1640	1.5 - 2.0	As	Hand Auger	6.7 NSJ
SV-L4A	SB5660	Soil	3-15-94/1640	1.5 - 2.0	As	Hand Auger	7.5 NSJ
SSV-U1	SB5654	Sediment	3-15-94/1520	0.4 - 0.8	As	Split Spoon	7.9 NSJ
SSV-L1	SB5655	Sediment	3-16-94/1030	1.0 - 1.4	As	Split Spoon	108.0
SW-U1	SB5641	Soil	3-15-94/1025	0.5 - 1.0	As	Hand Auger	127.0*
SW-L1	SB5642	Soil	3-15-94/1035	2.5 - 3.0	As	Hand Auger	2.3 BWNJ
SW-U2	SB5643	Soil	3-15-94/1030	0.5 - 1.0	As	Hand Auger	61.3*
SW-L2	BMN 21	Soil	3-15-94/1036	2.5 - 3.0	TCL	Hand Auger	18.0 NSJ
	MBNQ 21	Soil		2.5 - 3.0	TAL/CN		
SW-L2A	BMN 33	Soil	3-15-94/1036	2.5 - 3.0	TCL	Hand Auger	16.6 NSJ
	MBNQ 33	Soil		2.5 - 3.0	TAL/CN		
SW-U3	SB5646	Soil	3-15-94/1153	0.5 - 1.0	As	Hand Auger	20.5 SNJ
SW-L3	SB5647	Soil	3-15-94/1158	2.0 - 2.5	As	Hand Auger	76.1*
SW-U4	SB5648	Soil	3-15-94/1211	0.5 - 1.0	As	Hand Auger	77.9*
SW-L4	SB5649	Soil	3-15-94/1218	2.0 - 2.5	As	Hand Auger	131.0*
SSW-U1	SB5644	Sediment	3-15-94/1042	0.5 - 1.0	As	Split Spoon	8.8 SNJ
SSW-L1	SB5645	Sediment	3-15-94/1055	1.5 - 2.0	As	Split Spoon	15.7 SNJ
SX-U1	SB5630	Soil	3-15-94/0755	0.5 - 1.0	As	Hand Auger	22.7 SNJ
SX-L1	SB5631	Soil	3-15-94/0810	2.5 - 3.0	As	Hand Auger	4.7 SNJ
SX-U2	SB5632	Soil	3-15-94/0758	0.5 - 1.0	As	Hand Auger	48.9 SNJ
SX-L2	SB5633	Soil	3-15-94/0815	2.5 - 3.0	As	Hand Auger	16.6 SNJ
SX-U3	SB5636	Soil	3-15-94/0815	0.5 - 1.0	As	Hand Auger	355*
SX-L3	BMN 22	Soil	3-15-94/0830	2.5 - 3.0	TCL	Hand Auger	10.1 NSJ
	MBNQ 22	Soil		2.5 - 3.0	TAL/CN		

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SX-U4	SB5637	Soil	3-15-94/0908	0.5 - 1.0	As	Hand Auger	76.6*
SX-L4	SB5638	Soil	3-15-94/0922	2.2 - 2.6	As	Hand Auger	13.6 SNJ
SX-L4A	SB5639	Soil	3-15-94/0922	2.2 - 2.6	As	Hand Auger	6.9 SNJ
SSX-U1	SB5634	Sediment	3-15-94/0915	0.5 - 1.0	As	Split Spoon	4.2 SNJ
SSX-L1	SB5635	Sediment	3-15-94/0920	1.5 - 2.0	As	Split Spoon	4.2 SNJ
SY-U1	SB5621	Soil	3-14-94/1346	0.5 - 1.0	As	Hand Auger	2.7 SN* (R)
SY-L1	SB5622	Soil	3-14-94/1355	2.5 - 3.0	As	Hand Auger	2.8 SN* (R)
SY-U2	SB5623	Soil	3-14-94/1410	0.5 - 1.0	As	Hand Auger	595.0 *J
SY-L2	SB5624	Soil	3-14-94/1415	2.5 - 3.0	As	Hand Auger	6.2 SN* (R)
SY-U3	SB5627	Soil	3-14-94/1510	0.5 - 1.0	As	Hand Auger	203.0 *J
SY-L3	SB5628	Soil	3-14-94/1540	2.5 - 3.0	As	Hand Auger	1.2 BWN* (R)
SY-U4	SB5629	Soil	3-14-94/1450	0.5 - 1.0	As	Hand Auger	31.7 SNJ
SY-L4	BMN23 MBNQ23	Soil	3-14-94/1525	2.0 - 2.5	TCL TAL/CN	Hand Auger	6.9 NSJ
SSY-U1	SB5625	Sediment	3-14-94/1505	0.1 - 0.8	As	Split Spoon	8.0 SN* (R)
SSY-L1	SB5626	Sediment	3-14-94/1520	1.5 - 2.0	As	Split Spoon	11.2 SN* (R)
SZ-U1	SB5610	Soil	3-14-94/1100	0.5 - 1.0	As	Hand Auger	23.6 SN* (R)
SZ-L1	SB5611	Soil	3-14-94/1105	2.5 - 3.0	As	Hand Auger	1.3 BWN* (R)
SZ-U2	SB5612	Soil	3-14-94/1122	0.5 - 1.0	As	Hand Auger	267.0 *J
SZ-L2	SB5613	Soil	3-14-94/1135	1.2 - 1.5	As	Hand Auger	8.9 SN* (R)
SZ-U3	SB5615	Soil	3-14-94/1120	0.5 - 1.0	As	Split Spoon	152.0 *J
SZ-L3	SB5616	Soil	3-14-94/1130	2.5 - 3.0	As	Split Spoon	1.9 BWN* (R)
SZ-U4	SB5617	Soil	3-14-94/1204	0.5 - 1.0	As	Split Spoon	106.0 *J
SZ-L4	SB5618	Soil	3-14-94/1211	2.5 - 3.3	As	Split Spoon	0.85 BN* (R)
SZ-L4A	SB5619	Soil	3-14-94/1211	2.5 - 3.3	As	Split Spoon	4.4 SN* (R)
SSZ-U1	BMN24 MBNQ24	Sediment	3-14-94/1430	0.3 - 0.8	TCL TAL/CN	Split Spoon	4.1 NJ
SSZ-L1	SB5614	Soil	3-14-94/1440	1.5 - 2.0	As	Split Spoon	6.5 SN* (R)
SAA-U1	SB5851	Soil	3-23-94/0820	0.5 - 1.0	As	Hand Auger	37.0
SAA-L1	SB5852	Soil	3-23-94/0840	2.5 - 3.0	As	Hand Auger	6.1

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Fl. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SAA-U2	SB5853	Soil	3-23-94/0842	0.5 - 1.0	As	Hand Auger	177.0
SAA-L2	BMN25 MBNQ25	Soil	3-23-94/0842	2.0 - 2.5	TCL TAL/CN	Hand Auger	129.0
SAA-U3	SB5856	Soil	3-23-94/0920	0.5 - 1.0	As	Hand Auger	36.8
SAA-U4	SB5858	Soil	3-23-94/0925	0.5 - 1.0	As	Hand Auger	3.6
SSAA-U1	SB5854	Soil	3-23-94/0830	0.5 - 1.0	As	Split Spoon	2.0B
SSAA-L1	SB5855	Sediment	3-23-94/0840	1.5 - 2.0	As	Hand Auger	3.0
SBB-U1	SB5892	Soil	3-23-94/0945	0.5 - 1.0	As	Hand Auger	4.5
SBB-L1	SB5893	Soil	3-23-94/0950	2.5 - 3.0	As	Hand Auger	4.4
SBB-U2	SB5894	Soil	3-23-94/1000	0.5 - 1.0	As	Hand Auger	34.1
SBB-U3	SB5889	Soil	3-23-94/1030	0.5 - 1.0	As	Hand Auger	7.8
SBB-L3	BMN26 MBNQ26	Soil	3-23-94/1100	2.5 - 3.0	TCL TAL/CN	Hand Auger	2.9
SBB-U4	SB5888	Soil	3-23-94/1100	0.5 - 1.0	As	Hand Auger	4.9
SSBB-U1	SB5890	Sediment	3-23-94/1020	0.5 - 1.0	As	Split Spoon	2.7
SSBB-L1	SB5891	Sediment	3-23-94/1035	1.5 - 2.0	As	Split Spoon	3.7
SCC-U1	SB5886	Soil	3-22-94/1435	0.5 - 1.0	As	Hand Auger	10.0 SNJ
SCC-L1	SB5885	Soil	3-22-94/1450	2.5 - 3.0	As	Hand Auger	11.1 SNJ
SCC-U2	SB5884	Soil	3-22-94/1511	0.5 - 1.0	As	Hand Auger	6.3 SNJ
SCC-L2	SB5883	Soil	3-22-94/1520	2.5 - 3.0	As	Hand Auger	1.5 BWNJ
SCC-U3	SB5880	Soil	3-22-94/1445	0.5 - 1.0	As	Hand Auger	11.0 SNJ
SCC-L3	SB5879	Soil	3-22-94/1505	2.5 - 3.0	As	Hand Auger	16.8 SNJ
SCC-U4	SB5878	Sediment	3-22-94/1515	0.5 - 1.0	As	Split Spoon	6.6 +N
SCC-L4	BMN27 MBNQ27	Soil	3-22-94/1530	2.0 - 2.5	TCL TAL/CN	Hand Auger	4.5 J
SSCC-U1	SB5882	Sediment	3-22-94/1456	0.5 - 1.0	As	Split Spoon	15.7 SNJ
SSCC-L1	SB5881	Sediment	3-22-94/1509	1.0 - 1.4	As	Split Spoon	10.3 SNJ
SDD-U1	SB5877	Soil	3-22-94/1335	0.5 - 1.0	As	Hand Auger	10.5 SNJ
SDD-L1	SB5876	Soil	3-22-94/1340	2.5 - 3.0	As	Hand Auger	14.4 SNJ
SDD-U2	SB5875	Soil	3-22-94/1340	0.5 - 1.0	As	Hand Auger	28.3 SNJ

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SDD-U3	SB5872	Soil	3-22-94/1410	0.5 - 1.0	As	Hand Auger	21.2 SNJ
SDD-U4	SB5859	Soil	3-22-94/1420	2.0 - 2.5	As	Hand Auger	4.1 SNJ
SDD-L4	SB5860	Soil	3-22-94/1405	0.5 - 1.0	As	Hand Auger	8.7 SNJ
SSDD-U1	BMN28 MBNQ28	Sediment	3-22-94/1400	0.5 - 1.0	TCL TAL/CN	Split Spoon	5.6 SJ
SSDD-U1A	BMN39 MBNQ39	Sediment	3-22-94/1400	0.5 - 1.0	TCL TAL/CN	Split Spoon	13.3 SJ
SEE-U1	SB5861	Soil	3-22-94/1235	0.5 - 1.0	As	Hand Auger	10.5 SNJ
SEE-U1A	SB5865	Soil	3-22-94/1235	0.5 - 1.0	As	Hand Auger	7.5 SNJ
SEE-U2	SB5863	Soil	3-22-94/1230	0.5 - 1.0	As	Hand Auger	14.2 SNJ
SEE-L2	SB5864	Soil	3-22-94/1235	2.5 - 3.0	As	Hand Auger	26.6 SNJ
SEE-U3	SB5867	Soil	3-22-94/1245	0.5 - 1.0	As	Hand Auger	25.9 SNJ
SEE-L3	SB5868	Soil	3-22-94/1300	2.5 - 3.0	As	Hand Auger	7.7 SNJ
SEE-U4	SB5869	Soil	3-22-94/1310	0.5 - 1.0	As	Hand Auger	2.9 NJ
SEE-L4	SB5870	Soil	3-22-94/1315	2.5 - 3.0	As	Hand Auger	1.8 BNJ
SRC-U1	BMN32 MBNQ32	Soil	3-23-94/1000	0.2 - 0.7	TCL TAL/CN	Trowel	3.6
SRC-L1	SB5896	Soil	3-23-94/1005	2.5 - 3.0	As	Hand Auger	1.6 B
SRC-U2	SB5897	Soil	3-23-94/1105	0.0 - 0.5	As	Trowel	4.6
SRC-U2A	SB5898	Soil	3-23-94/1105	0.0 - 0.5	As	Trowel	5.1
SRC-U3	SB5899	Soil	3-23-94/1100	0.0 - 0.5	As	Trowel	10.6
SRC-U4	SB5900	Soil	3-23-94/0936	0.0 - 0.5	As	Trowel	2.1B
SRC-U5	SB5901	Soil	3-23-94/1112	0.0 - 0.5	As	Trowel	4.6
SRC-U5A	SB5902	Soil	3-23-94/1112	0.0 - 0.5	As	Trowel	5.7
SRC-U6	SB5903	Soil	3-23-94/1055	0.0 - 0.5	As	Trowel	2.0 B
SRC-U7	SB5004	Soil	3-23-94/1045	0.0 - 0.5	As	Trowel	3.7
SRC-U7A	SB5005	Soil	3-23-94/1045	0.0 - 0.5	As	Trowel	5.6
SRC-8	SB5006	Soil	3-23-94/0917	0.0 - 0.5	As	Trowel	4.7
SRC-9	SB5007	Soil	3-23-94/0910	0.0 - 0.5	As	Trowel	2.1 B
SRC-10	SB5008	Soil	3-23-94/0926	0.0 - 0.5	As	Trowel	3.3

TABLE 3-6 CONTINUED
ARSENIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (Ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
SRC-10A	SB5009	Soil	3-23-94/0926	0.0 - 0.5	As	Trowel	3.3
SEW-U1	SB5010	Soil	3-24-94/0934	0.2 - 0.6	As	Scoop	5.9 S*
SEW-U1A	SB5011	Soil	3-24-94/0935	0.2 - 0.6	As	Scoop	7.9 S*
SEW-U2	SB5012	Soil	3-24-94/0950	0.2 - 0.7	As	Scoop	5.0 S*
SEW-U3	SB5013	Soil	3-24-94/1009	0.3 - 0.6	As	Scoop	6.4 S*
SEW-U4	BMN31 MBNQ31	Soil	3-24-94/1000	0.2 - 0.7	TCL TAL/CN	Scoop	3.5 NJ
SEW-U5	SB5014	Soil	3-24-94/1025	0.2 - 0.6	As	Scoop	3.4 S*
SEW-U6	SB5015	Soil	3-24-94/1020	0.2 - 0.7	As	Scoop	5.2 S*
SEW-U6A	SB5016	Soil	3-24-94/1020	0.2 - 0.7	As	Scoop	5.6 S*
SEG-U1	SB5017	Soil	3-24-94/1107	0.0 - 0.5	As	Scoop	259.0 S*
SEG-U2	SB5018	Soil	3-24-94/1110	0.1 - 0.5	As	Scoop	4.3 *
SEG-U3	SB5019	Soil	3-24-94/1130	0.2 - 0.7	As	Scoop	4.5 *
SEG-U3A	SB5020	Soil	3-24-94/1135	0.2 - 0.7	As	Scoop	6.0 S*
SEG-U4	BMN29 MBNQ29	Soil	3-24-94/1113	0.0 - 0.5	TCL TAL/CN	Scoop	6.1 NJ
SEG-U5	SB5021	Soil	3-24-94/1119	0.1 - 0.6	As	Scoop	3.9 S*
SEG-U6	SB5022	Soil	3-24-94/1120	0.0 - 0.5	As	Scoop	3.2 S*
SEG-U7	SB5023	Soil	3-24-94/1125	0.0 - 0.5	As	Scoop	4.3 S*
SEG-U8	SB5024	Soil	3-24-94/1130	0.0 - 0.5	As	Scoop	4.3 S*
SEG-U9	SB5025	Soil	3-24-94/1140	0.0 - 0.5	As	Scoop	5.2 S*
SEG-U10	SB5026	Soil	3-24-94/1140	0.0 - 0.5	As	Scoop	5.1 S*
SEG-U11	BMN30 MBNQ30	Soil	3-24-94/1207	0.3 - 0.8	TCL TAL/CN	Scoop	3.8 NJ
SEG-U12	SB5027	Soil	3-24-94/1148	0.0 - 0.5	As	Scoop	6.3 S*
SEG-U13	SB5028	Soil	3-24-94/1145	0.2 - 0.7	As	Scoop	3.5 +*J
SEG-U14	SB5029	Soil	3-24-94/1200	0.2 - 0.7	As	Scoop	5.2 SNJ
SEG-U15	SB5030	Soil	3-24-94/1215	0.3 - 0.8	As	Scoop	4.5 NJ
SSCG-1	SB5031	Sediment	3-23-94/1340	0.5 - 1.0	As	Hand Auger	10.5
SSCG-2	BMN56	Sediment	3-23-94/1405	1.0 - 1.5	TCL	Split Spoon	29.6S

TABLE 3-6 CONTINUED
ARSENIC RESULTS – SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE IDENTIFICATION CODE	CLP SAMPLE CODE	MATRIX	SAMPLE DATE/TIME	SAMPLE DEPTH (ft. Bgs)	ANALYTICAL PARAMETERS	SAMPLE METHOD	ARSENIC CONCENTRATION (mg/kg)
	MBNQ56				TAL/CN		
SSCG-3	BMN57 MBNQ57	Sediment	3-23-94/1430	2.0 – 3.0	TCL TAL/CN	Split Spoon	8.0S
SSCG-4	SB5032	Sediment	3-23-94/1640	0.0 – 0.5	As	Hand Auger	2.1 B
SSCG-6	BMN58 MBNQ58	Sediment	3-23-94/1550	0.0 – 1.5	TCL TAL/CN	Split Spoon	4.3
SSCG-7	BMN59 MBNQ59	Sediment	3-23-94/1612	1.5 – 2.0	TCL TAL/CN	Split Spoon	1.7B
SSCG-8	BMN60 MBNQ60	Sediment	3-23-94/1455	0.0 – 0.5	TCL TAL/CN	Split Spoon	4.2
SSCG-9	SB5034	Sediment	3-23-94/1455	0.0 – 0.5	As	Hand Auger	9.0
SSCG-10	SB5035	Sediment	3-23-94/1455	0.0 – 0.5	As	Hand Auger	7.6

B – If the reported value was obtained from a reading that was less than the Contract Required Detection Limit (CRDL) but greater than or equal to the Instrument Detection Limit (IDL).

N – Spiked sample recovery not within control limits.

S – The reported value was determined by the Method of Standard Additions (MSA).

W – Post-digestion spike for Furnace AA analysis is out of control limits (85 – 115%), while sample absorbance is less than 50% of spike absorbance.

(R) – Unusable results.

* – Duplicate analysis not within control limits.

+ – Correlation coefficient for the MSA is less than 0.995.

Sample Identification Code Prefix – SEG denotes Edison Glenn

Sample Identification Code Prefix – SEW denotes Edison Wood

Sample Identification Code Prefix – SRC denotes Rodak Circle

TABLE 3-7
VOLATILE ORGANIC RESULTS - SOILS/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

	1	2	3	4	5	6	7	8	9	10	11	12	13
SAMPLE NUMBER	8A-U1	8BA-U1	8B-U2	8C-U3	8CC-U4	8DD-U1	8EQ-U4	8EQ-U11	8EW-U4	8SD-U1A	8B-U9	8B-U10	8E-U12
SAMPLE LOCATION NO.	TRANSECT A	TRANSECT A	TRANSECT B	TRANSECT C	TRANSECT CC	TRANSECT DD	EDISON GLEN	EDISON GLEN	EDISON WOODS	TRANSECT DD	BMN4	BMN50	TRANSECT E
CLP SAMPLE CODE	BMN01	BMN02	BMN03	BMN04	BMN07	BMN28	BMN29	BMN30	BMN31	BMN38	BMN4	BMN50	BMN05
DEPTH INTERVAL	0.5 TO 1.0 ft	0.5 TO 0.8 ft	0.5 TO 1.0 ft	1.0 TO 1.5 ft	2.0 TO 2.5 ft	0.5 TO 1.0 ft	0.0 TO 0.5 ft	0.3 TO 0.8 ft	0.2 TO 0.7 ft	0.5 TO 1.0 ft	FIELD BLANK	FIELD BLANK	2.5 TO 3.0 ft
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	ug/kg
DATE SAMPLE COLLECTED	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94
Chloromethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Bromomethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Vinyl Chloride	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Chloroethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Methylene Chloride	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Acetone	20 U	44 U	12 U	12 U	24 U	63 U	13 U	13 U	12 U	30 U	50 B	44 B	20 U
Carbon Disulfide	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,1-Dichloroethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,1-Dichloroethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,2-Dichloroethane (total)	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Chloroform	12 U	13 U	12 U	0.7 J	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,2-Dichloroethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
2-Butanone	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,1,1-Trichloroethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Carbon Tetrachloride	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Bromochloromethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,2-Dichloropropane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
cis-1,3-Dichloropropane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Trichloroethane	12 U	4 J	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Dibromochloromethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,1,2-Trichloroethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Benzene	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
trans-1,3-Dichloropropane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Bromodichloromethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
4-Methyl-2-Pentanone	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
2-Hexanone	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Tetrachloroethane	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
1,1,2,2-Tetrachloroethane	12 U	13 U	2 J	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Toluene	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	1 J	3 J
Chlorobenzene	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Ethylbenzene	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Styrene	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Xylene (total)	12 U	13 U	12 U	12 U	13 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U
Total VOCs		4 J	2 J	0.7 J		2 J					32 BJ	95 BJ	3 J
TKC Count		8	1	2								1	
Total TKC Concentration		51 JN	45 JN	96 JN								10 JN	

U = Analyte was not detected at the instrument detection limit given
J = Estimated Value
B = Analyte was detected in blank
0 = Estimated value due to matrix interference
0 = Determined after sample dilution
UJ = Reporting the value for presence of analyte; estimated quantity
R = Reported during data collection

TABLE 1-1 CONTINUED
VOLATILE ORGANIC RESULTS - SOILS/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

	14	15	16	17	18	19	20	21	22	23	24	25	26
SAMPLE NUMBER	60-L2	60-U1	68A-L2	68B-L2	68C-U1	68H-U1	68H-U1A	68H-U1	68H-U1	68H-U1	68H-U1	68H-U1	68H-U1
SAMPLE LOCATION NO.	TRANSECT O	TRANSECT O	TRANSECT AA	TRANSECT BB	NODAK CIRCLE	TRANSECT H	TRANSECT H	TRANSECT H	TRANSECT H	TRANSECT H	TRANSECT H	TRANSECT H	TRANSECT H
CLP SAMPLE CODE	BMN07	BMN12	BMN25	BMN26	BMN32	BMN35	BMN36	BMN37	BMN47	BMN49	BMN54	BMN57	BMN58
DEPTH INTERVAL	2.0 TO 2.5 ft.	0.3 TO 1.2 ft.	2.5 TO 3.0 ft.	2.5 TO 3.0 ft.	0.2 TO 0.7 ft.	0.5 TO 1.0 ft.	0.5 TO 1.0 ft.	0.5 TO 1.0 ft.	FIELD BLANK	FIELD BLANK	1.0 TO 1.5 ft.	2.0 TO 3.0 ft.	0.0 TO 1.5 ft.
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94
Chloromethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Bromomethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Vinyl Chloride	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Chloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Methylene Chloride	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Acetone	62 U	67 U	10 U	17 U	17 U	12 U	12 U	12 U	36 U	15 U	74 U	16 U	20 U
Carbon Dioxide	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,1-Dichloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,1-Dichloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,2-Dichloroethane (total)	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Chloroform	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,2-Dichloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
2-Butanone	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,1,1-Trichloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Carbon Tetrachloride	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Bromochloromethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,2-Dichloropropane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
cis-1,3-Dichloropropane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Trichloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Dibromochloromethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,1,2-Trichloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Benzene	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
trans-1,3-Dichloropropane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Bromodimethylsilane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
4-Methyl-2-Pentanone	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
2-Hexanone	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Tetrachloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
1,1,2,2-Tetrachloroethane	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Toluene	2 J	11 U	10 U	12 U	12 U	1 J	12 U	2 J	10 U	10 U	12 U	12 U	11 U
Chlorobenzene	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Ethylbenzene	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Styrene	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Xylene (total)	12 U	11 U	10 U	12 U	12 U	12 U	12 U	12 U	10 U	10 U	12 U	12 U	11 U
Total VOCs	2 J					1 J		2 J	40 B	22 BJ	12 J		
TIC Count	1				1								
Total TIC Concentration	7 J				15 JN				22 JN				

U = Analyte was not detected at the instrument level

J = Estimated value

B = Analyte was detected in blank

B = Estimated value due to matrix interference

B = Estimated value after sample dilution

BJ = Presumptive evidence for presence of analyte, as

R = Reported during data validation

TABLE 3-7 CONTINUED
VOLATILE ORGANIC RESULTS - SOIL SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

	27	28	29	30	31	32	33	34	35	36	37	38	39
SAMPLE NUMBER	SSOG-7	SSOG-8	ESK-U1	SI-L1	FB-06	SO-L4	SO-L1	SH-L2	ST-L4	SSU-U1	SV-L1	SW-L2	SX-L3
SAMPLE LOCATION NO.	BMN10	BMN09	TRANSECT K	TRANSECT I	BMN13	TRANSECT O	TRANSECT Q	TRANSECT R	TRANSECT T	TRANSECT U	TRANSECT V	TRANSECT W	TRANSECT X
CLP SAMPLE CODE	BMN10	BMN09	BMN09	BMN13	BMN13	BMN14	BMN15	BMN16	BMN16	BMN16	BMN20	BMN21	BMN22
DEPTH INTERVAL	1.5 TO 2.0 ft.	0.5 TO 0.5 ft.	0.5 TO 1.2 ft.	0.5 TO 1.0 ft.	FIELD BLANK	2.5 TO 3.0 ft.	2.0 TO 2.5 ft.	2.5 TO 3.0 ft.	2.5 TO 3.0 ft.	0.4 TO 0.6 ft.	2.5 TO 3.0 ft.	2.5 TO 3.0 ft.	2.5 TO 3.0 ft.
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/2/94	3/2/94	3/18/94	3/18/94	3/18/94	3/17/94	3/17/94	3/16/94	3/16/94	3/16/94	3/16/94	3/15/94	3/15/94
Chloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Bromomethane	13 UJ	12 UJ	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Vinyl Chloride	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Chloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 UJ	13 UJ	13 UJ	13 UJ	13 UJ	12 UJ	14 UJ
Methylene Chloride	13 U	12 U	11 U	13 U	7 J	10 J	15	13 U	13 U	13 U	31 U	12 UJ	14 U
Acetone	24 U	24 U	11 UJ	6 J	51 J	10 U	12 UJ	13 UJ	69 UJ	12 UJ	35 UJ	12 UJ	14 UJ
Carbon Disulfide	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	1 J	13 U	13 U	12 UJ	14 U
1,1-Dichloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
1,1-Dichloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
1,2-Dichloroethane (total)	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Chloroform	13 U	12 U	11 U	13 U	10 U	11 UJ	12 UJ	13 U	13 U	13 U	13 U	12 UJ	14 U
1,2-Dichloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
2-Substane	13 U	12 U	11 U	13 U	10 UJ	11 U	12 UJ	13 U	13 U	13 U	13 UJ	12 UJ	14 U
1,1,1-Trichloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Carbon Tetrachloride	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 UJ
Bromochloromethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 UJ	13 UJ	13 UJ	13 U	12 UJ	14 U
1,2-Dichloropropane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
cis-1,3-Dichloropropane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Trichloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Dibromochloromethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 UJ	13 UJ	13 UJ	13 U	12 UJ	14 U
1,1,2-Trichloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Benzene	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
trans-1,3-Dichloropropane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Bromocloroform	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
4-Methyl-2-Pentanone	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
2-Hexanone	13 U	12 U	11 U	13 U	10 U	11 U	12 UJ	13 U	13 U	13 U	13 UJ	12 UJ	14 U
Tetrachloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
1,1,2,2-Tetrachloroethane	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Toluene	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	2 J
Chlorobenzene	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Ethylbenzene	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Styrene	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Xylene (total)	13 U	12 U	11 U	13 U	10 U	11 U	12 U	13 U	13 U	13 U	13 U	12 UJ	14 U
Total VOCs				0 J	56 J	10 J	15		1 J				2 J
TIC Count			1									1	
Total TIC Concentration			12 J									12 JN	

U = Analyte was not detected at the instrument level
J = Validated Value
B = Analyte was detected in blank
B = Estimated value due to matrix interference
B = Determined after sample dilution
UJ = Presumptive evidence for presence of analyte, as
R = Reported during data validation

TAB. B3-7 CONTINUED
VOLATILE ORGANIC RESULTS - SOIL/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

	40	41	42	43	44	45	46	47	48	49	50	51
SAMPLE NUMBER	SV-L4	SO2-L1	SW-L2A	SSU-L1A	FB-01	FB-02	FB-03	FB-04	FB-05	SJ-L2	SM-L1	SL-U3
SAMPLE LOCATION NO.	TRANSECT Y	TRANSECT Z	TRANSECT W	TRANSECT U						TRANSECT J	TRANSECT M	TRANSECT L
CLP SAMPLE CODE	BMN23	BMN24	BMN33	BMN34	BMN41	BMN42	BMN43	BMN44	BMN45	BMN46	BMN48	BMN11
DEPTH INTERVAL	2.0 TO 2.5 ft	0.5 TO 0.8 ft	2.5 TO 3.0 ft	0.4 TO 0.8 ft	FIELD BLANK	FIELD BLANK	FIELD BLANK	FIELD BLANK	FIELD BLANK	2.0 TO 2.5 ft	2.5 TO 3.0 ft	0.5 TO 1.0 ft
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	ug/l	ug/l	ug/l	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94	3/1/94
Chloromethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Bromomethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Vinyl Chloride	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Chloroethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Methylene Chloride	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Acetone	20 U	13 U	12 U	37 U	4 J	54 J	63 U	1000 J	100 J	16 U	10 U	530 U
Carbon Disulfide	10 U	20	12 U	12 U	10 U	2 J	10 U	100 U	10 U	12 U	10 U	30 U
1,1-Dichloroethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
1,1-Dichloroethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
1,2-Dichloroethane (total)	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Chloroform	2 J	13 U	12 U	1 J	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
1,2-Dichloroethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
2-Butanone	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
1,1,1-Trichloroethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Carbon Tetrachloride	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Bromodichloromethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
1,2-Dichloropropane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
cis-1,3-Dichloropropane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Trichloroethene	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Dibromochloromethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
1,1,2-Trichloroethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Benzene	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
trans-1,2-Dichloropropane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Bromocloromethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
4-Methyl-2-Pentanone	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
2-Hexanone	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Tetrachloroethene	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
1,1,2,2-Tetrachloroethane	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Toluene	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Chlorobenzene	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Ethylbenzene	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Styrene	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Xylene (total)	10 U	13 U	12 U	12 U	10 U	10 U	10 U	100 U	10 U	12 U	10 U	30 U
Total VOCs	30 J	30 J		1 J	4 J	50 J	5 J	1000 J	100 J	12 U	10 U	30 U
TIC Count												
Total TIC Concentration												

U = Analyte was not detected at the instrument detect
J = Estimated Value
B = Analyte was detected by blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
N = Presumptive evidence for presence of analyte, so
R = Reported during data validation

TABLE 3-1
SEMI-VOLATILE ORGANIC RESULTS - SOIL SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	SA-U1	SA-U1	SB-U2	SC-L3	SC-L4	SSDD-U1	SEQ-U4	SEQ-U11	SEW-U4	SSDD-U11A	FB-08	FB-10	SE-12	SO-12
SAMPLE LOCATION NO.	TRANSECT A	TRANSECT A	TRANSECT B	TRANSECT C	TRANSECT C	TRANSECT D	EDISON GLEN	EDISON GLEN	EDISON WOODS	TRANSECT D			TRANSECT E	TRANSECT D
CLP SAMPLE CODE	BMN01	BMN02	BMN03	BMN04	BMN07	BMN08	BMN09	BMN10	BMN11	BMN13	BMN14	BMN15	BMN05	BMN07
DEPTH INTERVAL	0.5 TO 1.0 ft	1.5 TO 2.0 ft	0.5 TO 1.0 ft	1.0 TO 1.5 ft	2.0 TO 2.5 ft	0.5 TO 1.0 ft	0.5 TO 0.5 ft	0.5 TO 0.5 ft	0.2 TO 0.7 ft	0.5 TO 1.0 ft	FIELD BLANK	FIELD BLANK	2.5 TO 3.0 ft	2.0 TO 2.5 ft
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/22/94	3/22/94	3/22/94	3/22/94	3/22/94	3/22/94	3/24/94	3/24/94	3/24/94	3/24/94	3/22/94	3/24/94	3/24/94	3/24/94
Phenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
1,2-Dichloroethane	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2-Chlorophenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
1,3-Dichlorobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
1,4-Dichlorobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
1,2-Dichlorobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2-Methylphenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2,2'-methylenebis(1-Chloropropane)	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
4-Methylphenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
N-Nitroso-d-n-propylamine	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Hexachlorobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Nitrobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Isophorone	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2-Nitrophenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2,4-Dinitrophenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
1,2-Dichloroethane	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
1,4-Dichlorobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
1,2,4-Trichlorobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Naphthalene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
4-Chloroaniline	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Hexachlorocyclopentadiene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
4-Chloro-3-methylphenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2-Methylnaphthalene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Hexachlorocyclopentadiene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2,4,6-Trichlorophenol	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2,4,5-Trichlorophenol	890 U	2100 U	890 U	850 U	1000 U	930 U	940 U	930 U	940 U	860 U	25 U	25 U	1000 U	1000 U
2-Chloronaphthalene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2-Nitroaniline	890 U	2100 U	890 U	850 U	1000 U	930 U	940 U	930 U	940 U	860 U	25 U	25 U	1000 U	1000 U
Dimethylnaphthalene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Acenaphthylene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2,6-Dinitroaniline	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
3-Nitroaniline	890 U	2100 U	890 U	850 U	1000 U	930 U	940 U	930 U	940 U	860 U	25 U	25 U	1000 U	1000 U
Acenaphthene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2,4-Dinitrophenol	890 U	2100 U	890 U	850 U	1000 U	930 U	940 U	930 U	940 U	860 U	25 U	25 U	1000 U	1000 U
4-Nitrophenol	890 U	2100 U	890 U	850 U	1000 U	930 U	940 U	930 U	940 U	860 U	25 U	25 U	1000 U	1000 U
Dibenzofuran	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
2,4-Dinitroaniline	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Dimethylnaphthalene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
4-Chlorophenyl-phenylphosphonate	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U
Phenanthrene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	380 U	380 U	400 U	10 U	10 U	410 U	410 U

TABLE 3-4 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - SOIL SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	EA-U1	ESA-U1	EB-U2	EC-L3	ECC-L4	ESDD-U1	EEG-U11	EDG-U11	EDW-U1	ESDD-U1A	FB-08	FB-10	EE-L2	EO-L2
SAMPLE LOCATION NO.	TRANSECT A	TRANSECT A	TRANSECT B	TRANSECT C	TRANSECT C	TRANSECT D	EDISON GLEN	EDISON GLEN	EDISON WOODS	TRANSECT D			TRANSECT E	TRANSECT G
CLP SAMPLE CODE	BMN01	BMN02	BMN03	BMN04	BMN27	BMN28	BMN29	BMN30	BMN31	BMN38			BMN03	BMN07
DEPTH INTERVAL	0.5 TO 1.0 ft	1.5 TO 2.0 ft	0.5 TO 1.0 ft	1.0 TO 1.5 ft	2.0 TO 2.5 ft	0.5 TO 1.0 ft	0.0 TO 0.5 ft	0.3 TO 0.8 ft	0.2 TO 0.7 ft	0.5 TO 1.0 ft	FIELD BLANK	FIELD BLANK	2.5 TO 3.0 ft	2.0 TO 2.5 ft
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/22/94	3/22/94	3/22/94	3/22/94	3/22/94	3/22/94	3/24/94	3/24/94	3/24/94	3/22/94	3/22/94	3/22/94	3/22/94	3/22/94
4-Nitroanisole	950 U	2100 U	980 U	850 U	1000 U	830 U	940 U	930 U	940 U	960 U	25 U	25 U	1000 UJ	1000 UJ
4,6-Dinitro-2-methylphenol	890 U	2100 U	980 U	850 U	1000 U	830 U	940 U	930 U	940 U	960 U	25 U	25 U	1000 U	1000 U
N-Nitrosodiphenylamine (I)	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 U	410 U
4-Bromophenyl-phenylether	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 UJ	410 UJ
Heuchlorobenzene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 UJ	410 UJ
Pentachlorophenol	990 U	2100 U	980 U	850 U	1000 U	830 U	940 U	930 U	940 U	960 U	25 U	25 U	1000 U	1000 U
Phenanthrene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 U	410 U
Anthracene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 U	410 U
Carbazole	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 U	410 U
Di-n-butylphthalate	28 J	850 U	28 J	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 U	410 U
Fluoranthene	43 J	2400	1000	61 J	100 J	300 J	88 J	140 J	48 J	440	10 U	10 U	120 J	410 U
Pyrene	53 J	1800	1200	66 J	120 J	380 J	84 J	110 J	42 J	850	10 U	10 U	130 J	410 U
Butylbenzylphthalate	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 UJ	410 UJ
3,3'-Dichlorobenzidine	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 U	410 U
Benzo(a)anthracene	37 J	890	720	31 J	50 J	120 J	38 J	73 J	25 J	240 J	10 U	10 U	59 J	410 U
Chrysene	38 J	1000	720	41 J	82 J	150 J	59 J	72 J	31 J	270 J	10 U	10 U	81 J	410 U
ba(2-Ethylhexyl)phthalate	410 U	2300	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	2 BJ	3 BJ	130 J	64 J
Di-n-octylphthalate	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 UJ	410 UJ
Benzo(b)fluoranthene	53 J	1200	890	50 J	120 J	380 J	88 J	140 J	48 J	440	10 U	10 U	89 J	410 U
Benzo(k)fluoranthene	410 U	850 U	400 U	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	72 J	410 U
Benzo(a)pyrene	28 J	770 J	550	24 J	46 J	100 J	40 J	74 J	25 J	210 J	10 U	10 U	66 J	410 U
Indeno(1,2,3-cd)pyrene	24 J	720 J	440	380 U	46 J	87 J	37 J	61 J	390 U	140 J	10 U	10 U	410 U	410 U
Dibenz(a,h)anthracene	410 U	180 J	150 J	380 U	420 U	380 U	390 U	390 U	390 U	400 U	10 U	10 U	410 U	410 U
Benzo(g,h,i)perylene	28 J	890 J	440	380 U	43 J	84 J	38 J	61 J	390 U	170 J	10 U	10 U	410 U	410 U
Total SVOCs	310 J	15490 J	7185	306 J	657 J	1803 J	420 J	671 J	191 J	3229 J	2 BJ	3 BJ	809 J	64 J
TIC Count	20	26	17	18	18	18	13	16	16	16	1	1	16	16
Total TIC Concentration	11590 JN	84700 JN	9500 JN	24630 JN	8550 JN	8850 JN	8490 JN	4785 JN	8348 JN	10130 JN	11 JN	3 J	6448 JN	3238 JN

U = Analyte was not detected at the instrument detection limit given
J = Estimated Value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence of analyte; estimated quantity
R = Rejected during data validation
(I) = Cannot be separated from Diphenylamine

TABLE 3-8 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - SOILS/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	SSO-U1	SAA-L2	SBB-L3	SRC-U1	SH-U1	SH-U1A	SH-U1	FB-07	FB-08	SSCO-2	SSCO-3	SSCO-6	SSCO-7	SSCO-8
SAMPLE LOCATION NO.	TRANSECT O	TRANSECT AA	TRANSECT BB	ROAD CIRCLE	TRANSECT H	TRANSECT H	TRANSECT H							
CLP SAMPLE CODE	BMN12	BMN25	BMN26	BMN32	BMN35	BMN36	BMN37	BMN47	BMN48	BMN56	BMN57	BMN58	BMN59	BMN60
DEPTH INTERVAL	0.5 TO 1.2 ft	2.5 TO 3.0 ft	2.5 TO 3.0 ft	0.2 TO 0.7 ft	0.5 TO 1.0 ft	0.5 TO 1.0 ft	0.5 TO 1.0 ft	FIELD BLANK	FIELD BLANK	1.0 TO 1.5 ft	2.0 TO 3.0 ft	0.0 TO 1.5 ft	1.5 TO 2.0 ft	0.0 TO 0.5 ft
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94
Phenol	350 U	330 U	350 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
ba(2-Chlorophenyl)ether	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2-Chlorophenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
1,3-Dichlorobenzene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
1,4-Dichlorobenzene	350 U	330 U	380 U	430 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
1,2-Dichlorobenzene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2-Methylphenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2,2-dimethyl(1-Chloropropyl)ether	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
4-Methylphenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
N-Methyl-2-n-propylamine	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Hexachlorobenzene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Nitrobenzene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Isophorone	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2-Nitrophenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2,4-Dimethylphenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
ba(2-Chlorophenyl)methane	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2,4-Dichlorophenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
1,2,4-Trichlorobenzene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Naphthalene	350 U	60 J	380 U	400 U	380 U	21 J	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
4-Chloroaniline	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Hexachlorobutadiene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
4-Chloro-2-methylphenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2-Methylnaphthalene	350 U	60 J	380 U	400 U	380 U	21 J	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Hexachlorocyclopentadiene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2,4,6-Trichlorophenol	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2,4,6-Trichlorophenol	880 U	1300 U	950 U	880 U	830 U	830 U	830 U	25 U	25 U	830 U	830 U	830 U	1100 U	880 U
2-Chloronaphthalene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	880 U	880 U	880 U	430 U	400 U
2-Nitroaniline	880 U	1300 U	950 U	880 U	830 U	830 U	830 U	25 U	25 U	830 U	830 U	830 U	1100 U	880 U
Dimethylnaphthalene	880 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Acenaphthylene	350 U	180 J	380 U	400 U	380 U	28 J	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2,6-Dichloroaniline	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
3-Nitroaniline	880 U	1300 U	950 U	880 U	830 U	830 U	830 U	25 U	25 U	830 U	830 U	830 U	1100 U	880 U
Acenaphthene	350 U	49 J	380 U	400 U	380 U	880 U	380 U	10 U	10 U	880 U	880 U	880 U	430 U	400 U
2,4-Dichlorophenol	880 U	1300 U	950 U	880 U	830 U	830 U	830 U	25 U	25 U	830 U	830 U	830 U	1100 U	880 U
4-Nitrophenol	880 U	1300 U	950 U	880 U	830 U	830 U	830 U	25 U	25 U	830 U	830 U	830 U	1100 U	880 U
Dibenzofuran	350 U	83 J	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
2,4-Dichloroaniline	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Dimethylnaphthalene	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
4-Chlorophenyl-phenyl ether	350 U	330 U	380 U	400 U	380 U	380 U	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U
Fluorene	350 U	82 J	380 U	400 U	380 U	34 J	380 U	10 U	10 U	380 U	380 U	380 U	430 U	400 U

TABLE 3-9 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - SOILS/SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
POISON, NEW JERSEY

SAMPLE NUMBER	SSO-U1	BAA-L2	BBB-L3	SAC-U1	SH-U1	SH-U1A	SH-U1	FB-07	FB-08	SSCO-2	SSCO-3	SSCO-4	SSCO-7	SSCO-8
SAMPLE LOCATION NO.	TRANSECT D	TRANSECT AA	TRANSECT BB	ROAD CIRCLE	TRANSECT H	TRANSECT H	TRANSECT H							
CLP SAMPLE CODE	BMN12	BMN25	BMN26	BMN32	BMN35	BMN36	BMN37	BMN47	BMN48	BMN50	BMN57	BMN58	BMN59	BMN60
DEPTH INTERVAL	0.3 TO 1.2 H	2.5 TO 3.0 H	2.5 TO 3.0 H	0.2 TO 0.7 H	0.5 TO 1.0 H	0.5 TO 1.0 H	0.5 TO 1.0 H	FIELD BLANK	FIELD BLANK	1.0 TO 1.5 H	2.0 TO 3.0 H	0.0 TO 1.5 H	1.5 TO 2.0 H	0.0 TO 0.5 H
UNITS	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94
4-Nitroanisole	880 U	1300 U	950 U	960 U	930 U	930 U	930 U	25 U	25 U	930 U	930 U	870 U	1100 U	980 U
4,6-Dinitro-2-methylphenol	880 U	1300 U	950 U	960 U	930 U	930 U	930 U	25 U	25 U	930 U	930 U	870 U	1100 U	980 U
N-Nitrosodiphenylamine (1)	950 U	530 U	390 U	400 U	960 U	390 U	390 U	10 U	10 U	390 U	390 U	390 U	430 U	400 U
4-Bromophenyl-phenylether	350 U	530 U	390 U	400 U	390 U	390 U	390 U	10 U	10 U	390 U	390 U	390 U	430 U	400 U
Hexachlorobenzene	950 U	830 U	390 U	400 U	390 U	390 U	390 U	10 U	10 U	390 U	390 U	390 U	430 U	400 U
Pentachlorophenol	880 U	1300 U	950 U	960 U	930 U	930 U	930 U	25 U	25 U	930 U	930 U	870 U	1100 U	980 U
Phenanthrene	52 J	1400	390 U	23 J	180 J	420	110 J	10 U	10 U	73 J	180 J	5800	180 J	400 U
Anthracene	350 U	140 J	390 U	400 U	390 U	390 U	390 U	10 U	10 U	390 U	35 J	850	40 J	400 U
Carbazole	350 U	100 J	390 U	400 U	390 U	27 J	390 U	10 U	10 U	390 U	390 U	590	430 U	400 U
Di-n-butylphthalate	350 U	37 J	390 U	29 J	390 U	390 U	390 U	10 U	10 U	75 J	390 U	390 U	430 U	400 U
Fluorene	88 J	1800	390 U	53 J	290 J	450	140 J	10 U	10 U	140 J	290 J	5500	270 J	400 U
Pyrene	88 J	2400	21 J	60 J	250 J	480	180 J	10 U	10 U	180 J	390 J	4400	280 J	400 U
Dibutylphthalate	350 U	50 J	390 U	400 U	390 U	390 U	390 U	10 U	10 U	390 U	390 U	350 U	430 U	400 U
3,3'-Dichlorobenzidine	350 U	530 U	390 U	400 U	390 U	390 U	390 U	10 U	10 U	390 U	390 U	390 U	430 U	400 U
Benzofluorene	88 J	1500	16 J	31 J	130 J	220 J	80 J	10 U	10 U	88 J	180 J	2600	150 J	400 U
Chrysene	55 J	1800	23 J	42 J	150 J	290 J	91 J	10 U	10 U	100 J	180 J	2600	150 J	400 U
bis(2-Ethylhexyl)phthalate	170 J	320 J	390 U	400 U	110 J	90 J	90 J	1 BJ	1 BJ	390 U	390 U	410	430 U	400 U
Di-n-octylphthalate	350 U	530 U	390 U	400 U	390 U	390 U	390 U	10 U	10 U	390 U	390 U	48 J	430 U	400 U
Benzofluorene	48 J	2000	50 J	72 J	140 J	250 J	78 J	10 U	10 U	180 J	210 J	3500	130 J	400 U
Benzofluorene	58 J	1700	390 U	400 U	130 J	180 J	71 J	10 U	10 U	390 U	390 U	390 U	130 J	400 U
Benzofluorene	54 J	1400	22 J	38 J	130 J	210 J	74 J	10 U	10 U	94 J	140 J	2200	100 J	400 U
Indeno(1,2,3-cd)pyrene	47 J	590	390 U	26 J	92 J	140 J	390 U	10 U	10 U	59 J	70 J	740	160 J	400 U
Dibenz(h)anthracene	350 U	390 J	390 U	400 U	390 U	390 U	390 U	10 U	10 U	390 U	24 J	870 J	80 J	400 U
Benzofluorene	45 J	630	390 U	26 J	91 J	130 J	390 U	10 U	10 U	58 J	68 J	870	200 J	400 U
Total SVOCs	754 J	16570 J	132 J	404 J	1683 J	2950 J	914 J	1 BJ	1 BJ	1024 J	1167 J	32135 J	1930 J	
TIC Count	18	17	9	13	7	13	8			3	11	18	4	12
Total TIC Concentration	284 JN	7750 JN	1450 JN	2977	1789	2541 JN	1308 JN			6 JNB	728 JN	1918 JN	830 JN	3360 JN

U = Analyte was not detected at the test
J = Estimated Value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NI = Presumptive evidence for presence
R = Rejected during data validation
(1) = Cannot be separated from Dithienyl

TABLE 3-3 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - SOIL SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	88K-U1	M-U1		80-L4	80-L1	8R-L2	8T-L4	8SU-U1	8V-L1	8W-L2	8X-L3	8Y-L4	8SZ-L11	8W-L2A
SAMPLE LOCATION NO.	TRANSECT K	TRANSECT I		TRANSECT O	TRANSECT Q	TRANSECT R	TRANSECT T	TRANSECT U	TRANSECT V	TRANSECT W	TRANSECT X	TRANSECT Y	TRANSECT Z	TRANSECT W
CLP SAMPLE CODE	BMN09	BMN13	BMN16	BMN14	BMN15	BMN18	BMN18	BMN18	BMN20	BMN21	BMN22	BMN23	BMN24	BMN33
DEPTH INTERVAL	0.5 TO 1.2 ft	0.5 TO 1.0 ft		2.5 TO 3.0 ft	2.0 TO 2.5 ft	2.5 TO 3.0 ft	2.5 TO 3.0 ft	0.4 TO 0.8 ft	2.5 TO 3.0 ft	2.5 TO 3.0 ft	2.5 TO 3.0 ft	2.0 TO 2.5 ft	0.3 TO 0.8 ft	2.5 TO 3.0 ft
UNITS	ug/kg	ug/kg	ug/l	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/15/94	3/16/94		3/17/94	3/17/94	3/16/94	3/16/94	3/15/94	3/15/94	3/15/94	3/15/94	3/14/94	3/14/94	3/15/94
Phenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
ba(2)-Chlorophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2-Chlorophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
1,3-Dichlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
1,4-Dichlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
1,2-Dichlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2-Methylphenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,2'-oxybis(1-Chloropropene)	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
4-Methylphenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
N-Nerone-d-n-propylmhe	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Hexachlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Hexachlorocyclopentadiene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Hexachlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Hexachlorocyclopentadiene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2-Nitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,4-Dinitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
ba(2)-Chlorophenylmethane	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,4-Dichlorophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
1,2,4-Trichlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Naphthalene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
4-Chlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Hexachlorobutadiene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
4-Chloro-3-methylphenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2-Methylnaphthalene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Hexachlorocyclopentadiene	380 U	420 U	25 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,4,6-Trichlorophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,4,5-Trichlorophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2-Chloronaphthalene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2-Nitroniline	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Dinitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Acanaphthylene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,6-Dinitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2-Nitroniline	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Acanaphthylene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,4-Dinitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
4-Nitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Dibenzofuran	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
2,4-Dinitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Dinitrophenol	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
4-Chlorophenyl-phenyl ether	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U
Fluorene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	470 U	400 U

TABLE 3-3 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - SOILS SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER SAMPLE LOCATION NO. CLP SAMPLE CODE DEPTH INTERVAL UNITS DATE SAMPLE COLLECTED	6SK-U1 TRANSECT K BMN09 0.5 TO 1.2 ft ug/kg 3/18/94	6I-U1 TRANSECT I BMN13 0.5 TO 1.0 ft ug/kg 3/18/94	6M-U1 BMN16 ug/l	6O-L1 TRANSECT O BMN14 2.5 TO 3.0 ft ug/kg 3/17/94	6O-L1 TRANSECT O BMN15 2.0 TO 2.5 ft ug/kg 3/17/94	6R-L2 TRANSECT R BMN16 2.5 TO 3.0 ft ug/kg 3/18/94	6T-L4 TRANSECT T BMN18 2.5 TO 3.0 ft ug/kg 3/18/94	6SU-U1 TRANSECT U BMN18 0.5 TO 0.8 ft ug/kg 3/18/94	6V-L1 TRANSECT V BMN20 2.5 TO 3.0 ft ug/kg 3/15/94	6W-L2 TRANSECT W BMN21 2.5 TO 3.0 ft ug/kg 3/15/94	6X-L3 TRANSECT X BMN22 2.5 TO 3.0 ft ug/kg 3/15/94	6Y-L4 TRANSECT Y BMN23 2.0 TO 2.5 ft ug/kg 3/15/94	6Z-U1 TRANSECT Z BMN24 0.5 TO 0.8 ft ug/kg 3/15/94	6W-L2A TRANSECT W BMN33 2.5 TO 3.0 ft ug/kg 3/15/94
4-Nitroanisole	920 U	1000 U	28 U	810 UJ	990 U	1000 UJ	1000 U	1000 U	1000 U	930 U	1100 UJ	1300 U	1000 U	980 UJ
4,6-Dinitro-2-methylphenol	920 UJ	1000 UJ	28 U	810 U	990 UJ	1000 U	1000 U	1000 U	1000 U	930 U	1100 UJ	1300 U	1000 U	980 UJ
N-Nitrosodiphenylamine (I)	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	420 U	400 U
4-Bromophenyl-phenyl ether	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	420 U	400 U
Hexachlorobenzene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	420 U	400 U
Pentachlorophenol	920 U	1000 U	28 U	810 U	990 U	1000 U	430 U	1000 U	1000 U	930 U	1100 U	1300 U	1000 U	980 U
Phenanthrene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	420 U	400 U
Anthracene	380 U	420 U	10 U	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 U	540 U	420 U	400 U
Carbazole	380 U	420 U	10 UJ	380 U	410 U	420 UJ	24 J	420 U	430 U	380 U	450 U	540 U	420 U	400 U
Di-n-butylphthalate	380 U	420 U	0.5 J	380 U	24 J	420 U	430 U	420 U	23 J	48 J	450 U	540 U	420 U	28 J
Fluorene	74 J	420 U	10 U	200 J	740	420 U	430 U	420 U	85 J	240 J	200 J	44 J	220 J	180 J
Pyrene	74 J	420 U	10 U	190 J	710	420 U	430 U	420 U	82 J	210 J	540 J	42 J	210 J	400 J
Butylbenzylphthalate	380 U	420 U	10 U	24 J	410 U	420 U	430 U	420 U	34 J	380 U	450 UJ	540 U	420 U	400 UJ
2,2'-Dichlorobenzidine	380 U	420 U	10 UJ	380 U	410 U	420 U	430 U	420 U	430 U	380 U	450 UJ	540 U	420 U	400 UJ
Benzofluorenone	380 U	420 U	10 U	110 J	450	420 U	430 U	420 U	45 J	110 J	150 J	840 U	85 J	110 J
Chrysene	380 U	420 U	10 U	110 J	560	420 U	430 U	420 U	62 J	160 J	230 J	540 U	100 J	150 J
bis(2-Ethylhexyl)phthalate	340 U	420 U	10 U	350 U	410 U	420 U	430 U	420 U	8	600 U	450 UJ	540 U	420 U	420 UJ
Di-n-octylphthalate	380 U	420 U	10 U	3 J	410 UJ	34 J	430 U	420 U	430 U	380 U	450 UJ	540 UJ	420 UJ	400 UJ
Benzofluorenone	380 U	420 U	10 U	75 J	830	420 U	430 U	420 U	81 J	100 J	280 J	34 J	420 U	110 J
Benzofluorenone	380 U	420 U	10 U	200 J	480	420 U	430 U	420 U	430 U	130 J	450 UJ	29 J	420 U	140 J
Benzofluorenone	380 U	420 U	10 U	120 J	490	420 U	430 U	420 U	51 J	120 J	200 J	540 U	420 U	130 J
Indeno(1,2,3-cd)pyrene	380 UJ	420 UJ	10 U	140 J	480	420 U	430 U	420 U	38 J	90 J	180 J	540 U	420 U	85 J
Dibenz(h)anthracene	380 U	420 U	10 U	380 U	240 J	420 U	430 U	420 U	430 U	380 U	450 UJ	540 U	420 U	400 UJ
Benzofluorenone	380 UJ	420 UJ	10 U	150 J	600	420 U	430 U	420 U	44 J	100 J	250 J	540 U	420 U	99 J
Total SVOCs	145 J		0.5 J	1411 J	6476 J	34 J	24 J		585 J	1418 J	2220 J	148 J	1335 J	1587 J
TIC Count	10	15		6	12	14	17	15	22	22	18	20	5	18
Total TIC Concentration	1487 JN	3477 JN		9000 JN	16570 JN	17144 JN	8180	17799 JN	16440 JN	4480 JN	7367 JN	26510 J	1500	5826 JN

U = Analyte was not detected at the test
J = Estimated Value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence
R = Rejected during data validation
(I) = Cannot be separated from D(phenyl)

TABLE 3-4 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - BOLL'S SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	SSU-111A	GL-L2	SM-L1	SI-U3
SAMPLE LOCATION NO.	TRANSCT U	TRANSCT J	TRANSCT M	TRANSCT L
CLP SAMPLE CODE	BMN11	BMN08	BMN10	BMN11
DEPTH INTERVAL	0.4 TO 0.8 h	2.0 TO 2.5 h	2.5 TO 3.0 h	0.5 TO 1.0 h
UNITS	ug/kg	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/18/94	3/18/94	3/11/94	3/11/94
Phenol	110 U	10 U	10 U	10 U
Is(2-Chloroethoxy)ethanol	110 U	10 U	10 U	10 U
2-Chlorophenol	110 U	10 U	10 U	10 U
1,3-Dichlorobenzene	110 U	10 U	10 U	10 U
1,4-Dichlorobenzene	110 U	10 U	10 U	10 U
1,2-Dichlorobenzene	110 U	10 U	10 U	10 U
2-Methylphenol	110 U	10 U	10 U	10 U
2,2'-oxybis(1-Chloropropane)	110 U	10 U	10 U	10 U
4-Methylphenol	110 U	10 U	10 U	10 U
N-Nitro-o-d-n-propylene	110 U	10 U	10 U	10 U
Hexachlorocyclopentadiene	110 U	10 U	10 U	10 U
Hexachlorobenzene	110 U	10 U	10 U	10 U
Isophorone	110 U	10 U	10 U	10 U
2-Nitrophenol	110 U	10 U	10 U	10 U
2,4-Dimethylphenol	110 U	10 U	10 U	10 U
Is(2-Chloroethoxy)methane	110 U	10 U	10 U	10 U
2,4-Dichlorophenol	110 U	10 U	10 U	10 U
1,2,4-Trichlorobenzene	110 U	10 U	10 U	10 U
Naphthalene	110 U	10 U	10 U	10 U
4-Chloroaniline	110 U	10 U	10 U	10 U
Hexachlorobutadiene	110 U	10 U	10 U	10 U
4-Chloro-3-methylphenol	110 U	10 U	10 U	10 U
2-Methylnaphthalene	110 U	10 U	10 U	10 U
Hexachlorocyclopentadiene	110 U	10 U	10 U	10 U
2,4,6-Trichlorophenol	110 U	10 U	10 U	10 U
2,4,5-Trichlorophenol	1000 U	25 U	25 U	25 U
2-Chloronaphthalene	110 U	10 U	10 U	10 U
2-Nitroaniline	1000 U	25 U	25 U	25 U
Dimethylphthalate	110 U	10 U	10 U	10 U
Acenaphthylene	110 U	10 U	10 U	10 U
2,6-Dinitrotoluene	110 U	10 U	10 U	10 U
3-Nitroaniline	1000 U	25 U	25 U	25 U
Acenaphthene	110 U	10 U	10 U	10 U
2,4-Dinitrophenol	1000 U	25 U	25 U	25 U
4-Nitrophenol	1000 U	25 U	25 U	25 U
Dibenzofuran	110 U	10 U	10 U	10 U
2,4-Dinitrotoluene	110 U	10 U	10 U	10 U
Diethylphthalate	110 U	10 U	10 U	10 U
4-Chlorophenyl-phenylether	110 U	10 U	10 U	10 U
Fluorene	110 U	10 U	10 U	10 U

TABLE 3-2 CONTINUED
SEMI-VOLATILE ORGANIC RESULTS - SOIL SEDIMENTS
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER SAMPLE LOCATION NO. CIP SAMPLE CODE DEPTH INTERVAL (INCHES) DATE SAMPLE COLLECTED	SSU-U1A						SL-L2		SM-L1	SL-U3
	TRANSECT U						TRANSECT J		TRANSECT M	TRANSECT L
	BMN34						BMN08		BMN10	BMN11
	14 TO 08 IN						20 TO 25 IN		25 TO 30 IN	05 TO 10 IN
	ug/g	ug/g	ug/g	ug/g	ug/g	ug/g	ug/g	ug/g	ug/g	ug/g
	3/16/94	3/16/94	3/16/94	3/16/94	3/16/94	3/16/94	3/16/94	3/16/94	3/16/94	3/16/94
4-Nitroaniline	1000 U	25 U	25 U	25 U	25 U	25 U	990 U	1400 UJ	980 UJ	
4,6-Dinitro-2-methylphenol	1000 U	25 U	25 UJ	25 U	25 U	25 UJ	990 UJ	1400 U	980 U	
N-Nitrosodiphenylamine (1)	410 U	10 UJ	10 UJ	10 UJ	10 U	10 U	410 U	590 U	400 U	
4-Bromophenyl-phenylether	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	590 U	400 U	
Hexachlorobenzene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	590 U	400 U	
Pentachlorobenzene	1000 U	25 U	25 UJ	25 U	25 U	25 U	990 U	1400 U	980 U	
Phenanthrene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	210 J	370 J	
Anthracene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	32 J	92 J	
Carbazole	410 U	10 U	10 UJ	10 U	10 U	10 UJ	410 U	590 U	30 J	
Di-n-butylphthalate	29 J	10 U	10 UJ	10 U	10 U	10 U	410 U	62 J	400 U	
Fluoranthene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	450 J	600	
Pyrene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	500 J	550	
Butylbenzylphthalate	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	590 U	400 U	
2,3-Dichlorobenzidine	410 U	10 U	10 UJ	10 U	10 U	10 UJ	410 U	590 U	400 U	
Benzo(a)anthracene	410 U	10 UJ	10 UJ	10 U	10 U	10 U	410 U	240 J	400 J	
Chrysene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	270 J	270 J	
bis(2-Ethylhexyl)phthalate	410 U	09 J	10 UJ	1 J	23 J	05 J	410 U	590 U	400 U	
Di-n-octylphthalate	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	590 U	400 U	
Benzo(b)fluoranthene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	110 J	200 J	
Benzo(k)fluoranthene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	440 J	380 J	
Benzo(a)pyrene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	320 J	440	
Indeno(1,2,3-cd)pyrene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 UJ	180 J	110 J	
Dibenz(a,h)anthracene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 U	590 U	400 U	
Benzo(ghi)perylene	410 U	10 U	10 UJ	10 U	10 U	10 U	410 UJ	200 J	150 J	
Total EPOCs	29 J	09 J	1 J	23 J	05 J	14		3014 J	3616 J	
TIC Count	16	3	2	1	1	14		19	11	
Total TIC Concentration	3674 JN	9 JB	7 JB	5 J	2353 JN	14720 JN	10840 JN			

U = Analyte was not detected at the test
J = Estimated Value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence
R = Rejected during data validation
(1) = Cannot be separated from Diaphenyl

TABLE 2-9
PESTICIDES/PCBs RESULTS - SOILS/SEDIMENT
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	SA-U1	SSA-U1	SB-U2	SC-L3	SCC-L4	SSUD-U1	SED-U4	SED-L1	SEW-U4	SSUD-U1A		SE-L2	
SAMPLE LOCATION NO.	TRANSECT A	TRANSECT A	TRANSECT B	TRANSECT C	TRANSECT CC	--	EDISON GLEN	EDISON GLEN	EDISON WOODS	--		TRANSECT E	
LOCATION DESCRIPTION	BMN01	BMN02	BMN03	BMN04	BMN07	BMN20	BMN29	BMN30	BMN31	BMN39	BMN48	BMN50	
DEPTH INTERVAL	0.5 TO 1.0 ft	0.2 TO 0.9 ft	0.5 TO 1.0 ft	1.0 TO 1.5 ft	2.0 TO 2.5 ft	0.5 TO 1.0 ft	0.5 TO 0.5 ft	0.3 TO 0.6 ft	0.2 TO 0.7 ft	0.5 TO 1.0 ft		2.5 TO 3.0 ft	
UNITS	ug/l	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/l	ug/l	
DATE SAMPLE COLLECTED	3/22/94	3/22/94	3/22/94	3/22/94	3/22/94	3/22/94	3/24/94	3/24/94	3/24/94	3/22/94		3/22/94	
alpha-BHC	2.1 UJ	2.2 UJ	2.0 UJ	2.0 U	2.2 U	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 U	0.050 U	2.1 U
beta-BHC	2.1 UJ	2.2 UJ	2.0 UJ	R	2.2 U	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 U	0.050 U	2.1 U
delta-BHC	2.1 UJ	R	2.0 UJ	2.0 U	2.2 U	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 U	0.050 U	2.1 U
gamma-BHC (Lindane)	2.1 UJ	R	R	35	2.2 U	2.3 J	2.0 UJ	2.0 UJ	2.0 UJ	R	0.050 U	0.050 U	2.1 U
Heptachlor	2.1 UJ	2.2 UJ	2.0 UJ	2.0 U	2.2 U	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 U	0.050 U	2.1 U
Aldrin	2.1 UJ	2.2 UJ	2.0 UJ	2.0 U	2.2 U	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 U	0.050 U	2.1 U
Heptachlor epoxide	2.1 UJ	2.2 UJ	2.3 J	2.0 U	2.2 U	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 U	0.050 U	2.1 U
Endosulfan I	2.1 UJ	2.2 UJ	2.0 UJ	2.0 U	2.2 U	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 U	0.050 U	2.1 U
Dieldrin	0.1 J	0.5 JN	4.0 JN	13	4.2 U	0.1 J	4.0 J	3.0 UJ	3.0 UJ	4.5 J	0.10 U	0.10 U	0.5 J
4,4'-DDE	12 J	10 J	6.0 J	77 J	4.2 U	0.6 J	0.6	3.0 UJ	3.0 UJ	4.0 J	0.10 U	0.10 U	110 E
Endrin	4.1 UJ	4.0 JN	4.0 UJ	3.0 U	4.2 U	3.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.0 UJ	0.10 U	0.10 U	4.1 U
Endosulfan II	4.1 UJ	4.2 UJ	4.0 UJ	3.0 U	4.2 U	3.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.0 UJ	0.10 U	0.10 U	4.1 U
4,4'-DDD	50 J	100 J	0.5 JN	R	4.2 U	12 JN	11 J	4.3 J	3.0 UJ	4.4 JN	0.10 U	0.10 U	130 H
Endosulfan Sulfate	4.1 UJ	4.2 UJ	4.0 UJ	3.0	4.2 U	3.0 UJ	3.0 U	3.0 UJ	3.0 UJ	4.0 UJ	0.10 U	0.10 U	4.1 U
4,4'-DDT	20 J	5.0 J	25 J	570	4.2 U	17 J	10 J	7.0	3.0 UJ	6.0 JN	0.10 U	0.10 U	240
Methoxychlor	21 UJ	22 UJ	20 UJ	200 U	22 U	20 UJ	20 UJ	20 UJ	20 UJ	20 UJ	0.10 U	0.50 U	21 U
Endrin ketone	4.1 UJ	4.2 UJ	4.0 UJ	3.0 U	4.2 U	3.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.0 UJ	0.10 U	0.10 U	4.1 U
Endrin Aldehyde	4.1 UJ	4.2 UJ	4.0 UJ	3.0 U	4.2 U	3.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.0 UJ	0.10 U	0.10 U	4.1 U
alpha-Chlordane	4.1 J	0.3 J	2.0	7.3	2.2 U	2.3 U	2.3 J	2.7	2.0 UJ	2.3 J	0.050 U	0.050 U	6.0
gamma-Chlordane	4.4 J	0.3 J	2.0	7.7	2.2 U	2.7 J	1.0 J	2.7	2.0 UJ	2.0 UJ	0.050 U	0.050 U	6.1 J
Toxaphene	210 UJ	220 UJ	200 UJ	200 U	220 U	200 UJ	200 UJ	200 UJ	200 UJ	200 UJ	5.0 U	5.0 U	210 U
Aroclor-1016	41 UJ	42 UJ	40 UJ	39 U	42 U	38 UJ	39 UJ	38 UJ	39 UJ	40 UJ	1.0 U	1.0 U	41 U
Aroclor-1221	03 UJ	08 UJ	01 UJ	00 U	00 U	70 UJ	70 UJ	70 UJ	70 UJ	01 UJ	2.0 U	2.0 U	04 U
Aroclor-1232	01 UJ	42 UJ	40 UJ	39 U	42 U	38 UJ	39 UJ	38 UJ	39 UJ	40 UJ	1.0 U	1.0 U	41 U
Aroclor-1242	41 UJ	42 UJ	40 UJ	39 U	42 U	38 UJ	39 UJ	38 UJ	39 UJ	40 UJ	1.0 U	1.0 U	41 U
Aroclor-1248	41 UJ	42 UJ	40 UJ	39 U	42 U	38 UJ	39 UJ	38 UJ	39 UJ	40 UJ	1.0 U	1.0 U	41 U
Aroclor-1254	41 UJ	42 UJ	40 UJ	39 U	42 U	38 UJ	39 UJ	38 UJ	39 UJ	40 UJ	1.0 U	1.0 U	41 U
Aroclor-1260	41 UJ	42 UJ	40 UJ	39 U	42 U	38 UJ	39 UJ	38 UJ	39 UJ	40 UJ	1.0 U	1.0 U	41 U

U = Analyte was not detected at the instrument detection limit given

J = Estimated value

0 = Analyte was detected in blank

E = Estimated value due to matrix interference

D = Determined after sample dilution

NU = Presumptive evidence for presence of analyte; estimated quantity

P = There is a greater than 25% difference for detected concentrations between the two GC columns; the lower of the two values is reported.

R = Rejected during data validation

TABLE 3-6 CONTINUED
PESTICIDE/PCB RESULTS - SOILS/SEDIMENT
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	80-L2	850-U1	85A-L2	88B-L2	89C-U1	8H-U1	8H-U1A	8H-U1			85C0-2	85C0-3	850-8	85C0-7
LOCATION NO.	TRANSECT G	TRANSECT D	TRANSECT AA	TRANSECT BB	ROADK CIRCLE	TRANSECT H	TRANSECT H	TRANSECT H						
DEPTH INTERVAL	BM407	BM412	BM425	BM426	BM432	BM435	BM436	BM437						
UNITS	20 TO 25 H	0.5 TO 1.2 H	2.5 TO 3.0 H	2.5 TO 3.0 H	0.2 TO 0.7 H	0.5 TO 1.0 H	0.5 TO 1.0 H	0.0 TO 1.0 H			1.0 TO 1.5 H	2.0 TO 3.0 H	0.0 TO 1.5 H	1.5 TO 2.0 H
DATE SAMPLE COLLECTED	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	3/2/94	ug/l	ug/l	3/2/94	3/2/94	3/2/94	3/2/94
alpha-BHC	2.1 UJ	1.0 UJ	2.7 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	2.0 J	2.0 UJ	1.0 UJ	2.2 UJ
beta-BHC	2.1 UJ	1.0 UJ	2.7 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	2.0 UJ	2.0 UJ	R	2.2 UJ
delta-BHC	2.1 UJ	1.0 UJ	2.7 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	2.0 UJ	2.0 UJ	1.0 UJ	2.2 UJ
gamma-BHC (Lindane)	2.1 UJ	1.0 UJ	R	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	R	2.0 UJ	R	2.2 UJ
Heptachlor	2.1 UJ	1.0 UJ	2.7 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	2.0 UJ	2.0 UJ	1.0 UJ	2.2 UJ
Aldrin	2.1 UJ	1.0 UJ	2.7 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	2.0 UJ	2.0 UJ	1.0 UJ	2.2 UJ
Heptachlor epoxide	2.1 UJ	1.0 UJ	7.7 J	2.0 UJ	3.2	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	2.0 UJ	2.0 UJ	1.0 UJ	2.2 UJ
Endosulfen I	2.1 UJ	1.0 UJ	2.7 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.050 UJ	0.050 UJ	2.0 UJ	2.0 UJ	1.0 UJ	2.2 UJ
Dieldrin	4.1 UJ	3.0 UJ	67.0 J	3.0 UJ	4.0 UJ	10.0	14 J	3.0 UJ	0.10 UJ	0.10 UJ	5.0 J	7.0 J	3.0 UJ	4.3 UJ
4,4'-DDE	4.1 UJ	3.0 UJ	100 J	3.0 UJ	5.3 J	65	70	3.0 UJ	0.10 UJ	0.10 UJ	17 J	10 UJ	3.0 UJ	4.3 UJ
Endrin	4.1 UJ	3.0 UJ	56 JN	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	0.10 UJ	0.10 UJ	3.0 UJ	4.2 JN	3.0 UJ	4.3 UJ
Endosulfen II	4.1 UJ	3.0 UJ	5.3 UJ	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	0.10 UJ	0.10 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
4,4'-DDD	4.1 UJ	3.0 UJ	R	3.0 UJ	4.0 UJ	27 J	24 JN	3.0 UJ	0.10 UJ	0.10 UJ	57 JN	R	3.0 UJ	4.3 UJ
Endosulfen Sulfate	4.1 UJ	3.5 UJ	5.3 UJ	3.0 UJ	4.0 UJ	5.2	3.0 UJ	3.0 UJ	0.10 UJ	0.10 UJ	3.0 UJ	3.0 UJ	2.7 UJ	4.3 UJ
4,4'-DDT	4.1 UJ	3.5 UJ	67.0 J	2.0 UJ	0.0	81	250 J	3.0 UJ	0.10 UJ	0.10 UJ	0.1 J	0.7 J	3.7 J	4.3 UJ
Methoxychlor	2.1 UJ	1.0 UJ	2.7 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	2.0 UJ	0.10 UJ	0.10 UJ	2.0 UJ	2.0 UJ	0.0 J	2.2 UJ
Endrin ketone	4.1 UJ	3.5 UJ	5.4 J	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	0.10 UJ	0.10 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
Endrin Aldehyde	4.1 UJ	3.5 UJ	5.3 UJ	2.0 UJ	4.0 UJ	R	4.0 JN	3.0 UJ	0.10 UJ	0.10 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
alpha-Chlordane	2.1 UJ	3.3 J	2.3 J	2.0 UJ	2.1	4.7	5.2 JN	2.0 UJ	0.050 UJ	0.050 UJ	5.2 UJ	0.2 J	1.0 UJ	2.2 UJ
gamma-Chlordane	2.1 UJ	3.4 UJ	2.3 J	2.0 UJ	2.0 UJ	3.1 J	3.4 J	2.0 UJ	0.050 UJ	0.050 UJ	5.4 J	1.2 J	4.3 J	2.2 UJ
Toxaphene	2.1 UJ	100 UJ	270 UJ	30 UJ	200 UJ	200 UJ	200 UJ	200 UJ	5.0 UJ	5.0 UJ	200 UJ	200 UJ	100 UJ	2.0 UJ
Aroclor - 1016	4.1 UJ	3.5 UJ	5.3 UJ	0.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	1.0 UJ	1.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
Aroclor - 1221	04 UJ	72 UJ	110 UJ	3.0 UJ	0.1 UJ	78 UJ	78 UJ	78 UJ	2.0 UJ	2.0 UJ	78 UJ	78 UJ	73 UJ	08 UJ
Aroclor - 1252	4.1 UJ	3.5 UJ	5.3 UJ	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	1.0 UJ	1.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
Aroclor - 1242	4.1 UJ	3.5 UJ	5.3 UJ	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	1.0 UJ	1.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
Aroclor - 1248	4.1 UJ	3.5 UJ	5.3 UJ	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	1.0 UJ	1.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
Aroclor - 1254	4.1 UJ	3.5 UJ	5.3 UJ	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	1.0 UJ	1.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	4.3 UJ
Aroclor - 1260	4.1 UJ	3.5 UJ	5.3 UJ	3.0 UJ	4.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	1.0 UJ	1.0 UJ	3.0 UJ	3.0 UJ	3.0 UJ	74 P

U = Analyte was not detected at the instrument detection level
J = Estimated value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence of analyte
P = There is a greater than 25% difference for data between the two GC columns; the lower of the two is reported
R = Rejected during data validation

TABLE 3-2 CONTINUED
PESTICIDES/PCBs RESULTS - SOIL/SEDIMENT
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER	SSCO-8	SSK-U1	SI-U1		SO-L4	SO-L1	SR-L2	ST-L4	SSU-U1	SV-L1	SW-L2	SK-L3	SY-L4	SSZ-U1
SAMPLE LOCATION NO.	--	TRANSECT K	TRANSECT I		TRANSECT O	TRANSECT O	TRANSECT R	TRANSECT T	TRANSECT U	TRANSECT V	TRANSECT W	TRANSECT X	TRANSECT Y	TRANSECT Z
LOCATION DESCRIPTION	BMN00	BMN00	BMN13	BMN16	BMN14	BMN15	BMN16	BMN16	BMN19	BMN20	BMN21	BMN22	BMN23	BMN24
DEPTH INTERVAL	0.0 TO 0.5 h	0.5 TO 1.2 h	0.5 TO 1.0 h		2.5 TO 3.0 h	2.0 TO 2.5 h	2.5 TO 3.0 h	2.5 TO 3.0 h	0.4 TO 0.8 h	2.5 TO 3.0 h	2.5 TO 3.0 h	2.5 TO 3.0 h	2.0 TO 2.5 h	0.3 TO 0.8 h
UNITS	ug/kg	ug/kg	ug/kg	ug/l	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg
DATE SAMPLE COLLECTED	3/23/94	3/18/94	3/18/94		3/17/94	3/17/94	3/16/94	3/16/94	3/16/94	3/16/94	3/15/94	3/15/94	3/14/94	3/14/94
alpha-BHC	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	2.1 U	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
beta-BHC	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	2.1 U	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
delta-BHC	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	3.5 JN	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
gamma-BHC (Lindane)	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	2.1 U	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
Heptachlor	2.1 U	2.0 U	2.3 U	0.050 U	2.3	2.4 J	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
Aldrin	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	2.1 U	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
Heptachlor epoxide	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	2.1 U	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
Endosulfen I	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	2.1 U	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
Dieldrin	4.0 U	3.0 U	4.3 U	0.10 U	3.7 U	R	4.3 U	4.3 U	4.3 U	4.3 U	6.0 JN	4.5 U	5.0 J	4.3 U
4,4'-DDE	4.0 U	3.0 U	4.3 U	0.10 U	3.7 U	4.1 U	4.3 U	4.3 U	4.3 U	4.3 U	3.0 U	4.5 U	5.4 U	4.3 U
Endrin	4.0 U	3.0 U	4.3 U	0.10 U	3.7 U	4.1 U	4.3 U	4.3 U	4.3 U	4.3 U	3.0 U	4.5 U	5.4 U	4.3 U
Endosulfen II	4.0 U	3.0 U	4.3 U	0.10 U	3.7 U	4.1 U	4.3 U	4.3 U	4.3 U	4.3 U	3.0 U	4.5 U	5.4 U	4.3 U
4,4'-DDD	4.0 U	7.0 JN	4.3 U	0.10 U	3.7 U	17.0 JN	4.3 U	4.3 U	5.4 JN	3.0	2.2 JN	R	R	4.3 U
Endosulfen Sulfate	4.0 U	3.0 U	4.3 U	0.10 U	3.7 U	2.0 J	4.3 U	4.3 U	4.3 U	4.3 U	3.0 U	4.5 U	5.4 U	4.3 U
4,4'-DDT	4.0 U	1.5	7.7	0.10 U	3.7 U	1200 *	9.4	4.3 U	1.4	0.5 JN	120 *	7.6	3.5 J	8.1
Methoxychlor	4.0 U	2.0 U	2.3 U	0.050 U	3.7 U	2.1 U	2.2 U	2.2 U	2.2 U	2.2 U	2.0 U	2.3 U	2.0 U	2.2 U
Endrin ketone	2.1 U	3.0 U	4.3 U	0.10 U	1.0 U	4.1 U	4.3 U	4.3 U	4.3 U	4.3 U	3.0 U	4.5 U	5.4 U	4.3 U
Endrin Aldohyde	4.0 U	3.0 U	4.3 U	0.10 U	3.7 U	R	4.3 U	4.3 U	5.2	4.3 U	3.0 U	4.5 U	5.4 U	4.3 U
alpha-Chlordane	4.0 U	2.0 U	2.3 U	0.050 U	0.7	2.0 J	2.2 U	2.2 U	2.2 U	2.2 U	2.0	2.3 U	2.0 U	2.2 U
gamma-Chlordane	2.1 U	2.0 U	2.3 U	0.050 U	1.0 U	2.1 J	2.2 U	2.2 U	2.2 U	2.2 U	4.3 JN	2.3 U	2.0 U	2.2 U
Toxaphene	2.1 U	200 U	270 U	5.0 U	1.0 U	2100 U	220 U	220 U	220 U	220 U	200 U	230 U	200 U	200 U
Aroclor-1016	210 U	30 U	42 U	1.0 U	100 U	410 U	42 U	43 U	42 U	43 U	30 U	45 U	54 U	42 U
Aroclor-1221	40 U	77 U	86 U	2.0 U	37 U	640 U	85 U	87 U	86 U	88 U	78 U	92 U	110 U	85 U
Aroclor-1232	82 U	30 U	42 U	1.0 U	70 U	410 U	42 U	43 U	42 U	43 U	30 U	45 U	54 U	42 U
Aroclor-1242	40 U	30 U	42 U	1.0 U	37 U	410 U	42 U	43 U	42 U	43 U	30 U	45 U	54 U	42 U
Aroclor-1248	40 U	30 U	42 U	1.0 U	37 U	410 U	42 U	43 U	42 U	43 U	30 U	45 U	54 U	42 U
Aroclor-1254	40 U	30 U	42 U	1.0 U	37 U	410 U	42 U	43 U	42 U	43 U	30 U	45 U	54 U	42 U
Aroclor-1260	40 U	30 U	42 U	1.0 U	37 U	410 U	42 U	43 U	42 U	43 U	30 U	45 U	54 U	42 U

U = Analyte was not detected at the instrument de
J = Estimated value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence of analyte
P = There is a greater than 25% difference for data
between the two GC columns; the lower of the two
R = Rejected during data validation

TABLE 3-9 CONTINUED
PESTICIDES/PCBs RESULTS - SOILS/SEDIMENT
OFF-SITE INVESTIGATION
CHEMICAL INSECTICIDE CORPORATION SITE
EDISON, NEW JERSEY

SAMPLE NUMBER SAMPLE LOCATION NO. LOCATION DESCRIPTION DEPTH INTERVAL UNITS DATE SAMPLE COLLECTED	BW-L2A TRANSECT W BMN33 2.5 TO 3.0 ft ug/kg 3/15/94	BBU-L1A TRANSECT U BMN34 0.4 TO 0.8 ft ug/kg 3/15/94	BMN1 ug/l	BMN2 ug/l	BMN3 ug/l	BMN4 ug/l	BMN5 ug/l	SJ-L2 TRANSECT J BMN08 2.0 TO 2.5 ft ug/kg 3/15/94	SM-L1 TRANSECT M BMN10 2.5 TO 3.0 ft ug/kg 3/15/94	SL-U3 TRANSECT BMN11 0.5 TO 1.0 ft ug/kg 3/17/94
alpha-BHC	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.1 U	3.3 J	2.1
beta-BHC	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.1 U	38 U	
delta-BHC	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.1 U	24 J	4.6
gamma-BHC (lindane)	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.1 U	6.4 JN	2.1
Heptachlor	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.2	30 U	2.2
Aldrin	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.1 U	30 U	2.6
Heptachlor epoxide	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.1 U	30 U	2.1
Endosulfan I	2.1 U	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	2.1 U	30 U	2.1
Dieldrin	87	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	29	80	27
4,4'-DDE	17	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	72 *	230	68
Endrin	4.0 U	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	4.1 U	34 J	23
Endosulfan II	4.0 U	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	4.1	59 U	
4,4'-DDD	32 J	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	R	1500 *	160
Endosulfan Sulfate	4.0 U	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	4.1 U	59 U	4.6
4,4'-DDT	150 *	6.0 JN	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	360 *	360	570
Methoxychlor	21 U	21 U	0.50 U	0.50 U	0.50 U	0.50 U	0.50 U	21 U	300 U	21
Endrin ketone	4.0 U	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	4.1 U	59 U	
Endrin Aldehyde	4.0 U	4.1 U	0.10 U	0.10 U	0.10 U	0.10 U	0.10 U	4.1 U		4.0
alpha-Chlordane	5.6 J	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	6.6	120	23
gamma-Chlordane	47 JN	2.1 U	0.050 U	0.050 U	0.050 U	0.050 U	0.050 U	13 *	110 J	29
Toxaphene	210 U	210 U	5.0 U	5.0 U	5.0 U	5.0 U	5.0 U	210 U	3000 U	210
Aroclor-1016	40 U	41 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	41 U	590 U	40
Aroclor-1221	82 U	84 U	2.0 U	2.0 U	2.0 U	2.0 U	2.0 U	83 U	1200 U	82
Aroclor-1232	40 U	41 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	41 U	590 U	40
Aroclor-1242	40 U	41 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	41 U	590 U	40
Aroclor-1246	40 U	41 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	41 U	590 U	40
Aroclor-1254	40 U	41 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	41 U	590 U	40
Aroclor-1260	40 U	41 U	1.0 U	1.0 U	1.0 U	1.0 U	1.0 U	41 U	590 U	40

U = Analyte was not detected at the instrument de
J = Estimated value
B = Analyte was detected in blank
E = Estimated value due to matrix interference
D = Determined after sample dilution
NJ = Presumptive evidence for presence of analyte
P = There is a greater than 25% difference for data
between the two QC columns; the lower of the two
R = Rejected during data validation

TABLE 3 - 10
INORGANIC RESULTS - SOILS/SEDIMENTS
CHEMICAL INSECTICIDE CORPORATION SITE
OFF-SITE INVESTIGATION
EDISON, NEW JERSEY

	1	2	3	4	5	6	7	8
SAMPLE NUMBER	SA-U1	SSA-U1	SB-U2	SE-L2	SG-L2	SSO-U1	SAA-L2	SBB-L3
SAMPLE LOCATION NO.	TRANSECT A	TRANSECT A	TRANSECT B	TRANSECT E	TRANSECT G	TRANSECT O	TRANSECT AA	TRANSECT BB
CLP SAMPLE CODE	MBNQ01	MBNQ02	MBNQ03	MBNQ05	MBNQ07	MBNQ12	MBNQ25	MBNQ26
DEPTH INTERVAL	0.5 TO 1.0 ft.	1.5 TO 2.0 ft.	0.5 TO 1.0 ft.	2.5 TO 3.0 ft.	2.0 TO 2.5 ft.	0.3 TO 1.2 ft.	2.5 TO 3.0 ft.	2.5 TO 3.0 ft.
UNITS	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg
DATE SAMPLE COLLECTED	3/22/94	3/22/94	3/22/94	3/21/94	3/21/94	3/21/94	3/23/94	3/23/94
ALUMINUM	5700 J	5290 J	10800 J	14600	5270	3300	17000	8120
ANTIMONY	4.5 UNJ	4.7 UNJ	4.5 UNJ	4.7 UNJ	4.3 UNJ	4.4 UNJ	5.9 UNJ	4.6 UNJ
ARSENIC	21.3 J		11.0 J	15.4 S	0.23 BWJ	4.4	129	2.9
BARIUM	34.1 J	41.8 J	45.0 J	85.6	14.1 B	19.8 B	143	31.8 B
BERYLLIUM	0.3 J	0.88 J	0.37 BJ	0.43 B	0.24 B	0.29 B	0.80 B	0.49 B
CADMIUM	0.32 J	0.65 BJ	0.83 J	0.81 B	0.31 U	0.54 B	0.47 B	0.49 B
CALCIUM	1420 J	1750 J	1260 J	407 B	109 B	510 B	1020 B	588 B
CHROMIUM	10.5 JN	14.2 NJ	16.8 NJ	18.2 NJ	11.2 NJ	21.8 NJ	41.6 NJ	15.8 NJ
COBALT	2.5 JB	5.8 BJ	5.1 BJ	6.9 B	1.9 B	5.5 B	12.5	4.2 B
COPPER	19.5 J	31.9 J	31.1 J	27.3 J	3.1 B	11.3 J	82.0 J	6.6 J
IRON	9890 *J	13700 *J	16600 *J	15900 *	2580 *	12000 *	30500 *	13500 *
LEAD	25.9 N*J	74.9 N*J	44.4 N*J	63.6 N*J	7.0	36.1 N*J	234 N*J	18.8
MAGNESIUM	772 BJ	1350 J	1560 J	1160	727 B	924	2310	1280
MANGANESE	84.8 JN*J	108 N*J	102 N*J	311 N*J	16.9 N*J	99.2 N*J	242 N*J	64.2 N*J
MERCURY	0.16 J	0.13 J	0.13 J	0.13	0.06 U	0.09 B	0.32	0.06 U
NICKEL	6.6 J	18.3 J	12.2 J	12.2	5.4 B	10.9	19.4	9.9
POTASSIUM	520 BJ	530 BJ	974 J	648 B	723 B	410 B	1290	616 B
SELENIUM	0.53 BJ	0.35 BJ	0.34 BJ	0.71 BNJ	0.23 U	0.24 B	2.2 S	0.35 B
SILVER	0.32 UJ	0.40 BJ	0.44 BJ	0.34 U	0.31 U	0.32 U	0.42 U	0.33 U
SODIUM	52.9 BJ	1.47 BJ	76.4 BJ	57.5 B	41.6 B	95.4 B	178 B	144 B
THALLIUM	0.48 UWJ	0.51 UJ	0.49 UJ	0.51 UW	0.46 U	0.48 U	0.63 U	0.50 U
VANADIUM	15.5 J	19.4 J	26.0 J	29.9	7.0 B	13.9	47.1	19.6
ZINC	32.1 N*J	213 N*J	79.0 N*J	83.6 N*J	20.6 N*J	58.9 N*J	136 N*J	53.5 N*J
CYANIDE	0.18 UJ	0.19 UJ	0.18 UJ	0.37 B	0.17 U	0.38 B	0.54 B	0.19 U

U= Analyte was not detected at the instrument detection limit given

J= Estimated Value

B= Reported value is between the instrument detection limit and the contract required detection limit

E= Value is estimated due to interference

N= Spiked sample recovery was not within control limits

* = Duplicate analysis was not within control limits

S= Determined by Method of Standard Addition (MSA)

W= Post digestion spike for furnace AA analysis out of control limits, while sample absorbance is less than 50% of spike absorbance

+= Correlation coefficient for the MSA is less than 0.995

M= Duplicate injection precision criteria was not met

R= Rejected during data validation

TABLE 3 - 10
INORGANIC RESULTS - SOILS/SEDIMENTS
CHEMICAL INSECTICIDE CORPORATION SITE
OFF-SITE INVESTIGATION
EDISON, NEW JERSEY

	9	10	11	12	13	14	15	16
SAMPLE NUMBER	SSU-U1A						SSK-U1	S1-U1
SAMPLE LOCATION NO.	TRANSECT U						TRANSECT K	TRANSECT S
CLP SAMPLE CODE	MBNQ34	MBNQ41	MBNQ42	MBNQ43	MBNQ44	MBNQ45	MBNQ09	MBNQ13
DEPTH INTERVAL	0.4 TO 0.8 ft.						0.3 TO 1.2 ft.	0.5 TO 1.0 ft.
UNITS	mg/kg	ug/l	ug/l	ug/l	ug/l	ug/l	mg/kg	mg/kg
DATE SAMPLE COLLECTED	3/16/94						3/16/94	3/16/94
ALUMINUM	7060	20.8 BJ	20.8 UJ	20.8 UJ	20.8 UJ	20.8 UJ	7210 *	14000 *
ANTIMONY	7.0 UNJ	28.3 UJ	28.3 UJ	28.3 UJ	28.3 UJ	28.3 UJ	6.7 UNJ	7.5 UNJ
ARSENIC	9.8 SNJ	1.3 UJ	1.4 BJ	1.7 BJ	2.2 BJ	1.7 BJ	17.6 S*	8.1 S*
BARIUM	34.6 B	0.80 UJ	0.80 UJ	0.80 UJ	0.80 UJ	0.80 UJ	53.7	37.5 B
BERYLLIUM	0.94 B	0.20 UJ	0.20 UJ	0.20 UJ	0.20 UJ	0.20 UJ	1.9	0.62 B
CADMIUM	0.67 U	2.7 UJ	2.7 UJ	2.7 UJ	2.7 UJ	2.7 UJ	0.64 U	0.72 U
CALCIUM	1710	36.5 BJ	45.6 BJ	45.6 BJ	45.6 BJ	54.7 BJ	3010	1100 B
CHROMIUM	18.3 N*J	2.6 UJ	2.6 UJ	2.6 UJ	2.6 UJ	2.6 UJ	21.8	29.7
COBALT	6.5 B	2.9 UJ	2.9 UJ	2.9 UJ	2.9 UJ	2.9 UJ	18.5	8.9 B
COPPER	10.0 J	2.4 UJ	2.4 UJ	2.4 UJ	2.4 UJ	2.4 UJ	30.4 N*J	35.6 N*J
IRON	35800 *	53.2 BJ	65.2 BJ	13.7 BJ	39.1 BJ	36.7 BJ	37300 *J	51100 *J
LEAD	26.2 S	0.60 UJ	0.64 BJ	0.64 BJ	0.60 UJ	1.7 BJ	26.4 *J	55.3 S*J
MAGNESIUM	1880	26.8 UJ	26.8 UJ	26.8 UJ	26.8 UJ	26.8 UJ	1910	2660
MANGANESE	123 N*J	1.7 UJ	1.9 BJ	1.7 UJ	1.7 UJ	1.7 UJ	351 N*J	237 N*J
MERCURY	0.12 U	0.20 U	0.20 U	0.20 U	0.20 U	0.20 U	0.12 U	0.13 U
NICKEL	12.6	10.8 UJ	10.8 UJ	10.8 UJ	10.8 UJ	10.8 UJ	21.2 *	15.7 *
POTASSIUM	1340	348 UJ	348 UJ	348 UJ	348 UJ	348 UJ	892 B	2230
SELENIUM	0.30 U	1.2 UJ	1.2 UJ	1.2 UJ	1.2 UJ	1.2 UJ	0.28 U	0.46 B
SILVER	2.0 B	2.8 UJ	2.8 UJ	2.8 UJ	2.8 UJ	2.8 UJ	2.6 *J	3.5 *J
SODIUM	143 B	43.7 BJ	75.6 BJ	41.3 BJ	58.1 BJ	41.6 BJ	205 B	76.7 B
THALLIUM	0.25 U	1.3 BWJ	1.1 BJ	1.1 BWJ	1.6 BJ	2.2 BJ	0.42 B	0.67 B
VANADIUM	19.6 *J	2.3 UJ	2.3 UJ	2.3 UJ	2.3 UJ	2.3 UJ	20.2	46.0
ZINC	61.8 NJ	4.3 BJ	10.3 BJ	6.1 BJ	3.1 UJ	5.2 BJ	132 N*J	90.3 N*J
CYANIDE	0.62 UN	10.0 U	10.0 U	10.0 U	10.0 U	10.0 U	0.59 U	0.66 U

U= Analyte was not detected at the instrument detection limit given

J= Estimated Value

B= Reported value is between the instrument detection limit and the contract required detection limit

E= Value is estimated due to interference

N= Spiked sample recovery was not within control limits

* = Duplicate analysis was not within control limits

S= Determined by Method of Standard Addition (MSA)

W= Post digestion spike for furnace AA analysis out of control limits, while sample absorbance is less than 50% of spike absorbance

+ = Correlation coefficient for the MSA is less than 0.995

M= Duplicate injection precision criteria was not met

R= Rejected during data validation

TABLE 3 - 10
INORGANIC RESULTS - SOILS/SEDIMENTS
CHEMICAL INSECTICIDE CORPORATION SITE
OFF-SITE INVESTIGATION
EDISON, NEW JERSEY

	17	18	19	20	21	22	23	24
SAMPLE NUMBER		SCC-L4	SSDD-U1	SRC-U1	SH-U1	SH-U1A	SH-U4	
SAMPLE LOCATION NO.		TRANSECT CC	TRANSECT DD	RODAK CIRCLE	TRANSECT H	TRANSECT H	TRANSECT H	
CLP SAMPLE CODE	MBNQ46	MBNQ27	MBNQ28	MBNQ37	MBNQ35	MBNQ36	MBNQ37	MBNQ39
DEPTH INTERVAL		2.0 TO 2.5 ft.	0.5 TO 2.0 ft.	0.2 TO 0.7 ft.	0.5 TO 1.0 ft.	0.5 TO 1.0 ft.	0.5 TO 1.0 ft.	
UNITS	ug/l	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	
DATE SAMPLE COLLECTED		3/22/94	3/22/94	3/23/94	3/21/94	3/21/94	3/21/94	
ALUMINUM	20.8 UJ	13300 J	6590 J	14700	4290	4070	9500	8410 J
ANTIMONY	28.3 UJ	4.9 UNJ	4.3 UNJ	4.6 UNJ	4.3 UNJ	4.3 UNJ	4.5 UNJ	4.6 UNJ
ARSENIC	1.3 UJ	4.5 J	5.6 SJ	3.6	12.0 J	6.9 SJ	17.9 S	13.3 SJ
BARIUM	1.0 BJ	49.2 J	32.9 J	58.1	72.7	55.0	51.5	40.1 J
BERYLLIUM	0.20 UJ	0.49 BJ	0.56 BJ	0.41 B	0.22 B	0.29 B	0.37 B	0.80 BJ
CADMIUM	2.7 UJ	0.35 UJ	0.43 BJ	0.51 B	0.31 U	0.78	0.32 U	0.41 BJ
CALCIUM	35.9 BJ	929 J	1260 J	769 B	692 B	585 B	356 B	1200 J
CHROMIUM	2.6 UJ	22.6 NJ	16.6 NJ	14.4 NJ	9.3 NJ	8.2 NJ	14.5 NJ	26.6 NJ
COBALT	2.9 UJ	7.7 BJ	7.7 J	4.6 B	2.9 B	2.9 B	4.1 B	7.7 BJ
COPPER	2.4 UJ	16.5 J	84.9 J	19.2 J	20.3 J	29.1 J	17.5 J	21.9 J
IRON	15.7 BJ	23400 *	15700 *J	11600 *	7830 *	8100 *	15400 *	35300 *J
LEAD	0.99 BJ	103 N*J	59.9 N*J	35.5 N*J	46.9 N*J	65.7 N*J	28.6 N*J	58.3 N*J
MAGNESIUM	26.8 UJ	2680 J	2040 J	1420	625 B	554 B	1030	1830 J
MANGANESE	1.7 UJ	108 N*J	136 N*J	82.4 N*J	79.6 N*J	97.6 N*J	165 N*J	165 N*J
MERCURY	0.20 U	0.18 J	0.16 J	0.17	0.16	0.12	0.16	0.17 J
NICKEL	10.8 UJ	26.4 J	27 J	10.9	5.7 B	7.2	9.9	15.9 J
POTASSIUM	348 UJ	1170 J	754 BJ	768 B	437 B	343 B	641 B	863 J
SELENIUM	1.2 UJ	0.37 BWJ	0.25 BJ	0.59 B	0.51 B	0.47 B	0.56 BWJ	0.34 BJ
SILVER	2.8 UJ	0.35 UJ	0.31 UJ	0.33 U	0.31 U	0.31 U	0.59 B	0.33 UJ
SODIUM	104 BJ	465 BJ	192 BJ	77.9 B	61.2 B	47.9 B	45.5 B	152 BJ
THALLIUM	1.2 BJ	0.52 UJ	0.46 UJ	0.49 U	0.46 U	0.47 U	0.48 UW	0.49 UJ
VANADIUM	2.3 UJ	26.2 J	15.2 J	25.7	15.0	21.2	23.6	38.8 J
ZINC	3.1 UJ	82.6 N*J	189 N*J	33.8 N*J	46.8 N*J	67.9 N*J	34.7 N*J	94.6 N*J
CYANIDE	10.0 U	0.20 UJ	0.17 UJ	0.18 U	0.2 B	0.17 U	0.18 U	0.18 UJ

U= Analyte was not detected at the instrument detection limit given

J= Estimated Value

B= Reported value is between the instrument detection limit and the contract required detection limit

E= Value is estimated due to interference

N= Spiked sample recovery was not within control limits

* = Duplicate analysis was not within control limits

S= Determined by Method of Standard Addition (MSA)

W= Post digestion spike for furnace AA analysis out of control limits, while sample absorbance is less than 50% of spike absorbance

+= Correlation coefficient for the MSA is less than 0.995

M= Duplicate injection precision criteria was not met

R= Rejected during data validation

TABLE 3 - 10
INORGANIC RESULTS - SOILS/SEDIMENTS
CHEMICAL INSECTICIDE CORPORATION SITE
OFF-SITE INVESTIGATION
EDISON, NEW JERSEY

	25	26	27	28	29	30	31	32
SAMPLE NUMBER	SSCG-2	ST-L4	SSU-U1	SV-L1	SW-L2	SX-L3	SY-L4	SSZ-U1
SAMPLE LOCATION NO.		TRANSECT T	TRANSECT U	TRANSECT V	TRANSECT W	TRANSECT X	TRANSECT Y	TRANSECT Z
CLP SAMPLE CODE	MBNQ58	MBNQ18	MBNQ19	MBNQ20	MBNQ21	MBNQ22	MBNQ23	MBNQ24
DEPTH INTERVAL	1.0 TO 1.5 ft.	2.5 TO 3.0 ft.	0.4 TO 0.8 ft.	2.5 TO 3.0 ft.	2.5 TO 3.0 ft.	2.5 TO 3.0 ft.	2.0 TO 2.5 ft.	0.3 TO 0.8 ft.
UNITS	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg	mg/kg
DATE SAMPLE COLLECTED	3/23/94	3/16/94	3/16/94	3/15/94	3/15/94	3/15/94	3/14/94	3/14/94
ALUMINUM	4390	9440	7850	10500	8040	7690	15700	5380
ANTIMONY	4.4 UNJ	8.1 UNJ	7.1 UNJ	8.0 UNJ	7.2 UNJ	7.7 UNJ	9.3 UNJ	7.3 UNJ
ARSENIC	29.6 S	11.5 SNJ	5.2 NJ	4.1 SNJ	18.0 SNJ	10.1 SNJ	6.9 SNJ	4.1 NJ
BARIUM	19.0 B	54.9 B	34.3 B	70.9	53.8	43.6 B	141	30.3 B
BERYLLIUM	0.40 B	0.76 B	1.0 B	1.9 J	1.2 B	1.0 B	0.65 B	1.9 J
CADMIUM	0.31 U	0.77 U	0.68 U	0.76 U	0.69 U	0.74 U	0.89 U	0.7 U
CALCIUM	918	1800	2540	3170	925 B	521 B	1280 B	2530
CHROMIUM	12.8 NJ	25.9 N*J	21.5 N*J	19.0 N*J	18.2 N*J	42.4 N*J	24.6 N*J	17.9 N*J
COBALT	4.9 B	11.7 B	5.1 B	18.9	8.0 B	14.3	7.2 B	10.9 B
COPPER	13.3 J	74.5	49.7 J	23.1 J	36.1	29.7	47.0	9.5
IRON	16400 *	52800 *	34100 *	23700 *	31300 *	79300 *	18800 *	18200 *
LEAD	28.1 N*J	22.0 S	26.4 S	44.4 S	95.0 *	39.3 S	105 *	19.7 S
MAGNESIUM	1930	1960	2350	2370	1760	968 B	2200	2880
MANGANESE	125 N*J	289 N*J	130 N*J	429 N*J	115 N*J	243 N*J	80.8 N*J	236 N*J
MERCURY	0.08 U	0.14 U	0.13 U	0.14 U	0.13 U	0.14 U	0.16 U	0.13 U
NICKEL	12.7	28.7	14.9	20.9	11.1	15.6	12.8 B	30.5
POTASSIUM	670 B	1200 B	1530	1150 B	683 B	662 B	1050 B	1410
SELENIUM	0.30 B	0.34 U	0.30 U	0.34 U	0.38 B	0.33 U	0.39 U	0.31 U
SILVER	0.31 U	2.7 B	1.5 B	0.86 B	1.4 B	4.0	0.92 U	0.72 U
SODIUM	93.3 B	186 B	144 B	129 B	44.5 B	193 B	287 B	95.2 B
THALLIUM	0.47 U	0.54 B	0.47 BW	0.32 B	0.29 BWJ	0.80 B	0.59 BWJ	0.54 BWJ
VANADIUM	15.0	51.4 *J	26.1 *J	26.2 *J	22.7 *J	62.9 *J	31.7 *J	14.9 *J
ZINC	57.5 N*J	52.5 NJ	51.7 NJ	92.9 NJ	91.1 NJ	128 NJ	68.2 NJ	109 NJ
CYANIDE	0.17 U	0.72 UN	0.63 UN	0.70 UN	0.64 UN	0.68 UN	0.82 UN	0.65 UN

U= Analyte was not detected at the instrument detection limit given

J= Estimated Value

B= Reported value is between the instrument detection limit and the contract required detection limit

E= Value is estimated due to interference

N= Spiked sample recovery was not within control limits

* = Duplicate analysis was not within control limits

S= Determined by Method of Standard Addition (MSA)

W= Post digestion spike for furnace AA analysis out of control limits, while sample absorbance is less than 50% of spike absorbance

+ = Correlation coefficient for the MSA is less than 0.995

M= Duplicate injection precision criteria was not met

R= Rejected during data validation

TABLE 3 - 10
INORGANIC RESULTS - SOILS/SEDIMENTS
CHEMICAL INSECTICIDE CORPORATION SITE
OFF-SITE INVESTIGATION
EDISON, NEW JERSEY

	33	34	35	36	37	38	39	40
SAMPLE NUMBER	SW-L2A	SSGC-3	SSGC-8	SSGC-7		SC-L3	SEG-V4	SEG-11
SAMPLE LOCATION NO.	TRANSECT W					TRANSECT C	EDISON GLEN	EDISON GLEN
CLP SAMPLE CODE	MBNQ33	MBNQ57	MBNQ58	MBNQ59	MBNQ60	MBNQ04	MBNQ29	MBNQ30
DEPTH INTERVAL	2.5 TO 3.0 ft.	2.0 TO 3.0 ft.	0.0 TO 0.5 ft.	1.5 TO 2.0 ft.		1.0 TO 1.5 ft.	0.0 TO 0.5 ft.	0.3 TO 0.8 ft.
UNITS	mg/kg	mg/kg	mg/kg	mg/kg		mg/kg	mg/kg	mg/kg
DATE SAMPLE COLLECTED	3/15/94	3/23/94	3/23/94	3/23/94		3/22/94	3/24/94	3/24/94
ALUMINUM	6880	4550	5240	6310	10400	2940 J	10900 J	9070 J
ANTIMONY	6.8 UNJ	4.6 UNJ	4.5 UNJ	5.0 UNJ	4.6 UNJ	4.7 UNJ	4.5 UNJ	4.4 UNJ
ARSENIC	16.8 SNJ	8.0 S	4.3	1.7 B	4.2	6.1 NJ	6.1 NJ	3.8 NJ
BARIUM	53.9	24.1 B	65.3	64.7	46.5	16.9 BJ	43.9 J	46.3 J
BERYLLIUM	1.1 B	0.39 B	0.93	2.5	0.35 B	0.17 UJ	0.36 BJ	0.19 BJ
CADMIUM	0.63 U	0.35 B	0.42 B	0.36 U	0.33 U	0.34 UJ	0.32 UJ	0.58 BJ
CALCIUM	1350	1620	7150	4560	140 B	161 BJ	1060 J	1230 J
CHROMIUM	16.0 N*J	19.8 NJ	33.0 NJ	19.7 NJ	13.3 NJ	3.7 J	14.9 J	15.9 J
COBALT	5.9 B	5.1 B	8.5	8.4 B	7.2 B	1.9 UJ	6.6 BJ	4.9 BJ
COPPER	23.6	17.3 J	95.0 J	83.6 J	10.2 J	6.5 J	28.0 J	59.6 J
IRON	26000 *	23000 *	26200 *	29900 *	15500 *	1790 J	17000 J	16500 J
LEAD	33.1 S	42.1 N*J	137 N*J	36.9 N*J	16.0	13.2 SN*J	42.2 *J	23.4 *J
MAGNESIUM	950 B	1830	3780	2240	1350	266 BJ	2070 J	1740 J
MANGANESE	70.6 N*J	139 N*J	362 N*J	136 N*J	91.5 N*J	16.9 J	233 J	197 J
MERCURY	0.12 U	0.06 U	0.12	0.07 U	0.06 U	0.11 BJ	0.14 J	0.21 J
NICKEL	9.2 B	13.6	29.8	25.3	10.8	2.1 BJ	13.1 J	18.0 J
POTASSIUM	832 B	722 B	575 B	1400	450 B	225 BJ	777 BJ	1160 J
SELENIUM	0.28 U	0.24 U	0.33 BWJ	0.27 U	0.49 B	0.49 BWJ	0.66 BJ	0.27 BWJ
SILVER	1.5 B	0.33 U	0.46 B	0.36 U	0.37 B	0.72 BJ	0.74 BJ	0.31 UJ
SODIUM	34.3 B	107 B	385 B	135 B	52.9 B	43.5 BJ	80.8 BJ	77.0 BJ
THALLIUM	0.42 B	0.49 U	0.48 U	0.53 U	0.49 U	0.49 UWJ	0.47 UWJ	0.45 UJ
VANADIUM	22.2 *J	13.6	35.7	14.3	23.3	5.7 BJ	33.1 J	23.5 J
ZINC	59.1 NJ	73.0 N*J	322 N*J	218 N*J	68.3 N*J	8.0 J	72.4 J	43.9 J
CYANIDE	0.59 NR	0.26 B	0.37 B	0.29 B	0.29 B	0.19 UJ	0.41 BJ	0.53 BJ

U= Analyte was not detected at the instrument detection limit given

J= Estimated Value

B= Reported value is between the instrument detection limit and the contract required detection limit

E= Value is estimated due to interference

N= Spiked sample recovery was not within control limits

* = Duplicate analysis was not within control limits

S= Determined by Method of Standard Addition (MSA)

W= Post digestion spike for furnace AA analysis out of control limits, while sample absorbance is less than 50% of spike absorbance

+ = Correlation coefficient for the MSA is less than 0.995

M= Duplicate injection precision criteria was not met

R= Rejected during data validation

TABLE 3 - 10
INORGANIC RESULTS - SOILS/SEDIMENTS
CHEMICAL INSECTICIDE CORPORATION SITE
OFF-SITE INVESTIGATION
EDISON, NEW JERSEY

	41	42	43	44	45	46	47	48
SAMPLE NUMBER						SJ-L2	SM-L1	SL-U3
SAMPLE LOCATION NO.						TRANSECT J	TRANSECT M	TRANSECT L
CLP SAMPLE CODE	MBNQ31	MBNQ47	MBNQ48	MBNQ49	MBNQ50	MBNQ08	MBNQ10	MBNQ11
DEPTH INTERVAL	0.2 TO 0.7 ft.	FIELD BLANK	FIELD BLANK	FIELD BLANK	FIELD BLANK	2.0 TO 2.5 ft.	2.5 TO 3.0 ft.	0.5 TO 1.0 ft.
UNITS		mg/kg					mg/kg	mg/kg
DATE SAMPLE COLLECTED			3/24/94			3/18/94	3/17/94	3/17/94
ALUMINUM	11300 J	33.0 U	33.0 U	33.0 U	33.0 U	9560	15600	6190
ANTIMONY	4.4 UNJ	28.0 U	28.0 U	28.0 U	28.0 U	7.3 UNJ	25 NJ	7.5 UNJ
ARSENIC	3.5 NJ	1.0 U	1.0 U	1.0 U	1.0 U	85.6 NJ	96.3 NJ	130 NJ
BARIUM	42.7 J	1.0 B	1.0 U	1.0 U	1.0 U	45.6 B	157	130
BERYLLIUM	0.19 BJ	1.0 U	1.0 U	1.0 U	1.0 U	0.63 B	2.2 J	0.69 B
CADMIUM	0.32 UJ	2.0 U	2.0 U	2.0 U	2.0 U	0.70 U	1.1 B	0.72 U
CALCIUM	1420 J	75.7 B	42.5 B	54.0 B	37.0 U	475 B	612 B	2700
CHROMIUM	15.0 J	3.0 U	3.0 U	3.0 U	3.0 U	22.6 N*J	21.2 N*J	49.8 N*J
COBALT	5.0 BJ	11.0 U	11.0 U	11.0 U	11.0 U	14.1	11.1 B	9.0 B
COPPER	14.9 J	4.0 U	4.0 U	4.0 U	4.0 U	68.7	72.7	72.0
IRON	16200 J	24.1 B	11.0 B	146	10.0 U	23100 *	16900 *	28000 *
LEAD	20.4 *J	2.0 U	2.0 U	2.0 U	2.0 U	244 *	341 *	149 *
MAGNESIUM	1440 J	33.0 U	33.0 U	33.0 U	33.0 U	1490	1470 B	2160
MANGANESE	165 J	1.0 U	1.0 B	1.0 B	1.0 B	286 N*J	75.2 N*J	337 N*J
MERCURY	0.11 BJ	0.10 U	0.10 U	0.10 U	0.10 U	0.13 U	0.26	0.13 U
NICKEL	11.4 J	10.0 U	10.0 U	10.0 U	10.0 U	13.3	24.1	25.0
POTASSIUM	937 J	64.0 U	64.0 U	64.0 U	72.1 B	991 B	1040 B	815 B
SELENIUM	0.27 BJ	1.0 UW	1.0 U	1.0 U	1.0 U	0.62 B	1.9	0.32 U
SILVER	0.32 BJ	2.0 U	2.0 U	2.0 U	2.0 U	1.3 B	1.0 U	2.0 B
SODIUM	97.9 BJ	157 B	121 B	131 B	122 B	55.4 B	71.5 B	154 B
THALLIUM	0.45 UWJ	2.0 UW	2.0 UW	2.0 UW	2.0 U	0.69 B	0.67 BWJ	0.27 U
VANADIUM	25.7 J	2.0 U	2.0 U	2.0 U	2.0 U	52.9 *J	41.2 *J	22.0 *J
ZINC	38.6 J	13.1 B	8.4 B	10.3 B	8.4 B	58.2 NJ	290 NJ	650 NJ
CYANIDE	0.48 BJ	1.5 U	1.5 UJ	1.5 U	1.5 U	0.65 UN	0.93 UN	0.66 UN

U= Analyte was not detected at the instrument detection limit given

J= Estimated Value

B= Reported value is between the instrument detection limit and the contract required detection limit

E= Value is estimated due to interference

N= Spiked sample recovery was not within control limits

*= Duplicate analysis was not within control limits

S= Determined by Method of Standard Addition (MSA)

W= Post digestion spike for furnace AA analysis out of control limits, while sample absorbance is less than 50% of spike absorbance

+ = Correlation coefficient for the MSA is less than 0.995

M= Duplicate injection precision criteria was not met

R= Rejected during data validation

TABLE 3 - 10
INORGANIC RESULTS - SOILS/SEDIMENTS
CHEMICAL INSECTICIDE CORPORATION SITE
OFF-SITE INVESTIGATION
EDISON, NEW JERSEY

	49	50	51
SAMPLE NUMBER	SSO-U1	SQ-L1	SR-L2
SAMPLE LOCATION NO.	TRANSECT O	TRANSECT Q	TRANSECT R
CLP SAMPLE CODE	MBNQ14	MBNQ15	MBNQ16
DEPTH INTERVAL	0.3 TO 1.2 ft.	2.0 TO 2.5 ft.	2.5-3.0 ft.
UNITS	mg/kg	mg/kg	mg/kg
DATE SAMPLE COLLECTED	3/21/94	3/17/94	3/16/94
ALUMINUM	16400	8300	9600
ANTIMONY	6.8 UNJ	7.1 UNJ	7.4 UNJ
ARSENIC	5.6 SNJ	25.4 NJ	7.0 SNJ
BARIUM	60.5	38.3 B	32.1 B
BERYLLIUM	1.2 J	0.86 B	1.1 B
CADMIUM	0.63 U	0.68 U	0.71 U
CALCIUM	1260	3280	1770
CHROMIUM	41.0 N*J	21.7 N*J	15.8 N*J
COBALT	13.3	8.4 B	19.5
COPPER	24.9	32.2	10.5
IRON	39400 *	26800 *	22900 *
LEAD	37.8 S	47.8 S	26.7 S
MAGNESIUM	1680	2590	2830
MANGANESE	335 N*J	194 N*J	962 N*J
MERCURY	0.12 U	0.13 U	0.13 U
NICKEL	30.2	15.8	44.8
POTASSIUM	1320	1610	740 B
SELENIUM	0.28 U	0.30 U	0.32 U
SILVER	2.0 B	0.70 U	0.74 U
SODIUM	505 B	56.2 B	172 B
THALLIUM	0.37 B	0.50 BWJ	0.50 B
VANADIUM	39.3 *J	21.3 *J	12.8 B*J
ZINC	62.8 NJ	88.2 NJ	179 NJ
CYANIDE	0.58 UN	0.63 UN	0.66 UN

U= Analyte was not detected at the instrument detection limit given

J= Estimated Value

B= Reported value is between the instrument detection limit and the contract required detection limit

E= Value is estimated due to interference

N= Spiked sample recovery was not within control limits

* = Duplicate analysis was not within control limits

S= Determined by Method of Standard Addition (MSA)

W= Post digestion spike for furnace AA analysis out of control limits, while sample absorbance is less than 50% of spike absorbance

±= Correlation coefficient for the MSA is less than 0.995

M= Duplicate injection precision criteria was not met

R= Rejected during data validation

APPENDIX III

Administrative Record Index

ADMINISTRATIVE RECORD INDEX

DOCUMENT	DATE
1. Field Investigation Data	October 20, 1993 July 11, 1994
2. Draft Proposed Plan Comments, EPA's Environmental Impacts Branch	October 20, 1994
3. Draft Proposed Plan Comments, EPA's Air Programs Branch	October 21, 1994
4. Draft Proposed Plan Comments, EPA's Ground Water Management Division	October 27, 1994
5. Draft Proposed Plan Comments, EPA's Biological Technical Assistance Group	October 31, 1994
6. Draft Proposed Plan Comments, New Jersey Department of Environmental Protection (NJDEP)	November 2, 1994
7. Feasibility Study Memo	November 1994
8. Proposed Plan	November 1994
9. Proposed Plan Comments, Edison Glen Resident	November 22, 1994
10. Public Meeting Transcript	November 28, 1994
11. Proposed Plan Comments, Edison Wetlands Association/Technical Assistance Grant Committee	December 5, 1994
12. Proposed Plan Comments, Attorney Representing Property Owner	December 6, 1994
13. Proposed Plan Comments, Edison Department of Health and Human Resources	December 6, 1994
14. EPA Memo Requesting Internal Review of Draft Record of Decision (ROD)	January 13, 1995

[illegible]

APPENDIX IV

State Letter of Concurrence



State of New Jersey

Christine Todd Whitman
Governor

Department of Environmental Protection

Robert C. Shinn, Jr.
Commissioner

March 28, 1995

Mr. William Muszynski
Deputy Regional Administrator
USEPA - Region II
290 Broadway - Floor 19
New York, NY 10007-1866

Dear Mr. Muszynski:

The Department of Environmental Protection has evaluated and concurs with the Chemical Insecticide Corporation (CIC) Superfund Record of Decision (ROD) (see attached ROD dated March 9, 1995) which addresses contaminated soil and sediments located in residential and recreational areas.

The Department is aware that this ROD represents the second of three phases for the site. The first phase, which was implemented in September 1994, addressed contaminated runoff leaving the site. The third and final phase is expected to address on-site contaminated soil and associated groundwater contamination.

The specific components of the selected remedy for the second phase of the remediation as outlined in the ROD include the following:

- excavation of approximately 10,000 cubic yards of soil and sediment containing arsenic at levels greater than 20 parts per million;
- appropriate off-site disposal of contaminated soil and sediment; and
- restoration of the excavated areas to the extent practicable.

The State of New Jersey appreciates the opportunity to participate in the decision making process and looks forward to future cooperation with the USEPA.

Sincerely,

Robert C. Shinn, Jr.
Commissioner

attachment: CIC ROD

APPENDIX V

Risk Assessment Documents

Human Health Risk Assessment Summary Unnamed Creek and Mill Brook

Chemical Insecticide Corporation



Edison, New Jersey

EPA
Region 2

June 29, 1994

SUMMARY OF SITE RISKS

A risk assessment was conducted to estimate the human health risks associated with potential exposures to arsenic detected in the soils and sediments of the Unnamed Creek and Mill Brook downstream of the Chemical Insecticide Corporation site (CIC). The risk assessment estimated the human health risks which could result from the contamination if no remedial action is taken in the future.

Human Health Risk Assessment

A four-step process was utilized for assessing health risks according to a reasonable maximum exposure scenario: *Hazard Identification*--identifies the contaminants of concern and estimates the environmental concentrations of the contaminants. Previous studies determined that arsenic was the primary contaminant of concern in the soils and sediments of the Unnamed Creek and Mill Brook downstream of the CIC site. *Exposure Assessment*--estimates the magnitude of actual and/or potential human exposures, the frequency and duration of these exposures, and the pathways (e.g., ingesting contaminated creek sediments) by which humans are potentially exposed. *Toxicity Assessment*--determines the types of adverse health effects associated with chemical exposures, and the relationship between magnitude of exposure (dose) and severity of adverse effects (response). *Risk Characterization*--summarizes and combines outputs of the exposure and toxicity assessments to provide a quantitative assessment of site-related risks.

Previous studies at the CIC site determined that arsenic was the primary contaminant of concern in the soils and sediments of the Unnamed Creek and Mill Brook downstream of the site. The risk assessment began with a determination of a representative concentration of arsenic in the soils and sediments of the Unnamed Creek and Mill Brook. The concentrations of arsenic detected in the soils and sediments of the Unnamed Creek and Mill Brook ranged from <1 to 1100 ppm. The majority of the detected concentrations were below 20 ppm. The mean concentration was 43.5 ppm, and a conservative estimate of the mean used in the risk assessment was 49 ppm.

The risk assessment was conducted using a reasonable maximum exposure (RME) scenario. The RME scenario is intended to focus on the maximum plausible exposures to contamination at the Unnamed Creek and Mill Brook. By definition, this is a highly conservative estimate of exposure, which is likely to overestimate the health risks related to the Unnamed Creek and Mill Brook. This risk assessment identified adolescents playing at the Creek or the Brook as the most sensitive potential receptors to the contamination. Adolescents (ages 7-18) were assumed to play at the Unnamed Creek or Mill Brook once a week throughout the year, for 12 years. Exposures were assumed to occur primarily through incidental ingestion of soils and sediments contaminated with arsenic.

The dose response assessment utilized the latest information of the toxicity of arsenic from EPA's

Integrated Risk Information System (IRIS). Under current EPA guidelines, the likelihood of carcinogenic (cancer-causing) and noncarcinogenic effects due to exposure to site chemicals are considered separately. Noncarcinogenic risks were assessed using a hazard index (HI) approach, based on a comparison of expected contaminant intakes and safe levels of intake (Reference Doses). Reference doses (RfDs) have been developed by EPA for indicating the potential for adverse health effects. RfDs, which are expressed in units of milligrams/kilogram-day (mg/kg-day), are estimates of daily exposure levels for humans which are thought to be safe over a lifetime (including sensitive individuals). Estimated intakes of chemicals from environmental media (e.g., the amount of a chemical ingested from contaminated sediments) are compared to the RfD to derive the hazard quotient for the contaminant in the particular medium. The HI is obtained by adding the hazard quotients for all compounds across all media that impact a particular receptor population.

An HI greater than 1.0 indicates that the potential exists for noncarcinogenic health effects to occur as a result of site-related exposures. The reference dose for arsenic is 3×10^{-4} . The HI for noncarcinogenic effects from ingestion of arsenic in soils and sediments of the Unnamed Creek and Mill Brook (using the reasonable maximum exposure for adolescents) is 0.05, therefore, noncarcinogenic effects are highly unlikely to occur from the exposure scenario evaluated in the Risk Assessment.

Potential carcinogenic risks were evaluated using the cancer slope factor developed by EPA for arsenic. Cancer slope factors (SFs) have been developed by EPA's Carcinogenic Risk Assessment Verification Endeavor (CRAVE) for estimating excess lifetime cancer risks associated with exposure to potentially carcinogenic chemicals. SFs, which are expressed in units of $(\text{mg/kg-day})^{-1}$, are multiplied by the estimated intake of a potential carcinogen, in mg/kg-day, to generate an upper-bound estimate of the excess lifetime cancer risk associated with exposure to the compound at that intake level. The term "upper bound" reflects the conservative estimate of the risks calculated from the SF. Use of this approach makes the underestimation of the risk highly unlikely. EPA has classified arsenic as a known human carcinogen, and the SF for arsenic is $1.75 (\text{mg/kg-d})^{-1}$.

For known or suspected carcinogens, EPA considers excess upper-bound individual lifetime cancer risks of between 10^{-4} and 10^{-6} to be acceptable. This level indicates that an individual has not greater than a one in ten thousand to one in a million chance of developing cancer as a result of site-related exposure to a carcinogen under the specific exposure conditions at the site. The excess cancer risk for an adolescent exposed to arsenic in the soils and sediments of the Unnamed Creek and Mill Brook (using the reasonable maximum exposure scenario) is 5×10^{-6} , which is well within EPA's acceptable risk range.

CONCLUSIONS

The results of the risk assessment indicate that the health risks associated with contact arsenic in the soils and sediments of the Unnamed Creek and Mill Brook are well within EPA's acceptable risk range. The risk assessment utilized the reasonable maximum exposure scenario, which is likely to overestimate the health risks associated with the potential exposures to arsenic in the Unnamed Creek and Mill Brook.

Memorandum

DATE: November 3, 1994

SUBJECT: Arsenic Risk at Edison Glen Development

FROM: Mark Maddaloni *mm*

TO: Pat Evangelista

The remediation of soils in the dumpster area of the Edison Glen Housing Development is premised on the results of the risk assessment that was performed on the sampling data in that area.

As is indicated in Table 1, the excess lifetime cancer risk under a residential land-use scenario is $2.0 \text{ E-}04$. The corresponding Hazard Quotient (for a child receptor) is 3.2.

cc: Vince Pitruzello
janet Feldstein

CHEMICAL:

ARSENIC

Sample Number

Concentration
(mg/kg)

Q

Log of
Concentration
(mg/kg)

Frequency: 10/10

Average: 29.99

Average log: 1.90

UCL: 74.7

Maximum: 259

Sample Std. Dev. 80.47

Sample Std. Dev. 1.29

(log value)

1

4.3

1.5

2

4.3

1.5

3

3.9

1.4

4

4.5

1.5

5

6.1

1.0

6

5.2

1.6

7

259

5.6

8

3.2

1.2

9

4.3

1.5

10

5.1

1.0

11/2/04
59 IN-RES-CIC.XLS

TABLE 1

SOIL INGESTION PATHWAY
CHEMICAL INSECTICIDE CORPORATION
RISKS TO RESIDENTS

CARCINOGENS - REASONABLE MAXIMUM CASE SURFACE SOIL INGESTION EXPOSURE: Adults

Chronic Daily Intake= (mg/kg-day)	Soil Concentration	X	Ingestion Rate	X	Conversion Factor	X	Fraction Ingested	X	Exposure Frequency	X	Exposure Duration	X	$\frac{1}{\text{Body Weight}}$	X	$\frac{1}{\text{Averaging Time}}$						
	mg/kg	X	100 mg/day	X	$\frac{1 \text{ kg}}{1000000 \text{ mg}}$	X	1 (unitless)	X	350 days/year	X	24 years	X	$\frac{1}{70 \text{ kg}}$	X	$\frac{1}{25550 \text{ days}}$						
Chemicals	Soil Concentration		Ingestion Rate		Conversion Factor		Fraction Ingested		Exposure Frequency		Exposure Duration		Body Weight		Averaging Time		Chronic Daily Intake (CDI)		Slope Factor (SF)		RISK = (CDI*SF)
Arsenic	7.47E+01		100		1.0E-06		1		350		24		70		25550		3.5E-05		1.75E+00		6.1E-05

TOTAL RISK = 6.1E-05

CARCINOGENS - REASONABLE MAXIMUM CASE SURFACE SOIL INGESTION EXPOSURE: Children (0 - 6 years)

Chronic Daily Intake= (mg/kg-day)	Soil Concentration	X	Ingestion Rate	X	Conversion Factor	X	Fraction Ingested	X	Exposure Frequency	X	Exposure Duration	X	$\frac{1}{\text{Body Weight}}$	X	$\frac{1}{\text{Averaging Time}}$						
	mg/kg	X	200 mg/day	X	$\frac{1 \text{ kg}}{1000000 \text{ mg}}$	X	1 (unitless)	X	350 days/year	X	6 years	X	$\frac{1}{15 \text{ kg}}$	X	$\frac{1}{25550 \text{ days}}$						
Chemicals	Soil Concentration		Ingestion Rate		Conversion Factor		Fraction Ingested		Exposure Frequency		Exposure Duration		Body Weight		Averaging Time		Chronic Daily Intake (CDI)		Slope Factor (SF)		RISK = (CDI*SF)
Arsenic	7.47E+01		200		1.0E-06		1		350		6		15		25550		0.2E-05		1.75E+00		1.4E-04

TOTAL RISK = 1.4E-04

30-YEAR COMBINED RISK (ADULT + CHILD) = 2.0E-04

TABLE I

SOIL INGESTION PATHWAY
 CHEMICAL INSECTICIDE CORPORATION
 RISKS TO RESIDENTS

NONCARCINOGENS - REASONABLE MAXIMUM CASE SURFACE SOIL INGESTION EXPOSURE: Adults

Chronic Daily Intake= (mg/kg-day)	Soil Concentration	X	Ingestion Rate	X	Conversion Factor	X	Fraction Ingested	X	Exposure Frequency	X	Exposure Duration	X	$\frac{1}{\text{Body Weight}}$	X	$\frac{1}{\text{Averaging Time}}$			
	mg/kg	X	100 mg/day	X	$\frac{1 \text{ kg}}{1000000 \text{ mg}}$	X	1 (unitless)	X	350 days/year	X	24 years	X	$\frac{1}{70 \text{ kg}}$	X	$\frac{1}{8760 \text{ days}}$			
Chemicals	Soil Concentration		Ingestion Rate		Conversion Factor		Fraction Ingested		Exposure Frequency		Exposure Duration		Body Weight		Averaging Time	Chronic Daily Intake (CDI)	Reference Dose (RfD)	HQ= CDI/RfD
Arsenic	7.47E+01		100		1.0E-06		1		350		24		70		8760	1.0E-04	3.0E-04	3.4E-01

HAZARD INDEX = 3.4E-01

NONCARCINOGENS - REASONABLE MAXIMUM CASE SURFACE SOIL INGESTION EXPOSURE: Children (0-6 years)

Chronic Daily Intake= (mg/kg-day)	Soil Concentration	X	Ingestion Rate	X	Conversion Factor	X	Fraction Ingested	X	Exposure Frequency	X	Exposure Duration	X	$\frac{1}{\text{Body Weight}}$	X	$\frac{1}{\text{Averaging Time}}$			
	mg/kg	X	200 mg/day	X	$\frac{1 \text{ kg}}{1000000 \text{ mg}}$	X	1 (unitless)	X	350 days/year	X	6 years	X	$\frac{1}{15 \text{ kg}}$	X	$\frac{1}{2190 \text{ days}}$			
Chemicals	Soil Concentration		Ingestion Rate		Conversion Factor		Fraction Ingested		Exposure Frequency		Exposure Duration		Body Weight		Averaging Time	Chronic Daily Intake (CDI)	Reference Dose (RfD)	HQ= CDI/RfD
Arsenic	7.47E+01		200		1.0E-06		1		350		6		15		2190	9.6E-04	3.0E-04	3.2E+00

HAZARD INDEX = 3.2E+00

Lockheed ESAT Technical Review
ESAT Site-Specific Follow-up ERA for Chemical Insecticide Corp.

The following are the quick turn-around results of the ESAT desktop ecological risk assessment (ERA) using methodologies derived by ESAT, and data from Eisler, McVey, Persaud, and the "Draft Off-site Investigation Report." This report was dated June 24, 1994, and prepared by Roy F. Weston, Incorporated, for the Chemical Insecticide Corporation (CIC) site in Edison Township, Middlesex County, New Jersey.

Using a chronic toxicity benchmark for arsenic, the potential for risk to small mammals, represented by the deer mouse, is present from the CIC off-site stream area surface soils at maximum (680 mg/kg) and mean (50.3 mg/kg) soil concentrations of arsenic. The maximum and mean levels generate hazard indices (HIs) of 27.4 and 2.0, respectively. No chronic risk is anticipated at the median (14.2 mg/kg) concentration (HI = 0.6).

No acute toxicity is anticipated to the mouse at even the maximum exposure concentration (28.8 mg/kg/day), but exposure at this concentration does exceed the LD-0, or no effect, level of 10.4 mg/kg/day (Eisler). Therefore, using the LD-0 indicates that there is the potential for an acute effect to begin in the mouse population at soil concentrations exceeding 245 mg/kg. Only six in over 115 surface soil samples exceed this concentration, and only two more approach it. Therefore, acute effects to small mammals are unlikely, as the receptor would not likely forage exclusively at those locations.

To further support this assessment, acute risk was assessed for the eastern cottontail. No chronic risk was assessed because an appropriate benchmark could not be readily located. The potential for acute risk was found based on the maximum soil concentration (HI = 1.4). No potential for acute risk was found at the mean concentration (HI = 0.1). (Due to the low HI for the mean, the median level was not assessed.) Extrapolating from the maximum surface soil concentration and its corresponding HI, the potential for acute effects can be anticipated at surface soil levels exceeding 485 mg/kg. As only two surface soil levels exceed this concentration, and no others approach it, no acute effects are anticipated.

Calculating an effect level that would be anticipated based on the chronic exposure scenario for the mouse yields a soil level at which there is a potential for risk at 24.82 mg/kg, or approximately 25 mg/kg. Based strictly on this site-specific effect level, the soil invertebrate and plant effect levels referenced in Eisler (chronic microbiota effects beginning at 375 mg/kg; acute earthworm effects at 150 to 165 mg/kg; reduced plant productivity at 25 to 85 mg/kg), and the USEPA approved action level (20 mg/kg, CIC report, page 2-4), it would appear to be appropriate to conduct extensive removal activities in the off-site stream area. Restoration would then return a low value habitat that may improve over several decades. However, this area

currently exist as a mature habitat corridor in an otherwise heavily developed area. This increases the habitat value of this area, indicating that destruction of this habitat should be avoided. The increased value also increases the attraction of receptors into this potentially hazardous habitat. Therefore, a balance between the potential risk and the value of the habitat must be found.

It may be possible to affect a removal of the most grossly contaminated surface soils, yet mitigate the impacts by minimizing the areas to be removed. Using the graphical representation of the contaminant levels present in Figure 3-8, it is possible to group several sampling transects that contain the majority of the contamination. Removal of these contaminated soils should eliminate much of the potential risk, while attempting to minimize the elimination of mature habitat. Sediments from Transects A and B and surface soil from Transect A should be removed due to their heavy contamination and low habitat value. These transects will not be considered further in the data for this discussion. Transects I through N and Transects V through AA (both six transects, each approximately 700 to 900 feet in length) are recommended for removal. Removal of these twelve of twenty-nine transects (41%) eliminates significant contamination. The maximum, mean, and median soil concentrations in Transects I through N, Transects V through AA, and all remaining transects (excluding A and B) are:

Transects:	<u>I - N</u>	<u>V - AA</u>	<u>All remaining</u>
Maximum	680.0 mg/kg	595.0 mg/kg	224.0 mg/kg
Mean	104.5	107.6	18.2
Median	39.2	55.1	10.5

This would eliminate twenty-eight of forty surface soil locations (70%) containing levels exceeding 25 mg/kg. Extending the removal of Transects I through N to include P (containing the 224 mg/kg location) would reduce the maximum concentration of the remaining transects to 88.4 mg/kg, and reduce the mean to 15.0.

Removing surface and deep sediments exceeding the Ontario Guidelines (Persaud) from Transects I, K, and N would remove two of the four elevated surface sediment locations, as well as two of the five deep locations. Leaving some of the adjoining excavated soil areas unfilled may mitigate the threat of the transport of remaining elevated surface sediments, and reduce the potential for storm events to scour, expose, and then transport remaining elevated deep sediments. These small restorations of the flood plain may help to restore wetland areas that allow for flood storage. Thus, depending on the extent of excavation, this may help reduce the energy in the stream flow during heavy storms.

We hope these comments have been helpful. If you have any questions or comments, please feel free to contact us at (908) 417-2238.

REFERENCES

Eisler, R. 1988. Arsenic hazards to fish, wildlife, and invertebrates: a synoptic review. United States Fish Wildlife Service. Biological Report 85(1.12).

McVey et al. 1993. Wildlife Exposure Factors Handbook. United States Environmental Protection Agency. EPA/600/R-93/187a.

Persaud et al. 1992. Ontario Sediment Quality Guidelines.

APPENDIX VI

Responsiveness Summary

RESPONSIVENESS SUMMARY

CHEMICAL INSECTICIDE CORPORATION SITE

REMEDY FOR OFF-SITE AREAS

This community relations responsiveness summary is divided into the following sections:

Overview: This section discusses the U.S. Environmental Protection Agency's (EPA's) preferred alternative for remedial action.

Background: This section briefly describes community relations activities related to remediation of contaminated soil and sediment in off-site areas associated with the Chemical Insecticide Corporation (CIC) site.

Summary of Comments: This section provides a summary of commentors' major issues and concerns, and expressly acknowledges and responds to all significant comments raised by the local community. The local community includes residents, businesses, the municipality, public officials, and the Technical Assistance Grant Committee and its consultant.

OVERVIEW

At the initiation of the public comment period on November 7, 1994, EPA presented its preferred alternative for addressing contaminated soil and sediment in residential areas and areas in and immediately adjacent to the unnamed tributary and Mill Brook associated with the CIC site, located in Edison Township, New Jersey.

The selected remedy includes excavation and off-site disposal of contaminated soil and sediment within particular areas along the unnamed tributary and Mill Brook. In addition, contaminated soil in a grassy area behind Building 14 of the Edison Glen Condominium Complex will also be excavated for off-site disposal.

Approximately 10,000 cubic yards of soil and sediment will be removed and the remediated areas will be appropriately restored. This approach enables EPA to restore contaminated areas such that any long-term risk associated with these areas is removed and no property use restrictions will be required. By targeting specific contaminated areas in and near the unnamed tributary and Mill Brook, EPA is able to remove a large majority of the CIC

contamination while achieving a reasonable and acceptable balance in preserving a majority of the valuable ecology existing in these areas.

COMMUNITY RELATIONS BACKGROUND

The Proposed Plan and supporting documentation for this second cleanup phase associated with the CIC site were released to the public for comment on November 7, 1994. These documents were made available to the public in the administrative record repositories maintained at the EPA Region II office (formerly, 26 Federal Plaza and currently 290 Broadway, New York, New York), the Edison Township Municipal Complex (100 Municipal Boulevard, Edison, New Jersey), the Edison Library (340 Plainfield Avenue, Edison, New Jersey), and the Metuchen Library (480 Middlesex Avenue, Metuchen, New Jersey). A notice of availability for these documents was published in The Star-Ledger on November 7, 1994. A public comment period involving the documents was held from November 7, 1994 to December 7, 1994. In addition, a public meeting was held on November 28, 1994, at the Edison Township Municipal Building. At this meeting, representatives from EPA answered questions about the site and the remedial alternatives under consideration. Responses to the comments received during the comment period and at the public meeting are provided in this Responsiveness Summary.

COMPREHENSIVE SUMMARY AND RESPONSES TO SIGNIFICANT COMMENTS

This section provides a comprehensive response to all significant questions and comments raised by the local community during the public meeting and received during the public comment period.

A summary of these questions/comments and EPA's responses to them is provided as follows:

1. **A representative of the Edison Wetlands Association and the Technical Assistance Grant (TAG) Committee's consultant requested that EPA sample and analyze soil and sediment further downstream in the Mill Brook, south of the confluence of Mill Brook and the New Jersey Turnpike.**

EPA Response: EPA agrees that additional sampling further downstream is necessary to more accurately determine the extent of contamination associated with the CIC site. EPA plans to perform such additional sampling as part of the remediation process for the areas in and around the unnamed tributary and Mill Brook. This sampling will be conducted during the first phase of remediation activities. If necessary, additional areas will be designated for remediation.

2. **A representative of the Edison Wetlands Association requested that EPA provide access to the CIC site to perform a dye test to more clearly define the surface water drainage pathways leading from the site.**

EPA Response: EPA understands that the Edison Department of Health and Human Resources will provide the dye and experienced personnel to perform the dye test. As EPA has already stated at several recent Citizen's Advisory Committee meetings, the Agency is willing to participate in performing the dye test and providing supervised access to the site.

3. **A representative of the Edison Wetlands Association requested that EPA evaluate a specific data point [510 parts per billion (ppb) arsenic] generated by a former owner of a condominium in the Edison Glen Condominium Complex as a result of sampling a puddle in the pool area of the complex.**

EPA Response: EPA has evaluated the information provided by the Edison Wetlands Association representative. Based on this evaluation, EPA believes that an arsenic level of 510 ppb in a puddle in the pool area can be attributed to the naturally occurring levels of arsenic in soil. In addition, EPA believes that risk associated with exposure to this level of arsenic found in the puddled water would be acceptable and similar to that posed by exposure to naturally occurring arsenic levels [up to 20 parts per million (ppm)] in New Jersey soil.

4. **The TAG Committee's consultant and a representative of the Edison Wetlands Association requested that EPA, in consultation with state and local government, the Edison Wetlands Association and other stakeholders, consider appropriate forms of institutional control (in addition to the selected remedy) for the areas in and around the unnamed tributary and Mill Brook.**

EPA Response: EPA plans to use 20 ppm as a guideline in remediating the contaminated soils and sediment in residential areas and areas in and immediately adjacent to the unnamed tributary and Mill Brook. Although this is not a promulgated chemical-specific standard, and therefore not an applicable or relevant and appropriate requirement, it represents the upper limit of naturally occurring arsenic concentrations in New Jersey soils. Use of this guideline allows EPA to remove a large majority of the arsenic contamination while achieving a reasonable and acceptable balance in preserving a majority of the valuable ecology existing in these areas.

Based on EPA's human health and ecological risk evaluation,

the selected remedy is protective of human health and the environment and EPA does not believe that any additional institutional controls are warranted. In addition, because the average residual level of arsenic is expected to be below 20 ppm, the New Jersey Department of Environmental Protection (NJDEP) does not require any land use restrictions or form of institutional controls based on the New Jersey Industrial Site Recovery Act.

5. The TAG Committee's consultant requested that EPA identify the location of sample number SSCG-6, questioned if this location is part of one of the transects designated for cleanup, and, if not, suggested that the area where this sample was obtained be added to the cleanup.

EPA Response: Sample number SSCG-6 designates a sediment sample which was obtained from the most northern point of a Mill Brook tributary running parallel to the southern edge of the Edison Woods residential complex.

Although the concentrations detected at this location for benzo(a)anthracene, benzo(b)fluoranthene and benzo(a)pyrene (2.6 ppm, 3.5 ppm, and 2.2 ppm, respectively) exceed NJDEP's guidelines (as was indicated in the consultant's comment), any risk posed by these contaminants would be insignificant when compared to the risk posed by the primary contaminant of concern, arsenic, and would not change EPA's overall assessment of risk. Using the reasonable maximum exposure scenario described in this Record of Decision (ROD), the carcinogenic human health risk would remain within EPA's acceptable risk range and any adverse non-carcinogenic effects are still not likely to occur.

As is described in this ROD, the ecological risk assessment indicates that adverse ecological effects are likely to occur if no remedial action is taken. However, because the Mill Brook and its tributaries currently exist as a mature habitat corridor in an otherwise heavily developed area, the habitat value of this area is increased and its destruction should be avoided if possible. During the process of selecting areas to be remediated and achieving a balance between mitigating the potential risk to ecological receptors and preserving the value of the area's ecosystem (or minimizing destruction of the habitat through remedial activity), EPA determined that the area where sample number SSCG-6 was obtained does not require remediation.

6. **An attorney representing an owner of property located near or within the area(s) designated for remediation submitted to EPA written objection on behalf of the property owner to any "egress and ingress onto their private property for the purposes of a public cleanup of an adjacent or adjoining property."**

EPA Response: EPA may require future access to this property for the purpose of remediating an adjacent or adjoining property. A decision regarding access to this property will be made during the planning phase of implementation of the selected remedy. Each property owner will be formally notified by EPA in a timely manner in advance of the commencement of any field activities, and appropriate access arrangements will be made.

7. **A representative of the community inquired about the adverse effects of arsenic on the ecology and the food chain.**

EPA Response: As is described in the ROD, the results of the ecological risk assessment indicates that adverse ecological effects primarily from arsenic are likely to occur if no remedial action is taken. The results of soil samples taken in the vicinity of the unnamed tributary and Mill Brook indicate the presence of arsenic at levels sufficient to generate acute risk to soil invertibrates and chronic risk to soil microbiota and small mammals. In addition, surficial stream sediments in some areas along the unnamed tributary and Mill Brook contain sufficient arsenic to pose a risk to the benthic community. Risk to organisms at higher trophic levels via exposure through the food chain was also assessed. It was determined that no risk to such organisms via this pathway of exposure is anticipated.

8. **A representative of Congressman Pallone's office inquired about the process for cleaning the surface water, how far downstream EPA is willing to test and how EPA expects to actually clean up the areas in and around the unnamed tributary and Mill Brook.**

EPA Response: Although the selected remedy does not include active measures to clean the surface water, EPA believes that removal of contaminated soils and sediment will contribute to the improvement of surface water quality in the unnamed tributary and Mill Brook. The surficial cap which was recently installed at the CIC site will also contribute to the improvement of surface water quality by controlling the release of contaminated surface water runoff from the site.

EPA agrees that additional sampling further downstream is necessary to more accurately determine the extent of

contamination associated with the CIC site. A determination as to the extent of downstream sampling will be made based on an evaluation of sampling data. After collection of additional data, the results will be evaluated, along with existing data, to assess potential impacts to human health and the environment.

The cleanup of the unnamed tributary and Mill Brook will include excavation of contaminated soil and sediment and appropriate disposal of such material in a secure off-site landfill. Heavy equipment such as bulldozers, backhoes and dump trucks are expected to be used for the remediation. Temporary access roadways will be constructed to provide vehicular equipment access to areas to be remediated. EPA intends to minimize tree removal during the remedial activities. The Agency will work closely with the community during the remedial design phase to coordinate construction activities.

9. A concerned citizen inquired about the location of the off-site disposal facility, the method for remediating the contaminated soil (if incineration would be the remediation method) and if EPA would be using a particular stone to stabilize the remediated banks of the unnamed tributary and Mill Brook.

EPA Response: The location of the off-site disposal facility has not yet been selected. A secure off-site disposal facility will be identified prior to commencement of excavation activities. The contaminated material will be appropriately landfilled at the selected off-site facility. The material will not be incinerated. Depending on the material's ultimate waste classification prior to landfilling, some pretreatment (i.e., solidification) may be necessary before landfilling. If necessary, this would be performed at the off-site disposal facility.

During the restoration of the remediated areas in and along the unnamed tributary and Mill Brook, EPA will also stabilize the banks of these waterways. Stone may be used as a stabilizing material. The stabilization method(s) will be identified during the planning phase of the remediation.

10. A concerned citizen asked if EPA could repeat surface water sampling in the Spring of 1995 to assure that the cap over the CIC site has eliminated contaminated surface water runoff.

EPA Response: EPA does not believe that sampling the surface water and determining its quality will be indicative that the cap is fully working. Therefore, EPA does not plan to sample surface water to determine the effectiveness of

the cap. EPA believes that maintaining the integrity of the cap will assure that the cap continues to accomplish its intended goal, to control contaminated surface water runoff from the site. EPA will perform periodic site inspections and make any necessary repairs of the interim remedy (the cap and all its other components) to preserve the integrity of the remedy and ensure that rainwater or surface runoff does not contact contaminated materials on-site

11. A concerned citizen inquired about the maximum depth of soil and sediment sampling, the depth of arsenic contamination, and if an arsenic concentration gradient was established with depth of soil/sediment.

EPA Response: During EPA's investigation, soil/sediment samples were obtained at a maximum sample depth of 30-36 inches. Arsenic contamination was observed at the maximum soil depth sampled. Based on an evaluation of the data, EPA could not establish a clear relationship between soil depth and arsenic levels. In some instances, the arsenic levels increased with soil depth and in others, it decreased with depth.

12. A concerned citizen encouraged EPA to minimize disruption of the ecology in the areas of the unnamed tributary and Mill Brook by sampling and analyzing soil as the excavation activities proceed. The citizen recommended that areas which are confirmed clean can be left alone.

EPA Response: EPA agrees. In fact, EPA applies this process of sampling and analyzing soil during an excavation remedy as a standard operating procedure, and will minimize disruption to areas not requiring remediation.

13. A concerned citizen requested that EPA provide a landscaping plan which identifies which trees will be saved during implementation of the remedy and that any wetlands impacted by the remediation be restored.

EPA Response: EPA intends to minimize tree removal during the remedial activities. EPA plans to work closely with the community during the remedial design phase and to coordinate construction activities so that the community is kept fully informed. EPA will identify certain areas requiring tree removal (such as those areas where temporary access roadways will be constructed) prior to commencement of excavation activities. Other areas requiring tree removal will be identified during the excavation activities under EPA supervision.

Wetland areas disrupted by the remedial activities will be appropriately restored.

14. An Edison Councilman questioned why EPA ceased sampling immediately south of the confluence of Mill Brook and the New Jersey Turnpike by the Edison Woods residential complex and whether there is a concern about the potential for contaminated groundwater to continue to migrate and contaminate the waterways targeted for remediation.

EPA Response: EPA's standard operating procedure for performing field investigations is to take a phased sampling approach. An initial plan is developed targeting specific areas to be sampled. Based on the results of the initial sampling program, a determination regarding the need for additional sampling is made. In this case, additional sampling further downstream is necessary to more accurately determine the extent of contamination associated with the CIC site. EPA plans to perform such additional sampling as part of the remediation process for the areas in and around the unnamed tributary and Mill Brook.

Although contaminated groundwater may be discharging into the waterways designated for remediation, EPA does not believe that the groundwater is contaminated at levels which could contaminate soil and sediment to levels of concern. EPA believes that the contamination requiring remediation in these areas has resulted primarily from historical contaminated surface water runoff, discharged from the CIC site over the long period prior to the installation of the cap over the site. EPA further believes that the contribution, if any, of contaminated groundwater to the contamination in the areas to be remediated is insignificant.

15. A concerned citizen inquired about the naturally occurring level of arsenic in soil, the arsenic level which is deemed to be acceptable in groundwater and whether certain forms of arsenic are more harmful than others.

EPA Response: NJDEP has determined the upper limit of naturally occurring arsenic for New Jersey soils to be 20 ppm. The Federal acceptable drinking water level for arsenic is 50 ppb.

The arsenic that occurs naturally may occur in different forms with different toxicities or degrees of potential harm to human health and the environment. During EPA's risk assessment process, the Agency assumes that arsenic is in its most harmful or toxic form, thereby providing an additional level of assurance regarding protection of human and the environment.

16. A representative of the Edison Wetlands association requested access to data which may be generated from resampling the property adjacent to Prince Street, in Edison, New Jersey, during the course of remediation. The representative also asked if a developer sampled the property and found higher arsenic levels, would EPA review the area?

EPA Response: Although EPA does not anticipate further sampling of the subject property at this time, if EPA should generate any future data from resampling the property, the Agency would make it available for review.

If EPA were presented with data which conflicted with data the Agency collected, EPA would investigate and evaluate the situation further.

17. A concerned citizen asked whether the approximately 10% of total soil and sediment samples which were analyzed for other contaminants, including arsenic, were randomly selected and whether any of these samples were taken on Wayne Miller's former property?

EPA Response: These samples and their locations were randomly selected but were obtained from each of the four general areas which were sampled (Edison Glen, Edison Woods, Rodak Circle/Wilshire Road and the unnamed tributary/Mill Brook areas).

During EPA's most recent sampling effort, samples of this kind were obtained on or in the immediate vicinity of the property formerly owned by Wayne Miller. Similar samples have also been obtained from this property during previous EPA investigations.

18. A concerned citizen asked when the remediation activities would begin?

EPA Response: Based on EPA's ability to secure a contractor in a timely fashion, EPA believes that the remediation activities will commence during the Spring of 1995.

ROD FACT SHEET

SITE

Name : Chemical Insecticide Corporation Site
Location/State : Edison Township, Middlesex County, New Jersey
EPA Region : II
HRS Score (date): 47.53 (10/89)
Site ID # : NJD980484653

ROD

Date Signed: March 28, 1995
Remedy/ies: Excavation and Off-site Disposal
Operating Unit Number: OU-2
Capital cost: \$8,583,000 (in 1995 dollars)
Construction Completion: March 1997
O & M in 1995: none
1996: n/a
1997: n/a
1998: n/a
Present worth: n/a

LEAD

EPA Remedial
Primary contact (phone): Pat Evangelista (212) 637-4403
Secondary contact (phone): Janet Feldstein(212) 637-4417
Main PRP(s): Arnold M. Livingston
PRP Contact (phone): n/a

WASTE

Type: predominantly arsenic
Medium: soil and sediment
Origin: pesticides manufacturing
Est. quantity: 10,000 cu.yd.