Journal of Hazardous Substance Research

Volume 2

Article 6

1-1-1999

Environmental Site Characterization and Risk-Based Evaluation of a Site Contaminated with Tetrachloroethene (PCE) and Trichloroethene (TCE)

C. L. Quast Howard R. Green Company

M. G. Fisher Howard R. Green Company

A. H. Broughton Howard R. Green Company

Follow this and additional works at: https://newprairiepress.org/jhsr

Recommended Citation

Quast, C. L.; Fisher, M. G.; and Broughton, A. H. (1999) "Environmental Site Characterization and Risk-Based Evaluation of a Site Contaminated with Tetrachloroethene (PCE) and Trichloroethene (TCE)," *Journal of Hazardous Substance Research*: Vol. 2. https://doi.org/10.4148/1090-7025.1016

This Article is brought to you for free and open access by New Prairie Press. It has been accepted for inclusion in Journal of Hazardous Substance Research by an authorized administrator of New Prairie Press. For more information, please contact cads@k-state.edu.

ENVIRONMENTAL SITE CHARACTERIZATION AND RISK-BASED EVALUATION OF A SITE CONTAMI-NATED WITH TETRACHLOROETHENE (PCE) AND TRICHLOROETHENE (TCE)

C.L. Quast, P.E.; M.G. Fisher, R.E.M.; and A.H. Broughton Howard R. Green Company, 4250 Glass Road, PO Box 9009, Cedar Rapids, IA 52409-9009; Phone: (319) 395-7805; Fax: (319) 395-6410.

ABSTRACT

A healthcare company purchased property in eastern Iowa for a facility expansion. Before the purchase, a Phase I environmental site assessment (ESA) revealed that the property was the site of a former dry cleaning business. Phase II sampling and testing indicated that tetrachloroethene (PCE) and trichloroethene (TCE) had affected site soil and groundwater. Maximum concentrations of PCE and TCE in groundwater were 538 and 209 μ g/L, respectively, and 105 and 1.51 mg/kg in soil. Additional sampling delineated the vertical and horizontal extent of contamination in the soil. The concentrations of the chlorinated solvents in both the soil and groundwater were below levels of regulatory concern. However, the company was concerned that the Iowa statewide standard for PCE in soil (780 mg/kg) might not provide adequate protection for several exposure pathways and wanted to assess the risk to the public from the contamination at the site. The results of a receptor survey were used to develop site-specific target levels (SSTLs) for contamination, using accepted human health exposure factors, models, and chemical-specific toxicity values. The recommended remedial options allowed the company to minimize the human health risks posed by contamination at the site. *Key words: chlorinated, risk, dry cleaners, tetrachloroethene, trichloroethene*

INTRODUCTION

This paper presents the results of an environmental site characterization (ESC), consisting of Phase I and Phase II environmental site assessments (ESAs) and additional sampling, and the methods and results of a risk-based evaluation performed on a site impacted by tetrachloroethene (PCE) and trichloroethene (TCE). A healthcare company purchased several parcels of property in the downtown area of an eastern Iowa city. A Phase I ESA performed on the parcels before the purchase concluded that one of the parcels was the former location of a dry cleaning operation. The dry cleaners used a 3000-gallon underground storage tank (UST) for storage of dry cleaning chemicals. A previous owner removed the tank in the early 1980's but did not collect soil samples to evaluate whether the tank leaked.

A Phase II ESA with additional sampling conducted to evaluate the extent of contamination from PCE, TCE, and their degradation compounds, indicated that PCE and TCE were present in soil gas, soil, and groundwater below the site. Field personnel collected a total of 66 soil gas samples, 36 soil samples, and three groundwater samples from 35 locations across the one-half-acre site. The suspected location of the former UST and the dry cleaning chemical piping had the highest levels of contamination.

Groundwater at the site is about 55 feet below ground surface (bgs). An area approximately 120 feet southwest of the site contains a shallower groundwater unit at a depth of 15 to 20 feet, which is suspected to be a perched water table.

The Iowa Department of Natural Resources (IDNR) stated that contamination in groundwater at the site was not at a level of concern and that they would not require additional characterization of groundwater. The IDNR also said that although the site was not enrolled in Iowa's Land Recycling Program (LRP), a voluntary cleanup program, that the statewide standards of 780 mg/kg for PCE and 180 mg/kg for TCE from the LRP would apply for the soil. The statewide standards for soil are based on ingestion and do not take into account the potential for soil leaching to groundwater. Since concentrations of PCE and TCE in soil at the site were below the statewide standards, the IDNR did not require soil remediation. However, to ensure the safety of patients, employees, and their neighbors, the company proceeded to evaluate the potential human health risks posed by contamination at the site.

OBJECTIVES

The objective of this paper is to provide data from an ESC and demonstrate how those data were used to evaluate human health risks from contaminants in soil and groundwater. Three different sets of criteria were used to evaluate health risks at the site. The first set of criteria used the equations, model, and exposure factors from Iowa's UST regulations (IDNR, 1998, Appendix A-6). The second set of criteria also used the equations and model from Iowa's UST regulations but used exposure factors from IDNR's LRP (567 IAC 137.5(4)). These two sets of criteria are site specific. The third set of criteria, the most conservative of the three, was the statewide standards from the LRP regulations and are not site specific. The exposure factors from the UST and LRP regulations are in Table 1.

The UST regulations in Iowa are risk based. The IDNR has a published Tier 1 Lookup Table (IDNR, 1996, p. 10) which identifies the risk-based screening level (RBSL) or the maximum concentration of a petroleum constituent at a site for a specific exposure pathway at which the receptors for that pathway can be considered not at risk. A receptor is an enclosed space (like a basement), conduit, drinking or nondrinking water well, a protected groundwater source, surface water body, or public water supply system which, when impacted by chemicals of concern, may result in exposure to humans and aquatic life, explosive conditions, or other adverse affects on health, safety, or the environment (IDNR, 1998, Appendix H). Should levels of contamination exceed those in the Tier 1 Lookup Table, then modeling is performed to determine the site-specific target level (SSTL) for a chemical. The SSTL is the risk-based target level for a chemical of concern at the source in order to meet the target level at the receptor.

The LRP regulations are for non-petroleum contamination and offer three different approaches to determining cleanup criteria. The first is background standards (567 IAC 137.4) that compare concentrations of contaminants in soil and groundwater at a site with concentrations of contaminants that are naturally occurring or generally present and not related to a readily identifiable release. The second approach is to use the statewide standards (567 IAC 137.5), which represent concentra-

6-2 Volume Two https://newprairiepress.org/jhsr/vol2/iss1/6 DOI: 10.4148/1090-7025.1016

tions of contaminants in soil and groundwater that at normal exposure via ingestion are considered unlikely to pose a threat to human health. The statewide standards can be considered a screening concentration below which risk evaluation is not necessary. The third approach used by the LRP is site-specific standards (567 IAC 137.6) which are derived by applying exposure and risk assumptions applicable to the conditions at a particular site. For the purposes of this ESC, only the second and third approaches are discussed because the site was located in a downtown area near several contaminated sites.

Of the eight compounds identified in groundwater at the site- PCE, TCE, *cis* 1,2dichloroethene (DCE), *trans* 1,2-DCE, chloroform, vinyl chloride, 1,1-DCE and methylene chloride- this risk-based evaluation addressed only PCE and TCE. *Cis* 1,2-DCE, *trans* 1,2-DCE, chloroform, vinyl chloride, and 1,1-DCE were each below the maximum contaminant level (MCL) for these chemicals in drinking water. The reporting limit for methylene chloride was $10 \,\mu$ g/L, and the MCL is $5 \,\mu$ g/L. The evaluation for risk excluded methylene chloride since it was not only below the method detection limit, but was not found in elevated concentrations in the soil.

ENVIRONMENTAL SITE CHARACTERIZATION

Previous investigations

Historical review of the site indicated that it was exclusively in residential, single- and multifamily housing through 1913. By the 1940s, the site contained one residence and a dry cleaning operation. Between 1970 and 1982, the residence was demolished for the rerouting of a street, and between 1982 and 1987, the dry cleaners was demolished and the site paved for use as an automobile sales lot. During paving of the lot, a UST containing dry cleaning chemicals was discovered, emptied, and removed. No evidence existed of soil or groundwater sampling after the removal of the tank.

A Phase IESA first indicated the potential for contamination. The existence of a former dry cleaning facility and the discovery of the former UST that contained dry cleaning chemical constituted a "recognized environmental condition" and an area of risk. The ESA found that dry cleaning activities had occurred at the site as early as 1940 to as late as 1987. Limited soil gas sampling during the Phase II and on-site analysis conducted in the area believed to be the site of the former UST confirmed the presence of PCE and TCE in the soil at the site.

Physical setting

Continuous soil sampling, performed at several borehole locations, documented the geological setting of the site. The site still contained the buried remnants of the basement and foundation of the former dry cleaning facility and possibly other foundations or basements from the former residences. Sand filled some of the former basements. In other areas, the surface of the site consisted of a layer of fill material containing pulverized limestone road material and native soils. Below the fill material,

light brown, highly permeable sand existed with intermittent clay lenses. Refer to Figure 1 for locations of soil borings.

The bedrock at the site is Devonian Wapsipinicon formation. This formation consists of limestone on the top, then dolomite, shale, and clayey limestone, followed by dolomite limestone with dolomite at the base. The Wapsipinicon formation is part of the Silurian-Devonian aquifer. According to bedrock topography maps, bedrock in the area lies between 650 and 700 feet mean sea level (msl), which is approximately 60 to 120 feet bgs. Well logs from the Iowa Geological Survey Bureau (GSB) in Iowa City indicate bedrock in the area varies from 3 feet to 310 feet bgs.

A two-foot-thick silty clay layer at about 10 to 12 feet bgs was found below the sand. Water was not present above this clay unit on the site. However, in a monitoring well (MW-14) installed during a petroleum investigation (Advanced Environmental Services, Inc. 1998) on an adjacent property and located southwest of the site, groundwater was perched above the clay. Groundwater at the site was at 55 feet bgs within the sand unit. Soil borings advanced for the nearby petroleum contamination investigation on an adjacent property encountered bedrock at a range of 15 to 36 feet bgs and groundwater at about 55 feet bgs. The predominant groundwater flow direction, as determined by the petroleum site investigation, was generally to the south. At the site, however, there was no bedrock found within 105 feet of the ground surface.

Soil gas and soil sampling

During the additional sampling, Geoprobe[™] technology was used to draw soil gas samples from various depths at sampling locations most likely to contain chlorinated solvents. Sampling locations were determined based on the approximate location of the removed UST at the former dry cleaning establishment and the results of the Phase II soil gas samples. Analysis of soil gas samples took place on site using a gas chromatograph.

To confirm and correlate data obtained from soil gas sampling and to gather data on the underlying stratigraphy, field personnel collected 36 soil samples from 24 GeoprobeTM and soilboring locations as indicated in Figure 1. Areas both above and below the clay were sampled as well as deeper areas at several locations to determine the level of impact in the soil immediately above and within the water table.

Groundwater sampling

Two groundwater samples were collected during the Geoprobe[™] investigation from Geoprobe[™] location P-6. In addition, three monitor wells placed at two boring locations were used to investigate groundwater conditions at the site. Two wells (MW-6A and MW-31A) screened at shallow depths monitored water potentially located above the clay layer. A third well (MW-6B), placed near well MW-6A, monitored water at a depth of 92.5 to 102.5 feet bgs. Additionally, field personnel collected a groundwater sample from the existing monitoring well (MW-14) located on the southwest side of the block, as well as MW-4, MW-5, and MW-8

installed during the petroleum investigation on the adjacent property. Groundwater samples from MW-6B and MW-14 were analyzed for volatile organic compounds (VOCs) by EPA Method SW 8260B. MW-6A and MW-31A failed to produce water because there was no perched water above the clay layer at those locations.

Sampling results

The PCE results of the 66 soil gas samples collected ranged from non-detect to 19,485 μ g/L. Table 2 presents a summary of results for soil and soil gas between 10 and 25 feet bgs. The highest concentrations of PCE detected in soil gas were in the 12 to 25 bgs interval in the area of P-6. Actual soil sampling and testing in this area confirmed that it contained the highest concentrations of PCE in soil. Soil gas analyses also indicated elevated concentrations in the area of B-28 (2480 μ g/L) and B-31 (2361 μ g/L) at depths of 40 and 50 feet, respectively.

Results from the 36 soil samples collected and analyzed for VOCs using EPA Method 8260B ranged from non-detect in the outer perimeter samples to 105 mg/kg near the former UST location. The highest levels found were in the area of P-6 and P-40, both located near the suspected location of the former UST and piping runs. Elevated concentrations of PCE in the soil were present at a variety of depths but generally above the clay layer in the areas of P-6, MW-31, and P-34. Elevated concentrations were also present in an area suspected to be the dry cleaning facility's basement at P-43.

Laboratory analytical results indicated VOCs other than PCE in several samples. Table 3 contains a summary of the sample locations and compounds detected. Some of these compounds, such as cis 1, 2-dichloroethene (DCE), are suspect breakdown products of PCE and TCE, which are both used as dry cleaning solvents. Others, such as methylene chloride, may indicate other solvent use.

During GeoprobeTM sampling activities, analysis of a groundwater sample collected from a depth of 50 to 55 feet bgs in P-6 indicated groundwater from this depth contained 538 μ g/L PCE.

Two of the newly installed monitoring wells, MW-6A and MW-31A, screened from 10 to 20 feet and from 14 to 24 feet, respectively, never yielded enough water for sampling. Analysis of a sample collected from the third well, MW-6B, located near the suspected area of the UST and screened from 92.5 to 102.5 feet bgs, resulted in a concentration of 90.1 μ g/L of PCE. Monitoring well MW-14, screened at a depth of 15 to 20 feet bgs, had a PCE concentration of 5.4 μ g/L. Other contaminants found in groundwater samples in the area of P-6 included TCE, cis 1, 2-DCE, and chloroform.

RISK-BASED EVALUATION

Three different sets of criteria were used to evaluate the potential human health risk at the site. The first set of criteria used the equations, target risk, target hazard quotient, exposure frequency,

Journal of Hazardous Substance Research

and exposure duration from Iowa's UST regulations (567 IAC 135,IDNR, 1998, Appendix A-6). The second set of criteria also used the equations from these regulations but used exposure risk factors from Iowa's LRP regulations (567 IAC 137.5(3)). These first two sets of criteria are sitespecific. The third set of criteria is the statewide standards from the LRP regulations (567 IAC 137.5(4)) and is not site specific. Table 4 lists toxicity and chemical-specific human health risk factors that were used in the calculations referenced above.

IDENTIFICATION OF RECEPTORS

The identification of actual and potential receptors was the first step in the risk-based evaluation process. The receptor identification process included determining the presence of drinking and nondrinking water wells, protected groundwater sources, plastic (PVC) drinking water lines, enclosed spaces (i.e., basements), and surface water bodies. Table 6 lists the receptors surveyed.

The GSB provided information on drinking water wells within 1000 feet of the site. After hydraulic conductivity testing, a well search was requested from GSB for wells within one-half mile of the site. The search identified eight nondrinking water wells and no drinking water wells within one-half mile (2640 feet) of the site.

Field personnel performed three bail-down tests on the deep well installed at the site, logging the results with a datalogger. The test results, when analyzed by the Bouwer and Rice Method (Duffield, 1996), indicated hydraulic conductivities of 13.24 m/day, 8.75 m/day, and 84.57 m/day. The hydraulic conductivity used for the risk calculations was 13.24 m/day because it was the higher of the two closest tests. The timing of the last two tests (8.75 m/day and 84.57 m/day) was close together and may have affected the results. This hydraulic conductivity established the aquifer underlying the site as being in the IDNR category of a "protected groundwater source" since the hydraulic conductivity exceeded 0.44 m/day.

Plastic (PVC) drinking water lines in contact with contaminated soil or groundwater can provide a pathway for ingestion of contamination as some contaminants can seep through the piping or leaks in the piping into the drinking water. The city water department stated that no plastic drinking water lines were in service near the site.

The potential for vapor accumulation in enclosed spaces was a concern. Sanitary sewers can provide such an enclosed space as well as basements. One sanitary sewer ran along the northwest side of the site. An explosimeter survey alleviated concern that explosive levels of vapors were present in basements and other enclosed spaces near the site. Vapors coming from contaminated soil and groundwater and entering the basement of the future healthcare facility expansion were of concern. No surface water bodies were located near the site or subject to impact from contamination, requiring no further evaluation of this receptor pathway.

For each potential receptor identified, the Tier 2 Guidance, Appendix B-2 from the UST regulations (IDNR, 1998 567 IAC 135), provided the equations to determine the RBSL for

groundwater and soil to protect actual and potential receptors. The RBSL is the maximum contaminant concentration allowed at the point of exposure for a receptor at which the receptor can be classified as not at risk. For PCE and TCE, both classified as noncarcinogens, the RBSL for groundwater ingestion was determined by the following equation (IDNR, 1998, Appendix A-2):

$$RBSL_{w}\left[\frac{mg}{L-H_{2}O}\right] = \frac{THQ * R_{f}D_{o} * BW * AT_{n} * \frac{365days}{years}}{IR_{w} * EF * ED}$$

Where:

RBSL	= risk-based screening level for the contaminant in water (mg/L)
THQ	= target hazard quotient for individual constituents (unitless)
R _f D ₀	= oral chronic reference dose (mg/kg-day)
BW	= body weight (kg)
AT _n	= averaging time of exposure for noncarcinogens
IR	= daily water ingestion rate (L/day)
EF	= exposure frequency (days/year)
ED	= exposure duration (years)

For each receptor identified, the RBSL was calculated twice, once using risk and exposure factors from IAC 567 Chapter 135 and then again using risk and exposure factors from IAC 567 Chapter 137. After each RBSL for groundwater was calculated, the SSTLs at the source (C_s) for groundwater could be calculated from the contaminant transport equation provided in the Tier 2 Guidance (IDNR, 1998, Appendix B-1) using the RBSL as C(x):

$$C(x) = C_s \exp\left(\frac{x}{2\alpha_x} \left[1 - \sqrt{1 + \frac{4\lambda\alpha_x}{u}}\right]\right) erf\left(\frac{S_w}{4\sqrt{\alpha_y x}}\right) erf\left(\frac{S_d}{4\sqrt{\alpha_z x}}\right)$$

Where:

Х	= distance in the x direction downgradient from the source (m)
erf(f)	= error function
C(x)	= chemical concentration in groundwater at $x (ug/L)$
C _s	= source concentration in groundwater (groundwater concentration at x=0) (uglL)
S _w	= width of the source (perpendicular to x) (m)
S _d	= vertical thickness of the source (m)
Ů	= groundwater velocity (pore water velocity); $u=Ki/q_{e}$
K	= hydraulic conductivity (cm/yr)
Ι	= groundwater head gradient (cm/cm)
θ	= effective porosity
λ	= first-order decay coefficient, chemical specific (d^{-1})
$\alpha_{x}, \alpha_{y}, \alpha_{z}$	= dispersivities in the x, y, and z directions, respectively (m)

The same procedure was followed to determine the RBSL and SSTL for the soil pathways.

Groundwater ingestion pathway

In order to more accurately assess the potential impact of the contamination, field personnel collected groundwater samples for VOC analysis from monitoring wells located on neighboring properties, as well as collected a second round of groundwater samples from the Site. Table 5 presents groundwater sampling data summaries. The wells sampled included MW-6B, located at the site; MWs 4 and 5, installed southeast of the site during the earlier petroleum hydrocarbon investigation; and MW-8, installed west of the site during the same petroleum hydrocarbon investigation.

The high concentration of petroleum hydrocarbon in MW-5 necessitated that the laboratory make several dilutions before analysis. This resulted in an elevated reporting limit for PCE. The TCE concentration in MW-5, which was downgradient from the site, was higher than that at the site, potentially due to the proximity of other off-site sources of the chemical. These factors resulted in exclusion of the results from MW-5 from the risk-based evaluation. Investigation of potential off-site sources was outside the scope of this project, although historical evidence indicated that other drycleaners were at one time located in the vicinity.

Of the eight compounds identified in groundwater at the site- PCE, TCE, *cis* 1,2-DCE, *trans* 1,2-DCE, chloroform, vinyl chloride, 1,1-DCE and methylene chloride- this risk-based evaluation addressed only PCE and TCE. *Cis* 1,2-DCE, *trans* 1,2-DCE, chloroform, vinyl chloride, and 1,1-DCE were each below the maximum contaminant level (MCL) for these chemicals in drinking water.

For all groundwater calculations, the source of PCE and TCE was assumed to be P-6. A GeoprobeTM groundwater sample collected at that location yielded a PCE concentration of 538 μ g/L and a TCE concentration of 209 μ g/L.

Drinking water wells. A search by GSB for wells within one-half mile of the site yielded no drinking water wells. Using the transport equations from IDNR's UST regulations and guidance given above, the concentration of PCE at one-half mile from the site was calculated to be $4.3 \mu g/L$, based on the conservative assumption that no decay of PCE was taking place. This was below the statewide standard of $5.0 \mu g/L$. Groundwater flow direction (southerly), plume range (150 degrees), and hydraulic gradient (0.00176 ft/ft) were derived from a Tier 2 Site Cleanup Report prepared on an adjacent property (Advanced Environmental Services, Inc., 1998). Because the source concentration of TCE was less than that of PCE and the statewide standard for TCE was also $5.0 \mu g/L$, it was reasoned that the TCE plume at one-half mile from the site was also less than the statewide standard. Based on the above assumption, risk from exposure through this pathway was found to be minimal when measured against the site-specific requirements of IDNR's UST and LRP regulations, both of which require the receptor concentration not exceed 5.0 ug/L.

Statewide standards, however, as provided by IDNR's LRP regulations, require that the source concentration not exceed the statewide standard, meaning that PCE and TCE would be at unacceptable levels when compared to this standard. Since, however, it can be shown that no drinking water wells are present, site-specific standards can be developed.

Nondrinking water wells. The GSB identified eight nondrinking water wells within a onehalf-mile radius of the site. The well determined to be most at risk is located approximately onethird mile downgradient from the site (Well #1). Site-specific target levels (SSTLs) were calculated for Well #1 in the groundwater source area, based on a calculated RBSL at the well of 70 μ g/L PCE and 42 μ g/L TCE. Table 6 gives the SSTLs required to protect Well #1. The SSTLs for Well #1 were calculated twice, once using the UST exposure factors and once with LRP exposure factors. Both calculations used the UST risk equations given above. The most restrictive SSTLs of 3900 μ g/L PCE and 2100 μ g/L TCE were calculated using the exposure factors from IDNR's LRP regulations and equations from IDNR's UST regulations. The actual concentrations at the source for PCE and TCE were 538 μ g/L and 209 μ g/L, respectively, well below the SSTLs. No risk is indicated for Well #1 based on these calculations.

Calculations for RBSLs based on exposure factors in IDNR's UST regulations would permit up to 360 μ g/L of PCE and 210 μ g/L of TCE at Well #1. This would result in SSTLs of 20,200 μ g/L for PCE and 11,800 μ g/L for TCE at the groundwater source. Based on SSTLs calculated both ways, no risk from exposure through this pathway was found. Table 6 summarizes the SSTLs calculated from each set of exposure factors and the statewide standards.

Protected groundwater source. The site is a protected groundwater source by the IDNR definition. The PCE concentration of $538 \mu g/L$ at the source exceeded the two calculated SSTLs as well as the statewide standard. For calculations based on the IDNR's UST and LRP exposure factors, the SSTL for PCE at the source in protected groundwater was 360 and 70 $\mu g/L$, respectively, without an institutional control in place. An institutional control is a restriction on use or access to a site to eliminate or minimize exposure to contaminants (IDNR, 1998, Appendix H). An example of an institutional control applicable for this pathway would be a deed restriction to prevent the installation of wells on the property.

The TCE concentration at the source, 209 μ g/L, was less than the 210- μ g/L RBSL, based on IDNR's UST exposure factors. The TCE concentration at the source exceeded the RBSL of 42 μ g/L calculated, based on IDNR's LRP exposure factors. The TCE concentration at the source also exceeded the statewide standard, which requires a level less than 5.0 μ g/L at any point in a protected groundwater source. The concentrations of PCE and TCE present in the protected groundwater wells present, the hydraulic conductivity makes it a potential drinking water source.

Soil leaching to groundwater pathway

The concern with the soil leaching to groundwater pathway is that contaminants in the soil will continue to leach to the groundwater, increasing the chance of groundwater exceeding its target level for contamination.

The soil contamination source was assumed to be P-40, which had a PCE concentration of 105 mg/kg and a TCE concentration of 1.51 mg/kg.

Nondrinking water wells. The nearest well, Well #1, could be at risk for soil leaching PCE to groundwater under the most restrictive criteria, which was using the exposure factors from IDNR's LRP regulations (567 IAC 137.5(4)). To eliminate the risk from PCE for this pathway, the concentration in the soil would have to be less than 61.0 mg/kg. TCE in soil is already less than the 12.0 mg/kg required, making the risk to Well #1 negligible from TCE at the site. Using the less restrictive IDNR UST exposure factors (IDNR, 1998, Appendix A-6), PCE cleared this pathway with the SSTL being 314 mg/kg. The statewide standards for PCE and TCE in soil remain at 780 and 180 mg/kg, respectively. This pathway does not pose a risk using statewide standard criteria.

Protected groundwater source. Based on calculations using either the IDNR's UST or LRP exposure factors, the concentrations of PCE and TCE in the soil at the source exceeded the SSTLs calculated for the soil leaching to a protected groundwater source pathway. With an institutional control in place to prevent the installation of wells, the allowable PCE at the soil source was calculated at 1.9 mg/kg, and the allowable TCE was 0.41 mg/kg based on IDNR's LRP exposure factors (567 IAC 137.5(4)). Based on IDNR's UST exposure factors (IDNR, 1998, Appendix A-6), the SSTLs were 9.9 mg/kg and 2.1 mg/kg, respectively. The statewide standard for PCE in soil, as provided in IDNR's LRP, however, is 780 mg/kg, and the statewide standard for TCE is 180 mg/kg. The site was determined to be high risk for this pathway because of the combined risk for soil leaching into already contaminated groundwater.

Groundwater vapor. The groundwater vapor pathway requires evaluation of the potential for soil to leach to groundwater to such an extent that vapors from the groundwater become a health risk by migrating into enclosed spaces. The actual receptors at risk for soil leaching to groundwater vapor were the existing basement of the truck sales building on the site, the future basement of the new healthcare facility, and the sanitary sewer running along the alley to the north of the site. The SSTL for PCE in the soil was 33.5 mg/kg in order to protect the environment of the truck sales building basement using IDNR's LRP exposure factors (567 IAC 137.5(4)). Using the exposure factors from IDNR's UST regulations (IDNR, 1998, Appendix A-6), the allowable PCE in the soil was 160 mg/kg. Since the actual PCE concentration was 105 mg/kg, this pathway presented no risk for PCE using the exposure factors from the UST regulations but was high risk when using the factors from the LRP.

In order to protect the environment of the future facility basement, the SSTL for PCE was 56.6 mg/kg (IDNR's LRP exposure factors) or 270 mg/kg (IDNR's UST exposure factors). Again, the site was cleared of risk for this pathway for PCE using the exposure factors from IDNR's UST regulations but was determined to be high risk when using the factors from the LRP. Based on the location of the sanitary sewer, allowable concentrations of PCE in the soil were even higher still, so this was not calculated. Soil leaching TCE to groundwater vapor did not pose an unacceptable risk based on the low concentration of TCE in site soil.

Groundwater vapor to enclosed space pathway

Groundwater vapor accumulation in enclosed spaces from existing groundwater contaminant levels did not pose a risk to human health when SSTLs were calculated based on IDNR's UST exposure factors (IDNR, 1998, Appendix A-6). The SSTL for PCE in groundwater was 8100 μ g/ L, and for TCE it was 2000 μ g/L, well above the actual concentrations in groundwater. SSTLs for PCE and TCE in groundwater based upon the IDNR's LRP exposure factors (567 IAC 137.5(4)) were 1700 μ g/L and 1500 μ g/L, respectively. They were also well above the actual concentrations for this pathway. This pathway did not pose an unacceptable risk based upon the criteria.

Soil vapor to enclosed space pathway

The existing basement of the truck sales building was at risk for vapors from the soil due to PCE concentrations in the soil. The SSTLs based on exposure factors from IDNR's UST and LRP regulations were 52 mg/kg and 10.8 mg/kg, respectively. The PCE concentration in site soil (105 mg/kg) was above the RBSL. TCE concentrations did not pose a risk for either of the site-specific criteria. Due to its location, the future basement of the new facility was not at risk for exposure to unacceptable vapor concentrations of PCE or TCE from existing contamination.

CONCLUSIONS

These conclusions were based upon models, equations, and exposure factors found in IAC 567 Chapter 135 (IDNR's UST regulations) and IAC 567 Chapter 137 (IDNR's LRP regulations). The calculations were designed to project the potential for risk and do not predict future conditions such as the addition of wells or basements in the vicinity of the site. Chemical-specific parameters used were from public sources (U.S. Department of Energy, 1999).

Groundwater

Based upon site-specific evaluations of risk and using exposure factors from IDNR's UST and LRP regulations, concentrations of PCE and TCE in groundwater did not pose an unacceptable risk of exposure to the actual receptors evaluated, which included eight nondrinking water wells located within a one-half mile radius of the site.

The aquifer underlying the site is a protected groundwater source by the IDNR definition. PCE concentrations exceeded the calculated SSTLs, whether being calculated using IDNR's

Journal of Hazardous Substance Research

Volume Two

UST or LRP exposure factors. TCE concentrations exceeded the more stringent SSTL calculated with the exposure factors in IDNR's LRP regulations but passed using factors from IDNR's UST regulations.

The statewide standard for PCE or TCE in groundwater is $5.0 \,\mu$ g/L and not site specific. As mentioned above, the statewide standard provides a screening level below which risk posed by contaminants is considered to be non-existent and risk evaluation is not required. PCE and TCE concentrations both exceed $5.0 \,\mu$ g/L.

Soil

Concentrations of PCE and TCE in the soil were considered a risk due to the ability of the contaminant to leach into the groundwater, potentially putting the nondrinking water wells and the protected groundwater source at risk in the future. Calculations also indicated that contaminants leaching from the soil could elevate contaminant levels in the groundwater to an extent that vapors from the groundwater could become a problem in enclosed spaces in the vicinity of the truck sales building. These conclusions are from the site-specific criteria. Statewide standards do not address soil leaching.

Vapors moving directly from the soil into enclosed spaces also presented a risk to the sales building basement, according to the criteria used for this evaluation.

The statewide standards for PCE and TCE in soil are 780 mg/kg and 180 mg/kg, respectively. Contaminants are well below these concentrations.

RECOMMENDATIONS

Groundwater

Attaching an environmental easement to the property deed, and possibly on adjoining properties, that would prevent the installation of drinking and nondrinking water wells would provide protection against ingestion of contaminated groundwater.

Although the concentrations of PCE and TCE in the groundwater at the site exceeded statewide standards provided in IDNR's LRP regulations, the IDNR has stated that concentrations do not warrant further investigation of the site due to the depth and complexity of the hydrogeologic system and the potential for commingled plumes from off-site sources.

Soil

Excavation of an area approximately 15 feet square by 15 feet deep, centered on the location of sample P-40, would remove soil inside of the calculated 50-mg/kg soil plume. A level of 50 mg/ kg would clear the soil leaching to groundwater ingestion pathway for the nearest well and for soil leaching to groundwater vapor for the basement of the new hospital facility. In actuality, the concentration of remaining PCE in the soil will probably be much lower than 50 mg/kg, due to the way the model interpolated concentrations between data points. This should adequately protect the future basement from vapors.

After excavation, the only soil pathway which would remain affected by the soil contamination, according to risk calculations, would be the soil leaching to protected groundwater source and soil leaching to groundwater vapor to the existing basement of the truck sales building. With an institutional control in place to eliminate the extraction of groundwater near the site and the planned demolition of the truck sales building, adequate protection would be afforded to the public from contaminated soil at the site.

REFERENCES

- Advanced Environmental Services, Inc., 1998 Tier 2 Site Cleanup Report for Allen Motor Company.
- Duffield, Glenn M., HydroSOLVE, Inc, 1996, AQTESOLV, Version 1.17.
- Environmental Protection Commission, 1996, Chapter 135, "Technical Standards and Corrective Action Requirements for Owners and Operators of Underground Storage Tanks," 567 Iowa Administrative Code.
- Environmental Protection Commission, 1998, Chapter 137, "Iowa Land Recycling Program and Response Action Standards," 567 Iowa Administrative Code.
- Iowa Department of Natural Resource, 1996, Tier 1 Guidance, Version 1.0.
- Iowa Department of Natural Resources, 1998, Tier 2 Site Cleanup Report Guidance, Version 2.14.
- U.S. Department of Energy, Office of Environmental Management, Oak Ridge Operations Office and the DOE Center for Risk Excellence, 1999, Risk Assessment Tools, Toxicity Values and Chemical Specific Factors, <u>http://risk.lsd.ornl.gov/tox/tox_values.html</u>.

	Iowa's UST Regulations ^b	Iowa's LRP Regulations ^e
ATc-Averaging time for carcinogens (years)	70	70
ATn-Averaging time for non-carcinogens (years)	25	70
BW-Body weight (kg)	70	70
ED-Exposure duration (years)	25	70
EF-Exposure frequency (days/year)	250	365
TR-Target risk for carcinogens (unitless)	1x10 ⁻⁴ or 1x10 ^{-6d}	5x10 ⁻⁶
THQ-Target hazard quotient (unitless)	1.0	0.02 ^e or 0.20 ^f

Table 1. Exposure factors for non-residential zoning in a protected groundwater source^a.

^a A protected groundwater source is an aquifer with a hydraulic conductivity (K) of ³0.44 meters/day.

^b (Iowa Department of Natural Resources, 567 IAC 1351998) ^c (Environmental Protection Commission, 567 IAC 137 1998) ^d Pathway dependent, ^e For Cancer Group C chemicals such as 1,1-DCE.

^f For Cancer Groups D and E chemicals such as PCE and *cis* 1,2-DCE.

Depth (Feet)	P-2	P-4	P-5	P-6	P-7	P-9	B-20	B-21	P-22	B-23	B-24	P-25	B-26	B-27
10-15	5760ª	852ª	2017ª	89.3° 8173ª	4034ª	1595 ^ь	715 ^a 13 ^c <5.0 ^d	405ª	298ª		191ª		109ª	453ª
15-20				14935ª										
20-25				19485 ^a 13000 ^c <120 ^d			144ª	796 ^a	194ª	233 ^a ND ^c <5.0 ^d	279^{a} ND ^c $< 5.0^{d}$	380 ^a ND ^c <5.0 ^d	202 ^a ND ^c <5.0 ^d	49 ^a ND ^c <5.0 ^d

Table 2. Soil gas and soil PCE and TCE analytical results.

Depth (Feet)	B-28	B-29	B-31	B-32	MW-14	P-34	P-35	P-37	P-38	P-39	P-40	P-42	P-44
10-15	603ª	388ª	51.6° 8.1 ^d			14.6° 15.1° <5.0 ^d	ND ^c 6.3 ^c <5.0 ^d	ND ^c <5.0 ^d	ND ^c <5.0 ^d	ND ^c <5.0 ^d	$\frac{\text{ND}^{\circ}}{<5.0^{\text{d}}}$	ND ^c 10.3 ^c <5.0 ^d	ND ^c <5.0 ^d
15-20					5.4° <1.0 ^d			13.4° <5.0 ^d	362° 30.5 ^d	41.5° <5.0 ^d	105,000 ^c 1,510 ^d		37.8° <5.0 ^d
20-25	835 ^a 54.5 ^c <5.0 ^d	$\begin{array}{c} ND^a \ NC^c \ <5.0^d \end{array}$		459ª									

^a soil gas for PCE (mg/L), ^b soil sample, field analyzed for PCE (mg/kg), ^c soil sample, laboratory analyzed for PCE (mg/kg), ^d soil sample, laboratory analyzed for TCE (mg/kg), ND = not detected

	SOIL SAMPLE IDENTIFICATION AND DEPTH									
CONTAMINANT	P-6 22'-24' (μg/kg)	P-6 11.75' (μg/kg)	B28-A 24'-26' (µg/kg)	P38 15' (μg/kg)	P40 12.5' (μg/kg)	P40 15' (μg/kg)	P43 9.5' (μg/kg)			
Tetrachloroethene	13,000	89.3	54.5	362		105,000	503			
<i>n</i> -Butylbenzene	3,670	17.2		57.6		10,300				
sec-Butylbenzene	2,350			44.5		7,260				
Isopropylbenzene	207					1,060				
p-Isopropyltoluene	1,690	12.6		79.6	ND					
<i>n</i> -Propylbenzene	1,180					4,630				
1,2,4-Trimethylbenzene	2,900			20.7		5,710				
1,3,5-Trimethylbenzene	204					2,110				
Total Extractable Hydrocarbons as Diesel	1,000	530	12							
Methylene chloride		56.2			55.4					
Trichloroethene		214		30.5		1,510				
Chlorotoluene				5.3						
tert-Butylbenzene						4,730				
2-Chlorotoluene						394				
Ethylbenzene						398				
Xylenes, total						927				
cis 1,2-Dichloroethene										
	GOU	NDWATE	R SAMPLI	E IDENTII	FICATION	AND DE	РТН			
	Ρ-6 51' (μg/L)	Ρ-6 91' (μg/L)	MW-6B 92'-102' (µg/L)							
Tetrachloroethene	538	50.3	90.1							
Trichloroethene	209	13.7	18.5							
cis 1,2-Dichloroethene	5.5									
Chloroform	3.3									

Table 3. Other contaminant concentrations.

		CAS No.	Cancer Group ^b	SF _o ^c (mg/kg-day) ⁻¹	$\begin{array}{c} R_{f}D_{o}^{c}\\ (mg/kg-day)^{-1}\end{array}$	SF ^e _i (mg/kg-day) ⁻¹	$\begin{array}{c} R_{f}D_{i}^{f} \\ (mg/kg-day)^{-1} \end{array}$
Carcinogens	Chloroform	67663	B2 ^g	6.1e-3 ^g	1.0e-2 ^g	8.1e-2 ^h	none
	Vinyl chloride	75014	A^h	1.90 ^h	none	3.0e-1 ^h	none
	Methylene chloride	75092	B2 ^g	7.50e-3 ^g	6.0e-2 ^g	1.65e-3 ¹	8.57e-1 ^j
	Tetrachloroethene	127184	D	5.2e-2 ^k	1e-2 ^g	2.0e-3 ^k	1.70e-1 ^{j,k}
	Trichloroethene	79016	none	1.1e-2 ^k	6.0e-3 ^k	6.0e-3 ^k	none
Non-Carcinogens	cis 1,2-Dichloroethene	156592	D	none	1.0e-2 ^h	none	none
	trans 1,2-Dichloroethene	156605	none	none	2.0e-2 ^g	none	none
	1,1-Dichloroethene	75354	С	6.0e-1 ^g	9.0e-3 ^g	1.2 ^h	none

Table 4. Toxicity and chemical-specific human health risk factors^a.

^a(U.S. Department of Energy, 1999)

^bIDNR, 567 IAC Chapter 137, Iowa Land Recycling Program and Response Action Standards, Table 1. Standards for Groundwater, Iowa Land Recycling Program, (Current as of October 4, 1999)

^cOral Slope Factor

^dOral Reference Dose

^eInhalation Slope Factor

^fInhalation Reference Dose

^gThe U.S. Environmental Protection Agency's (EPA's) Integrated Risk Information System (IRIS), Internet: www.epa.gov/iris/ Current as of October 4, 1999.

^hEPA's Health Effects Assessment Summary Tables (HEAST), 9200.6-303 (97-1), EPA-540-R-97-036, PB97-921199, July 1997.

ⁱThe Inhalation Slope Factor was calculated from inhalation unit risk as described in Supplemental Guidance from RAGS: Region 4 Bulletins, Human Health Risk Assessment (Interim Guidance) (USEPA, November 1995).

^jInhalation Chronic RFC * (20/70). United States Environmental Protection Agency. November 1995. Supplemental Guidance to RAGS: Region 4 Bulletins, Human Health Risk Assessment (Interim Guidance). Waste Management Division, Office of Health Assessment.

^kThe Risk Assessment Program has contacted the Superfund Program and been given provisional values which should be used for DOE-ORR projects. This value should be clearly documented as provisional. For other projects, the Superfund Health Risk Technical Support Center should be contacted directly (513) 569-7300.

Boring/Well Number	Date	PCE (mg/L) Statewide Standard = 5ug/L	TCE (mg/L) Statewide Standard = 5ug/L
P-6 (91 feet bgs)	08/04/98	50.3	13.7
P-6 (51 feet bgs)	08/07/98	538	209
MW-14	08/11/98	5.4	<1.0
MW-6B	08/21/98	90.1	18.5
MW-6B	12/08/98	33.1	5.3
MW-5	12/08/98	<200	575
MW-4	12/08/98	<1.0	<1.0
MW-8	12/08/98	17.9	1.7

 Table 5. Groundwater sampling summary for PCE and TCE.

Table 6.Summary of source SSTLs.

MEDIA	PATHWAY	RECEPTOR		PCE		TCE				
			Chap. 135 Exp. Factors (UST)	Chap. 137 Exp. Factors (LRP)	Statewide Standards	Chap. 135 Exp. Factors (UST)	Chap. 137 Exp. Factors (LRP)	Statewide Standards		
Groundwater (µg/L) Actual PCE = 538 Actual TCE = 209	Groundwater	Well #1	20200	3900	5	11800	2100	5		
	Ingestion	Protected GW- No IC ^a	360	70	5	210	42	5		
		Protected GW With IC All	950	180	5	550	110	5		
Soil (mg/kg)	Soil Leaching to Groundwater	Well #1	314	61.0	780 ^b	65.0	12.0	180 ^b		
Actual PCE = 105 Actual TCE = 1.51		Protected GW- No IC	5.6	1.08	780 ^ь	1.2	0.23	180 ^b		
		Protected GW- With IC All	9.9	1.9	780 ^ь	2.1	0.41	180 ^b		
		GW Vapor- Truck Sales Basement	160	33.5	780 ^ь	14.0	10.5	180 ^ь		
		GW Vapor- New Facility	270	56.6	780 ^b	23.8	17.8	180 ^b		
	Soil Vapor to Enclosed Space	All	52	10.8	780 ^ь	4	3.4	180 ^b		

^a IC = institutional control

^b Statewide standards for soil are based on ingestion and do not address leaching or vapor. Bold values indicate an SSTL less than the current source concentration.



Figure 1. Sampling locations. B = Soil boring $P = Geoprobe^{TM} location$ MW = Monitoring well