Appendix F
Examples

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I. Maps and Cross-Sections Presentation of Data

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I. MAPS AND CROSS SECTIONS PRESENTATION OF DATA

All work related to site assessment and mitigation must be documented in a clear and concise manner. In geological and engineering activities, the use of maps and cross sections is a valuable tool in presenting simple-to-very-complex issues.

All site investigations and monitoring reports should incorporate maps and cross sections, including, but not limited to, the following information:

- Site location/vicinity map
- Adjacent land use map
- Utilities site map
- Site map
- Geological cross sections
- Groundwater gradient maps
- Groundwater sampling results map
- All sample locations and relevant sample results
SITE MAP AND SURROUNDING LAND USE

FIGURE 2

Scale (feet)

Date: June 4, 1995
APPENDIX F: EXAMPLES

SITE MAP

FIGURE 4

LEGEND

- Excavation Samples
- Monitoring Well
- Boring
- Vapor Extraction Well

Scale (feet)

Approximate location of Tank Excavation and Product Lines
Approximate location of Property Line

Date: June 4, 1995
GROUNDWATER GRADIENT -- March 23, 1995

FIGURE 6

LEGEND

MW-1 (190.50) Monitoring Well
Groundwater Elevation
Approximate location of Tank
Excavation and Product Lines
Approximate location of Property Line

Scale (feet)

0 20 40

Date: June 4, 1995
II. SAMPLE VAPOR PHASE RISK CALCULATIONS
EXAMPLES - VAPOR-PHASE MIGRATION AND RISK EVALUATION

The vapor-phase migration and risk evaluation are discussed in detail in Section 6. All the equations and tables referenced in this appendix can also be found in Section 6. Consultants are encouraged to use the VAPRISK 2000 Model (http://www.sdcounty.ca.gov/deh/water/sam_vapor_risk_assessment_2000.html) which is programmed with each of the equations provided in the following examples.

LEVEL 1 ANALYSIS

Site Description:

The site under evaluation is a neighborhood gasoline station that is surrounded by residential homes. This site experienced a release of gasoline from the underground storage tank system that was replaced in 1990. The site investigation identified the extent of soil and groundwater contamination. Contamination extends off-site with free product extending below a residential home that was built in the late 1960s.

Due to the presence of free-product beneath the residence, a preliminary vapor phase evaluation was warranted. The following are typical steps that should be taken to do a Level 1 Evaluation of the potential health risk.

VAPOR TRANSPORT

Step 1 – Review of site data

A review of the site investigation data indicated that in the area of the residence the subsurface soils were primarily medium to coarse sands. Free product ranging form 0.05 to 0.2 feet in thickness was observed in the area of the residence.

Step 2 – Field verify site conditions

Initially the consultant visited the site and performed a detailed visual evaluation of the residence and its construction. This inspection identified the residence was a structure built with a concrete slab on-grade and the house was ventilated passively with only a forced air heating system. The concrete slab was inspected to verify its condition. Field observations identified the building slab as being in good condition with no observed deterioration or cracking. Based on these observations it was concluded that the use of the 0.01 slab attenuation factor was acceptable for use in the health risk evaluation. Additionally, the residence had an interior room height of 8 feet (2.44 meters).
Step 3 – Calculation of soil gas concentration

Since free-product is present appropriate method to calculate the level of benzene in soils gas is using Equation 6-13 presented below:

\[ C_{sg} = \frac{VP \times MW \times MF}{R \times T} \]

Where:
- \( C_{sg} \) = the contaminant concentration in the soil vapor (mg/m\(^3\))
- \( VP \) = the contaminant vapor pressure at STP (atm)
- \( MW \) = the molecular weight of the compound of concern (mg/mole)
- \( MF \) = the mole fraction (dimensionless)
- \( R \) = the universal gas constant (atm-m\(^3\)/mole-°K)
- \( T \) = the temperature in degrees Kelvin (Standard temperature of 293°K)

Using the default values presented in Table 6-4 the soil gas concentration is calculated as follows.

\[ C_{sg} = \frac{0.13 \text{ atm} \times 78,110 \text{ mg/kg} \times 0.03}{0.000082 \text{ atm-m}^3/\text{mole-°K} \times 293 \text{ °K}} = 12,700 \text{ mg/m}^3 \]

Step 4 – Calculate Effective Diffusion Coefficient

To calculate the effective diffusion coefficient Equation 6-16, presented below, is used.

\[ D_e = \frac{D_a \times \theta_a^{3.33}}{\theta^2} \]

Where:
- \( D_e \) = the effective air diffusion coefficient (cm\(^2\)/sec)
- \( D_a \) = the diffusion coefficient of compound in air (cm\(^2\)/sec)
- \( \theta_a \) = the air filled porosity (dimensionless)
- \( \theta \) = the total soil porosity (dimensionless)

Since the soils identified in the area of concern (medium to coarse sands) have not been tested to determine the soils porosity and moisture content the default values for porosity and air filled porosity were used (Table 6-4). In reviewing Table 6-3

\[ D_e = \frac{0.088 \text{ cm}^2/\text{sec} \times 0.20^{3.33}}{0.30^2} = 0.0046 \text{ cm}^2/\text{sec} \]
Step 5 – Calculate Vapor Flux

To calculate vapor flux Equation 6-15 presented below is used.

\[
F_x = \frac{D_e \cdot C_{sg} \cdot 3,600 \text{ sec/hr}}{X \cdot 10,000 \text{ cm}^2/\text{m}^2}
\]

Where:
- \( F_x \) = the contaminant vapor flux (mg/hr-m^2)
- \( D_e \) = the effective air diffusion coefficient (cm^2/sec)
- \( C_{sg} \) = the contaminant concentration in the soil vapor (mg/m^3)
- \( X \) = the depth or distance to contamination in the vadose zone (m)

\[
0.0046 \text{ cm}^2/\text{sec} \cdot 12,700 \text{ mg/m}^3 \cdot 3,600 \text{ sec/hr} = \frac{4.6 \text{ m} \cdot 10,000 \text{ cm}^2/\text{m}^2}{4.56 \text{ mg/hr-m}^2}
\]

Step 6 – Calculation of Indoor Air Concentration

To calculate the indoor air concentration Equation 6-17 presented below is used.

\[
C_i = \frac{S_b \cdot F_x \cdot A}{V \cdot E} = \frac{S_b \cdot F_x}{R_h \cdot E}
\]

Where:
- \( C_i \) = the indoor air concentration (mg/m^3)
- \( S_b \) = the slab attenuation factor (dimensionless)
- \( F_x \) = the contaminant vapor flux (mg/hr-m^2)
- \( A \) = the room floor area (m^2)
- \( V \) = the room volume (m^3)
- \( E \) = the indoor air exchange rate per hour (hr\(^{-1}\))
- \( R_h \) = the room height (m)

\[
0.01 \cdot 4.56 \text{ mg/hr-m}^2 = \frac{0.0374 \text{ mg/m}^3}{2.44 \text{ m} \cdot 0.5 \text{ hr}^{-1}}
\]

**VAPOR RISK**

Steps 1 though 6 have calculated the indoor air concentration in overlaying residence. This is the vapor transport portion of the evaluation. This indoor air concentration, 0.0374 mg/m^3, is the air concentration that the occupants are exposed to through inhalation. The following steps calculate the human exposure and the potential health risk to these individuals. Since in this evaluation we are evaluating residential exposure then the default values for health risk exposure (Table 6-9) need to be used.
Step 7 – Calculating Human Exposure

To calculate human exposure through inhalation Equation 6-22, presented below, is used

\[
\frac{C_i \times IR \times ET \times EF \times ED}{BW \times AT} = IT
\]

Where:
- \(IT\) = the chemical intake (mg/kg-day)
- \(C_i\) = the indoor air concentration (mg/m\(^3\))
- \(IR\) = the inhalation rate (m\(^3\)/day)
- \(ET\) = the exposure time (hr/24hr)
- \(EF\) = the exposure frequency (days/yr)
- \(ED\) = the exposure duration (yr)
- \(BW\) = the body weight (kg)
- \(AT\) = the averaging time (days)

\[
\frac{0.0374 \text{ mg/m}^3 \times 20 \text{ m}^3/\text{day} \times 24\text{hr}/24\text{hr} \times 365 \text{ days/yr} \times 70 \text{ yr}}{70 \text{ kg} \times 25500 \text{ days}} = 0.0107 \text{ mg/kg-day}
\]

Step 8 – Calculation of Carcinogenic Risk

To calculate the carcinogenic risk Equation 6-23, presented below, is used.

\[
\text{Risk} = IT \times SF
\]

Where:
- \(Risk\) = the estimate of health risk (dimensionless)
- \(IT\) = the chemical intake (mg/kg-day)
- \(SF\) = the contaminant carcinogenic slope factor ([mg/kg-day]\(^{-1}\))

\[
\text{Risk} = 0.0107 \text{ mg/kg-day} \times 0.1 \text{ [mg/kg-day]}^{-1} = 1.07 \times 10^{-3}
\]

Based on this analysis, the incremental cancer risk is the inverse of the risk calculated above. This result indicates that there is a cancer risk of one in a population of 934 people. This result represents an unacceptable health risk. The acceptable level or risk is one in a population of 1,000,000 (one in a million).

Based on this result, the responsible party should either proceed with remediation or complete a higher level of investigation to collect site-specific information to support a Level 2 risk evaluation.
LEVEL 2 ANALYSIS

Site Description:

The site under evaluation is a commercial property that historically was operated as a dry cleaning business. Site investigation included the collection of soil and vapor data from beneath the concrete floor slab in the area of the former dry cleaning equipment. The investigation identified only tetrachloroethene (PCE) at a maximum concentration of 10 mg/kg in soil and 365 ug/l in soil vapor at approximately 1 foot below the floor slab.

Due to the elevated levels of PCE contamination, a vapor phase evaluation was warranted. The following are typical steps that should be taken to do a Level 2 Evaluation of the potential health risk.

VAPOR TRANSPORT

Step 1 – Review of site data

A review of the site investigation data indicated that the maximum soil vapor concentration was 265 ug/l at 1 foot below (0.33 meters) the floor slab. During the site investigation, the soils at the site were identified as Lindavista Formation-derived fill soils consisting of silty fine sands. Due to fine grained nature of the fill soils, the consultant obtained samples and did site-specific physical testing to determine the in situ soil porosity, moisture content and organic carbon content.

This additional testing provided the following physical properties:

<table>
<thead>
<tr>
<th>Property</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bulk density</td>
<td>1.9 gm/cm³</td>
</tr>
<tr>
<td>Total porosity</td>
<td>0.255 dimensionless</td>
</tr>
<tr>
<td>Water filled porosity</td>
<td>0.135 dimensionless</td>
</tr>
<tr>
<td>Air filled porosity</td>
<td>0.120 dimensionless</td>
</tr>
<tr>
<td>Total organic carbon content</td>
<td>0.01 dimensionless</td>
</tr>
</tbody>
</table>

Step 2 – Field verify site conditions

Additionally the consultant performed a detailed visual evaluation of the commercial space under consideration, in addition to the adjacent units, to evaluate the building construction and current condition. This inspection identified that the structure was built with an on-grade concrete slab within the past 10 years and the commercial unit was designed with a ventilation system that provided 1.0 air exchange per hour with the outside air.

The concrete slab was inspected to verify its condition. Field observations identified the building slab as being in good condition with no observed deterioration or cracking. Based on these observations, it was concluded that the use of the 0.01 slab attenuation factor was acceptable for use in the health risk evaluation. Additionally, the commercial space had a ceiling height of 8 feet (2.44 meters).
Step 3 – Calculation of soil gas concentration

Based on the site investigation, the maximum soil vapor concentration identified was 265 ug/l.

Since the vapor risk model must use units of mg/m³ for the soil gas, the maximum soil gas concentration needs to be converted from ug/l. The conversion table is presented in Section 6, Table 6.7.

In this example, the unit of ug/l is equal to that of mg/m³

\[ C_{sg} = 265 \text{ mg/m}^3 \]

Step 4 – Calculate Effective Diffusion Coefficient

To calculate the effective diffusion coefficient, Equation 6-16, presented below, is used.

\[
D_e = \frac{D_a \cdot \theta_a^{3.33}}{\theta^2}
\]

Where:
\[ D_e = \text{the effective air diffusion coefficient (cm}^2/\text{sec)} \]
\[ D_a = \text{the diffusion coefficient of compound in air (cm}^2/\text{sec)} \]
\[ \theta_a = \text{the air filled porosity (dimensionless)} \]
\[ \theta = \text{the total soil porosity (dimensionless)} \]

Since the soils were tested, the site-specific values presented in Step 1 are used along with the diffusion coefficient obtained from Table 6-2a.

\[
D_e = \frac{0.072 \text{ cm}^2/\text{sec} \cdot 0.120^{3.33}}{0.255^2} = 0.00095 \text{ cm}^2/\text{sec}
\]

Step 5 – Calculate Vapor Flux

To calculate vapor flux, Equation 6-15 presented below is used.

\[
F_x = \frac{D_e \cdot C_{sg} \cdot 3,600 \text{ sec/hr}}{X \cdot 10,000 \text{ cm}^2/\text{m}^2}
\]

Where:
\[ F_x = \text{the contaminant vapor flux (mg/hr-m}^2) \]
\[ D_e = \text{the effective air diffusion coefficient (cm}^2/\text{sec)} \]
\[ C_{sg} = \text{the contaminant concentration in the soil vapor (mg/m}^3) \]
\[ X = \text{the depth or distance to contamination in the vadose zone (m)} \]

\[
F_x = \frac{0.00095 \text{ cm}^2/\text{sec} \cdot 265 \text{ mg/m}^3 \cdot 3,600 \text{ sec/hr}}{0.33 \text{ m} \cdot 10,000 \text{ cm}^2/\text{m}^2} = 0.275 \text{ mg/hr-m}^2
\]
Step 6 – Calculation of Indoor Air Concentration

To calculate the indoor air concentration, Equation 6-17 presented below is used.

\[ C_i = \frac{S_b \times F_x \times A}{V \times E} = \frac{S_b \times F_x}{R_h \times E} \]

Where:
- \( C_i \) = the indoor air concentration (mg/m\(^3\))
- \( S_b \) = the slab attenuation factor (dimensionless)
- \( F_x \) = the contaminant vapor flux (mg/hr-m\(^2\))
- \( A \) = the room floor area (m\(^2\))
- \( V \) = the room volume (m\(^3\))
- \( E \) = the indoor air exchange rate per hour (hr\(^{-1}\))
- \( R_h \) = the room height (m)

\[ C_i = \frac{0.01 \times 0.275 \text{ mg/hr-m}^2}{2.44 \text{ m} \times 1 \text{ hr}^{-1}} = 0.00113 \text{ mg/m}^3 \]

**VAPOR RISK**

Steps 1 though 6 have calculated the indoor air concentration in overlaying commercial space. This is the vapor transport portion of the evaluation. This indoor air concentration, 0.000113 mg/m\(^3\), is the air concentration that the occupants are exposed to through inhalation. The following steps calculate the human exposure and the potential health risk to these individuals. Since in this evaluation we are evaluating commercial exposure, the default values for health risk exposure (Table 6-9) need to be used.

Step 7 – Calculating Human Exposure

To calculate human exposure through inhalation, Equation 6-22, presented below, is used

\[ IT = \frac{C_i \times IR \times ET \times EF \times ED}{BW \times AT} \]

Where:
- \( IT \) = the chemical intake (mg/kg-day)
- \( C_i \) = the indoor air concentration (mg/m\(^3\))
- \( IR \) = the inhalation rate (m\(^3\)/day)
- \( ET \) = the exposure time (hr/24hr)
- \( EF \) = the exposure frequency (days/yr)
- \( ED \) = the exposure duration (yr)
- \( BW \) = the body weight (kg)
- \( AT \) = the averaging time (days)
0.00113 mg/m³ * 20 m³/day * 12hr/24hr * 250 days/yr * 25 yr
IT = -------------------------------------------------------------
70 kg * 25500 days

IT = 3.94 x 10⁻⁵ mg/kg-day

**Step 8 – Calculation of Carcinogenic Risk**

To calculate the carcinogenic risk, Equation 6-23, presented below, is used. The slope factor for PCE is presented in Table 6-1.

\[
\text{Risk} = \text{IT} \times \text{SF}
\]

Where:
- \( \text{Risk} \) = the estimate of health risk (dimensionless)
- \( \text{IT} \) = the chemical intake (mg/kg-day)
- \( \text{SF} \) = the contaminant carcinogenic slope factor ([mg/kg-day]⁻¹)

\[
\text{Risk} = 3.94 \times 10^{-5} \text{mg/kg-day} \times 2.10 \times 10^{-2} \text{[mg/kg-day]^{-1}} = 8.28 \times 10^{-7}
\]

Based on this analysis, the incremental cancer risk is the inverse of the risk calculated above. This result indicates that there is a cancer risk of one in a population of 1,209,190 people. This result represents an acceptable health risk. The acceptable level or risk is one in a population of 1,000,000 (one in a million).