# INTERIM MEASURES WORK PLAN

# EXIDE ENVIRONMENTAL RESPONSE TRUST Frankfort Indiana Site 555 North Hoke Avenue EPA ID No. IND001647460

Prepared by:



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Montrose AGC Project # 2020-4123



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#### 1.0 OVERVIEW

This Interim Measures Work Plan (IMWP) has been prepared by Advanced GeoServices Corp. dba Montrose Environmental Solutions ("Montrose AGC") on behalf of the Exide Environmental Response Trust (Trust) to address impacts to groundwater from volatile organic compounds (VOCs) and chlorinated VOCs (CVOCs) at the former Exide Technologies (Exide) facility located at 555 North Hoke Avenue in Frankfort, Indiana (EPA ID# IND 001 647 460) (i.e., the Site).

Montrose AGC conducted a Comprehensive RCRA Facility Investigation (RFI) at the Site and presented the results of the RFI and Interim Corrective Measures (ICM) activities in a report titled *Comprehensive RCRA Facility Investigation (RFI) Report* dated February 24, 2021 (RFI Report). The RFI was completed as part of the RCRA Corrective Action (CA) process. Investigation of groundwater in May and July 2018 showed impact to groundwater in on-site monitoring wells from VOCs and CVOCs. The RFI Report recommended conducting focused investigation activities to define the movement of groundwater and groundwater contamination in the area north and east of MW-4, for the ultimate purpose of proceeding with interim measures for chlorinated VOCs. The focused investigation activities are detailed herein.

The investigation also includes the collection of representative soil gas samples using direct push methods from select locations (including in utility bedding, if possible) and vapor samples from sanitary and storm water manholes. This Interim Measures Work Plan (IMWP) was submitted as a draft document for USEPA review on May 5, 2021 by Montrose AGC on behalf of the Exide Environmental Response Trust. The following table outlines the date that USEPA provided comments to the IMWP and the date of the response to comments.

#### <u>Date Comments received from USEPA</u> <u>Date of Response to Comments</u>

May 21, 2021 July 6, 2021

July 16, 2021 August 6, 2021

August 12, 2021 August 19, 2021

August 20, 2021 August 26, 2021



On August 27, 2021, the USEPA responded by email stating that the responses were acceptable and requested that the revised IMWP reflecting the changes be submitted when available. This revised document is intended to fulfill that request.

#### 1.1 Facility Location and Description

The Site is located in central Indiana within Clinton County, approximately 50 miles northwest of Indianapolis (see Figure 1). The Site is bounded by North Hoke Avenue to the west, Kelley Avenue to the east, Michigantown Road to the north (also referred to as Washington Street on some maps), and Norfolk Southern railroad tracks to the south. The Site consists of eighteen (18) contiguous parcels now owned by the Trust which encompass approximately 13.7 acres. All but three of the parcels are located within a perimeter security fence. The majority of the area (12.1 acres) lies within the perimeter security, and with the exception of grass and a few shrubs along North Hoke Avenue is covered with former building pads, pavement or crushed stone. The facility was formerly a manufacturing plant that produced lead-acid automotive batteries. The plant was closed and the aboveground infrastructure was decontaminated and demolished January 2013.

#### 1.2 Reference Documents

The proposed focused investigation activities will be performed under the framework of the following site documents:

- 1. Supplemental Site Investigation Work Plan prepared for Excide Technologies by Advanced GeoServices Corp. dated December 21, 2018
- Quality Assurance Project Plan for the RCRA Facility Investigation, Excide Technologies Former Manufacturing Facility, Frankfort, Indiana, prepared for Excide Technologies by Advanced GeoServices Corp. dated October 12, 2017
- 3. Sampling and Analysis Plan for the RCRA Facility Investigation, Former Excide Manufacturing Facility, Frankfort, Indiana, prepared for Excide Technologies by Advanced GeoServices Corp. dated October 12, 2017



#### 1.3 Local Geology

The soil and unconsolidated materials in the subsurface of the Site consist of glacial till which is unsorted glacial sediment that washes off of retreating glaciers. The soil has been mapped as the by Fincastle-Crosby soils. This is a silty loam with slow infiltration rates (Class C) and is somewhat poorly drained. The hydraulic conductivity in these regional silt and clay deposits is low which makes these deposits semi-pervious.

Shallow groundwater represents local perched zones of saturation in sand and silt layers. The depth and thickness of the saturated layers varied from 4 to 10 feet bgs, and appear to be laterally discontinuous. The underlining aquiclude was encountered in all of the groundwater monitoring well locations and consisted of a very stiff to hard gray clayey Silt to silty Clay with trace amounts of sand and/or gravel. The perched groundwater near MW-4 is approximately 10 feet below ground surface and groundwater flows towards the north.



#### 2.0 FOCUSED INVESTIGATION ACTIVITES

#### 2.1 <u>Investigation Overview</u>

The following activities were proposed to assess the movement of groundwater and groundwater contamination in the area north and east of MW-4, for the ultimate purpose of proceeding with interim measures for chlorinated VOCs.

- Confirm the existence of the easterly flow direction of shallow groundwater between MW-4 and MW-9 suggested by groundwater surface elevations measured during the November 2019 groundwater sampling event.
- Collect detailed information regarding the types, sizes and depth of utilities in Kelley Avenue for the purpose of identifying potential preferential pathways for the VOC impacted groundwater observed in and around MW-4.
- Conduct targeted groundwater sampling using Hydropunch sampling techniques
  within and across Kelley Avenue to define the limits of VOC impacted groundwater.
  This should include targeted groundwater sampling within the backfill of utilities
  identified to be below the shallow groundwater table.
- Collect surface water samples for VOC and lead analysis at discharge point of site storm sewer to unnamed tributary along Michigantown Road, near Kelley Avenue.
- Complete passive in-situ microcosm study in MW- 1, 4, and 9, to collect microbial, chemical and geochemical data to determine if MNA, bio-enhancement or bioaugmentation are appropriate interim measures.
- Install additional monitoring wells at select locations selected based on the results of Hydropunch sampling in and across Kelley Avenue.



 Conduct one additional round of groundwater sampling at new and existing monitoring wells for VOCs.

In addition, groundwater, soil gas samples, and manhole vapor samples will be collected and analyzed on-Site using a Pace Analytical<sup>®</sup> Services mobile laboratory gas chromatograph (GC). The information obtained from the above activities should be sufficient to allow preparation of a detailed design for a selected Interim Measures to address chlorinated VOCs identified in the vicinity of MW-4 (and potentially into/across Kelley Avenue).

#### 2.2 Synoptic Water Levels

Well gauging data collected during the RFI suggest that the shallow groundwater gradient steepens with an easterly flow direction. Monitoring Wells MW-4 and MW-9 are both sand-packed and screened in approximately the same depth interval and should be representative of the water table aquifer. Groundwater occurs immediately below the Site exists as an unconfined water table aquifer, where the water level in the sediments can rise and fall in response to precipitation events. Depth-to-water will be measured by Montrose AGC in each well on the Site using an electronic water level indicator. The synoptic measurements will include the measurement of water levels and well depths in the monitoring wells in as short a time frame as possible to determine the potentiometric surface across the Site. The field personnel will measure the water levels in the wells to the nearest 0.01 foot using the surveyed point at the top of the inner well casing for reference. The goal of the well gauging event will be to confirm the previous water table conditions by preparing an updated groundwater contour map.

#### 2.3 Utility Investigation

Montrose AGC will contact the Frankfort's Street Department to identify whether utility maps are available for Kelley Avenue adjacent to MW-4. we will request copies of available plans for sewer lines and utilities in Hoke Avenue and Washington Avenue (aka Michigantown Road), and E. Morrison Street. We will review the plans for the depth and alignment of the sewer lines and utilities, and whether or not such features could represent a preferential pathway for groundwater contamination in this area of the site. If maps are not available, Montrose AGC will oversee a



focused subsurface utilities survey using geophysical techniques. The information will be utilized to evaluate whether or not utility trenches/backfill could represent a pathway for CVOC migration.

If needed, the subsurface survey will include the use of ground penetrating radar (GPR) and Radio Frequency (RF) transmitter and receiver operated by a competent geophysical contractor. GPR devices send radar pulses into the ground and detect reflections from subsurface structures. The RF transmitter emits a specific electromagnetic signal either indirectly induces, or with direct contact induces a signal on a metallic line. The RF receiver is configured to detect that signal on the ground surface to determine the path of the metallic line. The RF receiver can also be used in 'passive' mode, which can pick up any charged line, including electric lines. Nonmetallic lines can be traced by 'snaking' a metallic wire through it and then inducing a signal on that metallic wire. A map of the underground utilities will be generated.

#### 2.4 Targeted On-Site Groundwater Sampling

Hydropunch sampling is proposed to further investigate the Trichloroethene (TCE) detected in groundwater surrounding MW-4 and the area across Kelley Avenue near MW-9 to assess the limits of VOC impacted groundwater. If it can be done safely, Montrose AGC will sample groundwater within the backfill of utilities that are identified to be below the shallow groundwater table as discussed in Section 2.14. If needed, a pneumatic air knife could be used to excavate and install a temporary screen to sample close to underground utilities. In other areas, Hydropunch sampling will be performed using a direct-push drill rig to drive a sealed screen to the target depth. The screen is then exposed and a groundwater sample collected using a variety of methods: tubing with check valve, mini-bailer, or a small bladder pump. After sampling, the openings will be grouted closed. The On-site Hydropunch groundwater samples will be collected utilizing Hydropunch sampling techniques at Locations 7, 8, 9, 10 and 11, as shown on Figure 2. Water samples will be analyzed using a Pace Analytical® Services mobile laboratory GC for the following parameters.

- Benzene, Toluene, Ethyl Benzene, and xylenes (BTEX)
- cis-1,2-Dichloroethene (cis-1,2-DCE)
- Perchloroethylene (PCE)



- TCE
- Vinyl Chloride (VC)

Additional samples may be obtained based on the Decision Tree document contained in Appendix A. Basically, if the groundwater sample results are between 100 and 600 ug/L TCE, sampling will be deemed complete and no offset samples will be necessary. If the number is less than 100 ug/l, we will step in and install an offset boring closer to the source (MW-4). These locations are designated with an "A" on Figure 2. At a value of less than 100 ug/L, the extent of impact can be reasonably estimated employing multiple lines of evidence, including, but not limited to, analytical data; extrapolation or modeling based on existing data; application of a conceptual site model; or other means for determining the extent of the contamination for the purposes of interim remedial action. If the number is over 600 ug/L, we will step out and complete a step-out offset boring. These locations are designated with an "B" on Figure 2. Location 11 does not have an 11B, because groundwater well MW-3 will be utilized to interpret the 500 ug/L if the MW-11 sample is >600 ug/L.

#### 2.5 <u>Sediment and Surface Water Sampling</u>

A sediment sample and a surface water sample (if there is surface water flow from the outfall) will be collected at the discharge point of the Site storm water sewer to unnamed tributary along Michigantown Road, near Kelley Avenue. This is designated as Outfall Z on Figure 2. The sample will be sent to Pace Analytical for analyzed for VOC using EPA Method 8260 and lead analysis.

#### 2.6 Passive In-Situ Microcosm Study

Montrose AGC will conduct a passive in-situ microcosm study in monitoring wells MW- 1, 4 and 9. This entails collecting microbial, chemical and geochemical data to determine if monitored natural attenuation (MNA), bio-enhancement or bio-augmentation are appropriate interim measures. The in-situ study comprises Bio-Trap® Sampler and CENSUS analysis (Microbial Insights, Inc.). Bio-Trap® samplers are in-well study units that contain a sampling matrix favorable to colonization by microorganisms, which can later be harvested, quantified and assessed in the laboratory for microbial characterization and contaminant degradation potential



(i.e. CENSUS analysis). The CENSUS assessment tests for the presence and growth of Dehalococcoides, intrinsic and augmented, and the microbial reductase genes necessary for complete conversion of chlorinated VOCs (i.e., TCE) to non-toxic end products (i.e. ethene or ethane). Specifically, the parameters assessed with CENSUS and used as metrics for assessing biodegradation potential include the following:

- 1. Dehalococcoides population (>104 cells/milliliter groundwater benchmark for effective bioremediation strategy);
- 2. Expression of tceA Reductase gene (TCE degradation);
- 3. Expression of bvcA Reductase gene (VC degradation, prevents cis-DCE accumulation); and
- 4. Expression of vcrA Reductase gene (VC degradation, prevents cis-DCE accumulation).

In addition to the CENSUS analysis, Bio-Trap® samplers also collect groundwater for characterization of secondary lines of evidence indicative of microbial growth and contaminant degradation including dissolved gas production (ethane, ethene and methane), volatile fatty acids ("VFA") production, standard geochemical parameters and contaminant of concern concentration profiles. Detection of VFAs indicates active fermentation of intrinsic or added organic substrates (i.e. biostimulation) by microbial organisms has occurred and microbial growth is favorable. Geochemical parameters indicate if favorable conditions for microbial growth are present and are also applied for full-scale design assessment and modeling. Changes to the innate contaminant of concern concentration profile, with the production of intermediate degradation products (e.g., cis-1,2,-Dichloroethene and vinyl chloride) and non-toxic end products (e.g., ethene and ethane), provides a direct line of evidence that bioremediation is occurring. Lastly, the Bio-Trap® sampler includes sub-units that assess the benefit of carbon substrate addition, a critical factor in determining full-scale design parameters. Each Bio-Trap® unit will be deployed for approximately 3 months.



#### Slug Testing

Slug testing of wells MW-1, MW-3, MW-4, MW-7, MW-9, and MW-10 is proposed after retrieval of the microbial collection unit to assess the hydraulic properties of aquifer such as hydraulic conductivity. A falling-head test is proposed to be conducted by rapidly raising the water level in the monitoring well and subsequently measuring the falling water level. The test and resulting analysis will yield estimates of the average in-situ horizontal hydraulic conductivity of a small region surrounding the well screen. During the test, after first gauging the well, commercially available bottled drinking water is rapidly poured into the well. The rate at which the water level returns to at least 80% of its' initial static level is measured. The selected water-level measurement method depends upon how quickly water levels change in the well bore. The subsurface is sandy silt and clayey silt so the test may last for minutes to hours, indicating that the test data can be collected by hand using an electronic water level indicator. The resulting data will be plotted and analyzed using one or more of the three most-widely applied techniques for estimating horizontal hydraulic conductivity which are 1) Hvorslev (1951), 2) Bouwer and Rice (1976) (Bouwer, 1989), and 3) Butler's modification of the Bouwer and Rice approach (Butler and Healey, 1998; Butler et al., 2003).

#### 2.7 <u>Additional Monitoring Well Installation</u>

Based on the results of the of Hydropunch sampling in and across Kelley Avenue, Montrose AGC will install additional monitoring wells at select locations agreed upon with the USEPA. One flush-mount groundwater monitoring well will be installed at Location 12 on the south side of E. McClurg Street (new MW-11, approximately 50 feet east of MW-9) shown on Figure 2. A flush-mount monitoring well at Location 12 will help refine our understanding of the gradient between MW-4 and MW-10 and possible influences caused by utilities in Kelley Avenue.

Three new groundwater monitoring wells (MW-12, MW-13, and MW-14) will be installed at the fringe of the TCE plume around MW-4. The wells will be installed using hollow-stem auger techniques (direct push methods may also be utilized). The locations will be selected in the field based on the results of the Hydropunch sampling. The intent will be to locate one well south (upgradient), one well northeast and one well northwest of MW-4 where the TCE concentrations



in groundwater appear to decrease rapidly with distance from MW-4. The screened intervals will be selected to target the uppermost saturated intervals and are expected to be <5 feet in length.

During drilling, soil samples will be collected from the following intervals: 0 to 4 feet below ground surface (bgs), 4 to 8 feet bgs, and 8 to 12 feet bgs intervals. Soil conditions will be recorded and provided on boring logs. If the soil sample is collected from an unsaturated zone, the soil sample will be sent to Pace Analytical for analysis for VOC using EPA Method 8260 and lead analysis.

The monitoring wells will be finished as stick-up wells (flush mount may be considered depending on the conditions where they are installed). The wells will be constructed of 2-inch diameter Schedule 40 PVC screen and riser. A 5 to 10-foot long 0.010" slot screen will be installed in the first water bearing zone encountered to intercept the uppermost part of the aquifer. The bottom of the screen set is expected to be set approximately 15 to 18 feet bgs.

#### 2.8 Groundwater Sampling

One additional round of groundwater sampling at new and select existing monitoring wells will be conducted and the groundwater will be analyzed for VOCs. The additional round of groundwater sampling will be performed using low-flow purging methods at groundwater monitoring wells MW-1, MW-2, MW-3, MW-4, MW-7, MW-9, and MW-10 and the additional new wells MW-11, MW-12, MW-13, and MW-14 installed as described in Section 2.7. Groundwater samples will be analyzed for VOCs. Specific conductivity, dissolved oxygen, temperature oxidation-reduction potential (ORP), turbidity and pH will be measured in the field. Water levels will be obtained in all wells prior to sampling. An equipment blank will be collected from the sampling equipment for each day that non-disposable equipment is used for low flow sampling.

#### 2.9 Kelly Avenue Soil Gas Sampling

Montrose will conduct Soil Gas Screening along east side of Kelley Ave at Locations 1, 2, 3, 4, 5 and 6 shown on Figure 2. Soil Gas Probes will be installed as fixed/permanent flush-mount wells, with bolted lids. Probes will be located approximately 1 to 2 ft. above the groundwater surface



(est. 4 to 6 below ground level). Soil gas probes will be sampled and analyzed for BTEX, cis-1,2-DCE, PCE, TCE, and VC using a field GC operated by Pace Analytical.

Standard operating procedures (SOPs) for soil gas sampling are included in Appendix B. The soil gas probes will be constructed as permanent points. Sealing of each point will be accomplished using bentonite or grout in the annulus of the boring and the point will be finished with a stainless-steel capping fitting inside a flush-mount wellhead with a bolt-down "manhole" 12-inches or less in diameter. The water table is the presumed source of CVOCs in soil gas so the near-source soil gas samples will be collected within one (1) foot of the top of the capillary fringe. Nearby monitoring wells will be gauged to assess the appropriate depths of the soil gas probes. Based on the previous depth to water data for MW-3, 4, and 9, the probes are anticipated to be installed to a maximum depth of approximately 4 feet below grade. The tubing will be connected to the port sampler and a vacuum pump so that it can be checked for leakage by applying a vacuum to the line and monitoring an in-line pressure gauge for a loss of vacuum. An equilibration time of two hours will be allowed for direct push installations which will be increased to 48 hours if necessary to install probes using hand augured borings.

Before sampling, the sampling system will be leak checked by covering the sample port with a shroud that will be filled with helium. The shroud will allow two other ports where the leak check sample can be pulled and the helium concentration in the shroud can be tested. The shroud will be filled with helium (10-20%), while a soil gas sample is collected from the sample point and measured with a portable helium detector. A leak is occurring when the helium concentration is greater than 10% of the concentration within the shroud. In this case, the leak must be fixed and the leak check repeated. When the leak test is complete, the shroud, helium supply, and monitoring tubes will be removed, leaving the sample tube connected to the port for sampling.

Soil gas sampling will not be performed during rain events. Soil gas results will be compared to screening levels derived using the USEPA Vapor Intrusion Screening Level (VISL) calculator for "Near Source Soil Screening" HQ=1.0 and Target Risk 1x10<sup>-5</sup> which are shown on the following table:



USEPA Vapor Intrusion Screening Level (VISL)						
Near Source Soil Screening" HQ=1.0 and Target Risk 1x10⁻⁵						
	Target Sub-Slab and	Target	Is Target			
	Near-source Soil Gas	Groundwater	Groundwater			
	Concentration VISL	Concentration	Concentration			
	(ug/M³)	(ug/L)	< MCL?			
			(Cgw < MCL?)			
Trichloroethylene (TCE)	69.5	5.18	No (5)			
Tetrachloroethylene (PCE)	1,390	57.6	No (5)			
cis-1,2-dichloroethane (cis-1,2-DCE	) -	-	-			
Vinyl Chloride (VC)	55.9	1.47	Yes (2)			
Benzene	120	15.9	No (5)			
Toluene	174,000	19,200	No (1,000)			
Ethyl Benzene	374	34.9	Yes (700)			
Xylenes	3,480	385	Yes (10,000)			

#### Notes:

Results of the soil gas analysis will be provided to USEPA for further investigation as appropriate by the USEPA. This proposed scope of work does not include performing vapor sampling inside of the residential properties.

#### 2.10 Step-out Soil Gas Sampling

In addition, we will collect step out soil gas samples in the three locations shown on the attached map:

- 1. Location 13 Along the unpaved road north of East McCrurg Street (soil gas);
- 2. Location 14 Near the curve close to 1609 Goder Drive (soil gas); and
- 3. Location 15 East of North Kelley Ave. and north of MW-3 (soil gas and Hydropunch).

<sup>1.</sup> Output generated 27AUG2021:10:49:19

<sup>2.</sup> There is no VISL value for cis I,2-DCE on the EPA VISL website or in IDEM's 2021 Screening Level Vapor Exposure. Source: https://epa-visl.ornl.gov/cgi-bin/visl\_search



Soil Gas Probes will be installed as fixed/permanent flush-mount wells, with bolted lids. Probes will be located approximately 1 to 2 ft. above the groundwater surface (est. 4 to 6 below ground level). Soil gas probes will be sampled and analyzed for BTEX, cis-1,2-DCE, PCE, TCE, and VC using a field GC operated by Pace Analytical.

#### 2.11 <u>Targeted Off-Site Groundwater Sampling</u>

If soil gas concentrations for TCE and VC at Locations 1 and 2 are above VISL values, Hydropunch groundwater sampling will be performed utilizing a separate Geoprobe boring installed adjacent to the location with the highest TCE concentration. If soil gas results at Locations 1 and 2 are below the VISL values, a Hydropunch boring will be completed on west side of Kelley Avenue at Location 2A.

If a Hydropunch groundwater sample is collected at Location 2A and the corresponding TCE concentration is >600 ug/L, we will complete Hydropunch sampling adjacent to Location 2.

Hydropunch groundwater sampling will be conducted at Locations 3, 4, and 6 shown on Figure 2. The Hydropunch groundwater samples that are collected will be field analyzed for BTEX, cis-1,2-DCE, PCE, TCE, and VC using a field GC operated by Pace Analytical.

#### 2.12 Manhole Sampling on Kelley Avenue

Eight (8) grab vapor samples will be collected from sanitary manholes (B, E, F, G, H, and I) and storm sewer manholes (A and C) identified along Kelley Avenue as shown on the attached figure. Open grate storm sewer manholes will not be sampled. The grab vapor samples from manholes will also be analyzed for BTEX, cis-1,2-DCE, PCE, TCE, and VC using a field GC operated by Pace Analytical. In addition, the sanitary sewer gas samples will be analyzed for chloroform.



Each sewer manhole will be evaluated for accessibility and manhole type (i.e. closed manhole or open grate manhole) prior to sampling. If the sewer manhole is found to be inaccessible or, in the case of storm sewer manholes, is an open grate, a sewer gas sample will not be obtained from that particular manhole(s). The reason sewer gas vapor samples are not being obtained from open grate storm sewer manholes is because open grate storm sewer manholes readily vent directly to the atmosphere and a high mixing rate of sewer gas and ambient air occur in these locations. The tubing will be lowered into the manhole to the proper depth, and a syringe with a three-way valve will be used to purge the tubing prior to valving to the purge pump. A grab sample will be collected over 30 to 60 seconds in a Tedlar bag for analysis using the field GC.

#### 2.13 Manhole Sampling along Washington Avenue

We propose to collect VOC vapor grab samples from sanitary manholes J and L along Washington Avenue shown on Figure 3. In addition, we will attempt to locate storm water Manholes O, N, MH-10, and MH-11. If the manholes can be located and are assessable, and they are not an open grate, a vapor gas sample will be obtained for analysis. The grab vapor samples from manholes will also be analyzed for BTEX, cis-1,2-DCE, PCE, TCE, and VC using a field GC operated by Pace Analytical. In addition, the sanitary sewer gas samples will be analyzed for chloroform.

#### 2.14 Sampling from Utility Bedding

In addition, three (3) soil gas samples will be attempted to be collected from temporary soil gas probes selected based on the results of the utility information obtained from the City of Frankfort. The intention is to collect soil gas samples from immediately above the saturated granular bedding or backfill (if present) for the sanitary sewer and storm sewer.

Soil gas samples will be attempted to be collected from the temporary soil gas probes selected to target the granular backfill/bedding of the utilities will be screened against the VISL values



calculated for "Near Source Soil Screening" HQ=1.0 and Target Risk 1x10<sup>-5</sup> listed in Section 2.9. If the soil gas results are higher, then an additional soil gas sample will be collected from the same zone (i.e. immediately above saturated granular bedding or backfill) for the same utility at a location approximately 200 feet north of the failing location. Groundwater sampling will be attempted within the saturated granular bedding or backfill coinciding with the soil gas sample location. The technique for groundwater sample retrieval will be Hydropunch sampling or a soft dig technique selected to avoid damage to the utility.



#### 3.0 DATA ANALYSIS

#### 3.1 <u>Data Analysis</u>

A report presenting the findings and an analysis of the findings will be prepared. Given that the Bio-Trap® samplers described in Section 2.6 will be deployed for three months, Montrose will likely provide a preliminary report to USEPA containing the soil gas, utility vapor, and groundwater sampling analytical results followed by a final report that includes the Bio-Trap® sampler results and slug test analysis.

Targeted groundwater sampling data will be used to develop iso-concentration maps that will be used to further understand the extent and magnitude of the CVOC plume in groundwater. The IDEM VISL based definition 2021 Screening Level Vapor Exposure (SLVE) will be used to interpreting the overall extent of the VOCs plume:

Compound	Standard¹(ug/L)	2021 SLVE <sup>2</sup> (ug/ Residential Ind	
Benzene	5	28	120
Ethyl Benzene	700	-	-
Toluene	1,000	-	-
Xylenes	10,000	-	-
cis-1,2-DCE	70	-	-
Tetrachloroethene	5	6.5	28
TCE	5	9.1	38
Vinyl chloride	2	2.1	35

<sup>1.</sup> 2018 Remediation Closure Guide (RCG) GW Tap Limit, 2009 RISC TPH Closure Limits

<sup>&</sup>lt;sup>2.</sup> Indiana Department of Environmental Management 2021 Residential Screening Level Vapor Exposure



- The IDEM 2021 SLVE will be used to interpreting the overall extent of the VOC plumes to the extent that the data collected allows such an interpretation. When reporting the results of this investigation, the VI evaluation documentation will include a map using the data to interpret the extent of the VOCs groundwater plume that exceed the IDEM VI screening levels based the data collected. Maps will be provided for the contaminants of concern being assessed should they exceed VISLs, as appropriate. We will use the locations provided in the revised work plan document to assess on-Site and off-Site conditions. The extent of VOC impact may also be estimated using the analytical data, interpretation, and extrapolation.
- Well gauging data will be used to generate a groundwater contour map, which will be further enhanced with the additional data from the new groundwater monitoring wells.
- 4. Utility information will be used to further develop the Conceptual Site Model (CSM).
- 5. The passive in-situ microcosm study will provide microbial and geochemical data so that we can perform an evaluation of remedial options and their impermeability.
- 6. Slug testing and analysis will provide a measurement of horizontal hydraulic conductivity so the groundwater velocity and infiltration rates can be calculated for remedial planning.
- 7. Lastly the monitoring well sampling data from MW-1, MW-2, MW-3, MW-4, MW-7, MW-9, and MW-10 and the additional new wells MW-11, MW-12, MW-13, and MW-14 will provide current data and a greater understanding of the CVOCs in the subsurface.

The CSM was presented in our document titled *CVOC Remediation Evaluation* dated January 19, 2021, that outlined potential source areas, transport mechanisms, environmental media affected, potential exposure pathways, and potential exposure routes to potential receptors. Briefly, the source area of TCE impacted groundwater appears to be localized and contained by low-permeability soils around MW-4. Vapor intrusion (VI) typically is the main risk driver for non-



residential sites with chlorinated impacts. There are no on-site receptors and conducting the focused investigation proposed herein should allow further evaluation of potential off-site receptors. The results from the focused investigation activities will allow refinement of the CSM.



#### 4.0 CONCLUSION

The proposed focused study will provide the data necessary to design the interim remedial steps needed to address the CVOCs in groundwater on the Site. The main objective of any selected remedial measure is to prevent exposure to humans. The remedial action will provide for the protection of public health and the environment consistent with the intended property use, and comply with applicable environmental regulations and guidance. The timeline associated with the proposed focused investigation could take approximately six to seven months to complete.



#### **FIGURES**

Engineering for the Environment. Planning for People.<sup>TM</sup> 1055 ANDREW DRIVE, SUITE A, WEST CHESTER PA, 19380 tel 610.840.9100 fax 610.840.9199 www.advancedgeoservices.com

#### USGS TOPOGRAPHIC MAP

## EXIDE TECHNOLOGIES FRANKFORT, INDIANA

PROJECT ENGINEER:	PGS	SCALE:	NTS
CHECKED BY:	JSD	PROJECT NUMBER:	2011-2678
DRAWN BY:	KEZ	DATE:	FIGURE: 1

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EXISTING STORMWATER MANHOLE

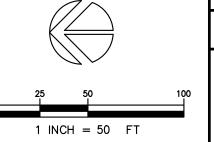
EXISTING SANITARY MANHOLE

EXISTING SANITARY LINE AND MANHOLES

EXISTING STORMWATER LINE AND MANHOLES

EXISTING WATER LINE AND HYDRANT

NOTE: LINE LOCATIONS ARE APPROXIMATE AND TAKEN FROM MAPS PROVIDED BY THE CITY OF FRANKFORT'S MUNICIPAL UTILITY SUPERINTENDENTS.



# PROPOSED MANHOLE VAPOR WASHINGTON AVENUE LOCATIONS



Scales	1" = 50"	~
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# APPENDIX A DECISION TREE DOCUMENT

# EXIDE TRUST – FRANKFORT INDIANA INTERIM MEASURES INVESTIGATION August 30, 2021

#### Purpose:

The purpose of the proposed field investigation is to further define the limits of TCE and other compounds in the vicinity of groundwater monitoring well MW-4 that will become the focus of interim measures intended to eliminate the suspected source area. In addition, the proposed field investigation will collect data to allow an evaluation of the potential for vapor intrusion at the nearby residential dwellings.

We propose to utilize a field gas chromatograph (GC) to provide real-time results for groundwater and soil gas samples, and guide the selection and number of sample locations. Analysis of soil samples will be performed at an off-site lab. Prior to the start of sampling activities, depth to groundwater measurements will be collected at existing groundwater monitoring wells.

#### **Approach to Sampling:**

Off-Site/Residential: The areas along the east side of Kelley Avenue are a predominantly residential, plus a small self-storage facility. The planned approach for Off-Site/Residential groundwater, soil gas and vapor sampling will be as follows:

- Conduct Soil Gas Screening along east side of Kelley Ave at Locations 1, 2, 3, 4, 5 and 6 (see attached Figure). In addition, we will collect step out soil gas samples in the three locations shown on the attached map: 1) Location 13 - Along the unpaved road north of East McCrurg Street (soil gas); 2) Location 14 - Near the curve close to 1609 Goder Drive (soil gas); and 3) Location 15 - East of North Kelly Ave. and north of MW-3 (soil gas and Hydropunch). Soil Gas Probes will be installed as fixed/permanent flush-mount wells, with bolted lids. Probes will be located approximately 1 to 2 ft. above the groundwater surface (est. 4 to 6 below ground level). Soil gas probes will be sampled and analyzed for trichloroethylene (TCE), vinyl chloride (VC), tetrachloroethylene (PCE), cis 1,2-dichloroethane (cis I,2-DCE), benzene, toluene, ethylbenzene, and xylene (BTEX). Analysis will be performed using a field gas chromatograph (GC) operated by Pace Analytical. Soil gas sampling will not be performed during rain events. Soil gas results will be compared to screening levels derived using the USEPA Vapor Intrusion Screening Level (VISL) calculator for "Near Source Soil Screening" HQ=1.0 and Target Risk 1x10<sup>-5</sup>. Results of the soil gas analysis will be provided to USEPA for further investigation as appropriate by the USEPA. This proposed scope of work does not include performing vapor sampling inside of the residential properties.
- If soil gas concentrations for TCE and VC at Locations 1 and 2 are above VISL values, hydro-punch GW sampling will be performed utilizing a separate geoprobe boring installed adjacent to the location with the highest TCE concentration. If soil gas results at Locations 1 and 2 are below the VISL values, a hydro-punch boring will be completed on west side of Kelley Avenue at Location 2A. The hydro-punch groundwater samples will be field analyzed for TCE and VC.

- If a hydro-punch GW sample is collected at Location 2A and the corresponding TCE concentration is >600 ug/L, complete hydro-punch sampling adjacent to Locations 2.
- Collect hydro-punch groundwater samples at locations 3, 4, and 6.
- Soil gas sampling and hydro-punch groundwater sampling will be performed at Location
   6.
- Eight (8) grab vapor samples will be collected from sanitary manholes (B, E, F, G, H, and I) and storm sewer manholes (A and C) identified along Kelley Avenue as shown on the attached figure. Open grate storm sewer manholes will not be sampled. The grab vapor samples from sanitary manholes will also be analyzed for the same VOCs plus chloroform. In addition, three (3) soil gas samples will be attempted to be collected from temporary soil gas probes selected based on the results of the utility information obtained from the City of Frankfort. The intention is to collect soil gas samples from immediately above the saturated granular bedding or backfill (if present) for the sanitary sewer and storm sewer.
- VOC vapor grab samples will be collected from sanitary manholes J and L along Washington Avenue. We will attempt to locate storm water Manholes O, N, MH-10, and MH-11. If the manholes can be located and are assessable, and they are not an open grate, a vapor gas sample will be obtained for analysis from these manholes.
- Soil gas samples collected from the temporary soil gas probes selected to target the granular backfill/bedding of the utilities will be screened against the VISL value calculated for "Near Source Soil Screening" HQ=1.0 and Target Risk 1x10<sup>-5</sup>. If the soil gas results are higher, then an additional soil gas sample will be collected from the same zone (i.e. immediately above saturated granular bedding or backfill) for the same utility at a location approximately 200 feet north of the failing location. Groundwater sampling will be attempted within the saturated granular bedding or backfill coinciding with the soil gas sample location. The technique for groundwater sample retrieval will be hydro-punch sampling or a soft dig technique selected to avoid damage to the utility.

On-Site: The on-site sampling will consist of hydro-punch sample taken from the first saturated permeable soil interval(s) at or below the depth corresponding to the depth to water in MW-4 (as corrected for well stickup). The purpose of the sampling is to define the limits of groundwater with TCE concentrations at or above 500 ug/L. The expectation is that any active remediation (thermal, excavation and removal, or enhanced biotreatment) will focus of the TCE source area.

- Groundwater samples will be collected utilizing hydro-punch sampling techniques at Locations 7, 8, 9, 10 and 11, as shown on the attached Google Earth image.
   Water samples will be analyzed for TCE and VC.
- If the groundwater sample results are between 100 and 600 ug/L TCE, sampling will be deemed complete and no offset samples will be necessary.

- If the TCE result is >600 ug/L an offset boring will be completed at a location further away from MW-4. These locations are designated with a "B" on the attached Google Earth image. (Please note that Location 11 does not have an 11B, because groundwater well MW-3 will be utilized to interpret the 500 ug/L if the MW-11 sample is >600 ug/L).
- If the TCE result is <100 ug/L an offset boring will be completed at location located closer to MW-4. These locations are designated with an "A" on the attached Google Earth image.

New Groundwater Monitoring Wells: Additional groundwater monitoring wells are planned in the vicinity of MW-4 to provide additional detail regarding groundwater gradients and to provide tighter data coverage around the perimeter of the MW-4 TCE plume.

- A groundwater monitoring well, designated as MW-11, will be installed at Location 12 on the south side of E. McClurg Street (approximately 50 feet east of MW-9).
   This will help refine our understanding of the gradient between MW-4 and MW-10 and possible influences caused by utilities in Kelley Avenue.
- Three new groundwater monitoring wells will be installed at the fringe of the TCE plume around MW-4. The locations will be selected in the field based on the results of the hydro-punch sampling. The intent will be to locate one well south (upgradient), one well northeast and one well northwest of MW-4 where the TCE in groundwater is <100 mg/L. The screened intervals will be selected to target the uppermost saturated intervals and are expected to be <5 feet in length. These wells will be designated as MW-12, 13, and 14.
- The new groundwater monitoring wells will be developed in accordance with the procedures in the RFI Work Plan.
- Slug tests will be performed at existing wells MW-1, 3, 4, 7, 9 and 10. In addition, the rates of recovery after development shall be measured for all of the new well locations.
- Groundwater sampling will be completed at all the existing wells MW-1, MW-2, MW-3, MW-4, MW-7, MW-9, and MW-10 and the additional new wells MW-11, MW-12, MW-13, and MW-14 four to six weeks after completion of development.



#### **APPENDIX B**

# STANDARD OPERATING PROCEDURES SOIL GAS SAMPLING

# STANDARD OPERATING PROCEDURES SOIL GAS PROBES AND VAPOR SAMPLING

Soil gas samples will be collected utilizing direct push drilling techniques with active soil gas sampling with Post Run Tubing (PRT). The probe depth will be selected based on lithology and PID reading observed in pilot holes completed adjacent to the proposed soil gas sampling location, or on the documented utility depths. The target depth for the soil gas probes is approximately 5 to 7 feet below the ground surface, while still remaining above the groundwater surface and avoiding utilities.

The direct push sampler, equipped with a PRT point sampler will be advanced to the target collection depth. The PRT tubing is connected to the PRT point sampler and a vacuum pump. The tubing is checked for leakage by applying a vacuum to the line and an in-line pressure gauge is monitored for a loss of vacuum. Once the tubing is determined to be air-tight, the drill rod is retracted approximately 6-inches to expose the PRT point sampler. Approximately 2.0 volumes of air are purged from the sample tube before a sample is collected using a syringe of at least 100 ml. The sample is injected directly into the GC for field analysis of chlorinated VOCs in accordance with USEPA Method 8260.

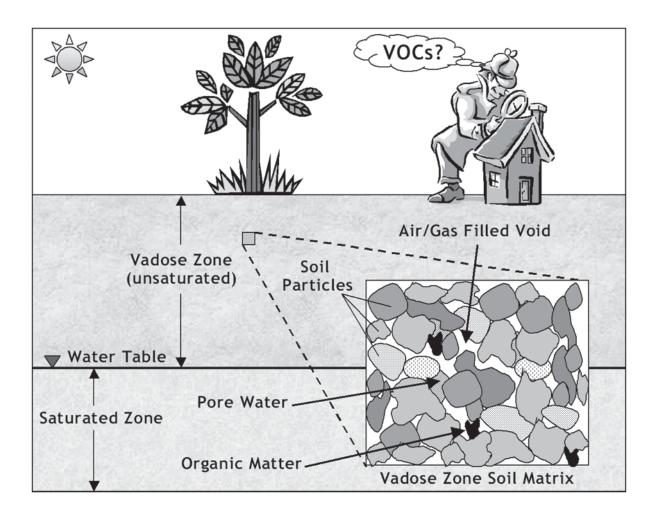
Specific details on the standard operating procedures for the Geoprobe® PRT sampling are provided Geoprobe Technical Bulletin MK3098 (May 2006) (attached). Soil samples corresponding to the soil gas sample horizon will be collected using Geoprobe® Macrocore sampler. These samples will be analyzed at an off-site fixed laboratory for VOCs. Soil samples designated for VOC analysis will be collected using EnCore® or equivalent sampling device following EPA Method 5035 sampling protocol.

Lithologic logs will be prepared for all probe locations. Boring holes will be sealed with bentonite.

Vapor samples will be collected from airspace within the interiors of select manholes, sewer cleanouts, pipes, conduit access boxes, etc. The airspace sample will collected using a Tedlar bag and vacuum box, or summa canister. New or decontaminated tubing will be lowered into space being sampled, taking care to avoid submersion in any water that may be present. If possible, the access location should be covered and allowed to sit/equilibrate for a minimum of 30 minutes between insertion of the tubing and sample collection to limit the introduction of atmospheric air.

# **Direct Push Installation of Devices for Active Soil Gas Sampling & Monitoring**

Technical Bulletin No. MK3098 Prepared: May, 2006



Geoprobe Systems®

### Section 1.0 Purpose

This document details the collection of representative soil gas samples with appropriate direct push (DP) methods to meet a range of data quality objectives, site-specific conditions, and regulatory requirements. New requirements for vapor intrusion investigations (U.S. EPA 2002, Cal. DTSC 2003, NYDOH 2005, etc.) establish higher sample data quality objectives to meet the needs for the human health risk assessment for this pathway. Downhole tools and sample collection methods needed to obtain the higher quality soil gas samples are reviewed.

### Section 2.0 Background

What is soil gas? Where is the vadose zone? The ASTM Standard Guide D5314 for soil gas monitoring gives us the following definitions:

**soil gas** = vadose zone atmosphere **vadose zone** = the hydrogeological region extending from the soil surface to the top of the principal water table

Soil gas is simply the gas phase (air) that exists in the open spaces between the soil particles in the unsaturated portion of the subsurface (Figure 1). Soil gas is normally comprised primarily of nitrogen and oxygen like the aboveground air it is in equilibrium with. However, when volatile organic compounds (VOCs) have been released in the subsurface due to spills or leaks, they will begin to evaporate from the fluid phase and become a part of the soil gas. Over time the VOCs can migrate through the soil and/or groundwater and present a potential hazard to human health or the environment. Intrusion of volatile contaminant vapors into an occupied building (home, office, school, etc.) can lead to exposure of the occupants to the potentially harmful contaminants (US EPA 2002).

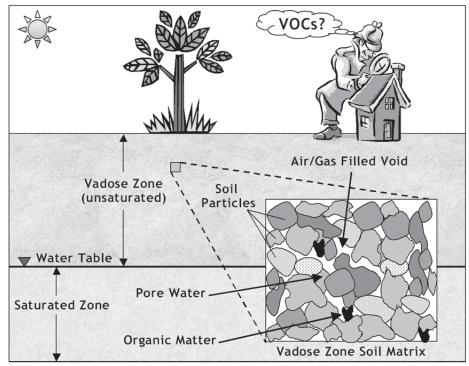


Figure 1: Schematic showing relationship of the vadose zone and saturated zone. Also exploded view showing the primary components of the soil matrix. VOCs will equilibrate with the solid, liquid, and gas phases of the soil matrix. Free product may also be present in the void space of the soil matrix.

Previously, soil gas sampling and monitoring were used primarily as cost- and time-effective means to track soil and groundwater contamination by VOCs and identify potential source areas (US EPA 1997). As our understanding of the vapor intrusion pathway develops, the use of soil gas data is being applied to evaluate this potential human health hazard. Because the vapor intrusion pathway deals with human health risk assessment, the sample and data quality objectives for soil gas samples are more stringent for this application than for simple plume tracking and source location activities. This technology overview will provide guidance on selection of the appropriate tools and methods for soil gas sampling and monitoring depending on the data quality objectives of the project.

It is useful to realize that VOCs will be partitioned between the different components (phases) of the vadose zone (Figure 1) (ASTM D5314, McCall et al. 2006) depending on their physical and chemical characteristics. The VOCs will be present in the soil gas phase, dissolved in any liquid water that may be present, and probably sorbed to the solid soil particles, especially any organic matter. Near the source of contamination, free product also may be present in the soil. The distribution of the contaminant(s) in different phases of the soil matrix is important to consider when sampling. If a strong vacuum is applied to the sample interval, VOCs that were partitioned into the liquid phase or sorbed to the solid matrix may be stripped and enter the vapor phase. This can result in an inaccurate measure of the amount of the contaminant actually in equilibrium with the vapor phase and lead to erroneous conclusions in a risk assessment.

### 3.0 VOCs and Other Soil Gases of Interest

The primary contaminants of interest in soil gas sampling and vapor intrusion studies are chlorinated volatile organic compounds (X-VOCs), aromatic compounds, and some landfill gases (Table 1). The volatility of a compound is primarily a function of its vapor pressure. The higher the compound's vapor pressure (Table 1) at ambient conditions the greater it's volatility. This simply means any spilled liquid will quickly evaporate and enter the gas phase. If spilled on soil or leaked into the subsurface, a compound will tend to enter the soil atmosphere. Don't forget these compounds also will be partitioned into the other phases of the soil matrix (Figure 1). In general, compounds with vapor pressures greater than about 0.5 mm Hg are considered to be detectable with active soil gas methods.

At many sites, VOC releases will contaminate the groundwater and then migrate down gradient with the flowing groundwater. Over time these groundwater contaminant plumes can become sources for VOC migration into the overlying soil gas that may potentially become a vapor intrusion source. The Henry's Law constant of a compound is used to describe its tendency to partition between water and vapor. One way to determine the Henry's Law constant of a compound is to divide the equilibrium concentration of the compound in air by it's concentration in water. This provides a unitless ratio. The higher the Henry's Law constant the more the compound tends to enter the gas phase. Usually compounds with Henry's law constants greater than 0.1 and sufficiently high vapor pressures are considered to be detectable with active soil gas sampling (EPA 1997).

### 4.0 Conceptual Site Model (CSM)

The conceptual site model (CSM) is a simplified picture and/or written description of what may be a complex system (U.S. EPA 2002). The purpose of the CSM is to provide the investigation team and other interested parties an initial idea about the subsurface conditions, possible contaminant migration pathways, potential receptors, local soils and geology, and potential sources for the volatile contaminants under investigation. The CSM should include preliminary site maps, geologic cross sections and other pertinent diagrams. This may include maps of sewer lines, water lines and other underground utility lines that may provide preferential conduits for migration of volatile contaminants to homes, schools, offices and other buildings where vapor intrusion could pose a health hazard. Sensitive environments and endangered species also should be evaluated.

A good CSM will help the investigation team identify additional information that will be required to complete the investigation and determine if the vapor intrusion pathway is complete. The CSM should be viewed as a dynamic tool that will be updated and modified as additional information, samples, and data are obtained that better characterize the site conditions. Soil samples may be required to verify local soil and geological conditions and identify potential source areas. Groundwater samples may be required to determine if a groundwater plume is acting as a source for volatiles across the site area and what the contaminant concentrations are. Measurements of hydraulic conductivity or soil permeability may be needed to evaluate the potential rate of contaminant migration. Several direct push methods are available to obtain these samples and other information needed to confirm or modify the CSM to accurately reflect site specific conditions. Direct push electrical conductivity logs, hydraulic profiling tool (HPT) logs, membrane interface probe (MIP) logs for volatiles, and CPT logs may provide methods to achieve a detailed site model in a relatively short time frame. In addition, ASTM Standards are available for particular methods and procedures for sampling and investigation that may provide useful insights and special knowledge. Some pertinent ASTM Standards particular to direct push methods include D6001, D6282, D6725, and D7242.

### Section 5.0 Data Quality Objectives (DQOs) and Selection of Sampling Methods

As mentioned previously, the data quality objectives will determine how samples are obtained in the field and analyzed. The project manager and field team will have to determine which downhole tools are required and the specific sample collection and analysis methods that should be used to meet the DQOs of the project. But first, what are data quality objectives? DQOs are basically a set of rules that define the amount of uncertainty that is acceptable in the results obtained from an environmental investigation. How confident are you that the sample you collected is representative? Are you sure the analytical method applied accurately identifies the contaminant and what its concentrations are in the sample?

One of the first steps in determining your DQOs is to clearly define the regulatory requirements that must be met to determine if the potential hazard (e.g. vapor intrusion) is an actual threat to the population of concern. Review the state or federal regulations and guidance documents (Cal RWQCB 1997,Cal.DTSC 2003,NYDOH 2005,U.S.EPA 2002,etc.) to be sure you know what is required and how the data is to be used. The following outline of objectives for three different projects may help clarify where and when different levels of data quality are required.

#### Project 1:

- Are VOCs present in soil gas?
- What are gross concentration ranges?
- What is approximate extent of the contamination?

#### Project 2:

- What are the specific types of VOCs present in the soil gas?
- What are approximate concentrations of each contaminant type at each location?
- Define and map the extent of the contaminant plume and locate source areas.

#### Project 3:

- Determine the specific VOC compounds present in the soil gas.
- Determine the concentration of each compound at each location with good precision and accuracy.
- Define and map the plume for each analyte and locate sources.
- Determine if the contaminants present a threat to human health or the environment (is the vapor intrusion pathway complete?).
- · Obtain data that can be used to determine appropriate remedial actions or if remedial actions are required.

Once you determine what level of data quality is required you can then determine the appropriate sampling tools, sample collection procedures, and analytical methods that are required to meet your objectives. Table 2 may be used to guide you in selecting the appropriate methods to use in the field to be sure the project manager, facility owners, and regulators are able to make the correct decisions. **Establishing project DQOs and selecting the appropriate field methods for sample collection and downhole tool systems is not a trivial process and should be completed well in advance of starting field work.** Details of the sample collection methods and downhole sampling systems listed in Table 2 are discussed in the following sections.

Of course the final data quality is controlled by the analytical method chosen and the analytical quality control that is used. You can collect a low quality sample and spend hundreds of dollars on an expensive analytical method with rigorous quality control, this would be a mistake. If you have low confidence in the sample quality (poorly representative) it will not matter how much you spend on the analysis, the final data will still be of low quality (poorly representative). You can't improve the quality of a bad sample by using expensive analytical methods. In the end the analytical result is only as good as the sample quality. For guidance on selection of the appropriate analytical methods review the appropriate federal or state guidance documents (Cal RWQCB 1997, Cal. DTSC 2003, NYDOH 2005, U.S. EPA 2002, etc.) and/or the SW846 manual (EPA 1986).

#### **5.1 Analytical Methods**

The EPA Draft Guidance for the vapor intrusion pathway includes a table listing more than a dozen analytical methods used to test for VOCs. The purge and trap methods such as 8260C were primarily written for soil and groundwater analysis. Purge and trap methods may be useful if you are conducting passive soil gas sampling and collecting VOCs in a sorbent trap. However, direct injection may be conducted with method 8260C and this provides a useful method for analysis of soil gas samples. Method TO-17 is also a method that requires the use of a sorbent trap for sample collection and later purge and trap gas chromatograph (GC) analysis. This is sometimes used for ambient air monitoring in buildings.

(continued on Page 6)

Table 1: Basic chemical properties ABGD of selected VOCs and gases of interest in soil gas investigations.

tics and Aliphatics         78.11         0.877         0.215         176°F           C <sub>e</sub> H <sub>e</sub> CH, CH <sub>e</sub> CH <sub>e</sub> 78.11         0.87         0.262         232°F           C <sub>e</sub> H <sub>e</sub> CH <sub>e</sub> CH <sub>e</sub> 106.2         0.87         0.262         232°F           C <sub>e</sub> H <sub>e</sub> CH <sub>e</sub> CH <sub>e</sub> CH <sub>e</sub> 106.2         0.88 to 0.86         0.304         269°F to 292°F           ther         CH <sub>e</sub> CH <sub>e</sub> CH <sub>e</sub> CH <sub>e</sub> CH <sub>e</sub> 114.2         0.70         258°F           ther         CH <sub>e</sub> CHCI         62.5         0.911         (1.1E-2)         258°F           ther         CH <sub>e</sub> CHCI         62.5         0.911         (1.1E-2)         13.3           thane         CHCI, CH <sub>g</sub> 99.0         1.18          133°F           whene         (CH <sub>g</sub> CH <sub>g</sub>	Chemical Name	Molecular Formula	Molecular Weight	Density (g/cc)	Henry's Law Constant Ratio <sup>c</sup> (atm·m³/mol) <sup>8</sup>	Boiling Point (°C)	Water Solubility %^ (g/L @ 25°C) <sup>8</sup>	Vapor Pressure (mm Hg @ 68°F)	Viscosity <sup>®</sup> (mPa·s @ 25°C, 1 atm)	lonization Potential (eV)
C_0H_0   P   P   P   P   P   P   P   P   P	Aromatics and	Aliphatics								
CeH,CH, Decomposition         92.1         0.87         0.262         232°F           Izene         CeH,CH,CH, CH, CH, CH, CH, CH, CH, CH, CH,	Benzene	C,H,	78.11	0.877	0.215	176°F	0.04% @ 54°F (1.77)	75	0.604	9.24
rzene C <sub>0</sub> H <sub>3</sub> CH <sub>3</sub> CH <sub>3</sub> rzene C <sub>0</sub> H <sub>3</sub> CH <sub>3</sub> CH <sub>3</sub>	Toluene	C <sub>6</sub> H <sub>5</sub> CH <sub>3</sub>	92.1	0.87	0.262	232°F	0.05% @ 61°F (0.53)	20 (at 65°F)	0.56	8.82
central parts         Central (CH <sub>2</sub> ) <sub>2</sub> CH <sub>3</sub> 106.2         0.88 to 0.86         0.304         269F to 292F           butylether         CH <sub>3</sub> (CH <sub>2</sub> ) <sub>2</sub> CH <sub>3</sub> 114.2         0.70         258°F           butylether         R8.15         0.74         0.018         55.2           Chlorinated VOCs         88.15         0.74         0.018         55.2           oride         CH <sub>2</sub> CHCl         62.5         0.911         (1.1E-2)         -13.3           oride         CHCl <sub>2</sub> CH <sub>2</sub> CH <sub>3</sub> 99.0         1.18          135°F           oroethylene         CHCl <sub>2</sub> CH <sub>3</sub> CH <sub>3</sub> 97.0         1.27         (3.0E-2)         118° to           hloroethane         CHCl <sub>2</sub> CH <sub>3</sub> CI         133.4         1.44          237°F           hloroethylene         CICH=CCl <sub>2</sub> 131.4         1.46          250°F         1           ethylene         CIC=CCl <sub>2</sub> 15.8         1.62          250°F         1           reachloride         CCl <sub>4</sub> 15.38         1.59          170°F           Landfill Gases         CO         AA         AA          16.04          -161.5	Ethyl Benzene	C <sub>6</sub> H <sub>5</sub> CH <sub>3</sub> CH <sub>2</sub>	106.2	0.87	0.322 (8.5E-3)	277°F	0.01% (0.16)	10 (at 79°F)	0.631	8.76
butyl ether  butyl ether  butyl ether  Chlorinated VOCs  Chlorioethene  Chlorioethene  ChCl, CH, CH, CH, CH, CH, CH, CH, CH, CH, CH	Xylenes (o,m,p isomers)	$C_6H_4(CH_3)_2$	106.2	0.88 to 0.86	0.304	269°F to 292°F	$\sim 0.01\%$ (0.16 to 0.18)	6/6/2	0.760, 0.581, 0.603	8.44 to 8.56
Chlorinated VOCs         88.15         0.74         0.018         55.2           Chlorinated VOCs         Chloroethene)         62.5         0.911         (1.1E-2)         -13.3         -13.3           oride         Chloroethene)         62.5         0.911         (1.1E-2)         -13.3         -13.3           oroethylene         CHCI, CH, CH         99.0         1.18          135°F           hloroethylene         CHCI, 2-DCE)         133.4         1.44          237°F           ethylene         CHCI, CH, CH         131.4         1.46          250°F         1           ethylene         CI, C=CCI, 2         15.8         1.62          250°F         1           etrachloride         CCI, 4         15.9          170°F           Landfill Gases         CH, 4         0.4228*2          -161.5	(n-)Octane	CH <sub>3</sub> (CH <sub>2</sub> ), <sub>C</sub> CH <sub>3</sub>	114.2	0.70		258°F	~insoluble	10	0.508	9.82
lorinated VOCs         CH2CHCI         62.5         0.911         (1.1E-2)         -13.3           CCH2CHCI         62.5         0.911         (1.1E-2)         -13.3           (chloroethene)         99.0         1.18          135°F           ylene         (CH) <sub>2</sub> Cl <sub>2</sub> 97.0         1.27         (3.0E-2)         118° to           thane         (CHC <sub>2</sub> CH <sub>2</sub> Cl         133.4         1.44          237°F           thane         CICH=CCl <sub>2</sub> 131.4         1.46          250°F           Iene         CI <sub>2</sub> C=CCl <sub>2</sub> 165.8         1.62          250°F           oride         CCl <sub>4</sub> 153.8         1.59          170°F           andfill Gases         CH <sub>2</sub> 16.04         0.4228 <sup>62</sup> -161.5	Methyl t-butyl ether (MTBE) <sup>D</sup>		88.15	0.74	0.018 (5.5E-4)	55.2	(51.2)	245		1
CH2CHCI         62.5         0.911         (1.1E-2)         -13.3           (chloroethene)         (chloroethene)         99.0         1.18          135°F           vlene         (CH) <sub>2</sub> Cl <sub>2</sub> 97.0         1.27         (3.0E-2)         118° to           vlene         (CH) <sub>2</sub> Cl <sub>2</sub> Cl         133.4         1.44          237°F           thane         CHCI <sub>2</sub> CH <sub>2</sub> Cl         131.4         1.46          189°F           inche         CI <sub>2</sub> C=CCl <sub>2</sub> 165.8         1.62          250°F           oride         CCl <sub>4</sub> 153.8         1.59          170°F           andfill Gases         CCl <sub>4</sub> 16.04         0.4228° <sup>62</sup> 161.5	Chlorinateo	IVOCs								
sine         CHCI2 CH3         99.0         1.18          135°F           ylene         (CH)2G2         97.0         1.27         (3.0E-2)         118° to           thane         (CHCI2CH2CI         133.4         1.44          237°F           thane         CHCI2CH2CI         131.4         1.46          237°F           iene         CICH=CCI2         131.4         1.66          250°F           iene         CI2C=CCI2         165.8         1.62          250°F           oride         CCI4         153.8         1.59          170°F           andfill Gases         CH4         16.04         0.4228°62           161.5	Vinyl Chloride	CH <sub>2</sub> CHCl (chloroethene)	62.5	0.911	(1.1E-2)	-13.3 (7°F)	~0.1% @ 77°F (2.8)	>760		66.6
ylene         (CH) <sub>2</sub> Cl <sub>2</sub> 97.0         1.27         (3.0E-2)         118° to           thane         (I,2-DCE)         133.4         1.44          237°F           thane         CHCl <sub>2</sub> CH <sub>2</sub> Cl         131.4         1.46          189°F           ne         CICH=CCl <sub>2</sub> 165.8         1.62          250°F           lene         Cl <sub>2</sub> C=CCl <sub>2</sub> 165.8         1.62          250°F           oride         CCl <sub>4</sub> 153.8         1.59          170°F           andfill Gases           CO         44.0	1,1 Dichloroethane (1,1-DCA)	CHCI <sub>2</sub> CH <sub>3</sub>	0.66	1.18	1	135°F	%9.0	230mm (at 77°F)	0.464	11.06
thane CHCI <sub>2</sub> CH <sub>2</sub> CI 133.4 1.44 237°F (TCA)  ne CICH=CCI <sub>2</sub> 131.4 1.46 189°F (TCE)  lene CI <sub>2</sub> C=CCI <sub>2</sub> 165.8 1.62 250°F (Perchloroethene)  oride CCI <sub>4</sub> 153.8 1.59 170°F (TCE)  andfill Gases  CO 440 161.5	1,2-Dichloroethylene	(CH) <sub>2</sub> Cl <sub>2</sub> (1,2-DCE)	97.0	1.27 (at 77°F)	(3.0E-2)	118° to 140°F	0.4%	180 to 264	Cis = 0.445 Trans = 0.317	9.65
ne CICH=CCI₂ 131.4 1.46 189°F (TCE) lene CI₂C=CCI₂ 165.8 1.62 250°F (Perchloroethene)  vride CCI₄ 153.8 1.59 170°F (TCE)  andfill Gases CCI₄ 16.04 0.4228 <sup>162</sup> 161.5	1,1,2-Trichloroethane	CHCI <sub>2</sub> CH <sub>2</sub> CI (TCA)	133.4	1.44	1	237°F	0.4% (4.4)	19		11.0
Iene         CI <sub>2</sub> C=CCI <sub>2</sub> 165.8         1.62          250°F           Oride         CCI <sub>4</sub> 153.8         1.59          170°F           andfill Gases         CH <sub>4</sub> 16.04         0.4228"62          -161.5	Trichloroethylene	CICH=CCI <sub>2</sub> (TCE)	131.4	1.46	1	189°F	0.1% @ 77°F	28	0.545	9.45
andfill Gases         CCI <sub>4</sub> 153.8         1.59          170°F           A4.0           -161.5	Tetrachloroethylene	Cl <sub>2</sub> C=CCl <sub>2</sub> (Perchloroethene)	165.8	1.62	1	250°F	0.02% @ 77°F (1.2)	14	0.844	9.32
andfill Gases  CH₄  16.04  0.4228 <sup>162</sup> -161.5  CO  AA 0   cublines	Carbontetrachloride	CCI⁴	153.8	1.59	1	170°F	0.05% (1.2)	91	0.908	11.47
CH <sub>4</sub> 16.04 0.4228 <sup>162</sup> 161.5	Landfill G	ases								
30 M OJ	Methane	CH₄	16.04	0.4228 <sup>-162</sup>	-	-161.5	Soluble	-	1	12.98
	Carbon dioxide	$CO_2$	44.0			sublimes	0.2% @ 77°F	>760		13.77

mm Hg = millimeters of mercury. mPa·s = millipascal second = centipoise. atm = atmosphere. eV = electron volt
A) NIOSH Pocket Guide to Chemical Hazards, 1990. B) CRC Handbook of Chemistry and Physics, 1995. C) U.S. EPA, 1997. D) www.epa.gov/opptintr/chemfact/s\_mtbe.txt g/cc = grams per cubic centimeter. atm·m3/mol = atomosphere cubic meter per mole. °C = degrees centigrade. °F = degrees Fahrenheit. g/L = grams per liter.

Table 2: Matrix for selection of field methods to achieve DQOs.

Downhole			Sample Colle	ction Metho	d
Sampling System		Syringe	Tedlar Bag	Glass Bulb	Summa Canister
	Increasing Quality				-
Direct Sampling		Low / Low			Low / High
PRT System					
Implants					
Gas Wells		High / Low			High / High

Dashed arrow shows general path for selection of methods that will result in sampling and collection with compatible quality procedures to achieve desired DQO level. If method pairs in the upper right or lower left corner are used the final sample analytical quality will be of low confidence regardless of the analytical method applied. To achieve intermediate DQOs for an initial survey the PRT system is often combined with collection in Tedlar bags or glass sampling bulbs and field analysis utilizing a mobile lab equipped with a gas chromatograph.

For active soil gas sampling you will probably be involved with one or more of the following methods:

Method #	Analytes	Instruments	Det. Limits
8260	Aromatic, Halogenated and other VOCs	GC/MS	50 ug/m³
8260 SIM	Aromatic, Halogenated and other VOCs	GC/MS	10 ug/m³
8021	Aromatic and Hologenated VOCs	PID/ELCD-GC	> 5 ug/m <sup>3</sup>
TO-15	VOCs in Air with Summa Canisters	GC/MS	1 to 10 ug/m <sup>3</sup>
TO-17	VOCs in Air with Sorbent Tubes	GC/MS	0.5 to 25 ppbv

 $\mathsf{GC} = \mathsf{Gas} \ \mathsf{Chromatograph} \ \ \mathsf{MS} = \mathsf{Mass} \ \mathsf{Spectrometer} \ \ \mathsf{PID} = \mathsf{Photo} \ \mathsf{Ionization} \ \mathsf{Detector}$ 

ELCD = Electrolytic Conductivity Detector

The 8021 Method is often modified for use in a mobile lab in the field. Air samples are direct injected on column for analysis. Sometimes an ECD (electron capture detector) is used instead of the ELCD( electrolytic conductivity detector) to detect the chlorinated VOCs. Field analysis with the 8021 method allows the field team to track/map plumes while in the field and locate source areas. The field analysis is also a useful way of screening samples to determine which are best submitted for more expensive off-site analysis. Knowing approximate sample concentrations also may be used to select the appropriate lab analysis.

For soil gas samples with elevated concentrations the standard 8260 method will provide sufficient sensitivity and detection limits. Samples with lower concentrations may be suitable for the 8260 SIM (selected ion monitoring) method. Low concentration samples requiring lower detection limits and high data quality may require the TO-15 method. Be aware that not all labs provide the same level of data quality for the same method. A cheaper price may mean less QA/QC. Be sure the final data package will meet your data quality requirements.

#### Section 6.0

## Downhole Soil Gas Sampling Methods for Preliminary Assessments and Plume Tracking

### 6.1 Methods to Advance Tool Strings

Several methods for soil gas sampling were developed in the 1980s and used to track VOC plumes and locate potential source areas. One of the earlier methods included manually driving galvanized steel pipe to the desired depth with a slam-bar or fence-post-driver. While physically demanding, these manual methods may still prove valuable in accessing delicate ground in residential areas (landscaped plant beds, etc.) or where access is otherwise limited. Improvements to this manual method have been made to improve sample quality and minimize the physical demands for driving and removing the probe rods (Figure 2) (Geoprobe Systems®, 2001). When site access conditions permit, soil gas sampling tools may be advanced into the subsurface with a Geoprobe® direct push machine (Figure 3) to minimize the physical effort required and speed up the sampling process.

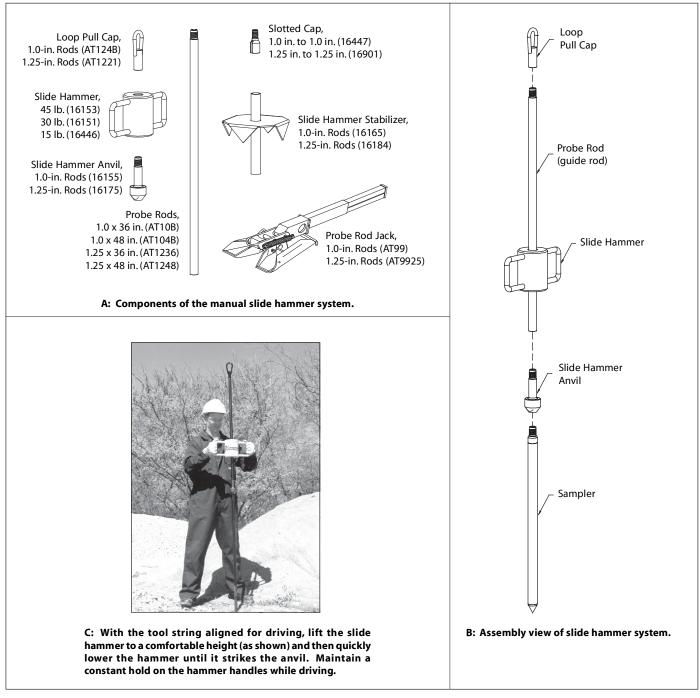


Figure 2: Components, assembly, and operation of the manual sampling tools.



Figure 3: Using a Geoprobe® direct push machine to advance soil gas sampling tools to depth. Note use of portable vacuum/volume system for purging and sampling.

### **6.2 Direct Sampling System**

Initially the collection of soil gas samples was conducted directly through the rods driven to depth (Figure 4). An expendable point was advanced to depth on the leading end of the drive rods. Then the rods were retracted a specified distance to release the expendable point and create an open void where soil gas could enter the rod string for sampling. A gas sampling cap with appropriate fitting was attached to the top of the drive rods and gas was purged and then collected for analysis. While this method is quick and cost effective there is potential for leaks at each rod joint that could compromise sample quality. This method also requires that the steel rods be thoroughly decontaminated between each use. In addition, some contaminants may be sorbed or degraded on the metal surface of the drive rods, especially if the rods are rusty.

### 6.3 PRT Sampling System

The Post Run Tubing (PRT) system was developed to help eliminate some of the problems with sampling directly through the steel rods. The PRT method uses an adapter and tubing to isolate the soil gas sample from the steel drive rods (Figure 5). This also eliminates possible leaks of ambient air from the rod joints into the gas being sampled. Tool configurations for soil gas sampling with the PRT system using different size probe rods are shown in Figure 6. PRT adapters (Table 3) must be selected to fit the desired tubing (Table 4) that will be used for sample collection.

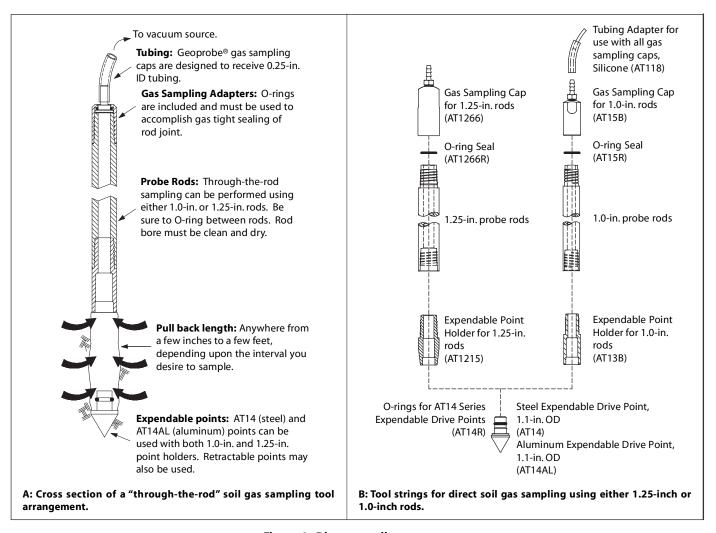


Figure 4: Direct sampling systems.

The basic steps to conduct soil gas sampling with the PRT system are as follows:

- An expendable point holder is attached to the lead rod.
- The expendable point is placed in the point holder, use O-ring.
- The tool string is advanced to the desired sampling depth. Additional probe rods are added as required.
- · Next, the stainless steel PRT adapter with appropriate sized barb is attached to the selected tubing cut to length.
- The drive cap is removed from the tool string and the PRT Adapter is lowered on the tubing down the tool string.
- The PRT adapter is threaded into the reverse thread fitting in the top of the PRT expendable point holder by manually rotating the tubing counterclockwise until the O-ring on the adapter seals against the point holder. The top end of the tube may be sealed to minimize gas exchange with the ambient air and possible loss of VOCs.
- A leak test of the PRT tubing system may be conducted at this time by applying vacuum to the tubing. An in-line gauge may be used to verify that leaks do not occur after the vacuum is applied.
- The direct push machine is then used to retract the probe rods a specified interval, usually 6 to 12 inches (or 150 to 300 mm). This provides an open cylinder in the soil through which the soil gas may be purged and sampled.
- The upper end of the tube is connected to the purging/sampling system and the sample is collected.
- The direct push machine is used to remove the probe rods. Once at the surface, the O-ring seal between the PRT adapter and the expendable point may be visually inspected to further verify sample integrity.

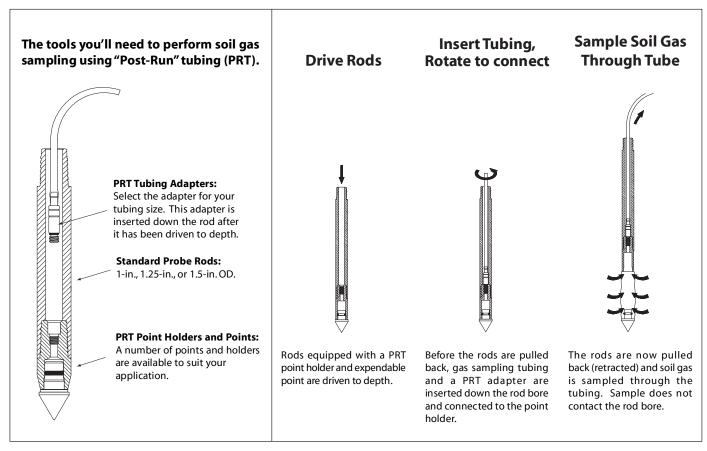


Figure 5: PRT sampling system.

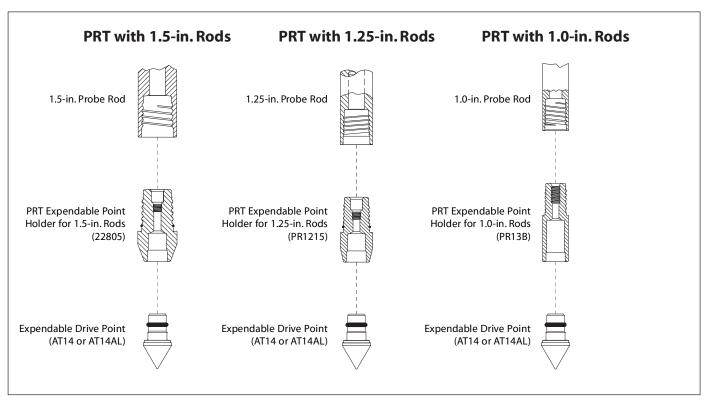


Figure 6: Tool configurations for PRT soil gas sampling.

Table 3: PRT adapter selection guide.

PRT Adapter Selection Guide			
PRT ADAPTER and PART NUMBER	RECOMMENDED TUBING SIZE (ID)	GEOPROBE® TUBING PART NO.	TUBING INTERNAL VOLUME
PR12S	1/8 in.	TB12T	2.4 mL/ft
	(3.2 mm)	Teflon®	(7.9 mL/m)
	0.17 in.	TB17L	4.5 mL/ft
	(4.3 mm)	LDPE	(14.6 mL/m)
PR17S	3/16 in.	TB17T	5.4 mL/ft
	(4.8 mm)	Teflon®	(17.8 mL/m)
PR25S	1/4 in.	TB25L	9.7 mL/ft
	(6.4 mm)	LDPE	(31.7 mL/m)
	5/16 in.	TB30T	15.1 mL/ft
	(7.9 mm)	Teflon®	(49.5 mL/m)
O-Rings for all PRT Adapters PR25R (pack of 25)			

**NOTE:** The sorption characteristics of certain tubing may not permit their use in all soil gas sampling applications.

Table 4: Tubing selection guide for PRT adapters and soil gas implants.

Part Number	Tubing Material	Inside Diameter, inches (mm)	Outside Diameter, inches (mm)	Wall Thickness, inches (mm)	Length, feet (m)	Tubing Internal Volume, mL/ft (mL/m	PRT Adapter Part Number	Implant Part Numbers <sup>1</sup>
TB17L	LDPE <sup>2</sup>	0.17 (4.8)	1/4 (6.4)	0.040 (1.0)	500 (152)	4.5 (14.6)	PR17S	AT8617S AT8717S
TB25L	LDPE <sup>2</sup>	1/4 (6.4)	3/8 (9.5)	1/16 (1.6)	500 (152)	9.7 (31.7)	PR25S	AT8625S AT8725S
TB37L	LDPE <sup>2</sup>	3/8 (9.5)	1/2 (12.7)	1/16 (1.6)	100 (31)	21.7 (71.2)	NA	AT9637 <sup>3</sup> AT9638 <sup>4</sup>
TB12T	Teflon®	1/8 (3.2)	1/4 (6.4)	1/16 (1.6)	50 (15)	2.4 (7.9)	PR12S	AT8612S AT8712S
TB17T	Teflon®	3/16 (4.8)	1/4 (6.4)	1/32 (0.8)	50 (15)	5.4 (17.8)	PR12S	AT8612S AT8712S
ТВЗОТ	Teflon®	5/16 (7.9)	3/8 (9.5)	1/32 (0.8)	50 (15)	15.1 (49.5)	PR30S	AT8630S AT8730S
TB45SL	Silicone	3/16 (4.8)	7/16 (11)	1/8 (3.2)	25 (8)	-	NA	NA
TB50TY	Vinyl	1/4 (6.4)	7/16 (11)	3/32 (2.4)	25 (8)	-	NA	NA
TB100TY	Vinyl	1/4 (6.4)	7/16 (11)	3/32 (2.4)	100 (31)	-	NA	NA
AT82	Stainless Steel	0.085 (2.2)	1/8 (3.2)	0.040 (1.0)	50 (15)	-	NA	AT86SW12

<sup>1 - 86</sup> series implants are 6 in. (152 mm) long and 87 series implants are 21 in. (533 mm) long.

<sup>2 -</sup> Low-density polyethylene.

<sup>3 - 96</sup> series implants are for use in 1.25-in. or 1.5-in. OD probe rods with an ID of 0.625 in. (15.9 mm). Screen is 12 in. (305 mm) long. Use with any plastic tubing with an ID ranging from 1/4 in. to 7/16 in. (6.4 mm to 11 mm).

<sup>4 -</sup> This implant is designed for air sparging only and is equipped with a porous polyethylene lining.

### 6.4 Vapor Implants / Small ID Rods

These simple devices allow the investigator to install a small screen at the desired depth to conduct long-term monitoring of soil gas in the subsurface. The implants may be installed through 1.0-,1.25-, or 1.5-inch outside diameter probe rods for many applications. However, when installed through these smaller rods it may not be possible to achieve the high-integrity seal required for human health risk assessments as in evaluation for the vapor intrusion pathway. High-integrity seals can be obtained by installing the implants through larger diameter rods as discussed later in Section 7.0.

The implants consist of a stainless steel wire mesh screen with a threaded fitting on the bottom for anchoring and a fitting at the top to connect to tubing (Figure7). The AT86 series screens are 6 inches (152 mm) long, the AT87 series screens are 21 inches (533 mm) long, and both have an OD of less than 0.5 inches (13 mm). The AT96 series screens are 12 inches (305 mm) long with an OD of less than 0.625 inches (16 mm). All have a pore diameter of approximately 0.006 inches (0.15 mm) in the stainless screens (Table 5).

The implants are installed through the drive rods after they are advanced to the desired depth (Figure 8). An expendable implant anchor/drive point at the lead end of the rod string has a threaded fitting at the upper end and the implant is anchored in place by threading in counterclockwise. As the rods are retracted with the direct push machine, fine glass beads (P/N AT84) may be poured in around the screen to prevent clogging. Once the rods are above the screen, a bentonite powder mix (P/N AT85) may be added to the rod annulus to seal the probe hole. Do not hydrate the bentonite until the rod string is completely removed from the ground. The tubing should be capped to prevent venting to the ambient atmosphere until ready for sampling. A simple PVC pipe with cap may be installed over the implant tubing at the surface for protection. Conventional well protectors (P/N WP1771 or WP1741) may be used if necessary.

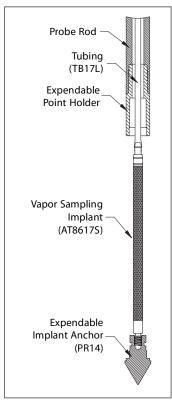


Figure 7: Schematic of a soil gas implant assembly.

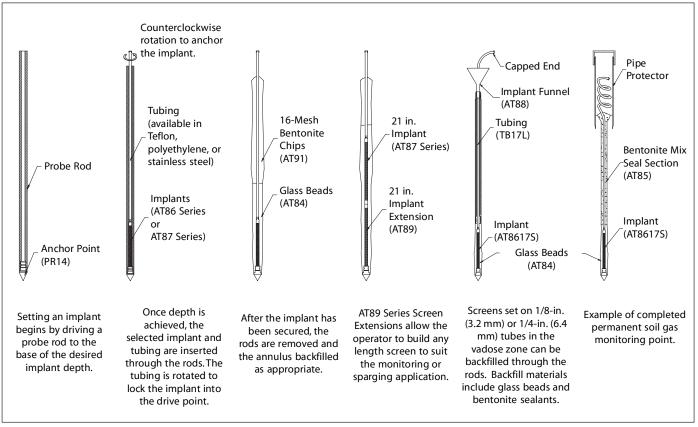


Figure 8: Installing implants through small-diameter probe rods.

Table 5: Implant selection guide.

END	SUGGESTED RECOMMENDED SAMPLING TUBING		GEOPROBE®	IMPLANT PART NUMBERS	
FITTING	USE	SIZE	TUBING PART NO.	6-in. (152 mm) SCREEN LENGTH	21-in. (533 mm) SCREEN LENGTH
Barbed	Soil Gas	1/8 in. ID (3.2 mm ID)	TB12T	AT8612S	AT8712S
	Soil Gas	0.17 in. ID (4.3 mm ID)	TB17L	AT8617S	AT8717S
Barbed	Jon das	3/16 in. ID (4.8 mm ID)	TB17T		
Barbed	Soil Gas, Shallow Groundwater	1/4 in. ID (6.4 mm ID)	TB25L	AT8625S	AT8725S
Barbed	Soil Gas, Shallow Groundwater	5/16 in. ID (7.9 mm ID)	TB30T	AT8630S	AT8730S
		1/8 in. OD (3.2 mm OD)	AT82 (1/8 in. stainless)	AT86SW12	_
	Soil Gas	3/16 in. ID (4.8 mm ID)		AT86SW17	
Swagelok™		1/4 in. OD (6.4 mm OD)	_	AT86SW25	AT87SW25
DDT Conduct	Soil Gas, Shallow Groundwater	Screen Extension (for use with AT87 Series Implants)			AT89
PRT Socket					

For 1.25-in. and 1.5-in. Probe Rods Only		RECOMMENDED TUBING SIZE	GEOPROBE® TUBING PART NO.	12-in. (305 mm) SCREEN LENGTH
Tapered Screw Thread	Soil Gas, Shallow Groundwater, Air sparging (AT9638)	Any plastic tubing 1/4 in. ID to 7/16 in. ID (6.4 mm to 11 mm ID)	TB37L	AT9637 (sampling) AT9638 (sparging)

AT86 Series: Screen is 0.25 in. ID  $\times$  6 in. long (6.4 mm  $\times$  152 mm) with 0.15 mm pore openings. AT86 implants will fit through 0.5-in. (12.7 mm) bore tooling.

AT87 Series: Screen is 0.25 in. ID x 21 in long (6.4 mm x 533 mm) with 0.15mm pore openings. AT87 implants will fit through 0.5-in. (12.7 mm) bore tooling.

AT9637 and AT9638 Implants: Screen is 3/8 in. ID x 12 in. long (9.5 mm x 305 mm) with 0.15mm pore openings. These implants require a minimum 0.625-in. (15.9 mm) rod bore for passage.

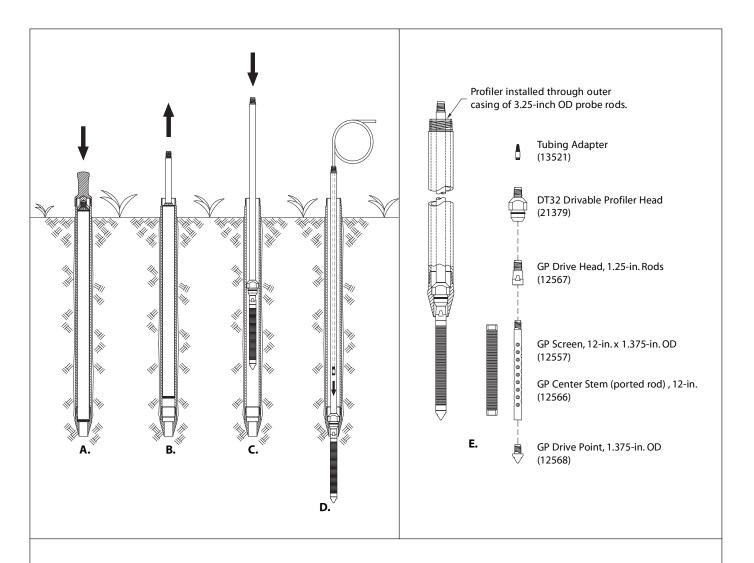
AT9638 implants are the only implants recommended for gas sparging (injection) as they have a porous polyethylene lining inside the implant screen. This interior filter prevents the implant from clogging with formation fines which may backflow into standard implants during periods of pressure shut down.

#### 6.5 Soil Gas Profiling with Dual Tube Systems

In some situations it may be necessary to conduct vertical profiles of soil gas to determine if there is a significant change in concentration or contaminant type with depth. The methods discussed in previous sections could be used to obtain profiles, but a separate advancement would be required for each depth interval. Conversely, dual tube systems can provide access to soil gas or ground water at multiple depths during a single advancement of the tool string. The DT325 soil sampling system can be combined with a simple screen to conduct profiles of soil gas or groundwater if necessary (Figure 9). A summary of the procedure for soil gas profiling follows:

- The equipment operator may conduct continuous soil sampling as the tool string is advanced or install a solid drive point (P/N 28509) if soil sampling is not required.
- Once the cutting shoe is immediately above the desired soil gas sampling interval, the sample sheath and soil sample or solid drive point is removed.
- The profiler screen assembly (Figure 9) with a 12-inch (305 mm) screen (P/N 14402) or 6-inch (152 mm) screen (P/N 14401) is then attached to the inner drive rods and lowered through the annulus of the 3.25-inch rods.
- The screen is pushed through the open cutting shoe and advanced into the formation until the O-ring on the profiler head (P/N 21379) seals in the throat of the cutting shoe.
- At this point the tubing adapter (P/N 13521) is attached to the selected tubing (Table 4) and lowered through the bore of the inner rods.
- The inner tube and adapter are rotated counterclockwise to thread the tubing adapter into the top of the screen drive head (P/N 12567 or 16913).
- This then provides a sealed sampling system for collection of soil gas samples.
- Once the soil gas sample is collected, the tubing is removed and the inner rods and screen are retracted and removed.
- To advance to the next depth, either the solid drive point is installed in the cutting shoe or a soil sample barrel is installed for the next interval. The tool string is advanced to the next desired depth (with removal of soil cores as necessary).
- Installation of the screen and tubing is repeated for the next soil gas sample (Figure 9, Steps C and D).

Once the soil gas profiling is complete, it is possible to install permanent implants at multiple depths as the 3.25-inch rod string is retracted. See Section 7.3 for further discussion of this procedure. Be sure to soil core through any intervals where soil gas implants are to be installed. This will help minimize compression of the formation and loss of gas permeability. If permanent implants will not be installed, abandon the probe hole as required by local regulations (see Section 9.0).



- A. Soil sampling as the tool string is advanced to minimize compression of the soil and loss of gas permeability.
- B. Extracting the soil sample.
- C. Installing the profiler screen through the rod bore.
- D. The selected tubing (Table 4) with attached tubing adapter (P/N 13521) are lowered through the bore of the inner rod string and threaded into the Screen Drive Head (P/N 12567). The soil gas is purged and sample is collected.
- E. Components of the profiler screen assembly for use with the DT325 system. The 12-inch (305 mm) screen assembly is shown here (P/N 14402). A 6-inch (152 mm) screen assembly (P/N 14401) is also available.

Figure 9: Soil gas profiling with the DT325 system.

#### Section 7.0

## Downhole Soil Gas Monitoring Methods for Evaluation of the Vapor Intrusion Pathway

Data obtained for evaluation of the vapor intrusion pathway (VIP) will generally be used to conduct a human health risk assessment (HHRA). Because the data will be used for a HHRA, the data quality objectives are very high and most of the soil gas sampling methods described in previous sections may not meet the sample quality requirements. To achieve the higher DQOs and to permit long-term monitoring of soil gas, most regulatory agencies are requiring the use of permanent soil gas monitoring wells or implants with filter media around the screens and high-integrity seals. In some situations, multilevel monitoring is required to define the vertical distribution of soil gas in the subsurface and evaluate the potential for contaminant degradation. The following sections outline some field methods that will enable the investigator to achieve the higher DQOs when either single-level or multilevel monitoring of soil gas is required.

### 7.1 Single-Depth High-Integrity Soil Gas Implants for Monitoring

If the installation of implants with smaller probe rods as described previously will not meet the higher DQOs for your project, the DT21 dual tube system may be required. This system uses 2.125-inch (54 mm) OD by 1.5-inch (38 mm) ID outer probe rods and 1-inch (25 mm) OD inner rods to provide access to the desired sampling interval either by use of a solid drive point or soil coring system (Figure 10, see also Technical Bulletin # 982100). To minimize compression of the formation and reduction of gas permeability, it is best to stop the DT21 system just above the interval where soil gas is to be sampled. The following steps (Figure 11) should be followed to install an implant for long-term soil gas monitoring:

- Once at the desired depth, the inner rod string and any soil sample is removed.
- A small thin-walled sample tube (P/N 19484) with drive head (P/N DT4055) is lowered through the outer rods and advanced ahead of the cutting shoe to remove a soil core where the implant will be installed (Figure 11). This sample tube (P/N 19484) is designed for low-density sands and finer-grained materials only.
- Once the soil core is removed a stiff brush (P/N BU600) may be lowered by extension rods (P/N AT671, etc.) into the open core hole in fine-grained or otherwise cohesive formations. The brush is rotated and moved up and down gently through the cored interval to relieve any smearing and reestablish natural gas flow into the void where the implant will be installed.
- The appropriate 6-inch (152 mm) or 12-inch (305 mm) long implant (Table 5) is attached to the selected tubing (Table 4) and lowered into the open core hole below the probe rods. Cap or cover the end of the tube at the surface to prevent clogging and/or loss of volatiles.
- Next, either environmental sand (P/N AT95) or fine glass beads (P/N AT93) are
  poured through the rod bore to fill the open hole around the implant screen.
  The probe rods should be retracted slowly to keep the filter media below the
  cutting shoe. Measure with suitable depth indicator to confirm the filter media
  extends at least 6 to 12 inches (152 to 305 mm) above the top of the implant.
- Grouting of the annulus may be accomplished in one of two ways.
  - 1. Fine bentonite chips (P/N AT91) are gravity poured through the rod bore as the rods are slowly retracted to keep chips below the cutting shoe. The chips are hydrated with water every 4- to 6-inch (102 to 152 mm) increment. Water should be added through a tube extending below the cutting shoe. This will prevent wetting the ID of the rods and possible bridging.
  - 2. A grout slurry of either bentonite powder (P/N AT92) or neat cement may be pumped by tremie tube using an appropriate grout machine to fill the bore hole as the drive rods are retracted.

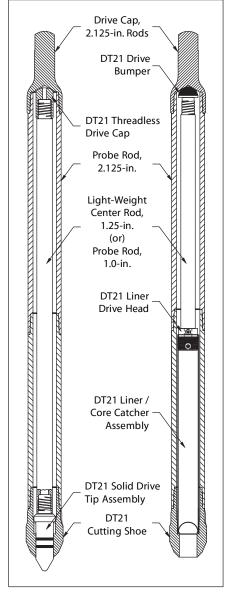
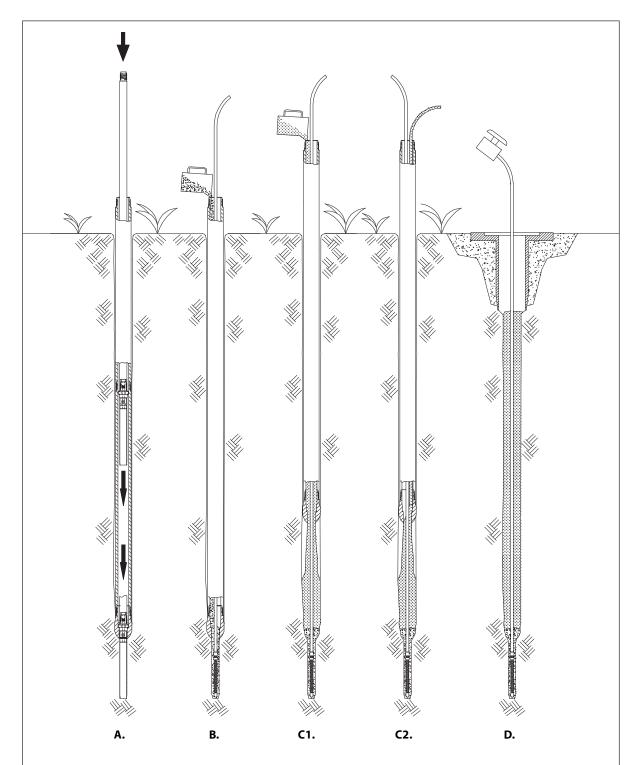


Figure 10: DT21 soil sampling tool string configurations.

• Once grouting is completed, suitable surface protection (P/N WP1771 or WP1741) should be installed to meet local needs and regulatory requirements. In remote locations such as landfills, a simple PVC pipe with slip-cap may provide suitable protection for short-term monitoring requirements.

The DT325 sampling system may also be used in a similar fashion to install high-integrity single-depth implants for soil gas sampling and long-term monitoring.



- A. Advance the DT21 system to depth and remove the inner rod string. The pre-core assembly is attached to the lead inner rod and lowered through the outer rod bore. The thin-walled tube is advanced into the virgin soil ahead of the drive rods to remove a soil core.
- B. The soil gas implant is assembled and lowered into the open core hole with the tubing. Fine sand or glass beads are added through the rods slowly to prevent bridging.
- C1. Fine bentonite chips are added through the drive rods. A tube with funnel at the surface should be used to add water to hydrate the bentonite incrementally as the rods are retracted.
- C2. Alternatively, grout slurry may be pumped through a tremie tube to fill the probe hole as the rods are retracted. Use appropriate pump (GS1000 or GS500) and nylon tremie tube (P/N 11633).
- D. The sample tube is sealed with a suitable airtight cap or valve. Well protection is installed to meet local requirements

Figure 11: Installation of single-level implants for soil gas monitoring with the DT21 system.

#### 7.2 Single-Depth, High-Integrity Soil Gas Wells

Under some conditions or to meet specific regulatory requirements, it may be necessary to install prepacked screen monitoring wells (Figure 12) for long-term soil gas monitoring. Smaller diameter wells (e.g. 0.5- or 0.75-inch ID) may be preferred for this option to minimize the amount of purging required before sampling is conducted. The DT21 system allows for this option and will provide high-integrity well construction with an excellent annular seal and well defined screened interval with filter media. This gas well installation process utilizes the same basic procedure as for the installation of prepacked screen wells for groundwater monitoring. See Geoprobe® Technical Bulletin No. 962000 for detailed discussion of this procedure or ASTM Standard Practice D6725. A different cap and seal mechanism will be required for the well to prevent venting of volatiles and permit for appropriate purging and sampling activities.

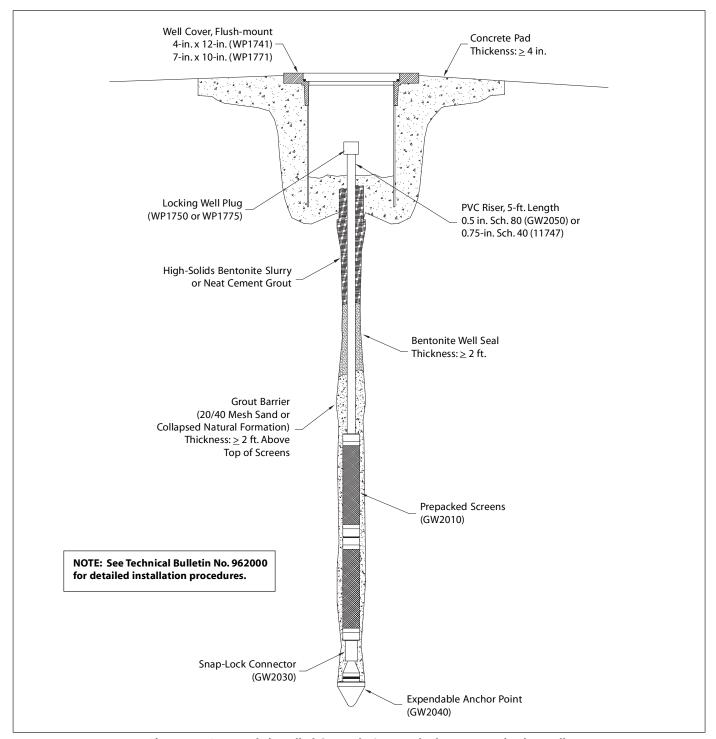


Figure 12: A properly installed Geoprobe® prepacked screen monitoring well.

#### 7.3 Multiple-Depth High-Integrity Implants

Soil gas profiling with the DT325 system was discussed previously in Section 6.5 and Figure 9. As noted, this system may be used to install multilevel soil gas implants as the 3.25-inch diameter rods are retracted. It is recommended that DT325 soil sampling be conducted through the entire interval where soil gas implants are to be installed. Removal of the soil cores will minimize compression of the formation and loss of gas permeability where implants are to be installed. Once soil coring has been completed to the maximum desired depth, the multilevel soil gas implant installation is conducted as follows (refer to Figure 13):

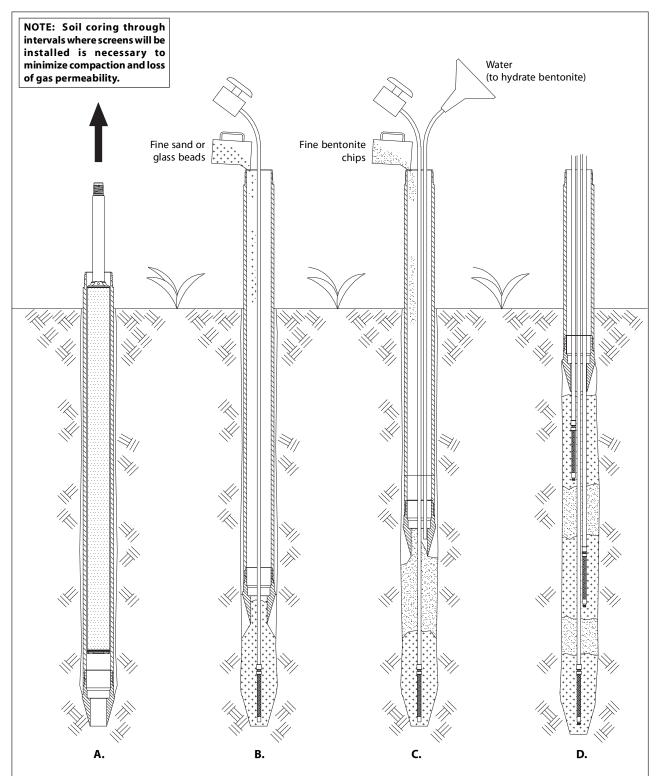
- The inner rod string and last soil core is removed.
- A soil core may be removed below the cutting shoe as described above (Figure 11) for installation of the deepest gas implant, if desired.
- The selected implant (Table 5) and tubing (Table 4) are assembled and lowered to the bottom of the bore hole.
- Filter media is poured through the rod bore as the rods are slowly retracted until filter media extends at least 6 to 12 inches (152 to 305 mm) above the implant. Measure depth to top of filter media and keep below rods and cutting shoe to prevent bridging.
- Fine bentonite chips are poured through the rod bore as the rods are slowly retracted. Water is added through a tube to hydrate each 4- to 6-inch (102 to 152 mm) increment of bentonite. Avoid wetting the interior of the rods to prevent bridging of sand and bentonite that could result in damage or loss of implants. (Alternately, grout slurry could be pumped by tremie tube and grout pump to install the seal at each level. See Figure 11, Part C2.)
- Repeat the last three steps for each soil gas implant as the rods are retracted.
- Grout from top of the filter pack for the uppermost implant to about 1 foot (0.3 m) below grade.
- Install surface protection as required.

Vapor monitoring implants installed following this procedure will enable you to achieve stringent data quality objectives for collection of soil gas samples to be used in human health risk assessments for the vapor intrusion pathway (Cal RWQCB 1997,Cal. DTSC 2003, NYDOH 2005, U.S. EPA 2002, etc.).

# Section 8.0 Direct Push Equipment Decontamination and Equipment Blanks

Without appropriate decontamination of downhole tooling and other sampling accessories there is significant opportunity for cross contamination and erroneous results. Various methods may be appropriate for equipment decontamination (decon) depending on the contaminant types and concentrations encountered (ASTM D 5088, Parker and Ranney, 2003). Most often downhole tools such as probe rods and point holders are cleaned with a soap-and-water wash and one or more clean water rinses. This may be most effective for VOCs if a heated pressure sprayer is used to conduct the decontamination. Measures for the control and containment of the spray and wastewater must be implemented. Appropriate health and safety protocols should be followed and adequate personal protective equipment should be worn. Refer to the project specific work plan and health and safety plan for guidance on appropriate safety protocols. Smaller parts may be washed and heated in an oven to drive off volatiles when appropriate.

When field analyses are conducted as sampling is performed in the field the analytical data can be used to guide decontamination requirements. Equipment used to collect samples that are nondetect will require less stringent decontamination measures than equipment used to collect samples from grossly contaminated areas. To assure that cross contamination is controlled equipment blanks should be collected on a regular basis (e.g. once a day or once every 10 samples, as appropriate to meet project DQOs). Equipment blanks should be collected from tool strings and sampling trains assembled just as they are for a typical sample location. The equipment blanks should be labeled and handled as other samples and submitted to the laboratory for the same analytical protocol.



- A. Advance DT325 system to depth with soil coring as needed. Remove last soil core.
- B. Install the selected implant and tubing. Gravity install fine sand or glass beads to provide a filter pack. Extend filter pack at least 4 to 6 inches (102 to 152 mm) above the top of the implant.
- C. Gravity install fine bentonite chips to create annular seal. Use funnel and tubing to incrementally hydrate the bentonite as the rods are retracted. Avoid wetting rod ID to prevent bridging and loss of boring. Alternatively, grout slurry can be installed with a grout pump and tremie tube.
- D. Repeat Steps B and C at specified intervals to obtain a multilevel monitoring system. Install well protection to meet local requirements.

Figure 13: Use of the DT325 system to install multilevel soil gas implants.

# Section 9.0 Abandonment and Grouting

To prevent migration of contaminants vertically along the probe hole to previously clean areas, proper abandonment and grouting techniques must be followed. For very shallow probe holes (e.g. < 5 ft (1.5 m)) in cohesive soils that remain open after the drive rods are removed, bentonite powder or fine bentonite chips may be added to the open hole incrementally and hydrated. In general, grout slurries should be pumped down the drive rods (Figure 14) or a tremie tube (Figure 11, Part C2) as the original drive rods are retracted from depths exceeding 5 feet (1.5 m). Pumps and accessories capable of injecting bentonite grout slurries or neat cement grout slurries through the small-diameter DP tools (Figure 15) will be required. Usually a 25% solids bentonite slurry is acceptable or a neat cement grout with 7 to 8 gallons (27 to 30 liters) of water per bag of cement may be required. Verify with state and local regulators the type and density of grout material necessary to meet local regulations. Proper abandonment and grouting is necessary to prevent substantial liability for contaminant migration if boreholes are left open.

# Section 10.0 Techniques for Collection of Soil Gas Samples

#### 10.1 DQOs Select the Technique

As discussed previously in Section 5.0, you must clearly define the DQOs before going to the field to collect soil gas samples. Samples that will be used for a human health risk assessment require much more stringent DQOs than samples that will simply be used to qualitatively determine if VOCs are present in the soil gas. A simple matrix (Table 2) will help you select the appropriate downhole tools paired with the correct sample collection methods to achieve the required sample quality. Remember, an expensive analysis does not make a poor quality sample better, it just wastes your money and time.

## 10.2 Sample Collection Flow Rate and Purge Volume

Another important consideration for sample collection is the actual flow

rate at which the sample is purged from the ground. Some guidance documents indicate that the purge and sample flow rate for soil gas sample collection should not exceed 200 ml/min. (Cal RWQCB 1997,Cal. DTSC 2003, NYDOH 2005, U.S. EPA 2002, etc.). If very high flow rates or strong vacuum is applied, volatiles that were sorbed to solid phases or liquids in the pore space may be stripped out and enter the gas phase for sampling. Thus, the use of high flow rates or strong vacuum for soil gas sampling may result in a strong positive bias. This could cause an over estimation of the risk to human health and may result in expensive remedial actions being applied that are not necessary.

For most soil gas sampling activities it is common to purge at least one to three volumes from the sampling system before the sample is collected. For direct sampling methods this will simply be the volume inside the drive rods multiplied by three. For PRT methods this will essentially consist of the volume inside the tubing and the open cylinder in the ground created by retraction of the rods. The internal volume of common tubing sizes by foot or meter of length is provided in Table 4.

For permanent soil gas implants or wells, the volume of void space in the filter pack around the screen should be included in the calculation along with the open volume inside the screen and tubing to the surface. For a clean sand filter pack a porosity of 30% is usually considered representative. The following example of purge volume calculation may be useful.

(continued on Page 23)

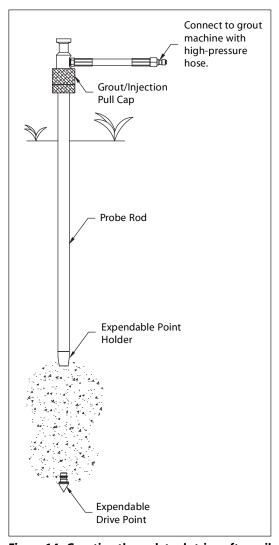
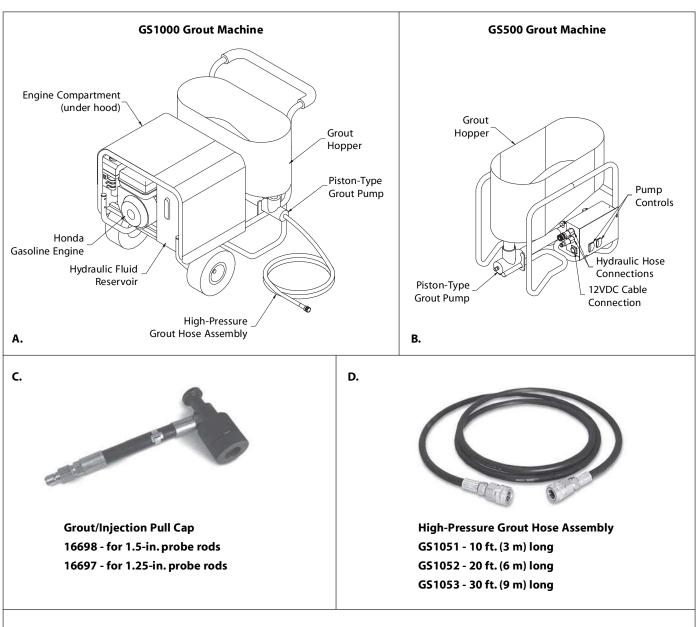


Figure 14: Grouting through tool string after soil gas sampling.



- A. Self-contained grouting system with gasoline engine and hydraulic fluid reservoir.
- B. Portable grout machine (65 lb / 30 kg) that is powered by the hydraulic and electrical systems of a direct push machine.
- C. Injection pull caps allow you to pump grout directly through the rod string during retraction.
- D. High-pressure grout hoses used to connect the grout pump to the injection pull cap and probe rod string.

Figure 15: Typical grouting equipment for direct push soil gas operations.

Volume of a cylinder =  $\pi$  x radius<sup>2</sup> x height, where  $\pi$  ~ 3.141

For a 2.125-inch (5.4 cm) diameter borehole 24 inches (61 cm) long, the total volume in milliliters would be:

$$3.141 \times (2.7 \text{ cm})^2 \times 61 \text{ cm} = 1397 \text{ cc} \text{ or approximately } 1400 \text{ ml}$$

For a 0.375-inch (0.95 cm) diameter implant 12 inches (30.5 cm) long, the volume would be:

$$3.141 \times (0.475)^2 \times 30.5 \text{ cm} = 21.6 \text{ cc} \text{ or approximately } 22 \text{ ml}$$

Now subtract the implant volume from the total probe hole volume and then multiply the remainder by 0.30 to determine the 30% pore space in the filter media. This will be:

$$(1400 \text{ ml} - 22 \text{ ml}) \times 0.30 = 413.3 \text{ ml}$$

Add back in the total volume of the void in the implant and you have:

$$413.3 \text{ ml} + 22 \text{ ml} = 435.3 \text{ ml}$$
 of void volume in the implant screen and filter media

If you have 12 feet (3.7 m) of 0.25-inch (6.4 mm) ID tubing, the volume per foot from Table 4 is 9.7 ml. One purge volume will be:

$$435.3 \text{ ml} + (12 \text{ ft } \times 9.7 \text{ ml/ft}) = 551.7 \text{ ml}$$

and three purge volumes would be about 1650 ml. Purging at 200 ml/min, it would require approximately

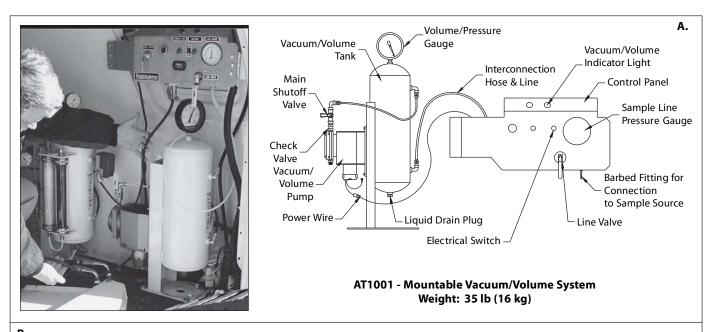
$$1650 \text{ ml} / 200 \text{ ml/min} = 8.25 \text{ minutes}$$

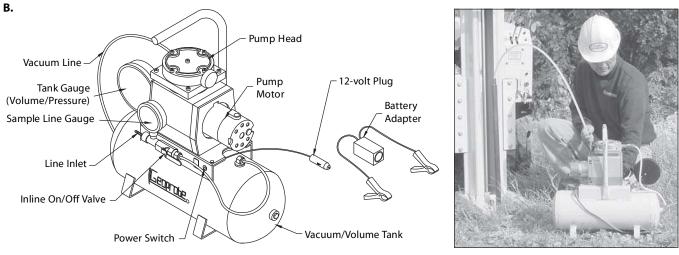
to complete the three volume purge for this implant system before you would be ready to collect your sample. So you would want to estimate at least ten minutes for sample collection in your cost proposal when similar implants are to be sampled.

The purge volumes and time required to purge using the PRT sampling system with three different probe rod diameters, using a one-foot (30.5 cm) retraction/void is summarized below in Table 6.

Table 6: Purge volume and time with PRT system and selected rod diameters.

Rod Diameter, inches (cm)	1.0 (2.54)	1.25 (3.18)	1.5 (3.81)
Void Radius (r), inches (cm)	0.5 (1.27)	0.625 (1.59)	0.75 (1.91)
Void Lenght (L), inches (cm)	12 (30.5)	12 (30.5)	12 (30.5)
Single Void Volume (mL) $(\pi \times r^2 \times L, \text{ where } \pi \sim 3.141)$	155	242	350
Tubing ID, inches (cm)	0.25 (0.64)	0.25 (0.64)	0.25 (0.64)
Tube Volume (mL/ft)	9.7	9.7	9.7
Tube Length (ft)	10	10	10
Total Tube Volume (mL)	97	97	97
Total System 1-volume (mL)	252	349	447
Approximate 3-System Volume (mL)	750	1050	1350
Required Purge Time at Flow Rate of 200 mL/min	3.75 min	5.25 min	6.75 min





- A. This system mounts in the back of a truck or van, often along with a Geoprobe® Model 5410 direct push machine.
- B. The portable 12V system may be powered from the vehicle 12V accessory outlet (cigarette lighter) or clipped directly to the vehicle battery.

AT1000 - Portable Vacuum/Volume System Weight: 40 lb (18 kg)

NOTE: Both systems include an 11-liter (2.9 gal.) tank and can provide up to 21 in. of Hg (70 centibars) of vacuum.

Figure 16: Vehicle-mounted and field-portable vacuum/volume systems for soil gas sampling.

#### 10.3 The Vacuum/Volume System

The vacuum/volume (V/V) system (Figure 16) has been used to good advantage for soil gas sampling during soil gas surveys, especially when large purge and sample volumes (liters) are required. The V/V system allows the field operator to perform three basic functions that are essential to successful soil gas sampling. These are measurement of the:

- volume of gas extracted at a sample point
- initial vacuum applied to a sample point
- amount of time for a sampling point to return to atmospheric pressure after a vacuum has been applied for sampling

The field operator may wish to incorporate an adjustable needle valve at the sample line inlet to control the flow rate to the desired level. The portable V/V system may be useful for periodic sampling of permanent wells or implants for monitoring purposes. Detailed operation instructions for the V/V system are available (Geoprobe® 2003).

### 10.4 Syringe sampling & purging

During initial surveys and plume tracking efforts, a mobile lab with onboard gas chromatograph (GC) may be used to conduct field analysis. This can significantly reduce analytical costs and the need for multiple mobilizations to determine the extent of the contaminant plume. When a van-mounted Geoprobe® direct push machine is equipped with a GC it is convenient to collect the soil gas sample directly from the PRT sample line with the syringe that will be used to inject the gas sample on the GC column (Figure 17). A short section of silicone tubing is placed in-line to allow for insertion of the GC syringe and collection of the sample. The sample is then immediately injected in the onboard GC for analysis. This collection and field analysis technique is especially amenable to use of the Triad approach for site characterization (EPA 2003).

When smaller purge volumes are required, a large syringe may be used to purge the sample tube. Gas sampling syringes made of glass or plastics are available from scientific supply warehouses in sizes ranging from 50 ml volume to 250 ml volume. This may be a particularly useful approach when shallow sampling is done in areas that are difficult to access with larger tools and equipment. Usually a small on/off valve is fitted to the end of the sample tube and is closed as the syringe is removed to be emptied and reattached for additional purging. The field tech will have to manually control the purge and sample flow rate. After purging the syringe may be used to inject a sample into a Tedlar bag, or glass bulb for on-site analysis. If off-site analysis will be conducted, a small summa canister may be used to collect the sample.

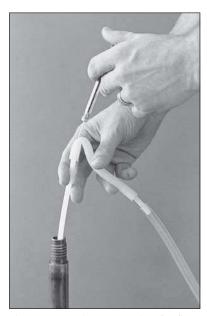


Figure 17: Syringe sampling for field analysis.

#### 10.5 Tedlar bags and glass sampling bulbs

These two sample collection options (Figure 18) have been widely used in soil gas surveys, especially when a mobile lab is stationed on-site to conduct the analyses as the samples are collected. These sample collection options also may be used when an off-site lab is to perform analyses but holding times for the samples may be limited. In addition, standard Tedlar bags and glass bulbs are transparent and exposure to sunlight is a concern. Some volatile organic contaminants may degrade when exposed to sunlight (especially ultraviolet light) so the sample containers must be stored out of the sunlight. A cooler or suitable box may be sufficient. Do not cool the samples if stored in a cooler as this may cause condensation and lead to biased sample results. Light-blocking Tedlar bags are available.

There are at least two ways to collect a sample in Tedlar bags. A large syringe may be used to extract the sample from the sample line and then it is injected into the Tedlar bag. Appropriate valves must be used on the syringe and bag to prevent ambient air contamination of the sample during transfer. Decontamination of the syringes between samples will be required to prevent cross contamination. An alternate method is to place the Tedlar bag inside a vacuum chamber with a port that allows connection to the sample line. A vacuum is applied outside the Tedlar bag and the bag expands and fills with gas from the sample line and downhole source. The downhole sample train must be purged before sample collection. Tedlar bags are generally considered a one-use disposable item, not to be decontaminated for reuse. Some regulatory agencies (Cal. EPA) discourage the use of Tedlar bags for soil gas sampling.

(continued on following page)

Alternatively, glass bulbs may be decontaminated for multiple reuse. However the glass bulbs are expensive and fragile, and decontamination requires additional equipment and man-hours. Blank analyses will also have to be performed periodically on the bulbs to verify no cross contamination occurs. The glass bulbs are usually placed in line during the purging process so they equilibrate with the gas being sampled. Smaller volume bulbs will require smaller system purge volumes and should still provide sufficient sample volume for duplicates and other QC measures if required. Decon may be accomplished by attaching the open bulbs to a vacuum pump and purging the bulbs while they are being heated in a small oven in the mobile lab on-site. A tubing manifold can connect several bulbs to one vacuum pump.

#### **10.6 Summa Canisters**

For evaluation of the vapor intrusion pathway and human health risk assessments, summa canisters (Figure 19) are preferred for sample collection and transport to a fixed lab facility for analysis. These stainless steel devices are usually supplied by the lab clean and ready for sample collection. The canisters are under vacuum and are simply attached to the sample line for collection of the soil gas. (The three-volume pre-sampling purge is conducted before the summa canister is attached to the sampling train.)

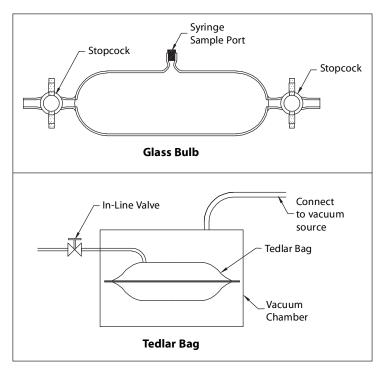


Figure 18: Tedlar bag and glass bulb for soil gas sampling.

A vacuum gauge and valve should be included on the canister inlet. The gauge will indicate when the canister is full and the valve can be used to control the sample flow rate from the downhole source at the 200 ml/min rate to prevent stripping of volatiles and biased sample results. Flow control valves may be set by the supplier at the requested flow rate. Smaller summa canisters (500 ml) may be preferable to minimize the sample collection time at the 200 ml/min flow rate.

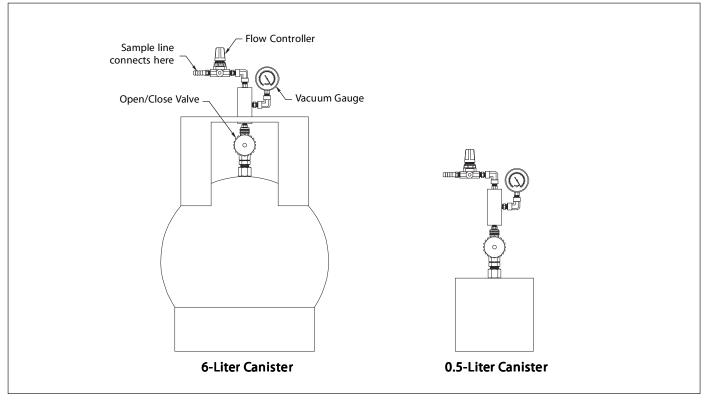


Figure 19: Summa canisters used for collection, storage, and transport of high-quality soil gas samples.

# Section 11.0 System Leak Tests for Field QC

Leak testing is a critical step to perform in the field to assure that system leaks do not degrade sample quality and cause biased results. There are at least three categories of leak testing that should be considered. These are:

- · Integrity of vacuum source
- Integrity of the sampling train
- Integrity of downhole tool system

The vacuum source may be as simple as a large syringe or as complex as a vacuum/volume system (see Figure 16). To verify that the vacuum supply itself does not have leaks, it should be tested periodically (e.g. once a day or every 10 samples). To test the integrity of your vacuum system the inlet valve should be closed or the inlet line should be plugged appropriately. The vacuum source is activated (syringe plunger retracted/vacuum pump turned on) and vacuum is applied. An in-line vacuum gauge should be included to quantitatively monitor loss of vacuum over time. Once the vacuum is applied, the system is "shut-in" so that loss of vacuum over time may be observed. For more complex systems the leak test may need to last 30 minutes to check for smaller leaks. Corrective action should be taken if leaks are detected.

Most often system leaks occur at aboveground plumbing connections where the sampling device is connected in-line between the vacuum supply and the sample line from the subsurface (sampling train). Probably the easiest and most effective way to determine if there are leaks during the sampling process is to apply a volatile compound around the fittings and analyze for that compound at the lab. One of the simplest and most effective ways to accomplish this is to wet paper towels with isopropyl alcohol and wrap them around the fittings and connections in the sampling train. Isopropyl alcohol may be purchased at the local pharmacy or department store. Remember, this material (liquid and vapor) is flammable and should be stored and handled appropriately.

Some downhole grab sampling systems can be tested for leaks before sampling is conducted. This may be accomplished for the direct sampling system (Figure 4) and the PRT sampling system (Figure 5). Expendable points with O-rings must be used so that an effective leak test can be accomplished. [Note: Neither monitoring nor grab sampling systems with downhole screens can be leak tested in this manner.] For the direct sampling system, the gas sampling cap with O-ring is installed on the drive rods and the tubing is connected between the cap and vacuum supply. Before the rods are retracted from the expendable point, a vacuum is applied to the sampling train and downhole system. The system is shut-in and the in-line vacuum gauge is observed for a period of time to determine if leaks are present.

For the PRT system, the PRT adapter (with fresh O-ring) is installed on the selected tubing. The adapter and tubing are lowered through the drive rods and the adapter is threaded into place. <u>Before</u> the drive rods are retracted off of the expendable point, vacuum is applied to the sampling train. The in-line vacuum gauge is observed to determine if leaks are present. Once the integrity of the sampling system is confirmed, the drive rods are retracted and soil gas purging and sampling is conducted.

**A note of caution:** System leak tests as described above will not allow you to determine if ambient air or soil gas from overlying zones is migrating into the interval being sampled. In order to prevent this problem, be sure that the expendable point and point holder are no larger in diameter than the outside diameter of the drive rods used in the tool string. Additionally, bentonite powder may be poured around the tool string at the ground surface and hydrated to help prevent ambient air intrusion during the sampling process.

# Section 12.0 Mapping Soil Gas Plumes

During the initial stages of a soil gas or VIP investigation it is wise to construct site maps that plot the location of each soil gas sample collection point. Then once the sample analyses are completed, the results may be plotted on the map and contours drawn to define the probable extent of the plume and areas of differing concentration (Figures 20 and 21). In many situations this will help locate sources for the contaminant plumes and define the general direction of contaminant migration. Often, soil gas plumes will develop over groundwater contaminant plumes and roughly mimic the extent of contamination in the local groundwater. In many urban and suburban areas there may be multiple plumes of different contaminants originating from different sources. Sometimes the source area(s) may be located outside the initial boundaries set for the investigation.

Use of on-site analysis with a mobile laboratory will optimize the investigation and plume definition. When field analyses are plotted on site maps daily as results are obtained, the field team can determine if new sample locations should be added to better define the plume. Just as important, nondetect samples can indicate that additional sampling locations planned for some areas may be deleted to minimize cost and time requirements. Application of the Triad Approach (EPA 2003) can be used to obtain the data needed while minimizing costs and multiple mobilizations to the field.

# Section 13.0 Health & Safety Considerations

All personnel working on an investigation where hazardous contaminants may be present must be OSHA 40 hour health and safety trained as required by 29 CFR 1910.120. The following discussion is provided as a brief overview of some of the health and safety issues that may be encountered during soil gas investigations. The field team should have a written health and safety plan that adequately addresses the possible physical and chemical exposure hazards that may be encountered during the investigation. Daily meetings to review health and safety conditions and any changes in the work environment should be conducted with the field team.

There are many types of health and safety issues to be considered during a soil gas investigation, or at any site where potentially hazardous contaminants may be present. These include physical hazards as well as potential exposure hazards to contaminants under investigation. Some of the physical hazards are associated with the use and operation of tools, equipment, and Geoprobe® direct push machines in the field. Basic precautions should be followed to prevent injury when operating tools and equipment such as wearing hard hats, steel-toe boots, appropriate gloves, eye protection, and hearing protection as necessary. Watch for pinch points on hydraulic equipment, machinery, and tools. Be familiar with the safety features of equipment being operated in the field, such as kill switches. Whenever subsurface sampling is to be conducted, all subsurface utilities must be professionally located and clearly marked to prevent accidents. Overhead utilities, especially high-voltage electrical lines, should be avoided. Adequate spacing between high-voltage lines and equipment masts or derricks should be maintained, check local requirements. When working in areas where vehicular traffic is a concern, wear appropriate high visibility clothing and use traffic cones to mark the work area.

The potential for exposure to hazardous contaminants is a reality whenever subsurface environmental investigations are conducted. The use of appropriate ambient air monitoring equipment is a must. Wearing appropriate clothing (e.g. tyvek, saranex, nitrile gloves) and air purification equipment (e.g. air purifying respirators (APR) or self contained breathing apparatus (SCBA)) for personal protection should be a routinely planned for and donned when necessary. The field team should have a written health and safety plan that reviews all these concerns and more that provides for appropriate measures when ambient air concentrations of contaminants exceeds the predetermined safety levels.

# Section 14.0 Summary

There are several tools and methods available to obtain soil gas samples with direct push techniques. Single tube methods are generally more time efficient and are often used during surveys to define the extent of the plume and help with location of source areas. These methods primarily provide sampling at a single-depth with each advance of the tool string. Conversely, dual tube methods are generally a little more time consuming but allow the operator the option to conduct soil gas profiling at multiple depths during one advancement of the tool string. The dual tube systems also allow the investigator to install high-integrity soil gas implants or monitoring wells. The placement of filter media in the screened interval and annular seals by the dual tube methods allows the sampler to achieve sample quality levels to meet the needs of human health risk assessments for the vapor intrusion pathway. Selection of the appropriate sample collection device and procedure is an integral part of achieving the required DQOs. While on-site analysis with a well-equipped mobile laboratory can expedite completion of the field work and plume definition, off-site analytical work may be required under some conditions to meet the more stringent requirements of human health risk assessment for the vapor intrusion pathway.

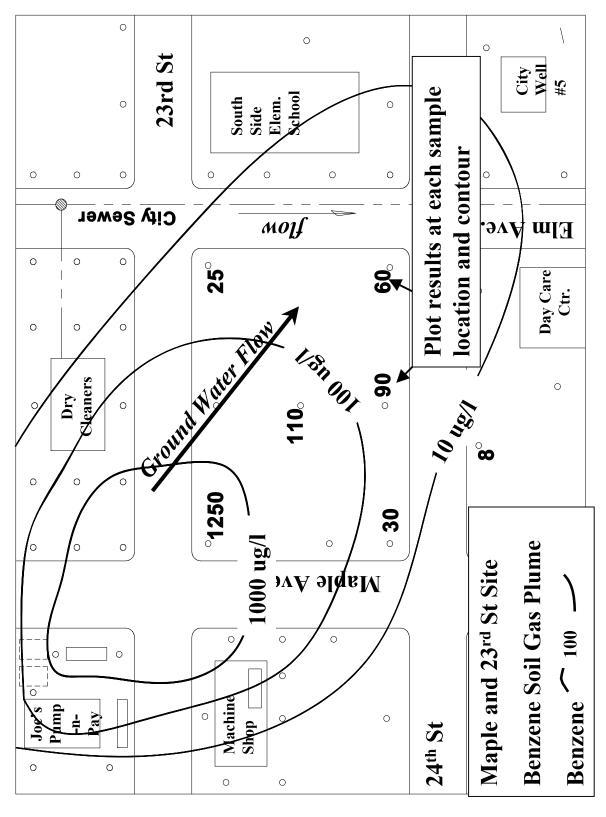


Figure 20: Mapping benzene plume for soil gas investigation.

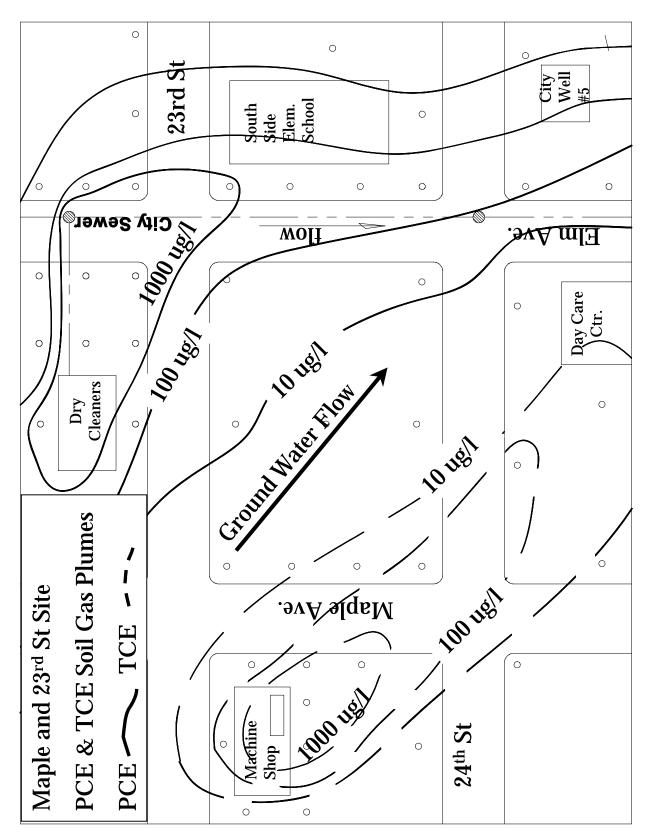


Figure 21: Mapping X-VOC plumes for soil gas investigation.

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