August 4. 1997

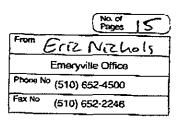
Comments on Polverosa Risk Assessment

Scott,

Since I found some errors in the hand calculation, they did some revisions and the updated calculations seems to be fine. On page 6, they summarized the results (SSTL's) based on the revised calculations. The concentrations on site for both soil (based on a maximum of 0.77) and groundwater (based on a maximum of .9 ppb) are below the calculated SSTL's for a risk of 10-5. Hopefully the maximum values they have stated in the risk assessment are acceptable to you based on site information. Also, in the revised calculation, they went back to using the default values for the air exchange rate as they did not have reference document to provide us. I left a message for Eric Nichols to go ahead and mail you a good copy (without scratches) of the risk assessment.

Madhulla

AUG



rections/reviewons

TO 93379335

LFR 1204.03

Risk-Based Corrective Action (RBCA) Evaluation for Polyorosa Business Park 1555 Doolittle Drive San Leandro, California

Based on a request from Alameda County Department of Environmental Health (ACDEH), Levine-Fricke-Recon (LFR) has prepared this Risk-Based Corrective Action (RBCA) evaluation for the Polvorosa Business Park site (hereafter referred to as "the Site"). This evaluation is derived from the Standard RBCA method applied to petroleum release sites (American Society for Testing of Materials, ASTM standard method E-1739-95). In the absence of specific state policy and guidance concerning RBCA, we have used the example policies presented in the Appendices of ASTM E-1739, modified as noted.

Step 1: Initial Site Assessment

The site investigation was conducted in several phases subsequent to UST removal in 1986. The groundwater characterization was completed by Levine-Fricke in 1988. The most current soil quality data is from samples collected by Groundwater Technology in October, 1986.

The fuel types of concern are gasoline- and diesel-range fuel hydrocarbons. The thickness of free-phase hydrocarbons on groundwater was measured when encountered. Groundwater quality samples were analyzed for TPHg, TPHd, benzene, toluene, ethylbenzene, and xylenes. Groundwater samples were not analyzed for the above analytes when free-phase hydrocarbons were detected. Soil samples (collected in 1986) were analyzed for motor fuels, benzene, toluene, and total xylenes.

Steo 2: Site Classification and Initial Response Action

Using Table 1 of ASTM Standard E 1739-95, the site falls under Classification 3 - Long-term threat to human health, safety, or sensitive environmental receptors. (This conclusion was drawn from both pre- and post-remediation conditions)

Interim Remedial Action

During UST removal activities in 1986, some unsaturated-zone hydrocarbon-affected soils in the immediate vicinity of the USTs were removed. However, the quantity of soils which

AUĢ

 were removed was not recorded. Based on available records, the volume of soils excavated may have been several hundred cubic yards.

From August, 1989 to August, 1993, a total fluids groundwater capture system was operated in the area where free-phase hydrocarbons were encountered. Approximately 766,000 gallons of total fluids were removed, including about 283 gallons of free-phase hydrocarbons.

Step 3: Tier 1 Evaluation

The following is a brief summary of the rationale used in screening of reasonable sources, pathways, and exposures for evaluation.

Primary Sources:

The tanks and piping were removed in 1986.

Secondary Sources:

Based on existing data, the following secondary sources are present at the site:

- impacted subsurface soils
- · dissolved groundwater plume
- free-phase liquid plume (limited extent based on observation of hydrocarbon sheen)

Transport Mechanisms:

The site is presently covered with asphalt and building structures. A one-story warehouse building overlies much of the more affected area. A landscaped area, without pavement, exists near the hydrocarbon-affected area, and is estimated at about 10% of the building footprint. Downgradient wells on the Site indicate that the plume has had little or no additional migration after the cessation of ground water capture.

The following transport mechanisms were judged to be significant:

- volatilization and atmospheric dispersion (through landscaped areas around the parking area)
- volatilization, migration of vapors through the building foundation, and enclosed-space accumulation

The following transport mechanisms were not considered significant as noted:

- wind erosion and atmospheric dispersion (because only a small portion of the site is unpaved)
- leaching and groundwater transport (because ground water is not extracted for human uses in the general vicinity of the Site)
- mobile free-liquid migration (because the source and mobile free-phase hydrocarbons were removed)
- stormwater/surfacewater transport (because the affected area does not include any affected surface soils or any large underground lines)

The receptors at the Site were characterized as:

- commercial/industrial
- construction workers

Applicable Risk-Based Screening Levels (RBSLs) from Tier 1 Look-up Table

Using the modified example ASTM RBCA table provided to us by ACDEH, the following is a comparison of applicable values for commercial/industrial sites:

SOIL-BASED EXPOSURE PATHWAYS RBSLS and available site data

Exposare Pathway	Larges Risk Level	Benzere jugiky	Totome ing/kg	Xylenes (tensi) (atg/kg)
Volatilization to outdoor air	Cancer 1x10 ⁴	0.133		
	Cancer 1x10 ⁻⁴	13.3		
	Chronic Hazard		RES	RES
Vapor intrusion into buildings	Cancer 1x10 ⁻⁶	0.00155		
"	Cancer 1x10 ⁻⁴	0.155		
	Chronic Hazard		54.5	RES
Max conc describe (1986)			0.27	1.2
Meanlover affected area @ 10 5 Bgs (1980)*		023	6,8	D 28

GROUNDWATER-BASED EXPOSURE PATHWAYS RBSLs and available site data

all

		My			
Exposure Pattings	Forget Biol Fryd (*)	Bonzene (16g/13)	(stant (ng/L)	Edwill-av Vinelij	
Volatilization to outdoor air	Cancer 1x10 ⁻⁶	5.34			
	Cancer 1x10 ⁻⁴	>5			
	Chronic Hazard		>S	>\$	>S
Vapor intrusion into buildings	Cancer 1x10 ⁻⁶	0.0069			
	Cancer 1x10 ⁻⁴	0.69			
	Chronic Hazard		54.5	· >S	RES
Max. conc detected (1995)		U 9809**	∠MDL	- <mde< td=""><td>SMDL.</td></mde<>	SMDL.
Estimated nizx bosed on free-phase HCs		5 #	83	20	198

EXPLANATION:

- **RES** the selected risk level is not exceeded for pure compound selected in any concentration
- >S the selected risk level is not exceeded for all possible dissolved levels
- <MDL the analyte was not detected above the lab minimum detection level bgs = below ground surface (depth)
- *the mean calculated was an arithmetic mean of soil sample data at the approximate depth of shallow ground water over the area and immediate vicinity of Building C
- ** the 0.0009 mg/L benzene concentration detected was in a well upgradient of the most affected area; well LF-12 was not sampled in May 1995 due to the measurement of 0.02 feet of free-phase hydrocarbons

The estimated maximum concentrations for benzene was based on a mole fraction of 0.03 and a pure-compound solubility of 1800 mg/L. The estimated maximum concentrations of toluene and xylenes were estimated from section X1.6 and Table X1.2 of ASTM E-1739.

 The lightly shaded cells of RBSL values are those exceeded at the Tier 1 screening level. also used shown

Step 4: Decision Tree/Comparison with RBSLs

Based on our conversations with ACDEH, we considered a (1x10 1 in ten thousand) excess cancer risk as the appropriate target risk level for the commercial/industrial receptors at the Site.

Chemical(s) of concern concentrations exceed RBSLs? - Yes, benzene (both soil and ground water into indoor air) and toluene (soil and ground water into indoor air). Both of the groundwater RBSLs were exceeded based on the assumption of free-phase hydrocarbons rather than the actual values detected.

Remediation to Tier 1 RBSLs practicable? - No

Interim remedial action appropriate? - Yes, but to further evaluate whether remaining concentrations pose an acceptable risk, we completed a Tier 2 evaluation to derive Site Specific Target Levels (SSTLs).

Step 5: Tier 2 Calculation of SSTLs

Based on the attached calculations, the groundwater-to-outdoor air pathway is well below the applicable commercial/industrial SSTL values using a combination of default and sitespecific assumptions.

Table 1 shows the Site-specific assumptions used to calculate the SSTLs. Other input parameters were taken from the default parameters presented in ASTM E1739.

Tables 2, 3 and 4 are the calculations used to derive the SSTLs for the indoor air exposure pathway. The SSTLs calculated are summarized below:

	Ben	Benzene				
Media	nsk=10-4	risk=10-5	HI=1			
Soil (mg/kg)	204	20.4	20,300.			
Groundwater	20.4	2.0	770.			
(mg/L)		200 ppb				

Step 6: Comparison with SSTLs

The calculated benzene and toluene concentrations in soil and groundwater necessary to exceed the indoor air SSTLs are well above those detected. The SSTLs calculated are also well above the soil and groundwater concentrations that would be expected adjacent to a

fresh liquid hydrocarbon sheen, based on equilibrium partitioning. A comparison of the calculated SSTL values above to the estimated maximum concentrations based on freephase hydrocarbons expressed in the above groundwater RBSL table supports this conclusion.

Conclusion

We recommend no further corrective action at the Site. It is our opinion that the possible exposure pathways have been considered and evaluated to pose an insignificant risk using Site data. This conclusion is consistent with existing current State of California guidelines, since benzene has not been detected at concentrations greater than 1 mg/L at the Site and the Site is greater than 750 feet from the nearest drinking water well.

	SHEET Table 1 p.1 OF 1 /alcs 1/8
-	JOB NO: 1204.00.03
	DATE: 18 March 1997
valuation	COMPUTED BY: J. STUTMEN
wters	CHECKED BY: E. Nithels

the building is relatively new (about 40 years) and was Constructed, with a polyethylene moisture barrier below the foundation Slab

PROJECT:

Symbol	Definition of Parameter	Tier 1 Default Value	Tjer 2 Sile- Specific Volve	Reason for Modification
O ās	volumetric air Confent vadose zone soils	0.26 (m3 oir	0.18 cmat	fine-grained soils under buildings are typically at least 50% Submited
Ows	volumetric unler content vadose zere soils	0.12 "	0.20 "	at at
O a crack	volumetric air content foundation cracks	0.26	0.18 "	foundation cracks are typically partially filled with soil and/or water
O w crack	Volumetric water content foundation cracks	0.12 "	0.20	ii h
hv	thickness of value zone	295 cm	305 cm	the approximate depth to ground under is about 9 feet at the site
Law	depth . to ground water	300 cm	310 cm	Law = hv + hcap
Ls	depth to subsurface soil	100 cm	170 cm	in 1986, some hydrocarbons were defected at 5.5 feet bgs
ER	enclosed-space air exchange rate	useQ as d	0.0028/sec 0.00014 efault, non 8/4/97	

0.01

0.001

areal fraction of cracks in foundation

η



PROJECT: Polvoresa Site RBCA Evaluation SUBJECT: Calculation of diffesion Calliciants SHEET Table 2, p. 1 OF 3, rules 2/8 JOB NO.: 1204.00.03 DATE: 18 March 1997 COMPUTED BY: J.STUTMAN CHECKED BY: E. Nichols

Diffusion Coefficient Soil - Vapor,
$$D_s^{eff}$$

$$D_s^{eff} = D_v^{air} \frac{\theta_{as}^{3,33}}{\theta_T^{2,0}} + D_v^{avet} \frac{1}{H} \frac{\theta_{ws}^{3,33}}{\theta_T^{2,0}}$$

For benzene, $D_{s}^{eff} = 0.093 \frac{(0.18)^{2.33}}{10.20)^{2.0}} + (1.1 \times 10^{-5}) \frac{1}{0.22} \frac{(0.18)^{3.33}}{(0.28)^{2.0}}$ $\stackrel{\simeq}{=} 0.093 \xrightarrow{0.0033} \frac{0.0033}{0.1444} + (11 \times 10^{-5}) (4.54) \frac{0.0033}{0.1444}$ $\stackrel{\simeq}{=} 0.021 + 1 \times 10^{-6}$ $\stackrel{\simeq}{=} 0.0021$

 $D_{s}^{eff} \cong 0.093 \frac{(0.18)^{3.33}}{(0.38)^{2.0}} + (9.4 + 10^{-6}) \frac{1}{0.26} \frac{(0.18)^{3.33}}{(0.38)^{2.0}} + 1 \times 10^{-6}$ = (02t 0.00Z1)

Diffusion Coefficient through foundation cracks, Direck $D_{crack}^{eff} \cong D^{air} \underbrace{\theta_{acrack}^{3,33}}_{\Phi^2} + D^{wat} \frac{1}{H} \underbrace{\theta_{ws}^{3,33}}_{\Phi^2}$

For benzene, Det = same as Det for benzene = 021

For toluene, Det = same as Diet for toluene = 0.21



PROJECT: Polvorosa Site RBCA Evaluation SUBJECT: Calculation of diffusion coefficients SHEET 1660 2, P. LOF 3, (AICS 3/8 JOB NO.: 1204.00.03 DATE: 18 March 1997 COMPUTED BY:) STUTMEN CHECKED BY E. Nichols

Diffusion Coefficient through capillary fringe,
$$D_{cap}^{eff}$$

$$D_{cap}^{eff} = D^{air} \frac{\theta_{acap}}{\theta_{\tau}^{2}} + D^{mat} \frac{1}{H} \frac{\theta_{wcap}}{\theta_{\tau}^{2,0}}$$

For benzere,

$$D_{Cap}^{eff} = 0.093 \frac{(0.038)^{3.33}}{(0.38)^{2.0}} + 1.1 \times 10^{-5} \frac{1}{0.22} \frac{(0.342)^{3.33}}{(0.38)^{2.0}}$$

$$= 0.093 \frac{0.000019}{0.1944} + 1.1 \times 10^{-5} \frac{1}{0.22} \frac{0.028}{0.1949}$$

$$= 1.2 \times 10^{-5} + 9.7 \times 10^{-6}$$

$$= 2.2 \times 10^{-5}$$

For toluene,

$$D_{(a)}^{eff} = 0.093 \frac{(0.038)^{3.33}}{(0.38)^{2.0}} + 9.4 \times 10^{-6} \frac{1}{0.26} \frac{(0.392)^{3.33}}{(0.38)^{2.0}}$$

$$= 0.093 \frac{0.000019}{0.1444} + 9.4 \times 10^{-6} \frac{1}{0.26} \frac{0.028}{0.1444}$$

$$= 1.2 \times 10^{-5} + 70 \times 10^{-6}$$

$$= 1.9 \times 10^{-5}$$

Diffusion Coefficient between ground water and soil surface Dws $D_{WS}^{eff} = \left(h_{cap} + h_{V}\right) \left(\frac{h_{csp}}{D^{eff}} + \frac{h_{V}}{D^{eff}}\right)^{-1}$

For benzene

$$D_{\nu S}^{eff} = \frac{(5+305)}{\left(\frac{5}{2.2\times10^{-5}} + \frac{305}{0.0021}\right)} = \frac{310}{2.27\times10^{5} + 1.45\times10^{5}} = \frac{1.36\times10^{-3}}{8.3\times70^{-4}}$$

ioceachers/revisions 8/4/97 Em

PROJECT: Polyorosa Ste BB(A Evaluation SUBJECT: Calculation of diffesion rectizion to SHEET Table 2, p.3 OF 3, Calcs 4/8 DATE: 18 Merch 1997 COMPUTED BY: 1. Sturmen CHECKED BY: E. Nickols

For toluene 7.63+10⁵+1.45+10³
7.6x co-4

corrections/revisions 8/4/at End

PROJECT: Polveresa Site RBCA Evaluation
SUBJECT: Calculation of Volatilization Factors

SHEET Table 3, p.1 OF 2, (a)cs 5/8

JOB NO.: 1204.00.03

DATE: 18 March 1997

COMPUTED BY:). Sturman

CHECKED BY: E Nichols

Volatilization Factor, Soil to indoor air VESESP

$$VF_{SESP} = \frac{H Ps}{[\Theta_{WS} + K_{S}P_{S} + H\Theta_{aS}]} \underbrace{\begin{bmatrix} D_{S}^{eff}/L_{S} \\ ER \cdot L_{B} \end{bmatrix}}_{ER \cdot L_{B}} \times 10^{3}$$

$$1 + \underbrace{\begin{bmatrix} D_{S}^{eff}/L_{S} \\ ER \cdot L_{B} \end{bmatrix}}_{ER \cdot L_{B}} + \underbrace{\begin{bmatrix} D_{S}^{eff}/L_{S} \\ D_{crrck}/L_{crack} \end{pmatrix}}_{Ocrrck} \times 10^{3}$$

For benzene, numerator = 0.22(1.7)terms = 0.20 + 0.38(1.7) + 0.22(0.18) 0.0028.300

= 0.42

0.0015 = 6.21×10/4

 $\frac{\text{denominator}}{\text{derms}} = 1 + \left[\frac{0.0028 \cdot 300}{0.0028 \cdot 300} \right] + \left[\frac{0.0021}{0.0021} \right] = 0.00$

 $1 + \begin{pmatrix} 0.0015 \\ 0.0015 \\ + 88.2 \\ = \begin{pmatrix} 0.0015 \\ -88.2 \\ - 88.2 \\ = \begin{pmatrix} 0.0015 \\ - 88.2$

 $VF_{SESP} = \frac{6.21 \times 10^{-4}}{89.2} \times 10^{3} = \frac{7.0 \times 10^{-3}}{m_{3/kg} soil}$

For tolvere,

numerator = $\frac{0.26(1.7)}{0.20 + 1.35(1.7) + 0.26(0.8)} = \frac{0.70}{0.0028/300}$

= 0.173

denominator = Same as benzena, above

siechans/ seuz 8/4/97 Em

0.0015 2.56+10-4

CONTRACTOR CONTINUES AND INJUNCTION OF THE										
	5 1			,	, .					

PROJECT: Polyonosa Site PBCA Evolution
SUBJECT: Calculation of Volatilization Factors

SHEET Table 3, p.2 OF 2 (alcs 6/8.

JOB NO.: 1204.00.03

DATE: 18 March 1997

COMPUTED BY: J. Sturman

CHECKED BY: E. Nichols

toluene
$$VF_{SESP} = \frac{2.56 \times 10^{-4}}{89.2} \times 10^3 = \frac{2.87 \times 10^{-5}}{9} \frac{mg/m^3.air}{mg/kg.soil}$$

Volatilization Factor, ground water to Indoor air, VFWESP

$$V_{LESP} = \frac{H \left(\frac{D_{ws}^{eff} / L_{SW}}{ER L_{B}} \right)}{1 + \left(\frac{D_{ws}^{eff} / L_{SW}}{ER L_{B}} \right) + \left(\frac{D_{ws}^{eff} / L_{Crock}}{D_{crock}^{eff} / L_{Crock}} \right) h}$$

$$\frac{1}{ER L_{B}} + \frac{D_{ws}^{eff} / L_{Crock}}{D_{ws}^{eff} / L_{Crock}} h$$

$$\frac{1}{ER L_{B}} + \frac{D_{ws}^{eff} / L_{Crock}}{D_{crock}^{eff} / L_{Crock}} h$$

$$\frac{1}{ER L_{B}} + \frac{1}{ER L_{B}} + \frac{$$

PROJECT: Polyarosa Site BBCA Evaluation SUBJECT: Calculation of SSTLS

SHEET Table 4, P.1 OF 2, Calcs JOB NO.: 1204.00.03 DATE: 18 March 1977 COMPUTED BY: J. Sturman CHECKED BY:

Calculation of Site-Specific Target Levels

Benzene

Using Cair = 143 x 10 mg/m3 as the muximum acceptable level based on the Tier 1 look-up table,

 $C_{T} = \frac{C_{air}}{VF_{SESP}} = \frac{1.43 \times 10^{1} \text{ ng/m}^3}{7.0 \times 10^{33}} = \frac{10^{-3} \text{ ng/m}^3}{\frac{mg/m^3 \text{ air}}{mg/kg \text{ soil}}} \times 10^{-3} \frac{mg}{mg}$ For soil

For ground water

$$C_{T}_{gw}^{Ben} = \frac{C_{air}}{V F_{WESP}}$$

Tolvene Cair = 5.84 × 102 Mg/m3 as above (HQ=1)

$$CT_{Soil} = \frac{Cair}{VF_{SESP}} = \frac{5.84 \, \text{F}/0^2 \, \text{Mg/m}^3 \, \text{air}}{2.87 \, \text{F}/0^{-35} \, \text{Mg/m}^3 \, \text{air}} \times 10^3 \, \text{Mg}$$

$$= 203 \text{ mg/kg Soi7} \\ 20300.$$

offections / revisions 8/4/97 End



March 21, 1997

97 MR 31 PM 4:00

LFR 1204.00.03

Mr. Scott Seery Hazardous Materials Specialist Local Oversight Program Alameda County Department of Environmental Health 1131 Harbor Bay Parkway Alameda, CA 94502-6577

Subject: Transmittal of Risk-Based Corrective Action (RBCA) Evaluation for Polvorosa Business Park Site, San Leandro, CA

Dear Mr. Seery:

As you requested, Levine-Fricke-Recon (LFR) prepared a Risk-Based Corrective Action (RBCA) evaluation for the underground storage tank release at the Polvorosa Business Park site in San Leandro, California. This RBCA evaluation was prepared in accordance with the American Society for Testing of Materials (ASTM) standard E-1739-95.

Based on the results of this evaluation, it is our recommendation that the fuel leak case be closed.

A copy of this transmittal was sent to you via fax earlier this week. We would appreciate a timely response to this evaluation. If you have any questions, please contact me at (908) 526-1000 extension 538 or Eric Nichols, P.E. at (510) 652-4500.

Sincerely,

John Sturman, P.E., R.G.

Senior Engineer

cc: Mr. Steven Chamberlain, Chamberlain Associates



March 19, 1997 LFR 1204.03

Risk-Based Corrective Action (RBCA) Evaluation for Polvorosa Business Park 1555 Doolittle Drive San Leandro, California

Based on a request from Alameda County Department of Environmental Health (ACDEH), Levine-Fricke-Recon (LFR) has prepared this Risk-Based Corrective Action (RBCA) evaluation for the Polvorosa Business Park site (hereafter referred to as "the Site"). This evaluation is derived from the Standard RBCA method applied to petroleum release sites (American Society for Testing of Materials, ASTM standard method E-1739-95). In the absence of specific state policy and guidance concerning RBCA, we have used the example policies presented in the Appendices of ASTM E-1739, modified as noted.

Step 1: Initial Site Assessment

The site investigation was conducted in several phases subsequent to UST removal in 1986. The groundwater characterization was completed by Levine-Fricke in 1988. The most current soil quality data is from samples collected by Groundwater Technology in October, 1986.

The fuel types of concern are gasoline- and diesel-range fuel hydrocarbons. The thickness of free-phase hydrocarbons on groundwater was measured when encountered. Groundwater quality samples were analyzed for TPHg, TPHd, benzene, toluene, ethylbenzene, and xylenes. Groundwater samples were not analyzed for the above analytes when free-phase hydrocarbons were detected. Soil samples (collected in 1986) were analyzed for motor fuels, benzene, toluene, and total xylenes.

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Using Table 1 of ASTM Standard E 1739-95, the site falls under Classification 3 - Long-term threat to human health, safety, or sensitive environmental receptors. (This conclusion was drawn from both pre- and post-remediation conditions)

Interim Remedial Action

During UST removal activities in 1986, some unsaturated-zone hydrocarbon-affected soils in the immediate vicinity of the USTs were removed. However, the quantity of soils which



were removed was not recorded. Based on available records, the volume of soils excavated may have been several hundred cubic yards.

From August, 1989 to August, 1993, a total fluids groundwater capture system was operated in the area where free-phase hydrocarbons were encountered. Approximately 766,000 gallons of total fluids were removed, including about 283 gallons of free-phase hydrocarbons.

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The following is a brief summary of the rationale used in screening of reasonable sources, pathways, and exposures for evaluation.

Primary Sources:

The tanks and piping were removed in 1986.

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Based on existing data, the following secondary sources are present at the site:

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- dissolved groundwater plume
- free-phase liquid plume (limited extent based on observation of hydrocarbon sheen)

Transport Mechanisms:

The site is presently covered with asphalt and building structures. A one-story warehouse building overlies much of the more affected area. A landscaped area, without pavement, exists near the hydrocarbon-affected area, and is estimated at about 10% of the building footprint. Downgradient wells on the Site indicate that the plume has had little or no additional migration after the cessation of ground water capture.

The following transport mechanisms were judged to be significant:

- o← ✓ volatilization and atmospheric dispersion (through landscaped areas around the parking
 - volatilization, migration of vapors through the building foundation, and enclosed-space accumulation



The following transport mechanisms were not considered significant as noted:

- wind erosion and atmospheric dispersion (because only a small portion of the site is unpaved)
- leaching and groundwater transport (because ground water is not extracted for human uses in the general vicinity of the Site)
- mobile free-liquid migration (because the source and mobile free-phase hydrocarbons were removed)
- stormwater/surfacewater transport (because the affected area does not include any affected surface soils or any large underground lines)

The receptors at the Site were characterized as:



• commercial/industrial

• construction workers



Applicable Risk-Based Screening Levels (RBSLs) from Tier 1 Look-up Table

Using the modified example ASTM RBCA table provided to us by ACDEH, the following is a comparison of applicable values for commercial/industrial sites:

SOIL-BASED EXPOSURE PATHWAYS RBSLS and available site data

Exposure Pathway	Target Risk Level	Benzene (mg/kg)	Toluene (mg/kg)	Xylenes (total) (mg/kg)
Volatilization to outdoor air	Cancer 1x10 ⁻⁶	0.133		
	Cancer 1x10 ⁻⁴	13.3		
	Chronic Hazard		RES	RES
Vapor intrusion into buildings	Cancer 1x10 ⁻⁶	0.00155		
	Cancer 1x10 ⁻⁴	0.155		
	Chronic Hazard		54.5	RES
Max conc. detected (1986)		0.77	0.27	1.2
Mean-over affected area @ 10.57 bgs (1986)*		0.23	6.11	1.23

- · Way detected benzene @ 10.5' BE (1986) exceeds 1E-4 RBSL for vapor intrusion into buildings from soil. However, fine grained soliments (silty day; gravely day, etc.) gonerally comprise the upper ~6' of soil column.
- . IE-4 RBSC for volatilization to outdoor air is not exceeded.



GROUNDWATER-BASED EXPOSURE PATHWAYS RBSLs and available site data

Exposure	Target Risk	Benzene	Toluene	Ethylbenzene	Xylenes
Pathway	Level	(mg/L)	(mg/L)	(mg/L)	(total)
Volatilization to outdoor air	Cancer 1x10 ⁻⁶	5.34			
	Cancer 1x10 ⁻⁴	>\$			
	Chronic Hazard		>S	>8	>S
Vapor intrusion into buildings	Cancer 1x10 ⁻⁶	0.0069			
	Cancer 1x10 ⁻⁴	0.69			
	Chronic Hazard		54.5	>\$	RES
Max, conc. detected (1995)		0.0009**	<mdl< td=""><td><mdl< td=""><td>≺MDL</td></mdl<></td></mdl<>	<mdl< td=""><td>≺MDL</td></mdl<>	≺MDL
Estimated max based on free-phase HCs		54		20	198

EXPLANATION:

RES — the selected risk level is not exceeded for pure compound selected in any concentration

- >S the selected risk level is not exceeded for all possible dissolved levels
- <MDL -- the analyte was not detected above the lab minimum detection level bgs = below ground surface (depth)
- *the mean calculated was an arithmetic mean of soil sample data at the approximate depth of shallow ground water over the area and immediate vicinity of Building C
- ** the 0.0009 mg/L benzene concentration detected was in a well upgradient of the most affected area; well LF-12 was not sampled in May 1995 due to the measurement of 0.02 feet of free-phase hydrocarbons

The estimated maximum concentrations for benzene was based on a mole fraction of 0.03 and a pure-compound solubility of 1800 mg/L. The estimated maximum concentrations of toluene and xylenes were estimated from section X1.6 and Table X1.2 of ASTM E-1739.



The lightly shaded cells of RBSL values are those exceeded at the Tier 1 screening level.

Step 4: Decision Tree/Comparison with RBSLs

Based on our conversations with ACDEH, we considered a $1x10^{-4}$ (1 in ten thousand) excess cancer risk as the appropriate target risk level for the commercial/industrial receptors at the Site.

Chemical(s) of concern concentrations exceed RBSLs? - Yes, benzene (both soil and ground water into indoor air) and toluene (soil and ground water into indoor air). Both of the groundwater RBSLs were exceeded based on the assumption of free-phase hydrocarbons rather than the actual values detected.

Remediation to Tier 1 RBSLs practicable? - No

Interim remedial action appropriate? - Yes, but to further evaluate whether remaining concentrations pose an acceptable risk, we completed a Tier 2 evaluation to derive Site Specific Target Levels (SSTLs).

Step 5: Tier 2 Calculation of SSTLs

Based on the attached calculations, the groundwater-to-outdoor air pathway is well below the applicable commercial/industrial SSTL values using a combination of default and site-specific assumptions.

Table 1 shows the Site-specific assumptions used to calculate the SSTLs. Other input parameters were taken from the default parameters presented in ASTM E1739.

Tables 2, 3 and 4 are the calculations used to derive the SSTLs for the indoor air exposure pathway. The SSTLs calculated are summarized below:

Media	Benzene	Toluene	
Soil (mg/kg)	2.04	203	_
Groundwater (mg/L)	17.0	663	

Step 6: Comparison with SSTLs

The calculated benzene and toluene concentrations in soil and groundwater necessary to exceed the indoor air SSTLs are well above those detected. The SSTLs calculated are also well above the soil and groundwater concentrations that would be expected adjacent to a



fresh liquid hydrocarbon sheen, based on equilibrium partitioning. A comparison of the calculated SSTL values above to the estimated maximum concentrations based on free-phase hydrocarbons expressed in the above groundwater RBSL table supports this conclusion.

Conclusion

We recommend no further corrective action at the Site. It is our opinion that the possible exposure pathways have been considered and evaluated to pose an insignificant risk using Site data. This conclusion is consistent with existing current State of California guidelines, since benzene has not been detected at concentrations greater than 1 mg/L at the Site and the Site is greater than 750 feet from the nearest drinking water well.

7

LEVINE, FRICKE

Symbol	Definition of Parameter	Tier 1 Default Value	Tier 2 Sile- Specific Whe	Reason for Modification
O _{as}	volumetric air Content vadose zone soils	0,26 Ca3 air	0.18 cmat	fine-grained soils under buildings are typically at 180st 50% salvated
Ows	volunatric unter content vadose zore soils	0.12 "	0,20 "	a di
to crack	volumetric air content foundation cracks	0,26	0.18	foundation cracks are typically partially folled with soil and/or water
Ow crack	Volumetric Water Content foundation cracks	0.12.	0.20 "	n y
hu	thickness of vaclose zone	295 cm	305 cm	the approximate depth to ground nater is about 9 feet at the Site
Low	depth to ground water	300 cm	310 cm	Law = hv + hcap
Ls	depth to subsurface soil	100 cm	170 cm	in 1986, some hydrocarbons were detected at 5.5 feet logs
ER	enclosed-space air exchange rate	0,00014/sec	0.0028/sec	an exchange vate of 10 volumes
		0;5	:10	perhour was used based on the use of the building as a worehouse with large doors open much of the time
7	areal fraction of cracks In foundation	0.01	0.001	the building is relatively new (about 40 years) and was Constructed with a polyathylene moisture barrier, below the foundation Slab

SUBJECT: PROJECT: Polvorosa Sike RBCA Evaluation Site-Specific Input Para meters

DATE: JOB NO.: SHEET TAKE 1 18 March 1204,00.03 10F1 1997

Calcs 1/8

68



PROJECT: Polvorosa Site RBCA Evaluation SUBJECT: <u>Calculation of diffesion Coefficients</u>

SHEET Table 2, p. 1 OF 3, Calcs 2/8 JOB NO .: 1204,00.03 DATE: 18 March 1997 COMPUTED BY: J.Sturman CHECKED BY: E, Nichols

For benzene,
$$\frac{\partial e^{3.33}}{\partial t} = \frac{\partial e^{3.33}}{\partial t} + \frac{\partial e^{3.33}}{\partial t} + \frac{\partial e^{3.33}}{\partial t}$$

For benzene,

For benzene, $D_s^{eff} \cong 0.093 \frac{(0.18)^{3.33}}{(0.38)^{2.0}} + (1.1 \times 10^{-5}) \frac{1}{0.22} \frac{(0.18)^{3.33}}{(0.38)^{2.0}} \cdot 02^{285}$ For benzene, $\stackrel{\simeq}{=} 0.093 \quad \frac{0.0033}{0.1444} + (1.1 \times 10^{-5}) \quad (4.54) \quad \frac{0.0033}{0.1444}$ $\stackrel{\simeq}{=} 0.21 \quad + 1 \times 10^{-6}$ $\stackrel{=}{=} 0.21 \quad cm/sec \quad = (.0021)$

00212

For toluene,
$$D_{s}^{eff} \cong 0.093 \frac{(0.18)^{3.33}}{(0.38)^{2.0}} + (9.4 \times 10^{-6}) \frac{1}{0.26} \frac{(0.18)^{3.33}}{(0.38)^{2.0}}$$

$$\cong 0.21 + 1 \times 10^{-6}$$

$$\cong 0.21 \text{ Cm}^{2}/\text{Sec}$$

Diffusion Coefficient through foundation cracks, Derack $D_{crack}^{eff} \cong D_{acrack}^{air} \oplus D_{acrack}^{3,33} \oplus D_{wat} \oplus D_{ws}^{3,33}$

For benzene, Deff = Same as Det for benzene = 0.21 cm²/sec

For tolvene. Derack = same as Diett for tolvene = 0.21 cm/sec



PROJECT: Polvorosa Site RBCA Evaluation
SUBJECT: Calculation of diffusion coefficients

SHEET Table 2, p.2 OF 3, Calcs 3/8

JOB NO.: 1204.00.03

DATE: 18 March 1997

COMPUTED BY: J. Styrman

CHECKED BY: E, Nichols

Diffusion Coefficient through capillary fringe,
$$D_{cap}^{eff}$$

$$D_{cap}^{eff} = D^{air} \frac{\theta_{acap}^{3,33}}{\theta_{T}^{2}} + D^{mat} \frac{1}{H} \frac{\theta_{wcap}^{3,33}}{\theta_{-}^{2,0}}$$

For benzene,

$$D_{Cap}^{eff} = 0.093 \frac{(0.038)^{3.33}}{(0.38)^{2.0}} + 1.1 \times 10^{-5} \frac{1}{0.22} \frac{(0.342)^{3.33}}{(0.38)^{2.0}}$$

$$= 0.093 \frac{0.000019}{0.1444} + 1.1 \times 10^{-5} \frac{1}{0.22} \frac{0.028}{0.1444}$$

$$= 1.2 \times 10^{-5} + 9.7 \times 10^{-6}$$

$$= 2.2 \times 10^{-5} \text{ cm}^{7/5ec}$$

For tolvene,

$$D_{cap}^{eff} = 0.093 \frac{(0.038)^{3.33}}{(0.38)^{2.0}} + 9.4 \times 10^{-6} \frac{1}{0.26} \frac{(0.342)^{3.33}}{(0.38)^{2.0}}$$

$$= 0.093 \frac{0.000019}{0.1444} + 9.4 \times 10^{-6} \frac{1}{0.26} \frac{0.028}{0.1444}$$

$$= 1.2 \times 10^{-5} + 7.0 \times 10^{-6}$$

$$= 1.9 \times 10^{-5} \text{ cm}^{2}/\text{sec}$$

Distrision Coefficient between ground under and soil surface D_{ws}^{eff} $D_{ws}^{eff} = \left(h_{cap} + h_{v}\right) \left(\frac{h_{cap}}{D_{cup}^{eff}} + \frac{h_{v}}{D_{s}^{eff}}\right)^{-1}$

For benzene

$$D_{VS}^{eff} = \frac{(5+305)}{\left(\frac{5}{2.2\times10^{-5}} + \frac{305}{421}\right)} = \frac{310}{2.27\times10^{5} + 1.45\times10^{3}} = 1.36\times10^{-3}$$



PROJECT: Polvorosa Ste RBCA Evaluation SUBJECT: Calculation of diffesion mellicents

SHEET Table 2, p.3 OF 3, Calcs 4/8 JOB NO.: 1204.00.03 DATE: 18 March 1997 COMPUTED BY: J. Sturman CHECKED BY: E, Nichols

For toluene

$$\frac{1}{1.9 \pm 10^{-5}} = \frac{(5 + 305)}{\frac{5}{1.9 \pm 10^{-5}} + \frac{305}{0.21}} = \frac{310}{2.63 \pm 10^{5} + 1.45 \pm 10^{3}} = 1.1 \pm 10^{3}$$

$$\frac{5}{1.9 \pm 10^{-5}} + \frac{305}{0.21} = \frac{310}{2.63 \pm 10^{5} + 1.45 \pm 10^{3}} = 1.1 \pm 10^{3}$$

$$\frac{5}{1.9 \pm 10^{-5}} + \frac{305}{0.21} = \frac{310}{2.63 \pm 10^{5} + 1.45 \pm 10^{3}} = \frac{1.1 \pm 10^{3}}{0.21}$$

$$= \frac{310}{2.63 \times 10^5 + 1.45 \times 10^3} = 1.1 \times 10^{-3}$$

$$\frac{310}{2.63 \times 10^5 + 1.45 \times 10^3} = 1.1 \times 10^{-3}$$

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PROJECT: Polvorosa Site RBCA Evaluation
SUBJECT: Calculation of Volatilization Factors

SHEET Table 3, p.1 OF 2, (alcs 5/8)
JOB NO.: 1204.60.03

DATE: 18 March 1997

COMPUTED BY: J. Styrman

CHECKED BY: E. Nichols

Volatilization Factor, Soil to indoorair VFSESP

$$VF_{SESP} = \frac{H\rho_{s}}{[\Theta_{WS} + K_{S}\rho_{s} + H\Theta_{as}]} \left[\frac{D_{s}^{eft}/L_{s}}{ER \cdot L_{B}} \right] \times 10^{3}$$

$$1 + \left[\frac{D_{s}^{eft}/L_{s}}{ER \cdot L_{B}} \right] + \left[\frac{D_{s}^{eft}/L_{s}}{(D_{crack}/L_{crack})_{R}} \right]$$

For benzene,

$$numerator = \frac{0.22(1.7)}{0.20 + 0.38(1.7) + 0.22(0.18)} = \frac{0.21/170}{0.0028.300}$$

 $0.646 + 0.0396$

$$\frac{0.377}{4.8856} = 0.42$$

$$= 0.000815$$

$$0.0015 = 6.21 \times 10^{-4}$$

$$6.3 \times 10^{-6}$$

denominator
terms =
$$1 + \left[\frac{0.021}{0.0028.300} \right] + \frac{0.021}{0.021/150} = \frac{21.515}{0.001}$$

$$VF_{SESP} = \frac{6.21 \times 10^{-16}}{89.2} \times 10^{3} = 7.0 \times 10^{-16} \frac{mg/m^{3} air}{mg/kg soil}$$

For tolvene,

numerator =
$$\frac{0.26(1.7)}{0.20 + 1.35(1.7) + 0.26(0.18)} = \frac{0.20/170}{0.0028/300}$$



PROJECT: Polvorosa Site PBCA Evaluation
SUBJECT: Calculation of Volatilization Factors

SHEET Table 3, p.2 OF 2 (alcs 6/8 JOB NO.: 1204.00.03

DATE: 18 March 1997

COMPUTED BY: J. Sturman

CHECKED BY: E. Nichols

Toluene
$$VF_{SESP} = \frac{2.56 \times 10^{-4}}{89.2} \times 10^3 = 2.87 \times 10^{-3} \frac{mg/m^3 \, air}{mg/kg \, soil}$$

Volatilization Factor, ground water to Indoor air, VFWESP

$$VF_{WESP} = \frac{H \left[\frac{D_{ws}^{eff}}{L_{gw}} \right]}{1 + \left(\frac{D_{ws}^{eff}}{L_{gw}} \right) + \left(\frac{D_{w$$



PROJECT: Polvorosa Site RBCA Evaluation
SUBJECT: Calculation of SSTLs

SHEET Table 4, P.1 OF 2, Calcs 7/8

JOB NO.: 1204.00.03

DATE: 18 March 1997

COMPUTED BY: J. Sturman

CHECKED BY: E. Nichols

Calculation of Site-Specific Target Levels

Benzene

Using Cair = 1.43 × 10' mg/m³ as the maximum acceptable level based on the Tier 1 look-up table,

For Soil

$$C_{T} = \frac{C_{air}}{V_{FSESP}} = \frac{1.43 \times 10^{1} \text{ ng/m}^{3}}{7.0 \times 10^{-3} \frac{\text{mg/m}^{3} \text{ air}}{\text{mg/kg soil}}} \times 10^{-3} \frac{\text{mg}}{\text{ng}}$$

$$= 2.04 \text{ mg/kg soil} \quad 4.33$$
For ground water
$$\frac{2.04 - 10^{-4}}{2.0-4 - 10^{-5}}$$

$$C_{7} = \frac{Cair}{yw} = \frac{1.43 \times 10^{1} \text{ ug/m}^{3}}{8.4 \times 10^{-4} \text{ mg/m}^{3} \text{ air}} \times 10^{-3} \text{ mg/m}^{3}$$

= 17.0 mg/L ground water

Toluene
For Soi?
$$Cair = 5.84 \times 10^2 \text{ Mg/m}^3$$
 as above $(HQ=1)$

$$C_{5iil} = \frac{Cair}{VF_{SESP}} = \frac{5.84 \times 10^2 \text{ Mg/m}^3}{2.87 \times 10^{-3} \frac{\text{Ng/m}^3 \text{ air}}{\text{Ng/kg soi?}} \times 10^3 \frac{\text{Mg}}{\text{Ng/kg soi?}}$$

= 203 mg/kg soil



PROJECT: Polvorosa Site RBCA Evalvation SUBJECT: Calculation of SSTLS

SHEET Table 4, p.20F2, Talcs 8/8 JOB NO .: 1204, 00.03 DATE: 18 March 1997 COMPUTED BY:), Styrman CHECKED BY: E. Nichols

Tolvene in ground water

$$CT_{GW}^{Tol} = \frac{Cair}{VF_{WESP}} = \frac{5.84 + 10^2 \text{ Mg/m}^3}{8.8 \times 10^{-4} \frac{\text{Mg/m}^3 \text{ air}}{\text{Mg/L} \cdot \text{Hz0}}} \times 10^{-3} \frac{\text{Mg}}{\text{Mg/L} \cdot \text{Hz0}}$$

= 663 mg/L. H2O ground water

Output Table 1

										7- 20 - 000	
	Site Name,	Polverosa	Jo	ob identification.			Software.	, GSI RBCA Spreadsheet			
	Site Location			ate Completed			Version				
	2.12 130001			Completed By.	madhulla Logan						
				, , .	··· == Ģ=···		NOTE, values	s which differ from Tier 1 default values are shown :	in bold italies and	d underlined	
	DEFA	ULT PARA	METERS								
Exposure	_		Residential		Commercia	//industrial	Surface			Commercia	al/industrial
Parameter	Definition (Units)	Adult	(1-6yrs)	(1-16 yrs)	Chronic	Constrctn	Parameters	_Definition (Units)	Residential	Chronic	Construction
ATC	Averaging time for carcinogens (yr)	70					t	Exposure duration (yr)	30	25	1
ΑTn	Averaging time for non-carcinogens (yr)	30	6	16	25	1	Α	Contaminated soil area (cm^2)	2.2E+06		1.0E+06
8W	Body Weight (kg)	70	15	35	70		W	Length of affected soil parallel to wind (cm)	1 5E+03		1.0E+03
ED	Exposure Duration (yr)	30	6	16	25	1	W.gw	Length of affected soil parallel to groundwater (c	1 5E+03		
EF	Exposure Frequency (days/yr)	350			250	180	Uarr	Ambient air velocity in mixing zone (cm/s)			
EF.Derm	Exposure Frequency for dermal exposure	350			250		delta	Air mixing zone height (cm)			
!Rgw	Ingestion Rate of Water (I/day)	2			1		Lss	Definition of surficial soils (cm)	1.0E+02		
IRs	Ingestion Rate of Soil (mg/day)	100	200		50	100	Pe	Particulate areal emission rate (g/cm^2/s)			
(Radi	Adjusted soil ing_rate (mg*yr/kg*d)	1 1E+02			9 4E+01			3			
IRa in	Inhalation rate indoor (m^3/day)	15			20		Groundwater	r Definition (Units)	Value		
iRa out	Inhalation rate outdoor (m^3/day)	20			20	10	delta gw	Groundwater mixing zone depth (cm)		-	
SA	Skin surface area (dermal) (cm^2)	5 8E+03		2.0E+03	5 8E+03	5 8E+03	1	Groundwater infiltration rate (cm/yr)			
SAadı	Adjusted dermal area (cm^2-yr/kg)	2 1E+03		· ·	1 7E+03		Ugw	Groundwater Darcy velocity (cm/yr)			
M	Soil to Skin adherence factor	1			-		Ugw tr	Groundwater Transport velocity (cm/yr)			
AAFs	Age adjustment on soil ingestion	FALSE			FALSE		Ks	Saturated Hydraulic Conductivity(cm/s)			
AAFd	Age adjustment on skin surface area	FALSE			FALSE		grad	Groundwater Gradient (cm/cm)			
tox	Use EPA tox data for air (or PEL based)	TRUE					Św	Width of groundwater source zone (cm)			
gwMCL?	Use MCL as exposure limit in groundwater?	FALSE					Sd	Depth of groundwater source zone (cm)			
goz.	and many and only and many and						BC	Biodegradation Capacity (mg/L)			
ĺ							BIO?	Is Bioattenuation Considered	FALSE		
ĺ							phi eff	Effective Porosity in Water-Bearing Unit			
							foc sat	Fraction organic carbon in water-bearing unit			
Matrix of Expo	sed Persons to	Residential			Commercia	l/Industrial					
	osure Pathways				Chronic	Constrctn	Soil	Definition (Units)	Value		
Groundwater							hc	Capillary zone thickness (cm)	5 0E+00	-	
GW i	Groundwater Ingestion	FALSE			FALSE		hv	Vadose zone thickness (cm)	3.0E+02		
GW v	Volatilization to Outdoor Air	FALSE			FALSE		rho	Soil density (g/cm²3)	17		
GW b	Vapor Intrusion to Buildings	FALSE			TRUE		foc	Fraction of organic carbon in vadose zone	0 01		
Soil Pathways							ph≀	Soil porosity in vadose zone	0.38		
Sv	Volatiles from Subsurface Soils	FALSE			FALSE		Ĺgw	Depth to groundwater (cm)	3.1E+02		
SSv	Volatiles and Particulate Inhalation	FALSE			FALSE	TRUE	Ĺs	Depth to top of affected soil (cm)	1,7E+02		
SS d	Direct Ingestion and Dermal Contact	FALSE.			FALSE	FALSE	Lsubs	Thickness of affected subsurface soils (cm)	1.3E+02		
SI	Leaching to Groundwater from all Soils	FALSE			FALSE		рH	Soil/groundwater pH	6.5		
Sb	Intrusion to Buildings - Subsurface Soils	FALSE			TRUE		•	- '	capillary	vadose	foundation
		-			•		phi.w	Volumetric water content	0.342	0.2	0.2
{							phia	Volumetric air content	0.038	<u>0.18</u>	0.1 <u>8</u>
							I				
							Building	Definition (Units)	Residential	Commercial	
	-t bl-t				0	ا - ماريمه - ا	Lb	Building volume/area ratio (cm)	2 0E+02	3.0E+02	
	eptor Distance		dential	_	Commercia		ER	Building air exchange rate (s^-1)	1 4E-04	2 3E-04	
and Location	on- or orr-site	Distance	On-Site		Distance	On-Site	Lork	Foundation crack thickness (cm)	1 5E+01		
\ .			F4:			E41.65	eta	Foundation crack fraction	<u>0.001</u>		
GW	Groundwater receptor (cm)		FALSE			FALSE					
S	Inhalation receptor (cm)		FALSE			FALSE	m =				
							Dispersive Ti			_	
Matrix of									Residential	Commercial	
Target Risks		Individual	Cumulative	_			Groundwater				
							ax	Longitudinal dispersion coefficient (cm)			
TRab	Target Risk (class A&B carcinogens)	<u>1.0E-05</u>					ay	Transverse dispersion coefficient (cm)			
TR¢	Target Risk (class C carcinogens)	1 0E-05					az	Vertical dispersion coefficient (cm)			
THQ	Target Hazard Quotient	1 0E+00					Vapor				
	Calculation Option (1, 2, or 3)	2					dcy	Transverse dispersion coefficient (cm)			
Opt Tier	RBCA Tier	2					dcz	Vertical dispersion coefficient (cm)			

য়ি Levine-Fricke-Recon

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FAX TRANSMISSION: This cover page plus	15	page(s)
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Time			
Date	20 March	199.7	
From	John Sturm		
LFR Project No.	1204.00.0	3	
Delive	er To _i	Name Of Firm	FAX Number
Mr. Stit See	ry	Alameda Country - Emiron Health	510-837-0335

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Comments:

This fax contains a RBCA evaluation of the Polvorosa site in San Leandro. Seven pages of text and eight pages of calculations are enclosed. The hard copy is being sent to you via regular mail. I'll give you a call in a couple weeks to check with you on your schedule for review. If you have any grestions, please call me at 908:526-1000, extension 538. Thank for

:forms/faxmast.doc

March 19, 1997

LFR 1204.03

Risk-Based Corrective Action (RBCA) Evaluation for Polyorosa Business Park 1555 Doolittle Drive San Leandro, California

Based on a request from Alameda County Department of Environmental Health (ACDEH), Levine-Fricke-Recon (LFR) has prepared this Risk-Based Corrective Action (RBCA) evaluation for the Polvorosa Business Park site (hereafter referred to as "the Site"). This evaluation is derived from the Standard RBCA method applied to petroleum release sites (American Society for Testing of Materials, ASTM standard method E-1739-95). In the absence of specific state policy and guidance concerning RBCA, we have used the example policies presented in the Appendices of ASTM E-1739, modified as noted.

Step 1: Initial Site Assessment

The site investigation was conducted in several phases subsequent to UST removal in 1986. The groundwater characterization was completed by Levine Fricke in 1988. The most current soil quality data is from samples collected by Groundwater Technology in October, 1986.

The fuel types of concern are gasoline- and diesel-range fuel hydrocarbons. The thickness of free-phase hydrocarbons on groundwater was measured when encountered. Groundwater quality samples were analyzed for TPHg, TPHd, benzene, toluene, ethylbenzene, and xylenes. Groundwater samples were not analyzed for the above analytes when free-phase hydrocarbons were detected. Soil samples (collected in 1986) were analyzed for motor fuels, benzene, toluene, and total xylenes.

Step 2: Site Classification and Initial Response Action

Using Table 1 of ASTM Standard E 1739-95, the site falls under Classification 3 - Long-term threat to human health, safety, or sensitive environmental receptors. (This conclusion was drawn from both pre- and post-remediation conditions)

Interim Remedial Action

During UST removal activities in 1986, some unsaturated-zone hydrocarbon-affected soils in the immediate vicinity of the USTs were removed. However, the quantity of soils which

were removed was not recorded. Based on available records, the volume of soils excavated may have been several hundred cubic yards.

From August, 1989 to August, 1993, a total fluids groundwater capture system was operated in the area where free-phase hydrocarbons were encountered. Approximately 766,000 gallons of total fluids were removed, including about 283 gallons of free-phase hydrocarbons.

Step 3: Tier 1 Evaluation

The following is a brief summary of the rationale used in screening of reasonable sources, pathways, and exposures for evaluation.

Primary Sources:

The tanks and piping were removed in 1986.

Secondary Sources:

Based on existing data, the following secondary sources are present at the site:

- impacted subsurface soils
- dissolved groundwater plume
- free-phase liquid plume (limited extent based on observation of hydrocarbon sheen)

Transport Mechanisms:

The site is presently covered with asphalt and building structures. A one-story warehouse building overlies much of the more affected area. A landscaped area, without pavement, exists near the hydrocarbon-affected area, and is estimated at about 10% of the building footprint. Downgradient wells on the Site indicate that the plume has had little or no additional migration after the cessation of ground water capture.

The following transport mechanisms were judged to be significant:

- volatilization and atmospheric dispersion (through landscaped areas around the parking area)
- volatilization, migration of vapors through the building foundation, and enclosed-space accumulation

The following transport mechanisms were not considered significant as noted:

- wind erosion and atmospheric dispersion (because only a small portion of the site is unpaved)
- leaching and groundwater transport (because ground water is not extracted for human uses in the general vicinity of the Site)
- mobile free-liquid migration (because the source and mobile free-phase hydrocarbons were removed)
- stormwater/surfacewater transport (because the affected area does not include any affected surface soils or any large underground lines)

The receptors at the Site were characterized as:

9085267886

- commercial/industrial
- construction workers

Applicable Risk-Based Screening Levels (RBSLs) from Tier 1 Look-up Table

Using the modified example ASTM RBCA table provided to us by ACDEH, the following is a comparison of applicable values for commercial/industrial sites:

SOIL-BASED EXPOSURE PATHWAYS RBSLS and available site data

	arget Risk evel	Benzene (mg/kg)	Toluené (mg/kg)	Xylenes (total (mg/kg)
Volatilization (to outdoor air	ancer 1x10 ⁻⁶	0.133		
C	ancer 1x10 ⁻⁴	13.3		
C	hronic Hazard		RES	RES
Vapor C intrusion into buildings	ancer 1x10 ⁻⁶	0.00155		
C	ancer 1x10 ⁻⁴	0.155		
	bronic Hazard	1100	54.5	RES
Mas cone Jelected 1986)		0,77	0.27	1.2
Assnerver fiscase area 2.10.5° bgs		0.23	0.11	0.28
1986)*				

GROUNDWATER-BASED EXPOSURE PATHWAYS RBSLs and available site data

Exposure Pathway	Farget Risk Level	Benzena (mg/L)	Foltrene (mg/L)	Ethylbenzem (mg/L)	Xylenes (total)
Volatilization to outdoor air	Cancer 1x10 ⁻⁶	5.34			
,	Cancer 1x10 ⁻⁴	>S			
2.4	Chronic Hazard		>\$	> S	>S
Vapor intrusion into buildings	Cancer 1x10 ⁻⁶	0.9069			
	Cancer 1x10 ⁻⁴	0.69			
	Chronic Hazard		54:5	>S	RES
Max. conc. detected (1995)		0.0009**	≺MDL	*MDL	< MD _L
Estimated max based on free-phase HCs		54	83	20	198

EXPLANATION:

- RES the selected risk level is not exceeded for pure compound selected in any concentration
- >S the selected risk level is not exceeded for all possible dissolved levels
- SMDL the analyte was not detected above the lab minimum detection level bgs = below ground surface (depth)
- *the mean calculated was an arithmetic mean of soil sample data at the approximate depth of shallow ground water over the area and immediate vicinity of Building C
- ** the 0.0009 mg/L benzene concentration detected was in a well upgradient of the most affected area; well LF-12 was not sampled in May 1995 due to the measurement of 0.02 feet of free-phase hydrocarbons
- The estimated maximum concentrations for benzene was based on a mole fraction of 0.03 and a pure-compound solubility of 1800 mg/L. The estimated maximum concentrations of toluene and xylenes were estimated from section X1.6 and Table X1.2 of ASTM E-1739.

The lightly shaded cells of RBSL values are those exceeded at the Tier 1 screening level.

Step 4: Decision Tree/Comparison with RBSLs

Based on our conversations with ACDEH, we considered a $1x10^4$ (1 in ten thousand) excess cancer risk as the appropriate target risk level for the commercial/industrial receptors at the Site.

Chemical(s) of concern concentrations exceed RBSLs? - Yes, benzene (both soil and ground water into indoor air) and toluene (soil and ground water into indoor air). Both of the groundwater RBSLs were exceeded based on the assumption of free-phase hydrocarbons rather than the actual values detected.

Remediation to Tier 1 RBSLs practicable? - No

Interim remedial action appropriate? - Yes, but to further evaluate whether remaining concentrations pose an acceptable risk, we completed a Tier 2 evaluation to derive Site Specific Target Levels (SSTLs).

Step 5: Tier 2 Calculation of SSTLs

Based on the attached calculations, the groundwater-to-outdoor air pathway is well below the applicable commercial/industrial SSTL values using a combination of default and site-specific assumptions.

Table 1 shows the Site-specific assumptions used to calculate the SSTLs. Other input parameters were taken from the default parameters presented in ASTM E1739.

Tables 2, 3 and 4 are the calculations used to derive the SSTLs for the indoor air exposure pathway. The SSTLs calculated are summarized below:

Media	Benzene	Toluene	
Soil (mg/kg)	2.04	203	
Groundwater (mg/L)	17.0	663	

Step 6: Comparison with SSTLs

The calculated benzene and toluene concentrations in soil and groundwater necessary to exceed the indoor air SSTLs are well above those detected. The SSTLs calculated are also well above the soil and groundwater concentrations that would be expected adjacent to a

fresh liquid hydrocarbon sheen, based on equilibrium partitioning. A comparison of the calculated SSTL values above to the estimated maximum concentrations based on free-phase hydrocarbons expressed in the above groundwater RBSL table supports this conclusion.

Conclusion

We recommend no further corrective action at the Site. It is our opinion that the possible exposure pathways have been considered and evaluated to pose an insignificant risk using Site data. This conclusion is consistent with existing current State of California guidelines, since benzene has not been detected at concentrations greater than 1 mg/L at the Site and the Site is greater than 750 feet from the nearest drinking water well.



PROJECT: Polivorosa Site RBCA Evaluation SUBJECT: Calculation of diffesion Calificients

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SHEET Table 2, p. 1 OF 3. Culcs 2/8 JOB NO .: 1204, 00.03 DATE: 18 March 1997 COMPUTED BY: J. Sturman CHECKED BY: F. Nichola

Diffusion Coefficient Soil - Vapor, Def
Deff = Duair
$$\frac{\theta_{as}^{3.33}}{\theta_{as}^{2.0}} + Duat \frac{1}{H} = \frac{\theta_{ws}^{3.33}}{\theta_{as}^{2.0}}$$

For benzene, $D_5^{eff} = 0.093 \frac{(0.18)^{3.33}}{(0.38)^{2.0}} + (1.1 \times 10^{-5}) \frac{1}{0.22} \frac{(0.18)^{3.33}}{(0.38)^{2.0}}$ $\stackrel{\simeq}{=} 0.093 \quad \frac{0.0033}{0.1444} + (1.1 \times 10^{-5}) \quad (4.54) \quad \frac{0.0033}{0.1444}$ $\stackrel{\simeq}{=} 0.21 \quad + 1 \times 10^{-6}$

= 0.21

For tolvene, $D_{5}^{\text{eff}} \cong 0.093 \frac{(0.18)^{3.33}}{(0.38)^{2.0}} + (9.4 \times 10^{-6}) \frac{1}{0.26} \frac{(0.18)^{3.33}}{(0.38)^{2.0}}$ € 20.21 + 1×10-6

Diffusion Coefficient through foundation cracks, Derack $D_{crack}^{ctf} \cong D^{dir} \underbrace{\theta_{acrack}^{3,33}}_{A^2} + D^{wat} \underbrace{\frac{1}{H}}_{H^2} \underbrace{\theta_{ws}^{3,33}}_{A^2}$ For benzene,

Deff = same as Deft for benzene = 0.21

For tolvene, Derack = same us Dief for tolvene = 0.21



PROJECT: Polvoresa Site RBCA Evaluation SUBJECT: Calculation of diffusion coefficients

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SHEET Table 2, p. 2 OF 3, Calcs 3/8 18 March 1997 COMPUTED BY: J. Sturmen CHECKED BY: E. Nichols

Diffesion Coefficient through capillary finge, Deap $\mathcal{D}_{cap}^{eff} = \mathcal{D}^{air} \frac{\theta_{acap}}{\theta_{+}^{2}} + \mathcal{D}^{wat} \frac{1}{H} \frac{\theta_{wcap}}{\theta_{-}^{2.0}}$

For benzene,

$$D_{cap}^{eff} = 0.093 \frac{(0.038)^{3.33}}{(0.38)^{2.0}} + 1.1 \times 10^{-5} \frac{1}{0.22} \frac{(0.342)^{3.33}}{(0.38)^{2.0}}$$

$$= 0.093 \frac{0.000019}{0.1444} + 1.1 \times 10^{-5} \frac{1}{0.22} \frac{0.028}{0.1449}$$

$$= 1.2 \times 10^{-5} + 9.7 \times 10^{-6}$$

$$= 2.2 \times 10^{-5}$$

For tolvena,

$$D_{cap}^{eff} = 0.093 \frac{(0.038)^{3.33}}{(0.38)^{2.0}} + 9.4 \times 10^{-6} \frac{1}{0.26} \frac{(0.342)^{3.33}}{(0.38)^{2.0}}$$

$$= 0.093 \frac{0.000019}{0.1444} + 9.4 \times 10^{-6} \frac{1}{0.26} \frac{0.028}{0.1449}$$

$$= 1.2 \times 10^{-5} + 7.0 \times 10^{-6}$$

$$= 1.9 \times 10^{-5}$$

Diffusion Coefficient between ground water and soil surface Dus $D_{WS} = \left(h_{Cap} + h_{V}\right) \left(\frac{h_{Cap}}{D^{aff}} + \frac{h_{V}}{D^{aff}}\right)^{-1}$

For benzene

$$D_{WS}^{eff} = \frac{(5+305)}{\left(\frac{5}{2.2\times10^{-5}} + \frac{305}{0.21}\right)} = \frac{310}{2.27\times10^{5} + 1.45\times10^{3}} = 1.36\times10^{3}$$

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PROJECT: Polyprosa Ste BBCA Evaluation SUBJECT: Calculation of diffesion calliciants

SHEET Table 2, p.3 OF 3, Calcs 4/8 JOB NO .: 1204.00.03 DATE: 18 Merch 1997 COMPUTED BY: 1. Sturmen CHECKED BY: E. Nichols

For tolvene

$$D_{ws}^{eff} = \underbrace{\frac{(5+305)}{5}}_{1.9\times10^{-5}} + \frac{305}{0.24} = \frac{310}{2.63\times10^{5}+1.45\times10^{3}} = 1.1\times10^{-3}$$



PROJECT: Polvorosa Site RBCA Evaluation SUBJECT: Calculation of Volatilization Factors

SHEET Table 3, p.1 OF 2, Calcs 5/8 JOB NO .: 1204. 60.03 DATE: 18 March 1997 COMPUTED BY: J. Sturman CHECKED BY: E. Niche/s

Volatilization Factor, Soil to indoorair VFSESP

$$VF_{SESP} = \frac{HPs}{\left[\frac{D_sef}{Ls}\right] + \left[\frac{D_sef}{ER Ls}\right]} \left[\frac{D_sef}{Ls}\right] \times 10^{ef}$$

$$VF_{SESP} = \frac{1 + \left[\frac{D_sef}{ER Ls}\right] + \left[\frac{D_sef}{D_{crack}/L_{crack}}\right] \times 10^{ef}}{1 + \left[\frac{D_sef}{ER Ls}\right] + \left[\frac{D_sef}{D_{crack}/L_{crack}}\right] \times 10^{ef}}$$

For benzene, numerator = $\frac{0.22(1.7)}{0.20 + 0.38(1.7) + 0.22(0.18)} = \frac{0.21/170}{0.0028.300}$

= 0.42

. 0.0015 = 6.21×10-4

denominator
Herms =
$$1 + \left[\frac{0.21/170}{0.0028.300}\right] + \frac{0.21/170}{0.21/15} (0.001)$$

 $1 + 0.0015 + 88.2 = 81.2$

 $VF_{SESP} = \frac{6.21 \times 10^{-4}}{89.2} \times 10^{3} = 7.0 \times 10^{-3} \frac{mg/m^{3} air}{mg/kg soil}$

For tolvere,

= 0.173

denominator = same as benzene, above



PROJECT: Polvorosa Site PBCA Evolvation
SUBJECT: Calculation of Volatilization Factors

SHEET Table 3, p.2 OF 2 (alcs 6/8 JOB NO.: 1204.00.03

DATE: 18 March 1997

COMPUTED BY: J. Sturman

CHECKED BY: F. Nichols

Toluene
$$VF_{SESP} = \frac{2.56 \times 10^{-4}}{89.2} \times 10^3 = 2.87 \times 10^{-3} \frac{mg/m3.ain}{mg/kg.soil}$$

Volatilization Factor, ground water to indoor air, VFWESP

$$VF_{WFSP} = \frac{H \left[\frac{D_{ws}}{ER} \right]_{LGW}}{1 + \left(\frac{D_{ws}}{L_{GW}} \right) + \left(\frac{D_{ws}}{D_{ws}} \right)_{LGW}} \times 10^{3}$$

$$1 + \left(\frac{D_{ws}}{L_{BW}} \right) + \left(\frac{D_{ws}}{D_{ws}} \right)_{LGW}}{\left(\frac{D_{eff}}{D_{crack}} \right)_{LCrack}} \times 10^{3}$$
For benzene,

$$1 + \left(\frac{1.36 + 10^{-3}}{0.0028.300} \right) + \frac{1.36 + 10^{-3}}{0.21 / 15.0001}$$

$$1 + \left(\frac{1.36 + 10^{-3} / 3.10}{0.0028.300} \right) + \frac{1.36 + 10^{-3} / 3.10}{0.21 / 15.0001}$$

$$1 + 5.2 + 10^{-6} + 0.31 = 1.31$$

$$VF_{WESP} = \frac{1.1 \times 10^{-6}}{1.31} \times 10^{3} = 8.4 \times 10^{-4} \frac{mg/m^{3} \text{ air}}{mg/L \cdot H_{2}0}$$

For tolvene,

Tunerator =
$$0.26 \frac{1.1 \times 10^{-3}/310}{0.0028 \cdot 300} = 1.1 \times 10^{-6}$$

denominator = $1 + \left(\frac{1.1 \times 10^{-3}/310}{0.0028 \cdot 300}\right) + \frac{1.1 \times 10^{-3}/310}{0.21/15 \cdot 0.001} = 1.25$
VF masp = $\frac{1.1 \times 10^{-6}}{1.25} \times 10^{3} = 8.8 \times 10^{-4} \frac{m_{Ym}^{3} air}{m_{g/1} \cdot H_{2}0}$



PROJECT: Polyerusa Site RBCA Evaluation SUBJECT: Calculation of SSTLS

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SHEET Table 4, P.1 OF 2, Calcs 7/8 JOB NO .: 1204.00.03 DATE: 18 March 1977 COMPUTED BY: J. Sturman CHECKED BY: E. Nichols

Calculation of Site-Specific Target Levels

Benzene

Using Cair = 143 × 10 1 mg/m3 as the maximum acceptable level based on the Tier 1 look-up table,

For soil $C_{7 \text{ Sen}} = \frac{C_{ast}}{VF_{5ESP}} = \frac{1.43 \times 10^{1} \text{ ug/m}^{3}}{7.0 \times 10^{-3} \frac{\text{mg/m}^{3} \text{ air}}{\text{mg/kg soil}}} \times 10^{-3} \frac{\text{mg}}{\text{mg/kg soil}}$ = 2.04 mg/kg soil

For ground water

Cyben =
$$\frac{\text{Cair}}{\text{VFWESP}} = \frac{1.43 \times 10^{4} \text{ ug/m}^{3}}{8.4 \times 10^{-4} \frac{\text{mg/m}^{3} \text{ air}}{\text{mg/L} \cdot \text{H}_{2}0}} \times 10^{-3} \frac{\text{mg}}{\text{mg/L}}$$

- 17.0 mg/L ground water

Tolvene Cair = 5.84 × 102 mg/m3 as above (HQ=1) $CT_{Siil}^{Tal} = \frac{Cair}{VF_{SESP}} = \frac{5.84 \times 10^{2} \, \text{Mg/m}^{3}}{2.87 \times 10^{-3} \, \frac{\text{Mg/m}^{3} \, \text{air}}{\text{Mg/kg siil}}} \times 10^{-3} \, \frac{\text{Mg}}{\text{Mg/kg siil}}$

= 203 mg/kg soil



PROJECT: Polyorosa Site RELA Evaluation SUBJECT: Calculation of SSTLS

SHEET Table 4, P.20F2, Calcs 8/8 DATE: 18 March 1997 COMPUTED BY: J. Styrmen CHECKED BY: E. Nichols

Tolvene in ground water

$$C_{TGW} = \frac{C_{air}}{VF_{wesp}} = \frac{5.84 \times 10^{2} \, Mg/m^{3}}{8.6 \times 10^{-4} \, Mg/L \cdot Hzp} \times 10^{-3} \, Mg/L \cdot Hzp$$

663 mg/L. H2O ground unter