REPORT OF INVESTIGATION CHEVRON SS 9-2582 7240 DUBLIN BOULEVARD DUBLIN, CALIFORNIA

Prepared for

Chevron U.S.A. Inc. San Ramon, California

Prepared by

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1. INTRODUCTION

1.1 SCOPE

At the request of Chevron U.S.A. Inc., EA Engineering, Science, and Technology, Inc. (EA) conducted a soil vapor contaminant assessment (SVCA) at Chevron Service Station 9-2582 in Dublin, California. This report describes the SVCA and presents the results.

1.2 SITE SETTING

Chevron Service Station (SS) 9-2582 is located at 7240 Dublin Boulevard, Dublin, Alameda County, California (Figure 1). It is on the southwestern corner of the intersection of Dublin Boulevard and Village Parkway (Figure 2). The neighborhood consists predominantly of commercial properties, but U.S. Interstate 680 and an unnamed canal border the site on the west, and there are residences to the northeast at a distance of less than one-quarter mile. There are no other known underground petroleum product storage tanks in the immediate area of the site.

The site is located in a relatively flat area at approximately 330 feet above mean sea level. The general topographic gradient is about 10 feet per mile (0.002), down to the south-southeast. There is significant topographic relief on the site itself: large planters on the north side of the station are approximately two feet higher than the paved area.

The Dougherty Hills are approximately one mile north-northeast of the site, and other hills of the Diablo Range rise from an elevation of 400 feet msl to 800 feet, beginning approximately one mile to the west. Other than the unnamed concrete channel that drains to the southeast, built to protect Highways 680 and 580 from storm runoff from the numerous intermittent streams in

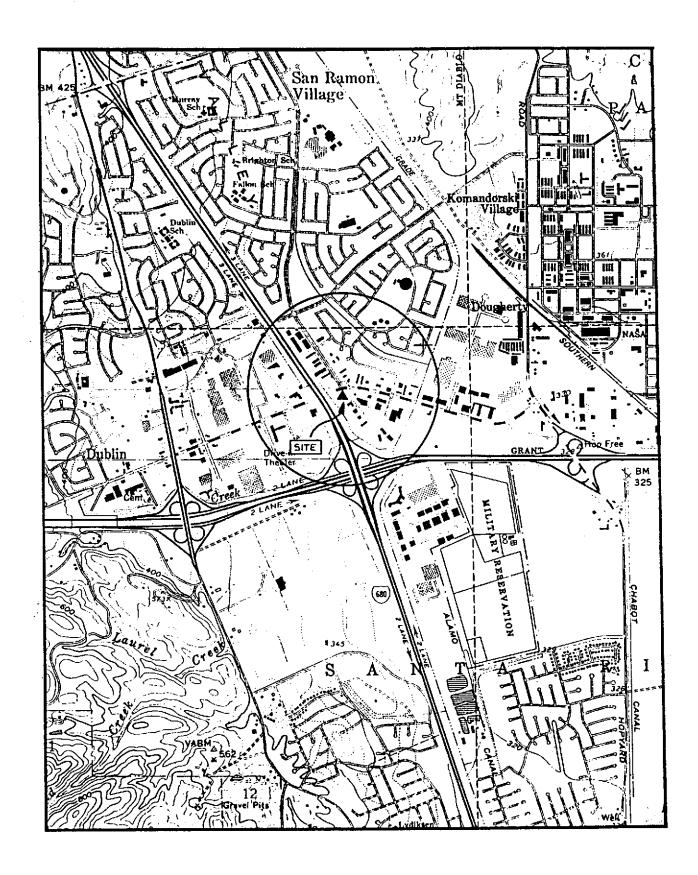


Figure 1. Location and topography of Chevron SS 9-2582, 7420 Dublin Blvd., Dublin, California

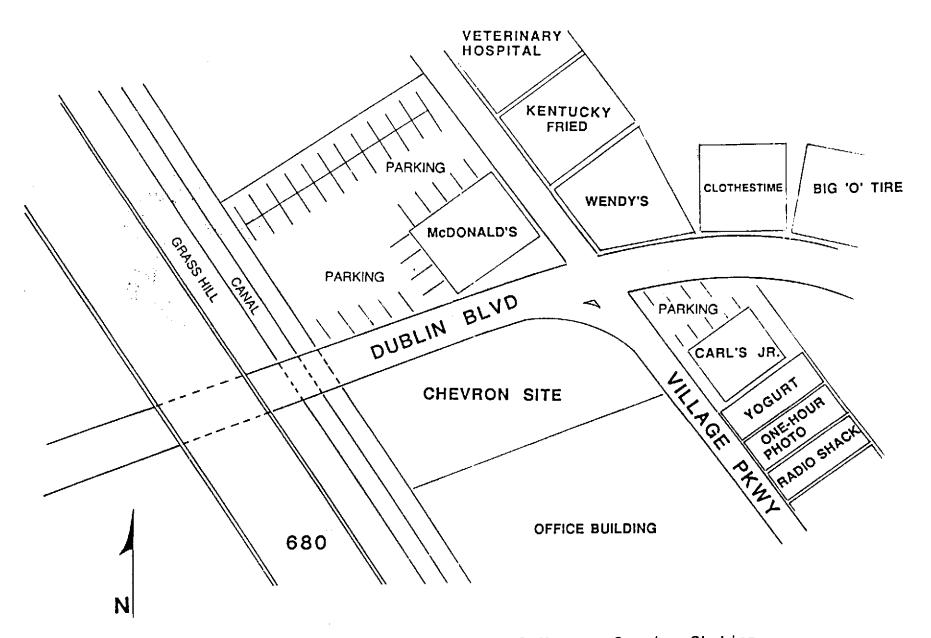


Figure 2. Land use in the vicinity of Chevron Service Station 9-2582, Dublin, California.

the hills, the nearest downgradient surface water is Laurel Creek, approximately 4,000 feet southwest of the site.

1.3 HYDROGEOLOGY AND WATER USE

The site is located on semi-consolidated and unconsolidated Quaternary alluvium which was deposited in lake, playa, and stream environments in the San Ramon Valley. The elevation of ground water in the vicinity is between 320 and 330 feet above sea level (Killingsted 1988, pers. comm.), which puts it at a depth of 10 feet or less.

No soil borings are available for this site, so the detailed local subsurface geology is unknown. The expected ground-water gradient is to the south (Killingsted 1988, personal communication).

The City of Dublin (Killingsted 1988, personal communication) has no records of public or private wells within one-half mile of the site, but because there are older businesses and homes in the area, there may be local wells not on record. The municipal water district uses reservoir water primarily, but one-third of its total water comes from wells in the region.

2. SOIL VAPOR CONTAMINANT ASSESSMENT

Following a subsurface gasoline release, as free product migrates downward towards the ground water some of the gasoline will be adsorbed to the soils, and some will vaporize. In the case of a spill of sufficient volume to exceed the soil binding capacity, free liquid will reach ground water, at which point it will float and may begin to vaporize and solubilize.

On the basis of these and other physicochemical properties and behaviors of hydrocarbon mixtures, described in Appendix A, it can be seen that associated with any ground-water, soil, or free-product contamination there is vapor phase contamination. The SVCA technique takes advantage of this, and through the collection and analysis of soil vapor permits rapid delineation of the extent of contamination.

2.1 SVCA SAMPLING

13

On 11 February 1988, EA conducted an SVCA at the Dublin site.

Samples of shallow soil gas were taken at 15 on-site locations

(Figure 3). Vertical profiles of soil gas hydrocarbon composition were made at

- . the southeastern corner of the tank field (V1, at 3, 5.5, 8, 10.5, and 13 feet)
- . the northwestern corner of the tank field (V2, at 3, 10.5 and 15.5 feet)
- . the southwestern corner of the tank field (V3, at 3 and 13 feet)
- . the northeastern planter (V4, at 3 and 15.5 feet)
- . the southern pump island (V7, at 3 and 5.5 feet)
- . the northern pump island (V8, at 3 and 5.5 feet).

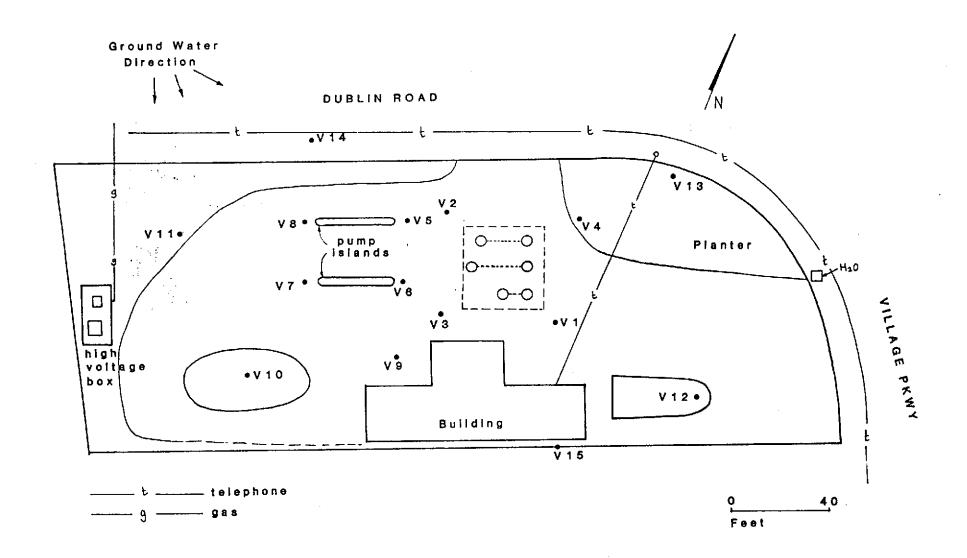


Figure 3. Site plan, with soil vapor sampling points, of Chevron SS 9-2582, Dublin, California, February 1988.

Before each sample was collected, a vacuum pump was used to purge previously collected vapor from the probe to ensure that the sample was not contaminated. The vacuum pressure reading on the purging apparatus was recorded. This vacuum pressure is related to the soil's gas permeability: a high vacuum reading suggests that there is a resistance to soil gas movement and a vacuum is being created between the tip of the probe and the pump. If this vacuum is maintained after the pumping period, a representative sample of the soil gas may not be obtained, and anomalously low hydrocarbon readings may result. In most situations, when the purging pump is turned off and the stopcock is closed the initial vacuum is released and the vacuum pressure reading drops. A low reading indicates that there is a free flow of soil gas from the vadose zone through the probe, and after purging the sample is assumed to be representative.

The samples were collected through a septum with a microsyringe and injected into an HNU 421 chromatograph for analysis. 421 is a laboratory-size, temperature-programmable gas chromatograph equipped with a flame ionization detector (FID). hydrogen-air flame ionizes compounds, generating an energy increase in the detector, which appears as an electrical signal. Vapor samples are injected into the gas chromatograph, separated on an analytical column, sensed by the detector, integrated, and reported as individual compounds on chromatograms. The instrument is operated isothermally at 60°C and the capillary column flow rate is 10 ml/min. These conditions ensure peak retention time stability and prevent contaminant build-up within the Blanks are run to verify that the system is free of contamination; as necessary, the instrument is re-calibrated by These data ensure system reproducibility. injecting standards.

The chromatograph yields a response in the form of an electrical signal, measured in volts; this is recorded and integrated across time by a Shimadzu C-R3A integrator. The peak area is expressed as volt-seconds (V-sec). The instrument is calibrated with a

multicomponent standard consisting of known concentrations of benzene, toluene, xylenes, and ethylbenzene. The integrator calculates and stores the response ratio, V-sec:ppm. The ratio for each component of the standard is used to quantify the concentrations of identifiable vapors in field samples according to their V-sec values.

The concentrations of unidentified compounds are calculated in a similar manner. Although petroleum hydrocarbons produce variable instrumental responses, the assumption may be made that all of the hydrocarbon constituents have response-to-concentration ratios approximately equivalent to that of benzene and that all quantifications may be based on the response ratio for benzene. In the table describing the results of the assessment, the column entitled "Peaks Prior to Benzene" represents the sum of the responses in V-sec for all peaks eluting prior to benzene, proportioned to the calibrated V-sec response for benzene. Similarly, the column entitled "Total Volatile Hydrocarbons" represents the sum of all V-sec responses, proportioned to that for benzene.

2.2 SVCA RESULTS

The results (Appendix B) of the SVCA conducted at the Dublin site are summarized in Table 1. The surfaces of the planters are about two feet above the paved surfaces, so for comparison of the vertical positions of constituent concentrations, the depths for points V4, V11, and V13 should be considered to be 2 feet less than those reported in Table 1. Isoconcentration contours of hydrocarbon constituents in the soil vapor are plotted in Figures 4 through 7. Moderate-to-high concentrations of hydrocarbons were detected at depths of 6 to 13 feet near the southern pump island, especially at point V7. Except for the very mobile compounds that elute prior to benzene, the hydrocarbon plumes appear to remain within the site boundaries.

TABLE 1 CONCENTRATIONS OF HYDROCARBON CONSTITUENTS IN SOIL VAPOR AT CHEVRON SS 9-2582, DUBLIN, CALIFORNIA 11 FEBRUARY 1988

Sample Location	Depth (ft)	Peaks Prior ^a to Benzene (ppm) ^b	Benzene (ppm)	Toluene (ppm)	o-Xylene (ppm)	m,p-Xylene (ppm)	Ethyl- benzene (ppm)	Peaks Not Otherwise Identified (ppm) ^b	Total Volatile Hydro- carbons (ppm) ^b
V1/A	3	. <1	< 1	<1	<1	<1	<1	6	6
V1/B	5.5	<1	<1	<1	<1	<1	<1	<1	<1
V1/C	8	84	<1	<1	<1	<1	<1	< 1	85
V1/D	10.5	Ċ1	<1	<1	<1	<1	<1	<1	<1
VI/E	13	330	1	<1,	<1	<1	<1	6	340
V2/A	3	<1	<1	<1	<1	<1	<1	<1	1
V2/B	10.5	<1	<1	<1	<1	< 1	<1	<1	<1
V2/C	15.5	. 1	<1	<1	<1	<1	<1	<1	1
V3/A	3	320	<1	<1	<1	<1	<1	14	330
V3/B	13	1	₹1	<1	<1	<1	<1	<1	1
V4/A	3	4	<1	<1	< 1	<1	<1	<1	5
V4/B	15.5	2,000	11	6	<1	<1	<1	120	2,100
v 5	3	1	<1	<1	<1	<1	<1	<1	1
V6	3	<1	<1	5	<1	<1	<1	1	6
V7/A	3	<1 '	<1	<1	<1	<1	<1	<1	<1
W7/B	5.5	7,500	150	200	<1	<1	3	1,800	9,700
V8/A	3	13	<1	<1	<1	<1	<1	<1	14
V8/B	5.5	1,100	65	140	<1	2	5	860	2,200
V9	4	15	<1	<1	<1	<1	<1	<1	15
V10	6.5	150	<1	<1	<1	<1	<1	<1	150
V11	6.5	<1	<1	<1	<1	< 1	<1	<1	<1
V12	13	280	₹1	<1	<1	<1	<1	<1	280
V13	15.5	3	<1	<1	<1	<1	<1	<1	3
V14	6.5	78	<1	<1	<1	<1	<1	<1	78
V15	13	<1	< 1	<1	<1	<1	<1	< 1	<1

a. Early peaks from blank data subtracted from total peaks prior to benzene.b. Quantification based on V-sec:ppm ratio for benzene (see text).

BLANK DATA

Time	Peaks Prior ^a to Benzene (ppm) ^b	Benzene (ppm)	Toluene (ppm)	o-Xylene (ppm)	m,p-Xylene (ppm)	Ethyl- benzene (ppm)	Not Otherwise Identified (ppm)D	Volatile Hydro- carbons (ppm)
1040	*** 1	<0.1	<0.3	<0.1	<0.1	<0.1	<0.1	1
1232*	1	<0.1				 `		
1528*	1	<0.1						 '

PERCENTAGE OF STANDARD RECOVERED

Test Time	Benzene (ppm)	Toluene (ppm)	o-Xylene (ppm)	m,p-Xylene (ppm)	Ethyl- benzene (ppm)
1010	100	100	100	100	100
133 7	104	107	137	115	126

GASOLINE STANDARD

Sample	Peaks Prior ^a to Benzene (ppm) ^b	Benzene (ppm)	Toluene (ppm)	o-Xylene (ppm)	m,p-Xylene (ppm)	Ethyl- benzene (ppm)	Peaks Not Otherwise Identified (ppm) ^D	Volatile Hydro- carbons (ppm)b
Regular	150,000	4,600	6,200	550	1,700	380	13,000	180,000

Note: * indicates an abbreviated blank chromatogram, used to determine peaks eluting prior to benzene.

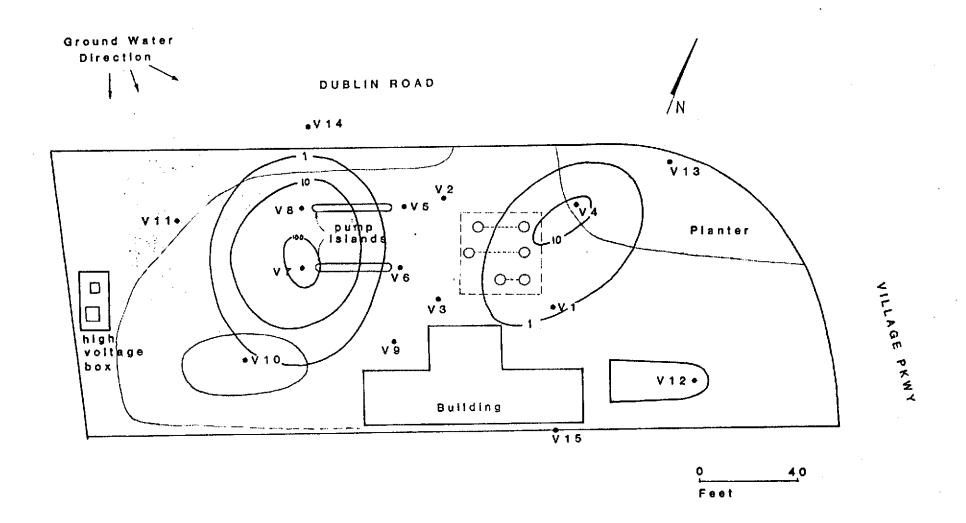


Figure 4. Isoconcentration contours (ppm) of henzene in the shallow soil gas at Chevron Service Station 9-2582, Dublin, California, February 1988.

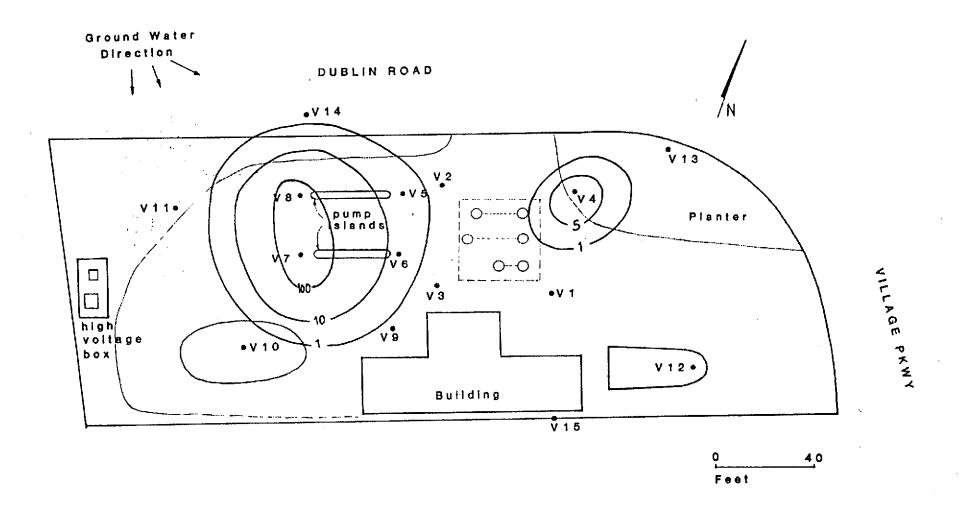


Figure 5. Isoconcentration contours (ppm) of **tolurne** in the shallow soil gas at Chevron Service Station 9-2582, Dublin, California, February 1988.

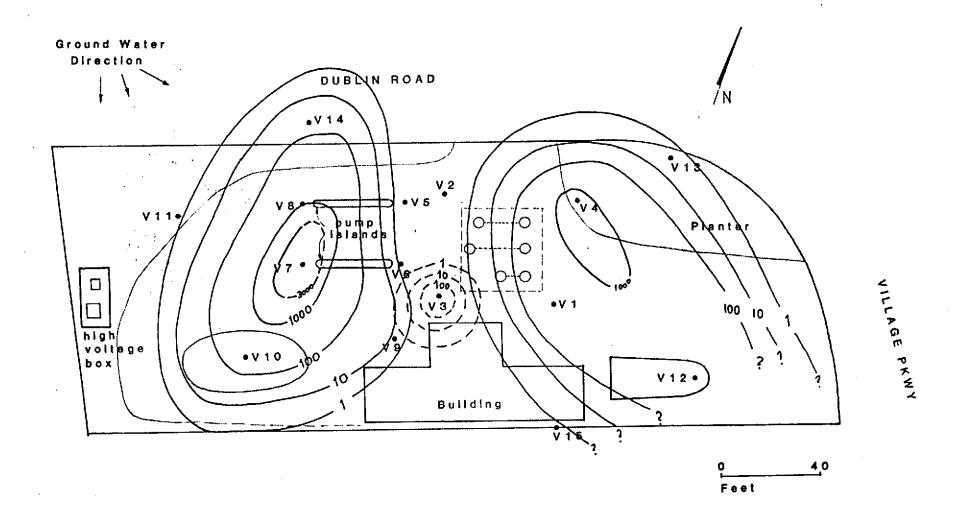


Figure 6. Isoconcentration contours (ppm) of compounds which elute prior to benzene in the shallow soil gas at Chevron Service Station 9-2582, Dublin, California, February 1988.

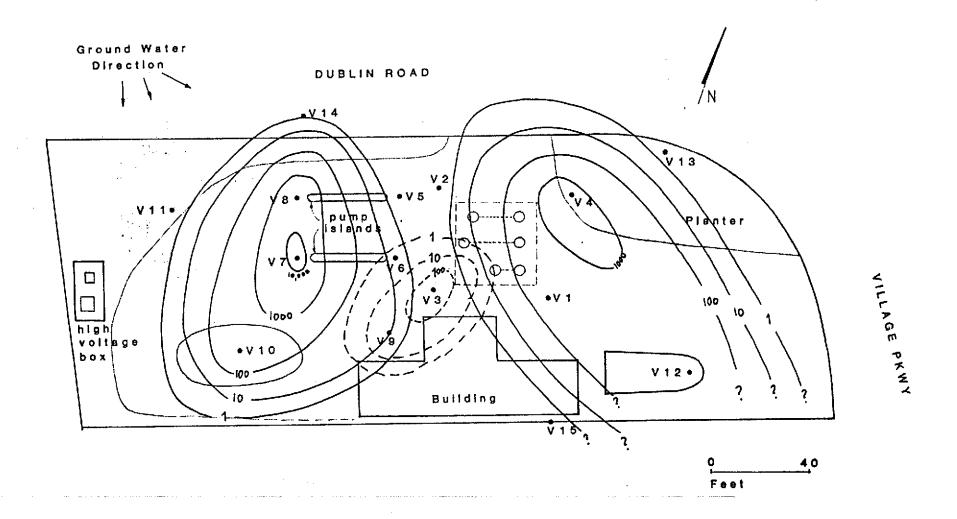


Figure 7. Isoconcentration contours (ppm) of total volatile hydrocarbons in the shallow soil gas at Chevron Service Station 9-2582, Dublin, California, February 1988.

Low levels of hydrocarbons were detected in the vicinity of the the northeastern planter and the tank field. The relative concentrations of the hydrocarbons suggest that these are the weathered residues of petroleum products rather than material of recent origin: there is little benzene and ethylbenzene relative to the concentrations of toluene and the small-molecular-weight compounds which elute prior to benzene. The extent of this contamination towards the southeast is unknown.

Finally, low-to-moderate concentrations of compounds which elute prior to benzene were found between the pump islands and the tank field at a relatively shallow depth (3-4 feet, see Figure 5). Neither benzene nor toluene was found at levels above the detection limit. It is unclear if this area has the same source as either of the other plumes, because V3 did not have any detected hydrocarbons at 13 feet (the depth at which hydrocarbons were found in the area of the tank field) and because V9, V6, and V5 (the locations between V3 and V7) did not have any detectable hydrocarbons.

2.3 DISCUSSION

The SVCA detected two (possibly three) areas of soil gas hydrocarbons at the Dublin site. Based on their chromatographic responses, most of the detected hydrocarbons in the eastern area of the site are low-boiling-point, low-molecular-weight compounds, probably alkanes such as pentane. Alkanes have a very low level of human toxicity (Sandmeyer 1981).

The highest detected concentrations of benzene (150 ppm) and toluene (200 ppm) were found at point V7 (see Table 1). These values can be used with Henry's Law to make order-of-magnitude estimates of potential ground-water concentrations. Inherent in this estimate is the assumption that product has reached ground water, which has not been confirmed for this site. A Henry's Law constant (H) is the ratio of a chemical's concentration in air to

its concentration in water at equilibrium (see Appendix A). It can be estimated by

$$H = C_{SV}/C_{W}$$

where

H = Henry's Law constant, atm'L/mole

C_{sv} = vapor concentration, atm

Cw = water concentration, mole/L.

For estimating ground-water concentrations, this equation can be rearranged as:

$$C_w = C_{sv}/H$$
.

To make the units compatible, the following conversions are used:

1 ppm soil vapor = 1x10⁻⁶ atmosphere; 1 mole = mole weight
expressed in milligrams. The molecular weights of benzene and
toluene are 78x10³ mg/mole and 92x10³ mg/mole, respectively (EPA
1986). Their Henry's Law constants are 5.6 atm'L/mole and 6.4
atm'L/mole, respectively (EPA 1986). Using these values, the
expected ground-water concentration of benzene can be estimated
as

For toluene, the ground-water concentration can be estimated by

On the basis of these equations, the order-of-magnitude estimates of dissolved benzene and toluene concentrations at the Dublin site would be 2 mg/L and 3 mg/L, respectively.

These levels can be put into perspective by comparing them to ground-water saturation values. A study sponsored by the San Francisco Bay Regional Water Quality Control Board (1985) found that gasoline-saturated ground water contained as high as 40 mg/L benzene and between 9 and 76 mg/L toluene. The estimated levels are an order of magnitude below those associated with water saturation.

3. CONCLUSIONS

The SVCA found two areas of hydrocarbon soil gas contaminination. One area seems to be associated with the tank field, although the highest measured levels of hydrocarbons were found in a planter northeast of the tank field. The levels of benzene and toluene within this plume are low (less than 10 ppm) and most of the detected hydrocarbons are suspected to be nontoxic alkanes. This suggests that the source of these hydrocarbons may be weathered gasoline.

A more significant area of contamination was detected in the vicinity of the pump islands. The relative concentrations of the hydrocarbons suggests that the source is fresher than the material discussed above. The benzene and toluene plumes in this area do not extend beyond the site boundaries. The SVCA results suggest that the levels of dissolved benzene and toluene in ground water under the site could be as high as 2 mg/L and 3 mg/L, respectively.

4. REFERENCES

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- SFBRWQCB (San Francisco Bay Regional Water Quality Control Board). 1985. Guidelines for Addressing Fuel Leaks. Available from SFBRWQCB, Oakland, California.

Appendix A

Principles and Application of Soil Vapor Contaminant Assessment

APPENDIX A: PRINCIPLES AND APPLICATION OF SOIL VAPOR CONTAMINANT ASSESSMENT

The soil vapor survey, or SVCA, technique takes advantage of the behavior of hydrocarbon mixtures and the physicochemical properties of the individual components in the subsurface. Following a subsurface gasoline release, free product will migrate downwards towards the ground water, some of the gasoline will volatilize, and some will adsorb to the soils. In the case of a spill of sufficient volume to exceed the soil binding capacity, free liquid will reach ground water, at which point it will float and may begin to vaporize and solubilize.

Like most hydrocarbon liquids, gasoline is a complex mixture of many compounds, each with its own physicochemical properties. The contaminants found in ground water located beneath a layer of floating hydrocarbon are generally less hydrophobic and are generally found in concentrations proportional to the hydrocarbon/water partition coefficient (i.e., the relative solubility of a given compound in the bulk hydrocarbon to its solubility in water) and to their percent composition in the gasoline. It may be noted that concentration of total benzene, toluene, and xylenes in product-saturated water may exceed 10-20 mg/L (API 1985a).

Hydrocarbons will also volatilize into the air- or gas-filled soil interstices. Volatilization is largely a function of vapor pressure. The natures of the contaminant mixtures, in terms of specific component mixtures, in either the aqueous or vapor phase, are distinctly different from each other and from the gasoline. That is, the more hydrophilic hydrocarbons will be more likely to move into ground water, while the more volatile compounds are more likely to move into the vapor phase, and the compounds that are both less volatile and more hydrophobic are more likely to remain in the free product or be adsorbed to soils (Hinchee and Reisinger 1987).

Hydrocarbons not remaining in the free product will partition into either ground water or soil vapor and migrate as the result of a variety of interacting forces. In ground water, contaminants will migrate with the ground-water flow, interacting with the rock or soil geological medium. As the contaminants pass through a medium, organic constituents in the medium interact with the contaminants, and some are adsorbed or bound to particle surfaces (Bruell and Hoag 1986). The result is a net retardation in the velocity of movement of those compounds relative to that of the ground water in which they are dissolved. The process is analogous to laboratory chromatography. The compound with the least affinity for the porous medium is least retarded and therefore moves most rapidly. This compound, then, is present at the leading edge of a contaminant plume.

The affinity of a compound for the soil porous medium is partly a function of the compound's hydrophobicity—that is, the more hydrophobic a compound the more likely it is to adsorb to the solid medium. Aqueous solubility is a good indicator of hydrophobicity: the more soluble a compound is, the less hydrophobic and more hydrophilic it is, and vice versa. Vapor pressure is a good indicator of volatility; compounds with higher vapor pressures are more volatile.

In determining the environmental fate of various hydrocarbon compounds in a hydrocarbon mixture such as gasoline, those which have a high vapor pressure are more likely to move into the vapor phase, or evaporate. Compounds with high solubility are more likely to move into ground water from the free product and, once in ground water, tend to move more rapidly. Compounds of low vapor pressure and low solubility tend to remain in the free product or be adsorbed to the solid matrix and remain relatively immobile.

Dissolved compounds will tend to volatilize from the aqueous phase. The Henry's Law constant is the equilibrium ratio of a

compound's concentration in the vapor phase to its concentration in the aqueous phase. The higher a compound's Henry's Law constant, the greater its tendency to volatilize from water into air.

Figure A-1 graphically illustrates the vapor pressure, aqueous solubility, and Henry's Law constants, and their relationships, for selected hydrocarbons typically found in gasoline. The Henry's Law constant is approximated here as the ratio of vapor pressure to solubility.

The Henry's Law constant is directly related to the tendency of compounds to volatilize, as opposed to solubilizing. Compounds with Henry's Law constants greater than 0.001 (atm m³/mole) volatilize from water into air very rapidly (Lyman et al. 1982); those with Henry's Law constants greater than 0.01 (atm m³/mole) are generally volatilized so rapidly that they are seldom found in gasoline-contaminated ground water. It may be observed (Figure A-1) that tetraethyl lead (TEL) has an extremely low solubility and a relatively low vapor pressure. As a result, this constituent would not be expected to solubilize and migrate in ground water, and although its low vapor pressure would indicate slow volatilization, its Henry's Law constant indicates that it may be more rapidly volatilized than solubilized. The fate of TEL would be expected to be long-term binding to the soil.

On the basis of these properties it can be seen that associated with any ground water, soil, or free-product contamination is vapor phase contamination. The SVCA technique takes advantage of this, and through the collection and analysis of soil vapor permits a rapid, cost-effective delineation of the extent of contamination.

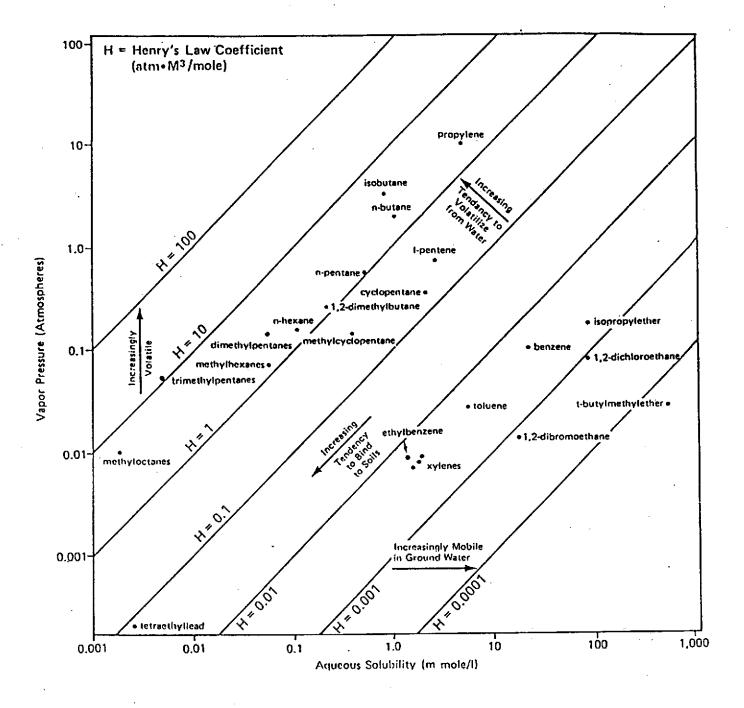


Figure A-1. Vapor pressures, solubilities, and corresponding Henry's Law constants for major constituents of gasoline.

Appendix B SVCA Data Sheets and Chromatograms



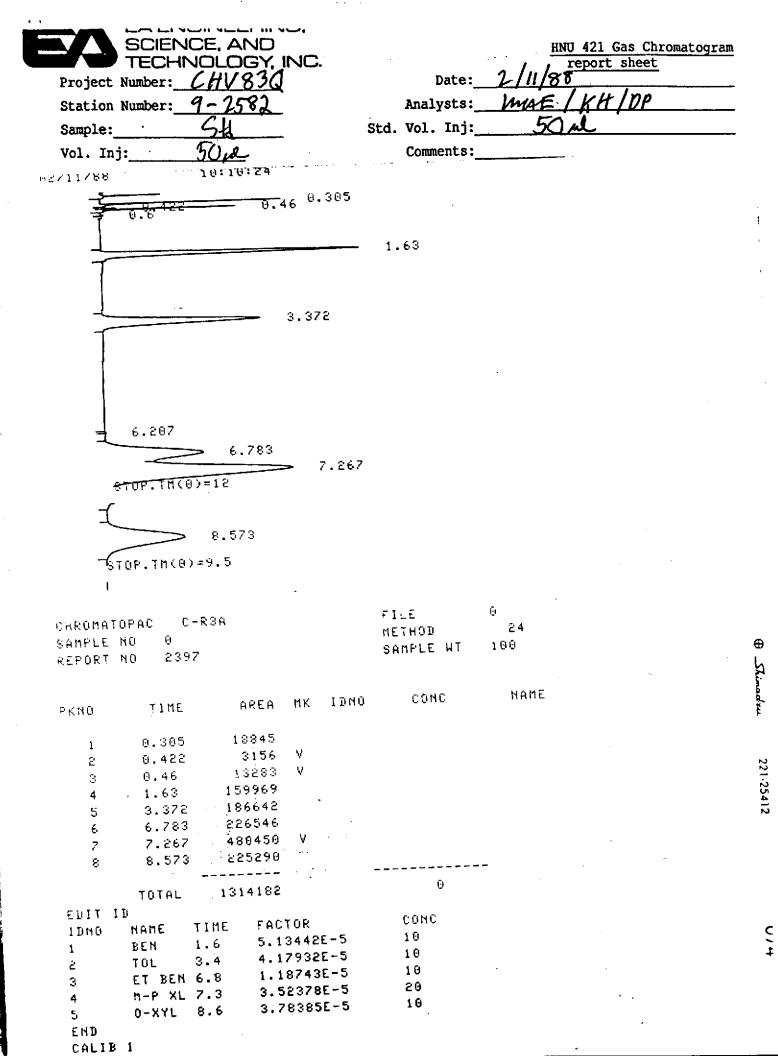
SVCA DATA SHEET

					,	HNU 421 Gas Chromatograph
Project No	mber:_	C4V8	3Q		Dat	te: 2/11/66
Station No	mber:	9-25	82		Analyst	The state of the s
•				ii.: A	_	
Site Locat		40			lud at	Glinge Parkway Dublin CA
Sample Location	Time	Depth	Purge Time (min)	Vacuum (in Hg)	Vacuum Release (min)	New RSLICO Copyllogy Columnic
5+1	1010	~		_		Rasslue
VIA	1023	3	10'	21	d	We Glor 10ml/mm EB from
VIB	1040	5.5		27		
·VZA	1055	3		7	0	(#3. % <u>2.3.</u>)
VIC	1110	8		21	Ø.5	
V28	1121	10.5		25		
VID	1136	10.5		27	.25	
VIC	1150	15.5		2.5	.25	
VIE	1204	13.0		25	,5	
V3A	1219	3		17	0	Odor 144 = 10 202
Blunk	1232			_		- Mtu=6 1000
ronge toot	1238			Ø	ō	No Olor.
<u> </u>	1246	3		25	.75	Nour lines]
<u> </u>	1255	3		28	25.	Sungled of 5" Vuc. 50,
V44	1308	3		28	60.	Sumpled it 27 Visc 500
<u> </u>	1321	13		28	.25	100
549	337		-			50.
	1348	3	10	27	1	160,
V4B	1359	15.5		28	10	S. Odor 50w
<u> </u>	1414	4.		27	1	No Glav- 100 m
V7B	1425	5.5		27	0	Odor 2011
V8A	1437	3		27	Ø	No Gor. 160M
V12	1501	13		25		Solil Kipple
Exstern block	a		_		_	
V13	1535	15.5	10	27	3	No Olor
V12	1552	_13		17	0.	5. Olor.
1/4 B	11.19	5.5		10		



SVCA DATA SHEET

	*					HNU 421 Gas Chro	xmatograpi
Project N	umber:_	CHV.	834		Dat	e:2/11/88	
Station N	umber:_	9-2	582		Analyst	s: MAE/KH/DP	
Site Loca	tion:_7	440	Dublin	Blud	(nt Villa	ige Karkway) Dubling Co	4
Sample Location	Time	Depth	Purge Time (min)	Vacuum (in Hg)	Vacuum Release (min)	Comments	
VIO	17/2	6.5	10	17	d	he Odor.	
V14	17-28	6.5		17	0	5. Odor.	140.1
V15	1749			27	2	No Odor.	
Chausen Regular	1803		V _			7 1 15	<u> </u>
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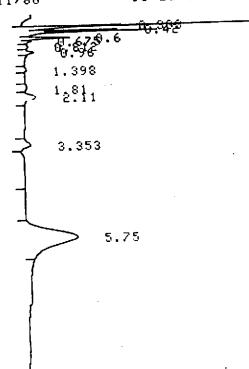
Station Number: 9-258

Sample:_

DON Vol. Inj:

START 02/11/88

10:23:54



C-R3A CHROMATOPAC 0 SAMPLE NO 2399 REPORT NO

FILE METHOD 100 SAMPLE NT

Date: Analysts:___

Std. Vol. Inj:

Comments:

PKNO	TIME	AREA	nК	IDMO	CONC	NAME
i	0.303	46942				
5	0.42	41824	A			
3	0.6	16671	٧			
4	0.675	3202	٧	•		
5	0.812	2084	, V			
6	0.96	3740	٧	• .		
7	1.398	1488	**			
8	1.81	1661		-		
9	2.11	13069			5 500	c T01
10	3.353	9940		2	8.532	6 TOL
11	5.75	159692				·
	-				0.532	6
	TOTAL	300312			8.336	. •

VOLUME INJECTED (UL) 100 DILUTION

HNU 421 Gas Chromatogram

report sheet



Project Number: CHV830

Station Number: 9-2582

Sample:___

Vol. Inj:

HNU 421 Gas Chromatogram report sheet Analysts: MAE Std. Vol. Inj:____

ALANGER - - - - - -START

02/11/88

18:49:40

593 Ma62, 303 .3.358 7.532

C-R3A CHROMATOPAC SAMPLE NO 2400 REPORT NO

FILE METHOD 24 100 SAMPLE NT

Comments:

PKNO	TIME	AREA	ńК	IDNO	CONC	NAME
i	0.303	27298				
2	0.42	7477	٧			
3	0.462	10767	٧			
4	0.593	3484		•		•
5	3.358	8714		5	0.4669	TOL
	TOTAL	57741	*** *		0.4669	

RUN

VOLUME INJECTED (UL)

? 100

DILUTION

? 1

BEN PBB ET BEN M,P-XYL -1.59321

ERROR 16: UNDEF'D STATEMENT IN

0-XYL TOL TVH IONG 0.23345 0.0389251

410

-1.32683

Sample: V2A

Vol. Inj: 100 W

HNU 421 Gas Chromatogram

Date: 2/11/8 report sheet

Analysts: WAE KH DP

Std. Vol. Inj: 50 sd

Comments:

START 02/11/88

10:55:34

0.595 0.858 0.858

- 7.684787

CHROMATOPAC C-R3A SAMPLE NO 0 REPORT NO 2402 FILE 0
METHOD 24
SAMPLE HT 100

NAME IDNO CONC AREA MΚ TIME PKNO 0.342 44261 0.43 29020 2 .8244 0.595 3 4448 7.787 0 85973 TOTAL

LIST

BASIC PROGRAM

1 B=1.05
2 PB=57000
3 EB=0
5 X2=0
7 X1=0
8 NI=0
10 T=0

077

HNU 421 Gas Chromatogram report sheet

Date:

Analysts:

Std. Vol. Inj:_

Comments:

Sample: Vol. Inj:_

Station Number: 9 - 2582

MERKOKA TOTOHNEL N STATEMENT

START

02/11/88 \ 11:10:19

		69639 0
1.165 1.405 1.64812 2.107	0.967	
3.333		_
9.032		

CHROMATOPAC C-R3A SAMPLE NO REPORT NO 2493

FILE 24 METHOD 100 SAMPLE NT

PKN0	TIME	AREA	МK	ONGI	CONC	NAME
1	0.317	55333 7				
2	0.372	54523	٧			
3	0.425	406697	٧			
4	. 0.465	495028	٧			
5	0.608	783343	٧			
6	0.678	156036	¥			
7	0.743	27094	V	•		
8	0.815	18041	٧			
9	0.967	169151	•			
19	1.057	47278	٧			
11	1.165	5590				
12	1.405	28832				
13	1.812	16456				
14	2.107	17493				
15	3.333	2062		2	0.110	5 TOL
	TOTAL	2780959		-	0.116	- 5

Project Number: CHV830

Station Number: 9-2582

Sample:

Vol. Inj:

HNU 421 Gas Chromatogram

report sheet

Analysts: WAE KH DP

Std. Vol. Inj:

Comments:

410

ERROR 16:UNDEFOD STHIEMEN: 18

START

02/11/88

11:21:28

0.592 0.₩2315 3.362

CHROMATOPAC C-R3A Θ SAMPLE NO REPORT NO 2404

FILE METHOD SAMPLE WT

TIME AREA MK IDNO CONC NAME PKNO 26510 0.315 23238 0.42 2 4614 V .0.592 3 0.1997 TOL 3.362 . 3727 0.1997 58089

TOTAL

VOLUME INJECTED (UL) ? 199

DILUTION

? 1

RUN

PBB

H,P-XYL

BEN ET BEN

TOL

0.0998312

0-XYL

HYT

0.0340342

Project	Number:	CHV	830

Station Number: 9-2582

Sample: VID

Vol. Inj: 100 w

HNU 421 Gas Chromatogram

report sheet

Analysts: MAE / KH / DP

Std. Vol. Inj: 50 M

Date: 2

Comments:

START

02/11/88

11:36:22

1.008 1.008

CHROMATOPAC C-R3A SAMPLE NO 0 REPORT NO 2405 FILE 0
METHOD 24
SAMPLE WT 100

KN0	TIME	AREA	MK	OMGI	CONC	NAME
1	0.182	1817				
2	0.308	22378				
3	0.427	6608	٧			
4	0.478	12829	۸.			
5	0.6	4340	٧			
	_			·		
	TOTAL	47973	·	•	0	

RUH

VOLUME INJECTED (UL)

? 100

DILUTION

? 1

PBB B M.P-XYL E

BEN ET BEN TOL PNOI

0 2.98023E-8 O-XYL TVH

-0.282154

4.10.

-Shimadeu

Project Number: CHV830

Station Number: 9-258

Sample:

Vol. Inj: 100 pg 4 kerkuk* 16: Unuer u Simienen

START

02/11/88

11:50:28

= 0.4793133.39

CHROMATOPAC C-R3A SAMPLE NO REPORT NO 2406

FILE METHOD SAMPLE WT 100

Date: 2

Std. Vol. Inj:

Comments:

Analysts: MAE

PKNO	TIME	AREA	MK	IDHO	CONC	NAME
1	0.313	27865				
2	0.43	18974	٧			
3	0.473	29170	٧			
4	0.61	10416	٧.			
5	0.675	2813	٧			
6	0.752	1067	٧	,		
7	0.973	1008				
8	3.39	6474		. 2	0.346	9 TOL
	-		,	- -		-
	TOTAL	97725			0.346	9

RUN VOLUME INJECTED (UL) ? 160 DILUTION

? 1

TOL PNOI 0-XYL HYT

HNU 421 Gas Chromatogram

report sheet

Project Number: <u>CHV83Q</u>

Station Number: 9-2582

VIE Sample:_ Vol. Inj:_ 100m

Std. Vol. Inj:___ Comments:

MERRORM 16: UNDER'D SIMILULUI START

12:04:24

02/11/88 1.42 1.655 2.125 8:383 3.388

Analysts:____

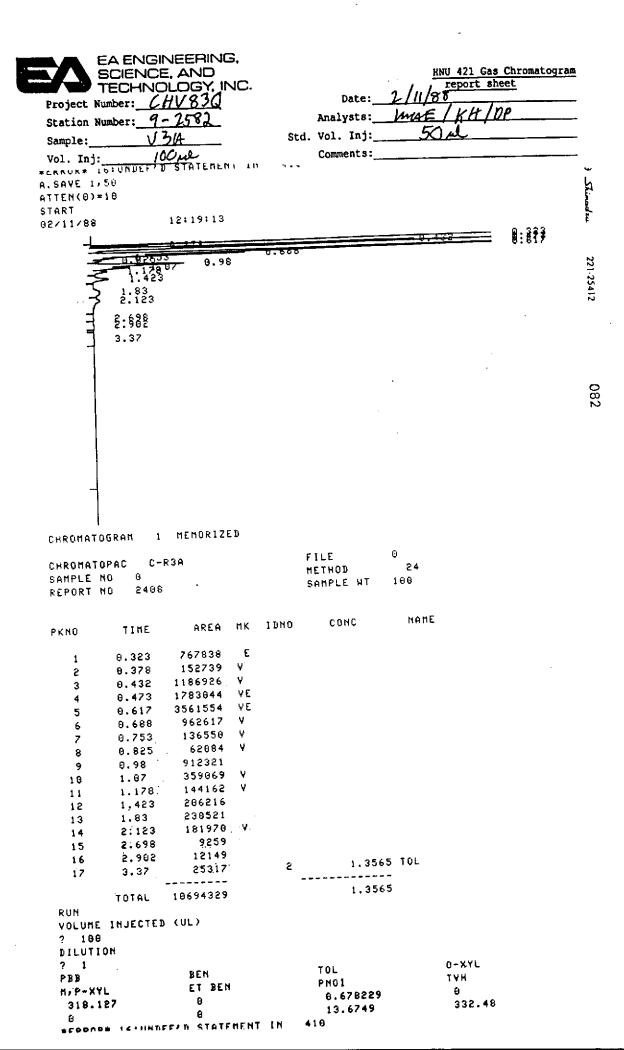
HNU 421 Gas Chromatogram

report sheet

CHROMATOPAC C-R3A SAMPLE NO 0 REPORT NO 2497

FILE METHOD SAMPLE WT 100

PKNO	TIME	AREA	ΜK	IBHO	CONC	NAME
			_			
ı	6.323	1499881	E			
2	0.377	2088 62	٧			
3	0.43	1405522	٧			
4	0.472	2163615	YΈ			
5	0.615	3114170	٧E			
6	0.685	790326	٧			
7	0.752	192337	٧			
8	0.822	67137	٧			
9	0.977	<i>77</i> 7400				
1 0	1.067	287368	٧			
1 i	1.175	9 3807	٧			
12	1.42	165857				
13	1.655	39173		1	2.4488	BEN
14	1.825	89764	٧			
15	2.125	72173				
16	2.703	7858				
17	2,912	6603				
18	3.388	25367	-	2	1.3591	TOL
				-		•
	TOTAL	11007214			3 .80 79	•
RUN		•				
UOL LIME	INJECTED	(UL)				



Project Number: CHV830

Station Number: 9-2582

Sample:

Vol. Inj: 100 Le

HNU 421 Gas Chromatogram report sheet

Analysts: MAE

Std. Vol. Inj:

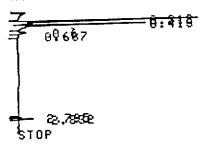
Comments:

Date:

START

02/11/88 12:32:38

ATTEN(0)=6



CHROMATOGRAM 2 MEMORIZED

CHROMATOPAC C-R3A SAMPLE NO REPORT NO 2409

FILE METHOD SAMPLE WT 100

PKN0	TIME	AREA	МK	IDHO	CONC	NAME
1 2 3	0.312 0.413 0.6	28626 18001 2981	V V			
	_					-
	TOTAL	49608			9	
RUN						

VOLUME INJECTED (UL)

? 100

DILUTION

? 1 PBB

BEN ET BEN M,P-XYL -0.231842

0-XYL TOL TVH PNOI

-0.231042



Project Number: CHV830

Station Number: 9 - 2583

Sample:

Vol. Inj:

Std. Vol. Inj: Comments:

ERROR 16:UNDER'D STRIEDERT IN

START

02/11/88

12:38:00

9.605.60423 0.525 1.415 3,407 \$TOP

CHROMATOGRAM 3 MEMORIZED

CHROMATOPAC C-R3A SAMPLE NO 9 REPORT NO 2410

FILE 24 METHOD SAMPLE WT

Date:

Analysts: _______

CONC NAME TIME AREA MK IDNO PKNO 0.307 30721 1 19817 2 0.423 3 0.468 6470 17797 0.525 5 0.603 21677 0.678 7501 7 3085 0.972 8 1.415 1750 3.407 8.2405 TOL

0.2405 112415 TOTAL

RUN

VOLUME INJECTED (UL).

? 100 DILUTION

2 1 PBB

BEN M,P-XYL ET BEN θ 1.59179

TOL PNOI

0.128227 0.0200462 0-XYL TVH

1.73206

HNU 421 Gas Chromatogram

report sheet

Project Number: CHV83Q

Station Number: 9-258

Sample: Vol. Inj:

START

82/11/88

12:46:47

report sheet Date: Analysts: MAE Std. Vol. Inj:__

HNU 421 Gas Chromatogram

Comments:

6.427 1.417 3.415 5.822 STOP

CHROMATOGRAM 4 MEMORIZED

CHROMATOPAC C-R3A Ü SAMPLE NO REPORT NO 2411

FILE 24 METHOD SAMPLE WT 100

NAME IDHO CONC AREA MK TIME PKNÖ 1669 0.273 1 31147 0.323 2 19680 0.427 3 19111 0.47 17841 . 0.605 5 0.685 . 5093 6 2370 0.975 7 0.4135 TOL 7718 8 3.415 0.4135 184629 TOTAL

RUN VOLUME INJECTED (UL) ? 108 DILUTION

BEN ET BEN TOL PNOI 0.206751 D-XYL TVH 9 1.4887

0.0344729

ET BEN

H,P-XYL

TVH

2.74423 XV

THE RESERVE OF THE PARTY OF THE

2.68898

8.435816 X

3.43 6.193

CHROMATOGRAM 6 MEMORIZED

CHROMATOPAC C-R3A FILE 0
SAMPLE NO 0 METHOD 24
REPORT NO 2413 SAMPLE WT 100

PKNO TIME AREA MK IDNO CONC NAME €.325 18777 0.43 2 48038 3 0.605 24534 SY: 4 0.892 1634 5 3.43 0.6009 TOL 0.6009 TOTAL 104197

RUN VOLUME INJECTED (UL) ? 50 DILUTION

PBB

BEH

TOL

D-XYI

086

			Ü
Project	Number:_	CHV83Q	

Station Number: 9-2582

Sample:

Vol. Inj:

START

02/11/88

13:21:40

0.692615 0.983 1.24 3.402 4.148

CHROMATOGRAM 7 MEMORIZED

C-R3A CHROMATOPAC SAMPLE NO Θ 2414 REPORT NO

FILE METHOD 24 SAMPLE WT 100

PKNO	TIME	AREA	пK	IDNO	CONC	NAME
1	0.328	29448				
2	9.432	40947	٧			
3	0.615	11745	٧.			
4	0.692	2596	٧			
5	0.983	1645	¥ .			
6	3.402	1932	-V	2	0.1035	TOL
	<u>-</u>		S			
	TOTAL	88312	•		0.1035	

VOLUME INJECTED (UL)

? 100

BILUTION

BEN ET BEN TOL PNOI

0.0517473 0.00862809 0-XYL TVH 8

6.978685

HNU 421 Gas Chromatogram

report sheet

Date: 2/11/88

Std. Vol. Inj: 50 M

Comments:

Analysts: WAE / KH / DP

Project Number: CHV83Q

Station Number: 9-25

Sample:

Vol. Inj:

HNU 421 Gas Chromatogram

report sheet

Date: Analysts: MAE

Std. Vol. Inj:

Comments:

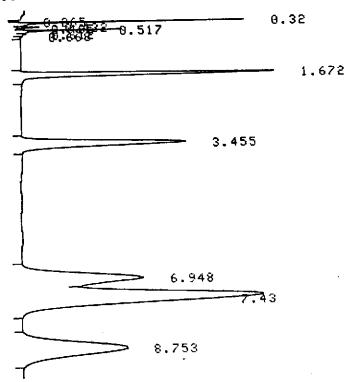
410

LKKOK 10:UNUEF U STATEMENT IN

START

88/11/88

13:37:58



8 MEMORIZED CHROMATOGRAM

CHROMATOPAC C-R3A SAMPLE NO 2415 REPORT NO

FILE METHOD SAMPLE WT 160

PKNO	TIME	AREA	мк	ONGI	CONC	NAME
1	0.32	23331	٧			
2	0.432	4897	٧.			
3	0.465	1950	٧			
4	0.517	14363	¥			
5	0.612	2475	w.Y			
6	1.672	168664		1	10.0434	
7	3,455	199145	•	2	10.6699	TOL
8	6.948	285593		3	12.6064	
9	7.43	622332	٧	4	25.9062	
10	8.753	310492	·	. 5	13.7819	0-XYL
	TOTAL	1624444			73.0078	

	Date:_	2/11/88
	Analysts:	MUSE / KH/DP
Std.	Vol. Inj:_	50 pl

/ report sheet

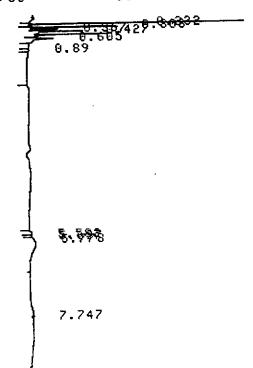
HNU 421 Gas Chromatogram

Comments:

02/11/88

Vol. Inj:_

13:48:87



CHROMATOGRAM 9 MEMORIZED

CHROMATOPAC C-R3A SAMPLE NO 0 REPORT NO 2416 FILE 0
METHOD 24
SAMPLE WT 100

KHO	TIME	AREA	MK	ONGI	CONC	NAME
1	0.232	12164				
2	0.308	18050				
3	0.367	6199	V			
4	0.427	23846	¥			
5	0.605	9462	, V			
				. - -		-
	TOTAL	69631	-4		Θ	

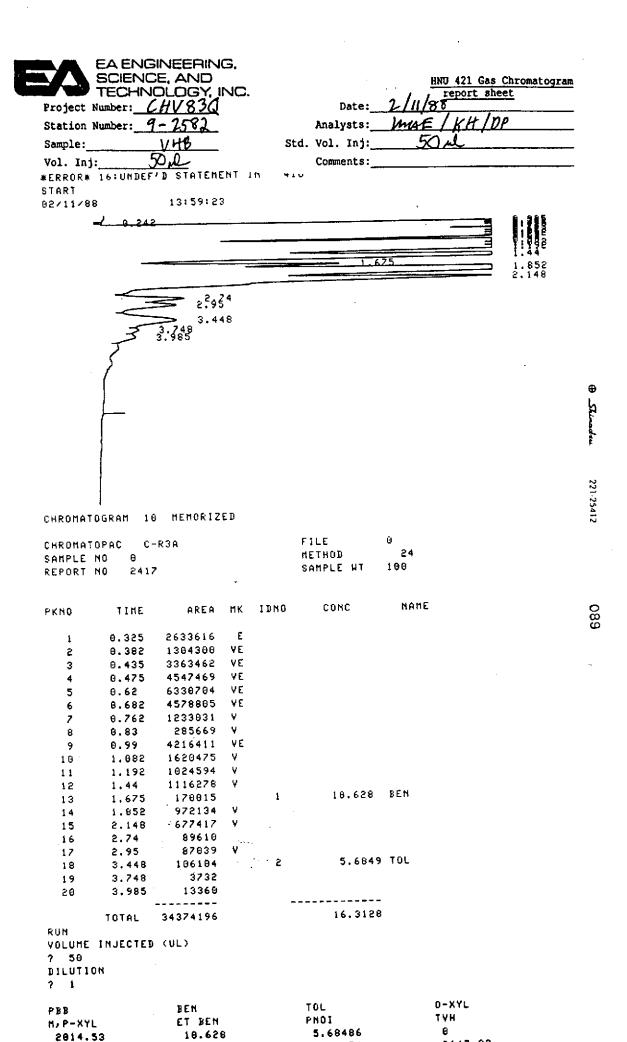
RUN VOLUME INJECTED (UL) ? 100

DILUTION ? 1

PBB BEN
M,P-XYL ET BEN
0.394794 0

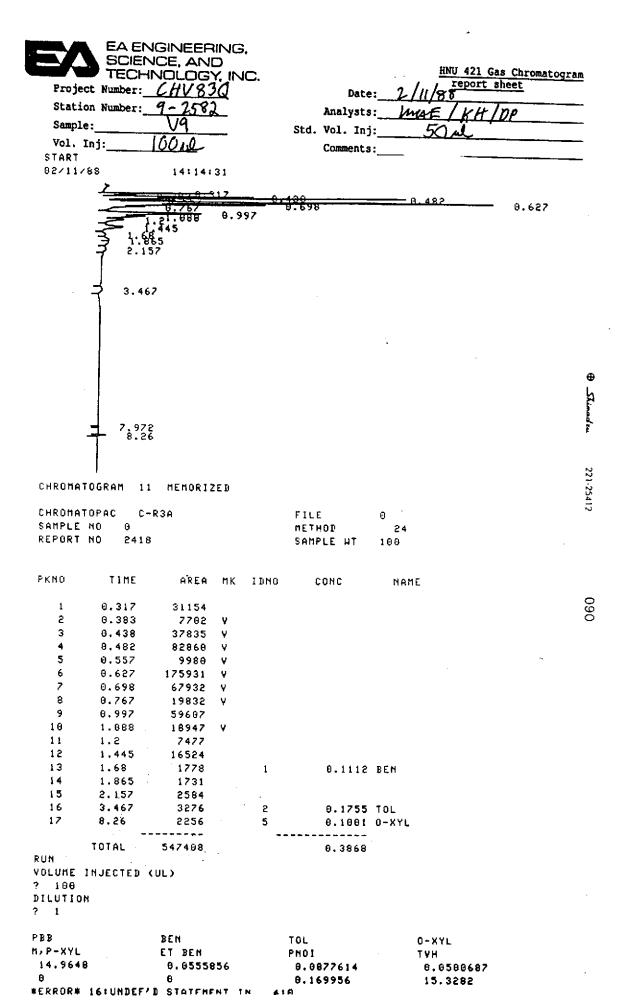
TOL 0-XYL
PNOI TVH
0 0
-2.98023E-8 0.394794

980

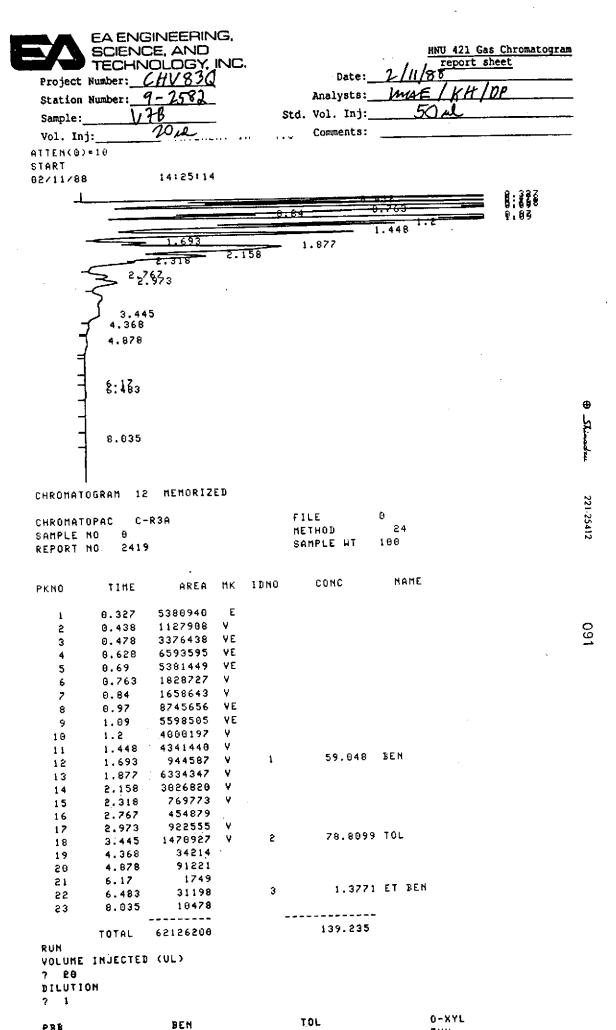


2147.62

1 5 6 1 2



1 42 0 3



PHOI

TVH

PBB

	Project Num Station Num Sample: Vol. Inj: TEN(0)=6 ART	A ENGINEER CIENCE, ANI ECHNOLOG ber: 64V83 ber: 9-2583 V8A	D Y. INC.	Std. Vol.	ysts: Mas	HNU 421 Gas Chroreport sheet	matogram
02	/11/88	14:37:4	8				
		ც გან					
		9.8355 0.59 1.1493988 0.99	12	Э	0.623		
	₹	1.443	, e			0.48	5
	7	2.157					
	}						
-]						
	1						
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	- 1						⊕
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							سيامهمناكمه
	1	•					i.
	1						
CHROM	ATOGRAM	13 MEMORIZET					22
		13 MEMORIZED					221-25412
SAMPLE		-R3A		FILE			5412
REPORT	: NO 6 ! NO 242	- ii	· ·	1ETHOD	0		
		•	ş	AMPLE WT	24 . 188		
PKNO	7140						
	TIME	. AREA MK	IDHO	CONC	N A 14 =		
1	0.325	449861		· · -	NAME		ο.
3 2	0.38	12341 v)92
4	0.433 0.485	56874 v					10
5	0.623	129433 V 98564 V					
6 7	0.697	33968 V				-	
8	0.76 0.835	8814 V					
9	0.993	4717 y 45878					
10	1.088	16831 y					
1 E	1.193	5627					
13	1.443 1.688	14543					
14	1.882	2329 14953	1	0.1456 B	FN		
15	2.157	4450					
	TOTAL			-			
KUN		493502		0.1456			
, 100 AOTAWE 14	IJECTED (U	L)		- -			•
DILUTION							·
? 1		-					
PBB							
H, P-XYL		BEN	TOL				
12.9642	E	T BEN A. AZZZOGO	PHOI		0-XYL		
Θ		0.0727998 0	θ		T V H 9		
==#### 16	INDEE'D	0 Statenfnt in	0.60	6441	13.6434		