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Air



Hazardous Waste Treatment, Storage, and Disposal Facilities (TSDF) — Air Emission Models

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Hazardous Waste Treatment, Storage and Disposal Facilities (TSDF) — Air Emission Models

Emission Standards Division

**U.S. ENVIRONMENTAL PROTECTION AGENCY
Office of Air and Radiation
Office of Air Quality Planning and Standards
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1.0 INTRODUCTION

1.1 BACKGROUND

This report was prepared for the Environmental Protection Agency's (EPA's) Office of Air Quality Planning and Standards (OAQPS) as part of the effort to develop air emission models for hazardous waste treatment, storage, and disposal facilities (TSDF). Basic to this effort is the determination of the means by which volatile organics (VO) escape to the environment from TSDF.

VO in surface impoundments, land treatment facilities, landfills, wastepiles, or wastewater treatment (WWT) plant effluents can depart through a variety of pathways, including volatilization, biological decomposition, adsorption, photochemical reaction, and hydrolysis. To allow reasonable estimates of VO disappearance, one must know which pathways predominate for a given chemical, type of waste site, and set of meteorological conditions.

Analytical models have been developed to estimate emissions of VO via various pathways from area emission sources at hazardous waste sites. Some of these models have been assembled into a spreadsheet that is included in this report as a floppy diskette for use on an IBM PC, or compatible, microcomputer. A user's guide for these models is included in the report as Appendix A. Area emission sources for which models are included on the diskette are as follows:

- Nonaerated impoundments, which include quiescent surface impoundments and open top WWT tanks
- Aerated impoundments, which include aerated surface impoundments and aerated WWT tanks
- Disposal impoundments, which include nonaerated disposal impoundments
- Land treatment
- Landfills.

These models can be used to estimate the magnitude of site emissions for regulatory purposes. Sample calculations using each model also are included in this report.

1.2 SCOPE

This report briefly describes the chemical and physical pathways for VO and discusses their importance for different types of sites and sets of conditions. Models developed for estimating the relative magnitude of environmental release in the presence of competing pathways are presented, and physical characteristics of the parameters that serve as inputs to the models are identified.

The models provide an estimate of the relative magnitude of VO pathways on a compound-specific basis. Models for aerated and nonaerated impoundments, lagoons, landfills, wastepiles, and land treatment facilities have been installed in an integrated spreadsheet program, CHEMDAT7, which allows a user to calculate the partitioning of VO among various pathways depending on the particular parameters of the facility of interest. The program is structured to allow new data (e.g., compounds and model facility parameters) to be added (see Appendix A for user's guide). The results of the calculated partitioning may be used to identify those characteristics that are important in determining relative VO loss rates.

Source variability will significantly influence the relative importance of the pathways. For highly variable sources, it may be possible to exclude insignificantly small pathways from consideration. The relative magnitude of these pathways then can be compared by applying the methodology to a model facility to determine relative differences among various compounds.

1.3 REPORT ORGANIZATION

Chapter 2.0 describes each of the potential pathway mechanisms that determine the fate of various chemical species. Chapter 3.0 discusses the importance of the pathways for surface impoundments and aerated and non-aerated WWT facilities, land treatment sites, and landfills/wastepiles. Chapters 4.0, 5.0, and 6.0 describe the emission models applicable to these sites. Models for estimating emissions from transfer, storage, and handling operations are described in Chapter 7.0. Chapter 8.0 compares emission model predictions with the field data that are available.

This report compares relative rates of VO destruction and volatilization to determine the most significant pathways. The rate of VO volatilization destruction for any one pathway is calculated so that it can be expressed as a fraction of the loss/destruction from all pathways.

Appendix B contains supplementary material, and Appendix C presents a comprehensive source list that includes pertinent literature in addition to that cited in the sections and appendixes of this report.

Properties of compounds of interest to TSDF pathways and emission estimation are presented in Appendix D. A subset of these compounds is a part of CHEMDAT7. The user's guide, Appendix A, describes the procedures that are used in estimating emissions using CHEMDAT7 and other procedures presented in the body of the report. The user's guide also contains instructions for modifying CHEMDAT7 to include additional compounds using the compound characteristics presented in Appendix D.

2.0 DESCRIPTION OF PATHWAYS

2.1 GENERAL

A pathway is considered here to be any process that removes volatile organics from a site. The removal may be physical (as in volatilization of a solvent from a surface impoundment) or chemical (as in oxidation of an alcohol in a wastewater treatment plant).

Pathways may be considered as rate processes, with rate often strongly dependent on concentration of the disappearing species and temperature of the system. Rates vary in order from zero to mixed, with perhaps first order predominating, that is:

$$\text{rate} = - \frac{dc}{dt} = k_v c \quad (2-1)$$

where

c = concentration of disappearing substance, g/L

t = time, s

k_v = volatilization constant, s^{-1}

Half-life, the time required for one-half of the substance to disappear, is a useful concept. It provides an easily visualized measure of the time required for disappearance. For a first-order rate process:

$$t_{1/2} = (\ln 2) k_v^{-1} = 0.693 k_v^{-1} \quad (2-2)$$

where

$t_{1/2}$ = half-life, s.

The half-life of a second-order equation is as follows:

$$t_{1/2} = k_v'^{-1} c_o^{-1} \quad (2-3)$$

where

k_v' = second-order volatilization constant, L/(g•s)

C_0 = initial concentration, g/L.

Note that first-order half-lives are independent of initial concentration while higher order half-lives are not.

Much of the following material is taken from ICF.¹ The pathways described are physical (volatilization, adsorption, migration, and runoff) and chemical (biological decomposition, photochemical decomposition, hydrolysis, oxidation/reduction, and hydroxyl radical reaction).

2.2 VOLATILIZATION

Volatilization occurs when molecules of a dissolved substance escape to an adjacent gas phase. The driving force for this process in nonturbulent liquids is molecular diffusion. Equation (2-1) shows the rate of volatilization of an organic chemical from water. For this case, the rate constant can be estimated:²

$$k_v = \frac{1}{L} \left[\frac{1}{k_1^0 (D_1^c/D_1^0)^m} + \frac{RT}{(10^6) H k_g^w (D_g^c/D_g^w)^n} \right]^{-1} \quad (2-4)$$

where

L = mixing depth of water, cm

k_1^0 = mass transfer coefficient of oxygen in water, cm/s

D_1 = diffusion coefficient of the chemical (c) or oxygen (o) in water, cm²/s

m = liquid turbulence exponent, 0.5 to 1, dimensionless, from Table 2-1

R = ideal gas constant, atm cm³/(mol•K)

T = temperature, K

H = Henry's law constant, atm m³/mol

k_g^w = mass transfer coefficient for water vapor in air, cm/s

D_g = diffusion coefficient of the chemical (c) or water (w) in air, cm²/s

n = gas turbulence exponent, 0.5 to 1.0, dimensionless, from Table 2-1.

Equation (2-4) requires values of diffusion coefficients and Henry's law constants. If tabulated values are not available, the following estimations can be used. For the diffusion of a chemical in air:³

$$D_g = 0.0067T^{1.5} (0.034 + M^{-1})^{0.5} M^{-0.17} [(M/2.5d)^{0.33} + 1.81]^{-2} \quad (2-5)$$

where

M = molecular weight of chemical, g/g mol

d = density of liquid chemical, g/cm³.

For diffusion coefficients in water:

$$D_l = 1.518 (10^{-4}) V_{cm}^{-0.6} \quad (2-6)$$

where

V_{cm} = molar volume of chemical, cm³/g mol.

This equation assumes the system temperature to be 300 K. For other temperatures, a more rigorous form of the equation should be used, as in Perry.⁴ Molar volume is estimated as the ratio of molecular weight to liquid density at room temperature.

If ideal gases and solutions are assumed, Henry's law constant can be estimated from:

$$H = P/(14.7s) \quad (2-7)$$

where

P = pure component vapor pressure, psia

s = solubility of chemical in water, g mol/m³.

Values for other terms in Equation (2-4) have been tabulated by ICF and are given in Table 2-1.

In general, equations are available to estimate volatilization from wastewater treatment systems and surface impoundments.^{6,7} In the case of land treatment and landfills, the models for volatilization are much less well developed and the supporting data are more limited than those of the

TABLE 2-1. VALUES OF CONSTANTS FOR USE IN EQUATION (2-4)⁵

Constant	Value	
	Rivers	Lakes
L (cm)	200	200
k_1^0 (cm•s ⁻¹)	0.0022	0.0005
m	0.7	1.0
T (K)	293	293
RT (m ³ •atm•mol ⁻¹)	2.40×10^{-2}	2.40×10^{-2}
k_g^w (cm•s ⁻¹)	0.58	0.58
n	0.7	0.7

aqueous systems. The rate of volatilization at a soil-air interface is a function of the concentration and properties of the escaping chemical, soil properties (moisture, temperature, clay, and organic content), and properties of the air at soil level (temperature, relative humidity, and wind speed).⁸

2.3 ADSORPTION

Adsorption takes place when molecules of a dissolved chemical (in a liquid-solid system) become physically attached to elements of the solid phase. Chemical bonding may also occur (chemisorption). An example of adsorption is molecules of solvent being sorbed by particles of silt in a surface impoundment. If the adsorptive capacity of the solid material is reached, no further net sorption will occur. With reductions in concentration in the bulk liquid of the chemical being sorbed (adsorbate), desorption may take place. The amount of material adsorbed depends on (1) the concentration of adsorbate, (2) the amount of solid phase (adsorbent), and (3) the temperature. For systems with constant adsorbent properties, primarily surface area per unit mass, the amount of material adsorbed at a particular concentration and temperature is proportional to the mass of adsorbent. For example, the Freundlich adsorption isotherm equation allows prediction of amount adsorbed as follows:

$$\frac{X}{m} = K_f C^{1/n} \quad (2-8)$$

where

X = mass of chemical adsorbed, g

m = mass of adsorbent, g

K_f = Freundlich adsorption coefficient, (g sorbate/g sorbent)/
(g sorbate/g solution)

C' = concentration of chemical in solution at equilibrium, g
sorbate/g solution

n = empirical constant, ranging from 0.7 to 1.1, typically 1.0 for
soils, dimensionless.

A Langmuir adsorption isotherm can be derived from a kinetic rate theory describing the adsorption and desorption rates. The rate of adsorption is proportional to the rate of collisions between adsorbate molecules

and free adsorbent surface. The rate decreases with lowering adsorbate concentration and with decreasing surface sites available for adsorbing molecules. The following rate equation applies:

$$\text{Rate of adsorption} = k_1 C' (1-f) \quad (2-9)$$

where

k_1 = rate constant for adsorption, g/s

f = fraction of adsorption sites occupied, dimensionless.

For desorption:

$$\text{Rate of desorption} = k_2 f \quad (2-10)$$

where

k_2 = rate constant for desorption, g/s.

At equilibrium the two rates are equal, and

$$f = \frac{k_1 C'}{k_1 C' + k_2} \quad (2-11)$$

Adsorption rates are usually rapid compared to the other processes discussed here. However, mass transfer limitations may reduce effective rates, especially for poorly mixed systems. Lack of sorbent and its saturation may also reduce the effectiveness of adsorption.

For estimating adsorption partitioning, a linear relationship is assumed ($n = 1$ or $k_1 C \ll k_2$). The equilibrium relationship for biomass is estimated from an equation of Matter-Muller,⁹ based on the logarithm of the octanol-water partition coefficient, LOW. For land treatment and landfills, the only partitioning of importance to fate predictions is gas-liquid partitioning.

2.4 MIGRATION

Migration occurs when chemicals applied to soils are transported through the soils to groundwater. Leaching and percolation are the mechanisms that physically remove chemical molecules from a point of deposit and

carry them toward a water table. Capillary flow is a resisting mechanism that moves the molecules upward through the soil. The leachability of a chemical is a function of soil texture and cation exchange capacity, amount of soil organic content, amount and intensity of rainfall, and mechanical placement and adsorptive properties of the chemical.¹⁰

2.5 RUNOFF

Chemicals at or near the soil may be washed away by rain. The rate depends on soil and chemical characteristics and on rainfall rates and frequency. Clark, Viessman, and Hammer¹¹ state that runoff in any drainage area is a function of climate and the physical characteristics of the area. Significant factors include precipitation type; rainfall intensity, duration, and distribution; storm direction; antecedent precipitation; initial soil moisture conditions; soil type; evaporation; transpiration; and, for a given drainage area, its size, shape, slope, elevation, directional orientation, and land use characteristics. If rainfall is heavy shortly after application of a chemical, runoff and erosion can physically remove it. The chemical may be dissolved in runoff water, carried along by it, or adsorbed on eroding soil particles that move with runoff. For pesticide applications, about 3 to 10 percent of the applied material appears in runoff water. Below a certain intensity, rainfall will promote leaching of nonadsorbed chemical into the ground rather than result in runoff.

2.6 BIOLOGICAL DECOMPOSITION

Biological decomposition takes place when microbes break down organic compounds for metabolic processes. The rate of decomposition depends on the structure of the compound and on the needs of the microbes. If the compound is present in excess, the rate of population increase is as follows:

$$dx/dt = Rx \quad (2-12)$$

where

x = concentration of biomass, g/L

R = specific growth rate coefficient, s^{-1} .

If the compound is present in limited amount, the rate becomes a hyperbolic saturation function of the compound (substrate) concentration:¹²

$$\frac{dx/dt}{x} = R_{\max} S / (K_s + S) \quad (2-13)$$

where

R_{\max} = maximum specific growth rate coefficient (where substrate is in excess), s^{-1}

S = concentration of substrate, g/L

K_s = substrate concentration at which the rate of substrate utilization is one-half of the maximum rate, g/L.

Because the microbial population increases at the expense of the compound, the growth rate is proportional to the compound's rate of disappearance. The rate process may be of zero, first, or mixed order depending on concentration of the substrate. In the presence of multiple substrates, kinetics become complex.

For the case of S much greater than K_s , the equation approaches zero order, and Equation (2-13) becomes:

$$\frac{dx/dt}{x} = R_{\max} \quad (2-14)$$

For S much less than K_s , the equation approaches first order:

$$\frac{dx/dt}{x} = SR_{\max}/K_s \quad (2-15)$$

with R_{\max}/K_s being the first-order rate constant.

For intermediate values of S , the equation is mixed order, with the order dependent on values of the constants R_{\max} and K_s .

2.7 PHOTOCHEMICAL DECOMPOSITION

Photochemical decomposition may occur in two ways. A chemical may absorb light and react (direct photolysis), or the chemical may react because of light absorption by surrounding elements (indirect photolysis).

For direct photolysis, the rate of reaction of a dilute solution of chemical in pure water is as follows:

$$K_p = b \Phi \Sigma \epsilon_\lambda I_\lambda [C], \quad (2-16)$$

where

K_p = rate of direct photolysis, g/(L s)

b = unit conversion constant, 3.8×10^{-21} g mol cm^4 /(L photon)

Φ = reaction quantum yield, dimensionless

ϵ_λ = light absorption coefficient at wavelength interval λ ,
L/(g mol \cdot cm)

I_λ = light flux at wavelength interval λ , photons/(cm 3 \cdot s)

C = concentration of the chemical in water, g/L.

Lyman¹³ refers to Zepp and Cline;¹⁴ Zepp;¹⁵ and Mabey, Mill, and Hendry¹⁶ for details of rate calculations in aquatic systems. In these systems, the rate constant K_p varies with the distribution of sunlight and its intensity. Time of day, season, cloud cover, and latitude all affect K_p so that a reference condition must be stated; e.g., a light flux of photons per second corresponding to a cloudless yearly average at a latitude of 40°N.

Reactions may be photocatalyzed. For example, a TiO_2 catalyst can be photoexcited by light at wavelengths less than 360 nm. Ollis¹⁷ examined the degradation of halogenated hydrocarbons with this catalyst and found a rate equation of the form:

$$\frac{1}{dC/dt} = \frac{1}{k_p} + \frac{1}{k_p K_b C} \quad (2-17)$$

where

k_p = photolysis rate constant, g chemical/(L \cdot s \cdot g catalyst)

K_b = apparent binding constant of a reaction intermediate adsorbed on the illuminated catalyst surface, L/g chemical.

For 11 halocarbons, values of k ranged from 5.8×10^{-8} to 2.3×10^{-6} g/L \cdot s \cdot g of catalyst, with most about 2.8×10^{-7} to 1.7×10^{-6} . A twelfth halocarbon had a k value of 2.3×10^{-4} . Values of K_b for the 12 compounds ranged from 2 to 20 L/g.

2.8 HYDROLYSIS

Hydrolysis occurs when a chemical reacts with water. For organic compounds, the reaction usually replaces a functional group (X) with a hydroxyl:18



Reaction rate constants may be pH-dependent; for a specific pH:

$$k_H = k_a [H^+] + k_n + k_b [OH^-] \quad (2-19)$$

where

k_H = first-order hydrolysis rate constant, s^{-1}

k_a = second-order rate constant for acid-promoted hydrolysis,
 $L/(g \text{ mol} \cdot s)$

$[H^+]$ = hydrogen ion concentration, $g \text{ mol}/L$

k_n = first-order rate constant for pH-independent neutral hydrolysis, s^{-1}

k_b = second-order rate constant for base-promoted hydrolysis,
 $L/(g \text{ mol} \cdot s)$

$[OH^-]$ = hydroxyl ion concentration, $g \text{ mol}/L$.

If

$$k_w = [H^+] [OH^-] \quad (2-20)$$

where

k_w = ionization constant for water $\approx 10^{-14} g \text{ mol}^2/L^2$.

Equation (2-19) can be transformed to:

$$k_H = k_a [H^+] + k_n + k_b k_w / [H^+] . \quad (2-21)$$

The rate constant k_H depends on system pH and on the relative values of k_a , k_b , and k_n .

2.9 OXIDATION/REDUCTION

Organic compounds in aquatic systems may be oxidized by oxygen (particularly as singlet oxygen, 1O_2) or other oxidants such as hydroxyl radicals (OH) and peroxy radicals (RO_2). The OH radicals tend to be very reactive, but present only in low concentrations. The RO_2 radicals are less reactive than the OH radicals, but are present in greater concentrations. Singlet oxygen is highly reactive, but also selective. It has an affinity for electron-rich structures such as dienes and substituted olefins.

The oxidation rate can be calculated as:¹⁹

$$\frac{dC}{dt} = C (k_{RO_2} [RO_2] + k_{SO} [^1O_2] + k_x [X]) \quad (2-22)$$

where

k_{RO_2} = rate constant for peroxy radicals, L/(g mol·s)

$[RO_2]$ = concentration of peroxy radicals, g mol/L

k_{SO} = rate constant for singlet oxygen, L/(g mol·s)

$[^1O_2]$ = concentration of singlet oxygen, g mol/L

k_x = rate constant for "other" oxidants, L/(g mol·s)

$[X]$ = concentrations of "other" oxidants, g mol/L.

In anaerobic environments, reduction reactions may take place. Organochlorines are particularly affected. The reduction rate can be calculated as:²⁰

$$\frac{dC}{dt} = C \sum_i k_i [R_i] \quad (2-23)$$

where

k_i = rate constant for reductant i, L/g mol·s

$[R_i]$ = concentration of reductant i, g mol/L.

2.10 HYDROXYL RADICAL REACTIONS

Hydroxyl radical reactions may occur through addition of a hydroxyl radical, abstraction of a hydrogen atom, or both. In the addition, reaction molecules with high electron density portions attract electrophilic hydroxyl radicals. Hydrogen abstraction takes place when a carbon-hydrogen bond in an organic molecule is easily broken; it is controlled by electronic configuration and number of hydrogen reactions in the molecule. The rate constant for the reaction is often in the range of 6 to 60×10^8 L/(g mol·s).

A hydroxyl radical reaction rate can be calculated as:²¹

$$\frac{dC}{dt} = k_{OH} [OH^\cdot] C \quad (2-24)$$

where

k_{OH} = rate constant for hydrogen abstraction or hydroxyl addition, L/(g mol·s).

2.11 REFERENCES

1. ICF, Inc. The RCRA Risk-Cost Analysis Model Phase III Report, Appendix E. Chemical and Physical Processes Affecting Decay Rates of Chemicals in Aquatic Environments. Draft. Economic Analysis Branch, U.S. Environmental Protection Agency Office of Solid Waste. Washington, DC. 1984.
2. Reference 1, p. E-18, Equation (14).
3. Spivey, J. J., C. C. Allen, D. A. Green, J. P. Wood, and R. L. Stallings. Preliminary Assessment of Hazardous Waste Pretreatment as an Air Pollution Control Technique. Research Triangle Institute. Research Triangle Park, NC. EPA Contract No. 68-03-3149, Task 12-5. 1984.
4. Perry, R. H., and C. H. Chilton. Chemical Engineers' Handbook, Fifth Edition. New York, McGraw-Hill. 1973.
5. Reference 1, p. E-18 - E-19.
6. Allen, C. C., D. A. Green, and J. B. White. Preliminary Assessment of Aerated Waste Treatment Systems at TSDFs--Phase I. Draft. Research Triangle Institute. Research Triangle Park, NC. EPA Contract No. 68-03-3149, Task 54-01F. 1985.

7. Farino, W., P. Spawn, M. Jasinski, and B. Murphy. Evaluation and Selection of Models for Estimating Air Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. GCA/Technology. EPA 450/3-84-020. 1984.
8. Hornick, S. B. In: Land Treatment of Hazardous Waste, Parr, J. F. (ed). Noyes Data Company. Park Ridge, NJ. 1982.
9. Matter-Muller, C., W. Gujer, W. Geiger, and W. Stumm. The Prog. Wat. Tech. (Toronto) 12:299-314. IAWPR/Pergamon Press, Ltd., Great Britain. 1980.
10. Reference 8.
11. Clark, J. W., W. Viessman, Jr., and M. J. Hammer. Water Supply and Pollution Control. Scranton, PA, International Textbook Company. 1971.
- 12. Reference 1, p. E-16, Equation (11).
13. Lyman, W. J., et al. Research and Development of Methods for Estimating Physicochemical Properties of Organic Compounds of Environmental Concern. Phase II, Part I. NTIS AD 11875A. 1981.
14. Zepp, R. G., and D. M. Cline. Rate of Direct Photolysis in Aquatic Environment. Environ. Sci. Technol. 11(4):359-366. 1977.
15. Zepp, R. G. Quantum Yields for Reaction of Pollutants in Dilute Aqueous Solution. Environ. Sci. Technol. 12(3):327-329. 1979.
16. Mabey, W. R., T. Mill, and D. G. Hendry. Photolysis in Water. In: Laboratory Protocols for Evaluating the Fate of Organic Chemicals in Air and Water. Draft. U.S. Environmental Protection Agency. EPA Contract 68-03-2227. 1980.
17. Ollis, D. F. Contaminant Degradation in Water. ES&T. 19(6):480-484. 1985.
18. Reference 13.
19. Reference 1, p. E-12, Equation (2).
20. Reference 1, p. E-12, Equation (3).
21. Reference 1.

3.0 IMPORTANCE OF PATHWAYS

3.1 INTRODUCTION

The importance of the nine pathways described in Chapter 2.0 for surface impoundment, open tanks, land treatment facilities, landfills, and wastepiles is described in this section. The discussion centers on the pathways used in the emission models described in subsequent sections. The pathways described in Chapter 2.0 are repeated below for convenience:

- Volatilization
- Adsorption
- Migration
- Runoff
- Biological decomposition
- Photochemical decomposition
- Hydrolysis
- Oxidation/reduction
- Hydroxyl radical reaction.

Section 3.2 presents the relative importance of these pathways based on the theoretical discussions appearing in Chapter 2.0, the data appearing in the literature, and engineering judgment. Section 3.3 summarizes in tabular form the results of the emission model analyses in Chapters 4.0 through 6.0 and the pathways forming the basis for the emission models.

3.2 THEORETICAL BASIS

The relative importance of the nine pathways for TSDF is discussed in the following text and summarized in Table 3-1. These data were used as

TABLE 3-1. PATHWAYS FOR HAZARDOUS WASTE AREA EMISSION SOURCES^a

Pathway	Surface impoundments	Wastewater treatment plants		Land treatment	Landfill
		Aerated	Nonaerated		
Volatilization	I	I	I	I	I
Biodegradation	I	I	I	I	S
Photodecomposition	S	N	N	N	N
Hydrolysis	S	S	S	N	N
Oxidation/ reduction	N	N	N	N	N
Adsorption	N	S	S	N	N
Hydroxyl radical reaction	N	N	N	N	N
Migration ^b	N	N	N	N	N
Runoff ^b	N	N	N	N	N

I = Important.

S = Secondary.

N = Negligible or not applicable.

^aIndividual chemicals in a given site type may have dominant pathways different from the ones shown here.

^bWater migration and runoff are considered to have negligible effects on ground and surface water in a properly sited, operated, and maintained RCRA-permitted hazardous waste treatment, storage, and disposal facility.

the basis for the emission models contained in CHEMDAT7. Results of exercising these models to identify pathways of importance are discussed in Chapters 4.0 through 7.0 and are summarized in Section 3.3. A short discussion of the theoretical basis for pathways selection follows. Appendix B presents a more detailed discussion.

3.2.1 Surface Impoundments

Data reported by ICF show predominant removal mechanisms and half-lives for 71 chemicals. Table 3-2 lists the mechanisms and statistics for six surface water pathways. Average half-lives range from about 1/2 to 8 days, with predominant mechanisms being volatilization and biodegradation. The rate of photodecomposition depends on the depth of the surface impoundment. The rate is negligibly low for depths as great as 3 meters and is indicated in Table 3-1 as S for a secondary effect.

3.2.2 Aerated and Nonaerated Wastewater Treatment

As in the case of the surface impoundments, volatilization and biodegradation are potentially significant mechanisms. The relative rates of these mechanisms depend on the particular component and treatment system. Photodecomposition is not expected to be a significant pathway due to the opacity of the system, the depth of the liquid, and the residence time of the processes. Adsorption is not expected to be significant except for large loadings of suspended solids and oils in the wastewater. The concentrations for many VO are expected to be roughly the same in the biomass as in the aqueous phase.

3.2.3 Land Treatment

Based on available emission data and literature sources, volatilization and biodegradation are expected to be important in land treatment.²⁻⁶ For highly volatile constituents, volatilization is expected to be the predominant pathway; for low volatile constituents, biodegradation is expected to be the predominant pathway. Adsorption of organic compounds onto organic carbon in the soil also occurs at land treatment sites. However, calculations of land treatment air emissions both with and without consideration of adsorption show a difference of only 10 percent. Therefore, adsorption is not considered a major pathway for organics removal.

The method of waste application and incorporation into the soil influence the importance of photochemical reactions in the degradation of

TABLE 3-2. STATISTICS FOR SURFACE WATER PATHWAYS

	Pathway					
	Vola-tiliza-tion	Biodegrada-tion	Photo-decompo-sition ^a	Hydrolysis	Oxida-tion/reduc-tion	Adsorption
Range of half-lives, days	0.9-15	0.04-96	0.04-900	0.0003-35	0.1-5	0.04-1.5
Average half-life	2.24	8.05	76.3 1.37	5.39	2.05	0.55
Standard deviation	2.85	19.4	259.0 1.82	10.8	2.40	0.83
Number of chemicals	38	26	12	11	4	3

^aStatistics are given for chemicals with and without an outlier.

TABLE 3-3. PATHWAYS FOR TSDF SITES

Type of facility	Pathways included in model
Quiescent storage and treatment impoundments	Volatilization
Mechanically aerated impoundments	Volatilization Biodegradation
Quiescent disposal impoundments	Volatilization
Land treatment facilities	Volatilization Biodegradation
Closed landfills	Volatilization (diffusion through cap) Barometric pumping
Active landfills	Volatilization (diffusion through waste)
Wastepiles	Volatilization

organic wastes in land treatment facilities.⁷ Photodecomposition can occur in land treatment between application and tilling (usually 24 hours), although exposure to sunlight is limited to daylight hours. While exact rates of photodegradation are not known, they are expected to be low. The oil in which the hazardous materials are suspended is semiopaque to sunlight, which would tend to keep photodecomposition low. After tilling, photodegradation is nonexistent because sunlight does not penetrate the soil surface.⁸ Consequently, photodecomposition is not expected to be significant.

3.2.4 Landfills

Volatilization is expected to be a primary VO pathway for landfills. Biodegradation is expected to be negligible for hazardous waste landfills. The toxic properties of the water are expected to inhibit biological processes and therefore biodegradation.⁹

Rates of diffusion in the gas phase may be important. Components can diffuse through unsaturated soils (air pockets present). Control of liquid infiltration into the landfill is expected to keep migration into the soil at a negligible level.

3.3 EMISSION MODELS

Based on the exercise of CHEMDAT7 in predicting and comparing pathways for TSDF processes, the pathways shown in Table 3-3 are used as the basis of the models. Insignificant emissions or inadequate data upon which to develop the model relationships are the principal reasons for limiting the models to the pathways shown in Table 3-3.

It should be noted that CHEMDAT7 includes provisions to activate the unused pathways should further investigations and field tests indicate the desirability of incorporating additional pathways in the emission models.

3.4 REFERENCES

1. ICF, Inc. The RCRA Risk-Cost Assessment Model Phase III Report, Appendix E. Chemical and Physical Processes Affecting Accurate Rates of Chemicals in Aquatic Environments. Draft. Economic Analysis Branch, U.S. Environmental Protection Agency Office of Solid Waste. Washington, DC. 1984.
2. American Petroleum Institute. Land Treatment--Safe and Efficient Disposal of Petroleum Waste. Undated.

3. Bossert, I., et al. Fate of Hydrocarbons During Oily Sludge Disposal in Soil. *Applied and Environmental Microbiology*. 47(4):763-767. 1984.
4. Pelter, P. Determination of Biological Degradability of Organic Substances. *Water Research*. 10:231-235. 1976.
5. Dupont, R. Ryon, and J. A. Reinemon (Utah Water Research Laboratory). Evaluation of Volatilization of Hazardous Constituents at Hazardous Waste Land Treatment Sites. Prepared for U.S. Environmental Protection Agency. Ada, OK. August 1986. 157 p.
6. Eklund, B. M., T. P. Nelson, and R. G. Wetherold (Radian Corporation). Field Assessment of Air Emissions and Their Control at a Refinery Land Treatment Facility. Prepared for U.S. Environmental Protection Agency. Cincinnati, OH. DCN 86-222-078-15-07. September 12, 1986. 330 p.
7. Kaufman, D. D. Fate of Toxic Organic Compounds in Land-Applied Wastes. In: *Land Treatment of Hazardous Wastes*, Parr, J. F., et al. (eds). Park Ridge, NJ, Noyes Data Corporation. 1983. p. 77-151.
8. Reference 7.
9. Shen, T. T. Estimation of Hazardous Air Emissions from Disposal Sites. *Pollution Engineering*. pp. 31-34. August 1981.

4.0 SURFACE IMPOUNDMENTS AND OPEN TANKS

This chapter discusses the approach used to estimate air emissions from surface impoundments and open top tanks. The emission models are described, model facilities are defined, and example calculations are presented.

4.1 NARRATIVE DESCRIPTION OF EMISSIONS AND MODEL UNITS

Emissions from surface impoundments and open tanks originate from the uncovered liquid surface that is exposed to the air. The model used to estimate emissions from the liquid surface is based on an overall mass transfer coefficient that incorporates two resistances to mass transfer in series--the liquid-phase resistance and the gas-phase resistance. Numerous correlations are available to estimate the individual mass transfer coefficients (or resistances), and they depend upon the compound's properties and the system's parameters. The recommended correlations and their applicability are described in subsequent sections. The emission estimating procedure also incorporates a flow model that describes the method of operation. For flowthrough systems, the impoundment's or tank's contents may be completely mixed, plug flow, or somewhere in between with varying degrees of backmixing or axial dispersion. Biologically active impoundments and aeration tanks can be designed for either completely mixed or plug flow, and both types of flow models are discussed for these types of systems. For disposal impoundments, the contents are assumed to be well mixed, and the bulk concentration is expressed as a function of time. An expression for biodegradation is incorporated for those units specifically defined for biodegradation, such as treatment impoundments or wastewater treatment tanks. For these units, the relative rates of air emissions and biodegradation are determined to assess the predicted extent of each mechanism.

The general approach that is used to estimate emissions compares the relative rates of air emissions, biodegradation, and removal with the effluent. Several different types of model units are presented and include mass transfer to the air from quiescent, mechanically aerated, diffused-air, and oil-film liquid surfaces. The other major difference among the types of model units is the type of flow model that is used. For flow-through systems, the degree of mixing can range from complete mixing to plug flow (no mixing), and both cases are presented. For disposal units with no flow out, emissions are a function of time, and average emissions are estimated for some specified time since disposal. The major difference in the emission equations is the liquid-phase concentration that is used for the driving force for mass transfer to the air. The simplest case is represented by well-mixed systems in which the driving force is represented by C_L , the liquid-phase concentration in the bulk liquid, which is also equal to the effluent concentration. Relative removal rates can be compared for this well-mixed case from a simple material balance.

For plug flow, integration is required because the driving force for mass transfer changes as the liquid flows through the system. This concentration is a function of location or time (which are equivalent in plug flow) and is expressed as C_t (denoting a dependence on time). The effluent from a plug flow system is denoted as C_e . For disposal impoundments, the driving-force concentration changes with time and is also denoted as C_t ; however, there is no effluent from a disposal impoundment. The integration required for plug flow is from $t = 0$, when the material first enters the unit, to $t =$ residence time, when the material leaves the unit. For disposal units, the integration is from $t = 0$, when the material is first placed in the unit, to $t =$ time since disposal, which must be specified to estimate average emissions. The integrated forms of these emission equations are very similar.

The well-mixed flow model is recommended and is the model used in the computer program accompanying this report. This flow model is more generally applicable than plug flow; the calculations are more straightforward, and the two types give similar results. The only exception is a flow-through impoundment with an oil film surface, which uses the plug flow

model because the oil film inhibits mixing. Both models yield an estimate of air emissions, biodegradation, and the quantity leaving with the effluent. It is important to recognize that the quantity leaving with the effluent may also eventually contribute to air emissions, especially for treatment units in series or for discharges to streams or publicly owned treatment works.

Equations are presented for estimating the various removal rates, and example calculations for different types of impoundments are also provided. Example calculations are not presented separately for open tanks because the procedure is analogous to that used for impoundments. In general, open tanks will have different input parameters that will account for differences in emission rates compared to impoundments. For example, the liquid surface area for open tanks will be less, and the fetch-to-depth (F/D) ratio will be much lower for tanks. If the open tank has a wind barrier to reduce the wind velocity, the reduced wind velocity can be used in the mass transfer correlations. In addition, the modeling approach accounts for the shorter retention times in tanks (on the order of hours) compared to impoundments (on the order of days). For open tanks, the mass transfer correlation of Springer is recommended for windspeeds less than 3.25 m/s, and the correlation of MacKay and Yeun is recommended for windspeeds greater than 3.25 m/s. Both are discussed in the following section.

4.2 QUIESCENT SURFACES WITH FLOW

4.2.1 Emission Model Equations

The primary focus on emissions from impoundments and wastewater treatment tanks is on aqueous solutions contaminated with organics because aqueous waste is the most common waste type handled in these facilities. For aqueous systems, the basic relationship describing mass transfer of a volatile constituent from the open liquid surface to the air is:

$$E = KAC_L \quad (4-1)$$

where

E = air emissions from the liquid surface, g/s

K = overall mass transfer coefficient, m/s

A = liquid surface area, m^2

C_L = concentration of constituent in the liquid phase, g/m^3 .

The overall mass transfer coefficient (K) is estimated from a two-phase resistance model that is based on the liquid-phase mass transfer coefficient (k_L in m/s), the gas-phase mass transfer coefficient (k_G in m/s), and Henry's law constant in the form of a partition coefficient (Keq). The two resistances act in series and the overall resistance is expressed as:

$$\frac{1}{K} = \frac{1}{k_L} + \frac{1}{k_G Keq} \quad (4-2)$$

where

K = overall mass transfer coefficient, m/s

k_L = liquid-phase mass transfer coefficient, m/s

k_G = gas-phase mass transfer coefficient, m/s

Keq = equilibrium constant or partition coefficient, concentration in gas phase/concentration in liquid phase where both concentrations are in the same units.

Henry's law constant (H in $atm \cdot m^3/g \text{ mol}$) is estimated for the constituents of interest by dividing the constituent's vapor pressure (in atmospheres) by its solubility in water (in $g \text{ mol}/m^3$). The equilibrium constant is estimated by:

$$Keq = H/RT \quad (4-3)$$

where

H = Henry's law constant, $atm \cdot m^3/g \text{ mol}$

R = universal gas constant, $8.21 \times 10^{-5} \text{ atm} \cdot m^3/g \text{ mol} \cdot K$

T = temperature, K .

For a standard temperature of $25^\circ C$, the expression for Keq reduces to:

$$Keq = 40.9 \times H \quad (4-4)$$

The units associated with K_{eq} in Equation (4-4) are the ratio of gas-phase to liquid-phase concentrations and require that both be expressed in the same units of mass/volume.

Several mathematical models have been developed to estimate the individual liquid- and gas-phase mass transfer coefficients. The models are based on different systems, constituents, and sometimes different theoretical considerations. Many of these models yield similar results. The procedures used in this section to estimate the individual mass transfer coefficients rely primarily on existing mass transfer correlations that are believed to be generally applicable.

The liquid-phase mass transfer coefficient (k_L) has been shown to be a function of the constituent's diffusivity in water, windspeed, and liquid depth.^{1,2} Work performed at the University of Arkansas by Springer et al.³ confirmed these effects and resulted in the correlations given in Table 4-1. Springer used simulation studies in a wind tunnel water tank of a constant fetch (2.4 m) and variable depth (4.7 cm to 1.2 m). Fetch is defined as the linear distance across the liquid surface in the direction of the wind flow, and the F/D ratio is defined as the fetch divided by the depth of the impoundment. Ethyl ether was used as the volatile component in the desorption experiments, in which the wind velocity and F/D ratio were varied. Springer's results shown in Table 4-1 yield three different correlations for k_L that depend upon the combination of windspeed and F/D ratio of interest. Springer's model implies that k_L is constant for windspeeds of 0 to 3.25 m/s. Although Springer examined only the mass transfer of ethyl ether, his results are extrapolated to other compounds by the ratio of the compound's and ether's diffusivities in water to the $2/3$ power. The windspeed in Springer's correlation is defined as the windspeed 10 m above the liquid surface. For practical application of his correlation, typically reported values of windspeed are used. Springer's model does not include the case in which the F/D ratio is less than 14 and the windspeed is greater than 3.25 m/s. For this specific case, k_L was estimated from MacKay and Yeun's correlation shown in Table 4-1.^{7,8} MacKay and Yeun⁹ did not address the effect of depth; however, their correlation is based on data from 11 organic compounds in a well-mixed system, the

TABLE 4-1. EQUATIONS FOR CALCULATING INDIVIDUAL MASS TRANSFER COEFFICIENTS FOR VOLATILIZATION OF ORGANIC SOLUTES FROM QUIESCENT SURFACE IMPOUNDMENTS

Liquid phase

Springer et al.⁴ (for all cases except $F/D < 14$ and $U_{10} > 3.25$ m/s):

$$k_L = 2.78 \times 10^{-6} \left(\frac{D_w}{D_{\text{ether}}} \right)^{2/3} \quad (0 < U_{10} < 3.25) \text{ (m/s)} \\ \text{(All F/D ratios)}$$

$$k_L = [2.605 \times 10^{-9} (F/D) + 1.277 \times 10^{-7}] U_{10}^2 \left(\frac{D_w}{D_{\text{ether}}} \right)^{2/3} \quad (U_{10} > 3.25) \text{ (m/s)} \\ (14 < F/D < 51.2)$$

$$k_L = 2.611 \times 10^{-7} U_{10}^2 \left(\frac{D_w}{D_{\text{ether}}} \right)^{2/3} \quad (U_{10} > 3.25) \text{ (m/s)} \\ (F/D > 51.2)$$

where

U_{10} = windspeed at 10 m above the liquid surface, m/s

D_w = diffusivity of constituent in water, cm^2/s

D_{ether} = diffusivity of ether in water, cm^2/s

F/D = fetch-to-depth ratio (fetch is the linear distance across the impoundment).

Gas phase

MacKay and Matasugu (in Hwang⁵):

$$k_G = 4.82 \times 10^{-3} U^{0.78} Sc_G^{-0.67} d_e^{-0.11} \text{ (m/s)}$$

where

U = windspeed, m/s

Sc_G = Schmidt number on gas side = $\frac{\mu_G}{\rho_G D_a}$

μ_G = viscosity of air, $\text{g}/\text{cm}\cdot\text{s}$

(continued)

TABLE 4-1 (continued)

ρ_G = density of air, g/cm³

D_a = diffusivity of constituent in air, cm²/s

d_e = effective diameter of impoundment = $\left(\frac{4A}{\pi}\right)^{0.5}$, m

A = area of impoundment, m².

Liquid phase

Mackay and Yeun⁶ (for $F/D < 14$ and $U_{10} > 3.25$ m/s):

$k_L = 1.0 \times 10^{-6} + 34.1 \times 10^{-4} U^* Sc_L^{-0.5} (U^* > 0.3)$ (m/s)

$k_L = 1.0 \times 10^{-6} + 144 \times 10^{-4} U^{*2.2} Sc_L^{-0.5} (U^* < 0.3)$ (m/s)

where

U^* = friction velocity (m/s) = $0.01 U_{10} (6.1 + 0.63 U_{10})^{0.5}$

U_{10} = windspeed at 10 m above the liquid surface, m/s

Sc_L = Schmidt number on liquid side = $\frac{\mu_L}{\rho_L D_w}$

μ_L = viscosity of water, g/cm•s

ρ_L = density of water, g/cm³

D_w = diffusivity of constituent in water, cm²/s.

compounds represent a broad range of Henry's law constants, and their general correlation is applicable for the case described above that is not covered by Springer's correlation.

The gas-phase coefficient (k_g) was estimated from the correlation of MacKay and Matasugu as shown in Table 4-1.¹⁰ This correlation was developed from experiments on the evaporation of isopropyl benzene, gasoline, and water into air. These researchers verified that previous work, which assumed that the wind velocity profile follows a power law, could be used to quantify the rate of evaporation from a smooth liquid surface. The result was a correlation that expressed k_g as a function of windspeed and the fetch or effective diameter of the liquid surface.

The individual mass transfer coefficients estimated from the correlations in Table 4-1 are used in Equation (4-2) to estimate the overall mass transfer coefficient. The equilibrium constant for a constituent dissolved in water at 25 °C is estimated from Equation (4-4). However, an estimate of the concentration in the liquid phase (C_L) is needed in Equation (4-1) to estimate emissions.

The concentration C_L in Equation (4-1) is the driving force for mass transfer. For an impoundment that is instantly filled with waste, the driving force (C_L) is the initial concentration in the waste. However, this concentration will decrease with time as the constituent is lost to the air, which suggests that emissions may also decrease with time (assuming constant K and A). For flowthrough systems, the concentration may be cyclical if the loading of the process is cyclical. Continuous flowthrough systems may attain some equilibrium concentration.

The flow model assumed for quiescent impoundments and tanks with no biodegradation is that the contents of the system are well mixed and that the bulk concentration (driving force) in the system is equal to the effluent concentration. A material balance around this system yields:

$$QC_0 = KAC_L + QC_L \quad (4-5)$$

or

$$C_L = QC_0 / (KA + Q) \quad (4-6)$$

where

Q = volumetric flow rate, m^3/s

C_0 = initial concentration in the waste, g/m^3

C_L = equilibrium or bulk concentration in the impoundment, g/m^3

K = overall mass transfer coefficient, m/s

A = liquid surface area, m^2 .

The well-mixed assumption is made for the sake of simplicity and assumes that bulk convection and wind-induced eddies combine to mix the basin contents. Axial dispersion in the flow direction is also possible, and some systems may be designed specifically for plug flow (e.g., some biological treatment tanks). An assumption of plug flow instead of well-mixed flow would yield slightly higher estimates of emissions; however, the difference is small. Calculations presented by Thibodeaux for an aerated basin that was well-mixed or had plug flow showed that the plug-flow assumption yielded estimates that were higher by 11 percent for acetaldehyde, 5 percent for acetone, and 0 percent for phenol.¹¹

The approach described to estimate emissions from quiescent impoundments with no biodegradation includes the following steps:

1. Estimate the individual mass transfer coefficients from Table 4-1.
2. Estimate the equilibrium constant from Equation (4-3).
3. Estimate the overall mass transfer coefficient from Equation (4-2).
4. Estimate the liquid-phase concentration from Equation (4-6).
5. Estimate emissions from Equation (4-1).

The major assumptions associated with this procedure are:

- The two-resistance model and the correlations for the individual mass transfer coefficients are applicable to the system of interest.
- The impoundment's contents are well mixed.
- There is no significant removal by biodegradation, seepage, adsorption, or other forms of degradation.

- The waste material of interest is aqueous waste with no separate organic phase.
- The estimate of Henry's law constant (equilibrium partitioning between the vapor and liquid) is reasonably accurate.

The recommended procedure for quiescent impoundments is to assume that the liquid is well mixed. This assumption is used in the computer model accompanying this report and is illustrated in the example calculations. However, impoundments and tanks with quiescent surfaces can also be designed for plug flow with the use of baffles or other design techniques to reduce the extent of backmixing. In a plug-flow system, the rate of air emissions at any point in the system changes as the material flows through the system. There is no uniform liquid concentration within the plug-flow unit as there was in the well-mixed system, and the lowest concentration occurs in the effluent (i.e., there is no backmixing of the effluent with the influent). For plug flow, the rate of disappearance of a compound by air emissions is given by:

$$\frac{-d C_t (V)}{dt} = K A C_t \quad (4-7)$$

where

C_t = concentration after the plug has traveled t seconds

t = time, s

V = volume, m^3

and with the other symbols as previously defined.

Rearranging Equation (4-7) yields:

$$\frac{d C_t}{C_t} = (-KA/V)dt \quad (4-8)$$

Integrating Equation (4-8) from $C_t = C_0$ at $t = 0$ to $C_t = C_e$ at $t = V/Q$ (one residence time) gives:

$$C_e/C_0 = \exp (-KA/Q) \quad (4-9)$$

where

C_e = effluent concentration, g/m³

and with the other symbols as previously defined.

The residence time, τ in seconds, equals V/Q and $V = AD$ (area times depth); consequently, $A/Q = \tau/D$. Substituting into Equation (4-9) yields an equivalent expression:

$$C_e/C_o = \exp (-K\tau/D) \quad . \quad (4-10)$$

The ratio C_e/C_o represents the fraction removed with the effluent; therefore, $1 - C_e/C_o$ represents the fraction that is emitted (f_{air}) from the plug-flow system:

$$f_{air} = 1 - C_e/C_o = 1 - \exp (-K\tau/D) \quad . \quad (4-11)$$

The average emission rate is calculated from:

$$E = f_{air} Q C_o \quad (4-12)$$

where

E = emissions, g/s

f_{air} = fraction emitted from Equation (4-11)

Q = flow rate, m³/s

C_o = influent concentration, g/m³.

4.2.2 Model Plant Parameters for Quiescent Impoundments

A model facility was developed for quiescent impoundments to illustrate the emission estimating procedure. A 1981 survey compiled by Westat¹² showed that the median surface area for storage impoundments was approximately 1,500 m² and that the median depth was 1.8 m. Detention times ranged from 1 to 550 days, with over half of the values at 46 days or less. For this example, a detention time of 20 days was chosen. The area and depth yield a total volume of 2,700 m³, and the detention time of 20 days yields a flow rate of 1.6 L/s (0.0016 m³/s).

Meteorological conditions are also needed as input parameters for the emission models. For this emission estimate, a standard temperature of 25 °C and a windspeed of 4.47 m/s (10 mi/h) were used. Benzene was chosen

as an example constituent at a concentration of 10 g/m^3 (10 ppm) to estimate emissions from the model facility. The properties of benzene that are used include Henry's law constant ($5.5 \times 10^{-3} \text{ atm}\cdot\text{m}^3/\text{g mol}$), diffusivity in air ($0.088 \text{ cm}^2/\text{s}$), and diffusivity in water ($9.8 \times 10^{-6} \text{ cm}^2/\text{s}$). Table 4-2 lists the input parameters for the estimate of emissions given in Section 4.2.3.

4.2.3 Example Calculation for Storage Impoundments

This section presents a step-by-step example calculation for emissions from storage impoundments. The equations described in Section 4.2.1 are used with the model unit parameters given in Section 4.2.2 to estimate emissions from an aqueous waste containing 10 g/m^3 of benzene.

- a. Calculate liquid-phase mass transfer coefficient, k_L . Use Springer's model (see Table 4-1):

$$\text{Effective diameter} = \left(\frac{\text{Area}}{\pi} \right)^{0.5} \times 2 = \left(\frac{1,500}{\pi} \right)^{0.5} \times 2 = 43.7 \text{ m}$$

$$F/D = \text{Effective diameter/depth} = \frac{43.7}{1.8} = 24.3 .$$

$$\begin{aligned} \text{Windspeed} &= 4.47 \text{ m/s} \quad (U_{10} \geq 3.25 \text{ m/s}) \\ F/D &= 24.3 \end{aligned}$$

$$k_L = [2.605 \times 10^{-9} (F/D) + 1.277 \times 10^{-7}] U_{10}^2 \left(\frac{D_w}{D_{\text{ether}}} \right)^{0.67} \text{ m/s}$$

where

$$U_{10} = \text{windspeed} = 4.47 \text{ m/s}$$

$$D_w = 9.8 \times 10^{-6} \text{ cm}^2/\text{s} \text{ (benzene)}$$

$$D_{\text{ether}} = 8.5 \times 10^{-6} \text{ cm}^2/\text{s} \text{ (ether)}$$

$$F/D = 24.3.$$

Then

$$k_L = [2.605 \times 10^{-9} (24.3) + 1.277 \times 10^{-7}] (4.47)^2 \left(\frac{9.8 \times 10^{-6}}{8.5 \times 10^{-6}} \right)^{0.67}$$

TABLE 4-2. INPUT PARAMETERS--STORAGE IMPOUNDMENT

Area	1,500 m ²
Depth	1.8 m
Volume	2,700 m ³
Retention time	20 days
Flow	0.00156 m ³ /s
Temperature	25 °C
Windspeed	4.47 m/s
Constituent	Benzene in water
Concentration	100 g/m ³
Henry's law constant	5.5 x 10 ⁻³ atm•m ³ /g mol
Diffusivity in air (benzene)	0.088 cm ² /s
Diffusivity in water (benzene)	9.8 x 10 ⁻⁶ cm ² /s
Diffusivity in water (ether)	8.5 x 10 ⁻⁶ cm ² /s
Viscosity of air	1.81 x 10 ⁻⁴ g/cm•s
Density of air	1.2 x 10 ⁻³ g/cm ³

$$k_L = [2.605 \times 10^{-9} (24.3) + 1.277 \times 10^{-7}] (4.47)^2 (1.1)$$

$$k_L = 4.2 \times 10^{-6} \text{ m/s}$$

- b. Calculate gas-phase mass transfer coefficient, k_G . Use MacKay and Matasugu (see Table 4-1):

$$k_G = 4.82 \times 10^{-3} U^{0.78} Sc_G^{-0.67} d_e^{-0.11} (\text{m/s})$$

where

$$U = \text{windspeed, } 4.47 \text{ m/s}$$

$$Sc_G = \frac{\text{Schmidt No. for gas}}{\text{viscosity of gas}} = \frac{\text{viscosity of gas}}{(\text{gas density})(\text{diffusivity of } i \text{ in gas})}$$

$$\text{Gas} = \text{air}$$

$$\text{Viscosity (air)} = 1.81 \times 10^{-4} \text{ g/cm}\cdot\text{s}$$

$$\text{Density (air)} = 1.2 \times 10^{-3} \text{ g/cm}^3$$

$$\text{Diffusivity (benzene in air)} = 0.088 \text{ cm}^2/\text{s}$$

$$Sc_G = \frac{1.81 \times 10^{-4} \text{ g/cm}\cdot\text{s}}{(1.2 \times 10^{-3} \text{ g/cm}^3)(0.088 \text{ cm}^2/\text{s})} = 1.71$$

$$d_e = \text{effective diameter} = 43.7 \text{ m}$$

Then

$$\begin{aligned} k_G &= (4.82 \times 10^{-3}) (4.47)^{0.78} (1.71)^{-0.67} (43.7)^{-0.11} \\ &= 7.1 \times 10^{-3} \text{ m/s} \end{aligned}$$

- c. Calculate overall mass transfer coefficient (K) from Equation (4-2):

$$\frac{1}{K} = \frac{1}{k_L} + \frac{1}{K_{eq} k_G}$$

where

$$K_{eq} = \frac{H}{RT} = \frac{5.5 \times 10^{-3} \text{ m}^3 \cdot \text{atm/mol}}{(8.21 \times 10^{-5}) \left(\frac{\text{atm} \cdot \text{m}^3}{\text{mol} \cdot \text{K}} \right) (298 \text{ K})} = 0.225$$

Then

$$\frac{1}{K} = \frac{1}{4.2 \times 10^{-6}} + \frac{1}{(0.225)(7.1 \times 10^{-3})} = 2.39 \times 10^5$$

$$K = 4.2 \times 10^{-6} \text{ m/s} .$$

d. Estimate emissions for a well-mixed system:

$$QC_o = KC_L A + QC_L \text{ (from material balance of Equation (4-5))}$$

$$C_L = \frac{QC_o}{KA + Q}$$

$$\text{Detention time} = 480 \text{ h}$$

$$\text{Volume} = 2,700 \text{ m}^3$$

where

$$Q = \text{flow rate} = \left(\frac{2,700 \text{ m}^3}{480 \text{ h}} \right) \left(\frac{1 \text{ h}}{3,600 \text{ s}} \right) = 0.00156 \text{ m}^3/\text{s}$$

$$C_o = 10 \text{ g/m}^3$$

$$K = 4.2 \times 10^{-6} \text{ m/s}$$

$$C_L = \frac{(0.00156 \text{ m}^3/\text{s})(10 \text{ g/m}^3)^3}{(4.2 \times 10^{-6} \text{ m/s})(1,500 \text{ m}^2) + (0.00156 \text{ m}^3/\text{s})} = 19.8 \text{ g/m}^3$$

$$A = 1,500 \text{ m}^2 .$$

$$\text{Air emissions} = KC_L A \text{ (Equation 4-2)}$$

$$\begin{aligned} &= (4.2 \times 10^{-6} \text{ m/s})(19.8 \text{ g/m}^3)(1,500 \text{ m}^2) = 0.012 \text{ g/s} \\ &= 3.9 \text{ Mg/yr} . \end{aligned}$$

e. Estimate emissions for a plug-flow system:

$$f_{\text{air}} = 1 - \exp(-K\tau/D) \text{ (Equation 4-11)}$$

$$K = 4.2 \times 10^{-6} \text{ m/s (Step c)}$$

$$\tau = 480 \text{ h} = 1.73 \times 10^6 \text{ s}$$

$$D = 1.8 \text{ m}$$

$$f_{\text{air}} = 1 - \exp(-4.2 \times 10^{-6} \text{ m/s} \cdot 1.73 \times 10^6 \text{ s} / 1.8 \text{ m}) = 0.98$$

$$E = f_{\text{air}} Q C_o \text{ (Equation 4-12)}$$

$$f_{\text{air}} = 0.98$$

$$Q = 0.00156 \text{ m}^3/\text{s}$$

$$C_o = 100 \text{ g/m}^3$$

$$E = (0.98)(0.00156 \text{ m}^3/\text{s})(100 \text{ g/m}^3)$$

$$E = 0.15 \text{ g/s} = 4.7 \text{ Mg/yr}$$

4.3 BIODEGRADATION

This section identifies some of the major design features of biological treatment processes, such as activated sludge units and impoundments designed for biodegradation. Mathematical models for biodegradation are also presented and incorporated into predictive fate models.

4.3.1 Description of Biological Active Systems

The activated sludge process is an aerobic biological treatment in which the pollutants are degraded by microorganisms suspended uniformly in the reaction tank. Oxygen is introduced by mechanical means, and the microorganisms are maintained by recycling the activated sludge that is formed. In most units, the sludge is removed by settling in a separate unit, a portion of the sludge is recycled, and a small portion is wasted (removed from the system) on a continuous basis. Oxidation or stabilization impoundments and aerated impoundments are used to treat entire plant wastes as well as to polish the effluent from other treatment processes. Solids usually settle out in the impoundment or are removed in a separate vessel. Generally, the solids are not recycled; however, if the solids are returned, the process is the same as a modified activated sludge process.¹³

Typical design parameters for an activated sludge process are given in Table 4-3. Two of the most commonly used parameters are the food-to-microorganism (F/M) ratio and residence time. The F/M ratio describes the organic loading on the biological system and is calculated as the weight of BOD₅ (biochemical oxygen demand from a 5-day test) that enters the system in a 24-hour period divided by the total weight of biological solids in the system. The biological solids may be roughly estimated from the mixed

TABLE 4-3. DESIGN PARAMETERS FOR ACTIVATED SLUDGE PROCESSES¹⁴

Process	F/M, ^a kg BOD/kg biomass•day	Loading, kg BOD/m ³ •day	MLSS, ^b g/L	Retention time, h
Conventional ^c	0.2-0.4	0.3-0.6	1.5-3.0	4-8
CSTR ^d	0.2-0.6	0.8-2.0	3.0-6.0	3-5
Contact stabilization	0.2-0.6	1.0-1.2	1.0-3.0 ^e 4.0-10 ^f	0.5-1 ^e 3-6 ^f
Extended aeration	0.05-0.15	0.1-0.4	3.0-6.0	18-36
O ₂ systems	0.25-1.0	1.6-3.3	6.0-8.0	1-3

^aF/M = Food to microorganism ratio.

^bMLSS = Mixed liquor suspended solids.

^cPlug flow design.

^dCSTR = Continuous stirred-tank reactor.

^eContact unit.

^fSolids stabilization unit.

liquor suspended solids (MLSS) if substantial quantities of inorganics (such as silt) are not present. If inorganic solids are present, the biological solids may be better approximated by the mixed liquor volatile suspended solids (MLVSS).¹⁵ For municipal wastewater systems, the volatile solids comprise about 60 to 80 percent of the total suspended solids in the sludge; consequently, in the absence of a direct measurement of MLVSS, the biological solids in municipal wastewater can be estimated as 60 to 80 percent of the total suspended solids.¹⁶ Conventional plants, which use an activated sludge process that has long and narrow basins designed to approach plug flow, operate with an F/M ratio of 0.2 to 0.4, but values as low as 0.05 are not unusual. High F/M values indicate a high loading, as from a sudden influx of organics or the loss of biological solids, and will lead to a deterioration in effluent quality.¹⁷

Aeration tanks are usually constructed of reinforced concrete, are open to the atmosphere, and are usually rectangular in shape. Treatment plants may consist of several tanks, operated in series or parallel. Some of the largest treatment plants may contain 30 to 40 tanks arranged in several groups or batteries.¹⁸

Typical parameters associated with biologically active impoundments are given in Table 4-4. The loading parameter is expressed in terms of area or volume, and typical retention times in aerated impoundments range from 7 to 20 days. The level of suspended solids in these impoundments is over an order of magnitude less than the level in activated sludge processes. Although the parameters in Table 4-4 are listed as "typical," large variations exist among real facilities, and at a single facility the values may change with time. For example, a study conducted over 12 months at an aerobic impoundment used to treat municipal wastewater reported suspended solids levels of 0.02 to 0.1 g/L and volatile suspended solids of 0.01 to 0.06 g/L.²¹ Another study of eight quiescent impoundments at four different sites with confirmed biological activity estimated active biomass concentrations from the rate of oxygen consumption that ranged from 0.0014 to 0.22 g/L with an average of 0.057 g/L.²²

The biomass concentration is an important parameter in estimating biodegradation rates. The best value to use for a specific site is a

TABLE 4-4. IMPOUNDMENTS DESIGNED FOR BIODEGRADATION^{19,20}

Type	Application	Typical daily loading, kg BOD ₅ /m ³ day	Retention time, d	Typical depth, m	Suspended solids, g/L
Facultative	Raw municipal wastewater Effluent from primary treatment, trickling filters, aerated ponds, or anaerobic ponds	0.0011 - 0.0034 ^a	25-100	1.2-2.5	0.11-0.40
Aerated	Industrial wastes Overloaded facultative ponds Situations where limited land area is available	0.008 - 0.32	7-20	2-6	0.26-0.30
Aerobic	Generally used to treat effluent from other processes, produces effluent low in soluble BOD ₅ and high in algae solids	0.021 - 0.043 ^b	10-40	0.3-0.45	0.14-0.34
Anaerobic	Industrial wastes	0.16 - 0.80	20-60	2.5-5	0.08-0.16

^aBased on a typical depth of 2 m.

^bBased on a typical depth of 0.4 m.

direct measurement such as volatile suspended solids for the system of interest. In the absence of site-specific data, a number may be chosen from the ranges for suspended solids given in Tables 4-3 and 4-4. Alternatively, typical or default values for biomass concentration given in Table 4-5 may be used.

The major mechanisms of organic removal in biologically active systems include biodegradation, volatilization, removal with the effluent, and removal by adsorption on the waste sludge. A study by Petrasek et al. of purgeable volatile organics in a pilot-scale wastewater treatment system showed that less than 0.4 percent (generally less than 0.1 percent) of the volatiles were found in the waste-activated sludge.²³ Bishop, in a study of municipal wastewater treatment, concluded that only a modest amount of purgeable toxics were transferred to the sludge.²⁴ Hannah et al.²⁵ found that the concentrations of volatile organics in sludges from pilot-scale systems were generally comparable to or less than the corresponding concentrations in the process effluent. This indicated that volatile organics do not have a high affinity for wastewater solids and do not concentrate in the sludges. Kincannon and Stover found that 0 to 1 percent of three compounds (1,2-dichloroethane, phenol, and 1,2-dichlorobenzene) was adsorbed on the sludge.²⁶ Melcer, in a review of biological removal studies, concluded that polycyclic aromatic hydrocarbons, pyrene, anthracene, fluoranthene, and chrysene were the most commonly occurring priority pollutants found in sludges.²⁷ These studies suggest that the compounds most likely to be emitted to the air (volatiles) do not concentrate on sludges; however, some of the relatively nonvolatile organics may be adsorbed. Consequently, the modeling approach presented in this section assumes that the removal of volatile organics with the waste sludge is not significant. The major removal mechanisms that are considered include volatilization, biodegradation, and removal with the effluent.

4.3.2 Rate of Biodegradation

Numerous models have been proposed for the removal of organic compounds by biodegradation and include design equations for activated sludge systems and stabilization or oxidation impoundments.^{28,29} There is general agreement in the literature that, for high organic loadings relative to

TABLE 4-5. TYPICAL OR DEFAULT VALUES FOR
BIOMASS CONCENTRATION^a

Unit	Biomass concentration, g/L
Quiescent impoundments	0.05 ^b
Aerated impoundments	0.25 ^c
Activated sludge units	4.0 ^d

^aThese values are recommended for use in the emission equations when site-specific data are not available.

^bBased on the range (0.0014 to 0.22) and average (0.057) from actual impoundments as discussed in the text.

^cFrom the data in Table 4-4 for aerated impoundments. Assumes biomass is approximated by the suspended solids level. Range is typically 0.05 to 0.30.

^dMidrange value from Table 4-3 for CSTR based on mixed liquor suspended solids.

biomass, the biodegradation rate is zero-order with respect to concentration (i.e., the rate is independent of organic concentration). For lower residual levels, the rate becomes first order with respect to concentration or follows Monod-type kinetics.^{30,31,32} The Monod-type biodegradation rate equation can be written as follows:

$$r_B = VK_{\max}b_iC/(K_S+C) \quad (4-13)$$

where

r_B = biodegradation rate, g/s

V = volume, m³

b_i = biomass concentration, g/m³

K_{\max} = maximum rate constant, g/s-g biomass

C = component concentration, g/m³

K_S = half saturation constant, g/m³.

The Monod model was originally developed to describe microbial growth rates for a single microbial population based upon a single, rate-limiting substrate. A yield coefficient was subsequently employed to determine the utilization rate of that substrate. For convenience of use, the biodegradation rate model given in Equation (4-13) has been written directly for component disappearance in terms of overall biomass concentration. It is assumed that Equation (4-13) applies to each organic constituent in the waste (although the rate constants will be different for each constituent), and that the biodegradation of any one constituent is independent of the concentrations of other constituents. Subsequent references to the Monod or the Monod-type model in this report refer specifically to Equation (4-13). The significant features of this model are:

1. At high concentrations (specifically, $C \gg K_S$), C dominates the denominator and can, therefore, be eliminated from Equation (4-13). The biodegradation rate is then independent of (i.e., zero order with respect to) the component concentration.

2. At low concentrations ($C \ll K_S$), K_S dominates the denominator, and the biodegradation rate becomes directly proportional (i.e., first order with respect) to the component concentration. The apparent first-order rate constant is:
 $K_1 = K_{\max}/K_S$.

Theoretical Monod curves for several different compounds are presented in Figure 4-1 to illustrate these features.

A literature review was conducted to determine appropriate rate constants for the Monod model. References that served as primary sources of biodegradation rate data included: Pitter,³³ Kincannon et al.,³⁴ Petrasek et al.,³⁵ and Hannah et al.³⁶ Data obtained from each reference included rate constants as reported, influent concentrations, effluent concentrations, biomass concentration, retention time (RT), and fraction of the amount of component removed by biodegradation (F_B). Using this information and field data collected during specially designed biodegradation rate studies, Coburn et al. developed a base of component-specific biological removal rates that contains nearly 500 entries and removal data for 90 different organic constituents.³⁷

For most compounds, there were inadequate biodegradation rate data to determine the Monod rate constants using traditional methods (e.g., Lineweaver-Burke plot). However, when reported, values for K_S were generally between 1 and 10 mg/L for a variety of different compounds. Thus, the Monod constant, K_{\max} , was calculated from organic removal data when high concentrations ($C > 10$ mg/L) were employed by assuming strict zero-order kinetics as follows:

$$K_{\max} = F_B(C_0 - C_L)/[(RT)b_i] \quad (4-14)$$

where

F_B = the fraction of component removal attributed to biodegradation

C_0 = inlet concentration, g/m³

C_L = bulk liquid and effluent concentration, g/m³

(RT) = residence time, s.

Note that, with zero-order kinetics, Equation (4-14) applies to both continuous, well-mixed systems and to plug-flow and batch systems.

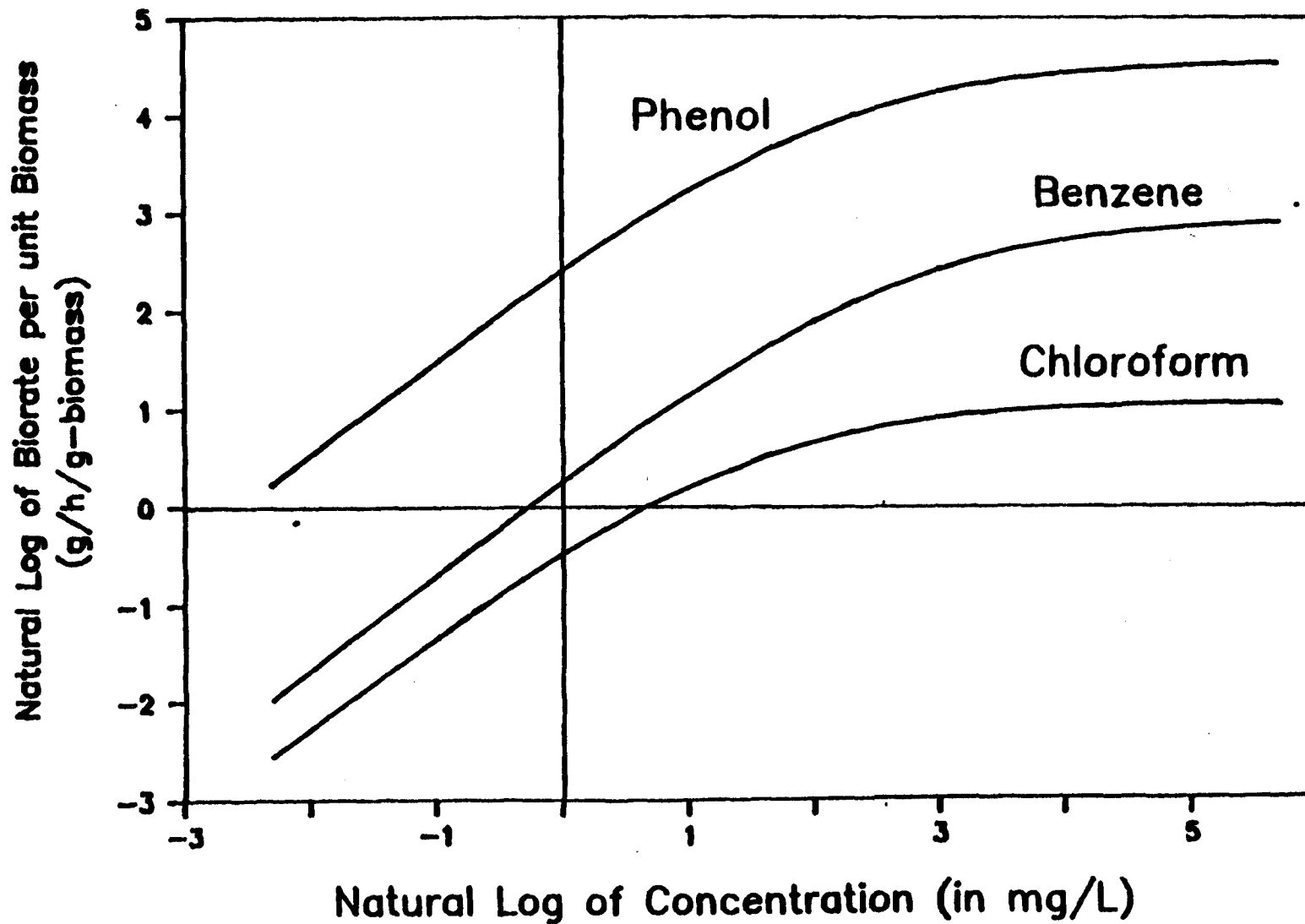


Figure 4-1. Theoretical relationship between concentration and biodegradation rates normalized by the amount of biomass as predicted using the Monod model for phenol, benzene, and chloroform.

The half-saturation constant K_s was estimated (knowing K_{max}) from the apparent first-order rate constants when low concentrations ($C < 1 \text{ mg/L}$) were present (specifically, $K_s = K_{max}/K_1$). The equation used to calculate the apparent first-order rate, K_1 , depends on the type of experimental system that was employed. For continuous, well-mixed systems, K_1 was calculated as follows:

$$K_1 = F_B(C_0 - C_L)/[(RT)b_iC_L] \quad (4-15)$$

For batch systems and for continuous, plug-flow systems, the equation used to calculate K_1 was:

$$K_1 = F_B \ln(C_0/C_L)/[(RT)b_i] \quad (4-16)$$

Using this approach, rate constants for specific compounds in the biodegradation rate data base were determined. These rate constants are provided in Appendix D, Table D-1. Upon evaluating the biodegradation rate data from several different laboratory and field studies, it is recognized that biodegradation rates can vary widely from site to site. Therefore, the following priority schedule is provided as guidance in determining the appropriate biodegradation rate constants to be employed in the emission models:

- Use site-specific biodegradation rate data in experiments controlled for air emissions where available.
- Use the rate constants suggested in Appendix D, Table D-1, as available.
- Estimate the biodegradation rate constants using the following methodology:
 - Approximate K_{max} from available data for K_{max} for compounds of similar structure and/or functional groups; and
 - Approximate K_1 either by using the correlation:

$$K_1 = 3.75 \times 10^{-8} K_{ow}^{0.38} \quad (4-17)$$

where

K_{ow} = octanol-water partitioning coefficient,

or by using the default (average) value for K_1 , which is: $K_1 = 1 \text{ L/h/g}$ ($2.78 \times 10^{-7} \text{ m}^3/\text{s/g}$), and then calculate K_s as: $K_s = K_{\max}/K_1$.

The correlation provided in Equation (4-17) was developed based upon the assumption that biodegradation was primarily an intracellular phenomenon. As such, the first-order biodegradation rate can be limited either by the rate of the internal reaction or by the rate of diffusion of the chemical through the cell membrane and into the cell. If the internal component concentrations are assumed to be proportional to the concentration of components absorbed onto the cell membranes, then, regardless of what limits the first-order biodegradation rate, the limiting first-order biodegradation rate will be directly proportional to the concentration of constituent absorbed onto the membrane. Because the octanol-water partitioning coefficient has been used to correlate the absorption partitioning of organic chemicals onto biomass,^{38,39} it follows that the octanol-water partitioning coefficient may also be used to correlate the limiting first-order biodegradation rate constant since the observed biodegradation rate is based on bulk liquid concentrations. To that end, the limiting first-order rate constants for a variety of compounds were plotted versus their corresponding octanol-water partitioning coefficient. The results, presented in Figure 4-2, indicate a fair correlation between the octanol-water partitioning coefficients and the limiting first-order rate constants for most compounds. The primary discrepancies are for ionizable or polar compounds.

Assuming continuous, steady-state operation for a system that is well-mixed, a mass balance on the system can be written as follows:

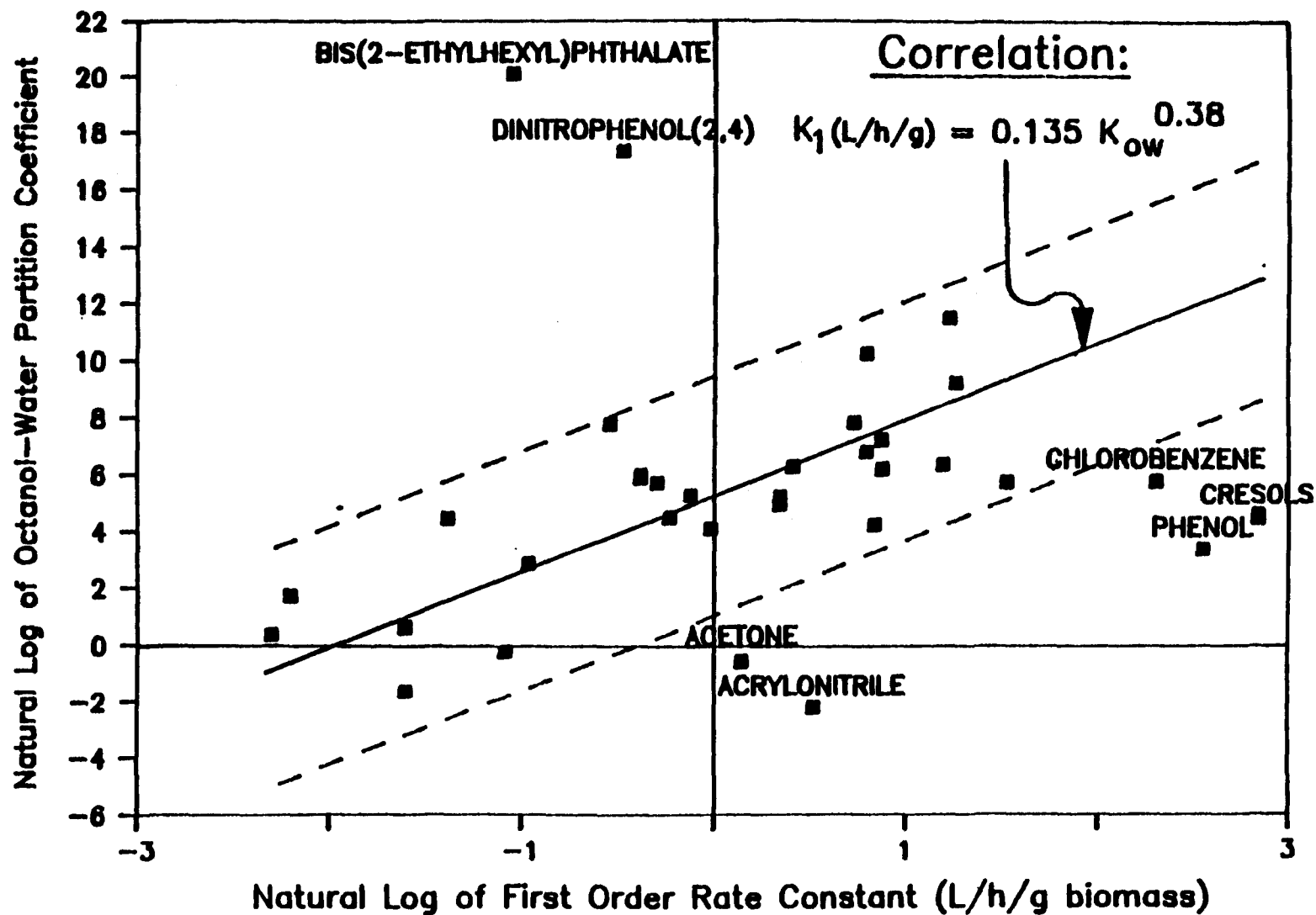
$$QC_0 = QC_L + VK_{\max}b_iC_L/(K_s+C_L) + K_{\text{other}} V C_L \quad (4-18)$$

where

Q = flow rate, m^3/s

K_{other} = sum of apparent first-order rate constants for competing mechanisms, $1/\text{s}$.

and the other symbols are as previously defined.



Note: Dashed lines represent a factor of 5 error.

Figure 4-2. Correlation of limiting first-order rate constant with the octanol water coefficient.

Note that Equation (4-18) was written in a general fashion so that, if desired, the rate of removal via adsorption onto biomass solids can be included. For most volatile organics, however, the adsorption pathway is negligible so that K_{other} is dominated by the volatilization rate. Consequently,

$$K_{other} = KA/V \quad (4-19)$$

where

K = overall mass transfer coefficient, m/s

A = area, m^2 .

To determine the fraction of VO emitted or biodegraded using the Monod model, one first has to solve for the effluent concentration. The effluent concentration can be determined by rearranging Equation (4-18) as follows:

$$K' C_L^2 + [K_S K' + (V/Q) K_{max} b_i - C_0] C_L - K_S C_0 = 0 \quad (4-20)$$

where

$K' = (K_{other})(V/Q) + 1$, dimensionless.

Equation (4-20) is easily solved using the quadratic formula as follows:

$$C_L = [-b + (b^2 - 4ac)^{0.5}] / 2a \quad (4-21)$$

where

$a = K' = (K_{other})(V/Q) + 1$

$b = K_S K' + (V/Q) K_{max} b_i - C_0$

$c = -K_S C_0$.

The plus sign is selected in Equation (4-21) to ensure positive effluent concentrations. Note that, because all of the rate constants and concentrations must have positive values, the constant, c , must be negative so that the quadratic equation always has real, positive roots.

Once the effluent concentration is calculated, the fraction of the component feed emitted to the air (f_{air}) is:

$$f_{air} = K A C_L / Q C_0 . \quad (4-22)$$

Emissions (E , g/s) are calculated from:

$$E = f_{air} Q C_0 . \quad (4-23)$$

Similarly, the fraction of the component feed biodegraded (f_{bio}) is:

$$f_{bio} = V K_{max} b_i / [(K_S + C_L) Q C_0] \quad (4-24)$$

If the biological system is operated with plug flow, the treated wastewater does not mix with the influent. The biodegradation rate and air emission rate change as the treatment progresses toward completion. For plug flow, the rate of disappearance of a compound by biodegradation and air emissions is given by:

$$\frac{-d C_t (V)}{dt} = \frac{V K_{max} b_i C_t}{(K_S + C_t)} + K A C \quad (4-25)$$

where

C_t = concentration at time = t

t = time, s

and with the other symbols as previously defined. Due to the nonlinear nature of the biodegradation rate term, Equation (4-25) cannot be directly integrated. Therefore, it is further assumed that first-order kinetics dominates the system's biodegradation. Equation (4-25) can then be rearranged as follows:

$$\frac{d C_t}{C_t} = (-K_1 b_i - K A / V) dt \quad (4-26)$$

where

$K_1 = K_{max} / K_S$, m^3/g biomass.

Integrating Equation (4-26) from $C_t = C_0$ at $t = 0$ to $C_t = C_e$ (effluent concentration) at $t = V/Q$ (one residence time) gives:

$$C_e/C_0 = \exp (-K_1 b_i V/Q - KA/Q) \quad (4-27)$$

The ratio C_e/C_0 represents the fraction leaving with the effluent; consequently, $1 - C_e/C_0$ represents the sum of the fractions that are biodegraded and emitted to the air. The fractions of component feed emitted to the air and biodegraded are calculated from their relative rates:

$$f_{air} = (1 - C_e/C_0)(KA)/(KA + K_1 b_i V) \quad (4-28)$$

$$f_{bio} = (1 - C_e/C_0)(K_1 b_i V)/(KA + K_1 b_i V) \quad (4-29)$$

The average emissions rate (E , g/s) is:

$$E = f_{air} QC_0 \quad (4-30)$$

4.3.3 Example Calculation for Quiescent Impoundments

The application of the biodegradation model to quiescent impoundments is presented in the form of an example calculation. The calculation is based on the quiescent impoundment's operating parameters from Table 4-2. For other types of impoundments, the application of the biodegradation model is illustrated in subsequent sections.

The waste stream for the example calculation is defined as containing benzene at 10 ppm with a total organic content of 250 ppm (0.25 g/L). The resultant organic loading on the impoundment on a daily basis is 12.8 kg/1,000 m³. The active biomass is assumed to be 0.05 g/L from a reported range from eight quiescent impoundments of 0.0014 to 0.22 g/L.

- a. Calculate the effluent concentration of benzene for a well-mixed system from Equation (4-21):

$$C_L = [-b + (b^2 - 4ac)^{0.5}] / 2a$$

where

$$a = K' = (KA/V) (V/Q) + 1 = KA/Q + 1$$

$$b = K_S K' + (V/Q) K_{max} b_i - C_0$$

$$\begin{aligned}
c &= -K_S C_0 \\
K &= 4.2 \times 10^{-6} \text{ m/s (Section 4.2.3, Step c)} \\
A &= 1,500 \text{ m}^2 \\
Q &= 0.00156 \text{ m}^3/\text{s} \\
K_{\max} &= 19 \text{ mg/g/L} = 5.28 \times 10^{-6} \text{ g/g/s (from Appendix D, Table D-1)} \\
K_S &= 13.6 \text{ mg/L} = 13.6 \text{ g/m}^3 \text{ (from Appendix D, Table D-1)} \\
b_i &= 0.05 \text{ g/L} = 50 \text{ g/m}^3 \\
V &= 2,700 \text{ m}^3 \\
C_0 &= 100 \text{ ppm} = 100 \text{ g/m}^3 \\
KA &= (4.2 \times 10^{-6} \text{ m/s})(1,500 \text{ m}^2) = 6.3 \times 10^{-3} \text{ m}^3/\text{s} \\
a = K' &= (6.3 \times 10^{-3} \text{ m}^3/\text{s}) / (0.00156 \text{ m}^3/\text{s}) + 1 = 5.0 \\
b &= (13.6 \text{ g/m}^3)(5.0) + (2,700 \text{ m}^3 / 0.00156 \text{ m}^3/\text{s})(5.28 \times 10^{-6} \text{ g/g/s})(50 \text{ g/m}^3) - (100 \text{ g/m}^3) \\
&= 425 \text{ g/m}^3 \\
c &= -(13.6 \text{ g/m}^3)(100 \text{ g/m}^3) = -1,360 \text{ g}^2/\text{m}^6 \\
C_L &= \{-[425 \text{ g/m}^3] + [(425 \text{ g/m}^3)^2 - 4(5.0)(-1,360 \text{ g}^2/\text{m}^6)]^{0.5}\} / [2(5.0)] \\
&= (-425 \text{ g/m}^3 + 517.6 \text{ g/m}^3) / 10 \\
&= 3.08 \text{ g/m}^3
\end{aligned}$$

- b. Calculate the fraction emitted for a well-mixed system from Equation (4-22):

$$f_{\text{air}} = KAC_L / (QC_0)$$

where

$$\begin{aligned}
f_{\text{air}} &= (6.3 \times 10^{-3} \text{ m}^3/\text{s})(3.08 \text{ g/m}^3) / [(0.00156 \text{ m}^3/\text{s})(100 \text{ g/m}^3)] \\
f_{\text{air}} &= 0.124
\end{aligned}$$

- c. Calculate benzene emissions for well-mixed system:

$$\begin{aligned}
E(\text{g/s}) &= f_{\text{air}} Q C_0 \\
&= (0.124)(0.00156 \text{ m}^3/\text{s})(100 \text{ g/m}^3) \\
&= 1.93 \times 10^{-2} \text{ g/s} = 0.61 \text{ Mg/yr}
\end{aligned}$$

- d. For a plug-flow system, calculate fraction removed with the effluent from Equation (4-27):

$$C_e/C_o = \exp (- K_1 b_i V/Q - KA/Q)$$

where

$$K_1 = 1.4 \text{ L/g-h} = 3.89 \times 10^{-7} \text{ m}^3/\text{g-s} \text{ (from Appendix D, Table D-1)}$$

$$b_i = 0.05 \text{ g/L} = 50 \text{ g/m}^3$$

$$V = 2,700 \text{ m}^3$$

$$Q = 0.00156 \text{ m}^3/\text{s}$$

$$C_o = 10 \text{ ppm} = 10 \text{ g/m}^3$$

$$K = 4.2 \times 10^{-6} \text{ m/s}$$

$$A = 1,500 \text{ m}^2$$

$$K_1 b_i V = (3.89 \times 10^{-7} \text{ m}^3/\text{s/g biomass})(50 \text{ g/m}^3)(2,700 \text{ m}^3) \\ = 5.25 \times 10^{-2} \text{ m}^3/\text{s}$$

$$KA = (4.2 \times 10^{-6} \text{ m/s})(1,500 \text{ m}^2) = 6.3 \times 10^{-3} \text{ m}^3/\text{s}$$

$$C_e/C_o = \exp \left[\frac{-5.25 \times 10^{-2} \text{ m}^3/\text{s}}{1.56 \times 10^{-3} \text{ m}^3/\text{s}} - \frac{6.3 \times 10^{-3} \text{ m}^3/\text{s}}{1.56 \times 10^{-3} \text{ m}^3/\text{s}} \right]$$

$$C_e/C_o = \exp (-37.7) = 0.00$$

e. Calculate fraction emitted from Equation (4-28):

$$f_{air} = (1 - C_e/C_o)(KA)/(KA + K_1 b_i V)$$

$$f_{air} = (1 - 0)(6.3 \times 10^{-3} \text{ m}^3/\text{s}) / (6.3 \times 10^{-3} \text{ m}^3/\text{s} + 5.25 \times 10^{-2} \text{ m}^3/\text{s})$$

$$f_{air} = 0.107$$

f. Calculate benzene emissions for plug flow:

$$E(\text{g/s}) = f_{air} Q C_o$$

$$= (0.107)(0.00156 \text{ m}^3/\text{s})(10 \text{ g/m}^3)$$

$$= 1.67 \times 10^{-2} \text{ g/s} = 0.53 \text{ Mg/yr}$$

4.4 MECHANICALLY AERATED IMPOUNDMENTS AND ACTIVATED SLUDGE UNITS

Some impoundments and tanks are mechanically agitated to improve mixing or to transfer air to the liquid (e.g., treatment tanks designed for biodegradation). The agitation creates a turbulent liquid surface that

enhances mass transfer to the air. A significant difference from the approach for quiescent surfaces discussed in Section 4.2 is the appropriate correlations for the individual mass transfer coefficients.

4.4.1 Emission Model Equations

The calculation of the overall mass transfer coefficient for mechanically aerated systems considers that the liquid surface is composed of two zones, quiescent and turbulent. The individual mass transfer coefficients for the turbulent zone are based on the correlations of Thibodeaux⁴⁰ and Reinhardt.⁴¹ Thibodeaux's model was developed from accepted interphase mass transfer concepts, published rate coefficient correlations, and existing operating data on 13 aerated basins at 11 pulp and paper mills. The basins represented a wide range of design and operating parameters, in spite of being from only one industry type. The simulation employed 11 organic chemical species common to industrial wastewater.

Reinhardt absorbed ammonia in aqueous sulfuric acid to measure the gas-phase mass transfer coefficient associated with flat-blade surface agitators in developing his correlation to calculate the gas-phase mass transfer coefficient.⁴²

Table 4-6 summarizes the correlations developed by Thibodeaux and Reinhardt. These correlations are used to estimate the individual mass transfer coefficients for the turbulent portion of the liquid surface. The individual coefficients are then used in Equation (4-2) to calculate an overall mass transfer coefficient for the turbulent zone. An overall mass transfer coefficient for the quiescent zone is calculated as described in Section 4.2. The two overall coefficients are combined to obtain a single coefficient for the system based on the relative areas of the turbulent and quiescent zones. For example, if 25 percent of the surface of the impoundment is turbulent, the overall coefficient would be the sum of 25 percent of the value for the turbulent area coefficient plus 75 percent of the value for the quiescent zone.

The model for mechanically aerated systems also incorporates biodegradation as a competing mechanism. The extent of biodegradation is difficult to predict in a generally applicable form because it is very dependent upon the constituent of interest, the waste matrix, the design and operation of

TABLE 4-6. EQUATIONS FOR CALCULATING INDIVIDUAL MASS TRANSFER COEFFICIENTS FOR VOLATILIZATION OF ORGANIC SOLUTES FROM TURBULENT SURFACE IMPOUNDMENTS

Liquid phase

Thibodeaux:43,44

$$k_L = [8.22 \times 10^{-9} J (\text{POWR})(1.024)^{t-20} O_t 10^6 \text{ MW}_L / (V a_v \rho_L)] (D_w / D_{O_2,w})^{0.5} \text{ (m/s)}$$

where

J = oxygen transfer rating of surface aerator, lb O₂/h•hp

POWR = total power to aerators, hp

T = water temperature, °C

O_t = oxygen transfer correction factor

MW_L = molecular weight of liquid

V = volume affected by aeration, ft³

a_v = surface-to-volume ratio of surface impoundment, ft⁻¹

ρ_L = density of liquid, g/cm³

D_w = diffusivity of constituent in water, cm²/s

D_{O₂,w} = diffusivity of oxygen in water = 2.4 x 10⁻⁵, cm²/s.

Gas phase

Reinhardt:45,46

$$k_G = 1.35 \times 10^{-7} R_e^{1.42} \rho^{0.4} S_{CG}^{0.5} .F_r^{-0.21} D_a \text{ MW}_a / d \text{ (m/s)}$$

where

R_e = d²wρ_a/μ_a = Reynold's number

d = impeller diameter, cm

w = rotational speed of impeller, rad/s

(continued)

TABLE 4-6 (continued)

ρ_a = density of air, g/cm³

μ_a = viscosity of air, g/cm•s

$$= 4.568 \times 10^{-7} T(^{\circ}\text{C}) + 1.7209 \times 10^{-4}$$

$p = P_I g_c / (\rho_L d^{*5} w^3) = \text{power number}$

P_I = power to impeller, ft•lb_f/s

$= 0.85 (\text{POWR}) (550 \text{ ft•lb}_f/\text{s•hp}) / \text{number of aerators,}$
where 0.85 = efficiency of aerator motor

g_c = gravitation constant, 32.17 lb_m•ft/s²/lb_f

ρ_L = density of liquid, lb/ft³

d^* = impeller diameter, ft

Sc_G = Schmidt number on gas side = $\mu_a / \rho_a D_a$

$Fr = d^* w^2 / g_c = \text{Froude number}$

D_a = diffusivity of constituent in air, cm²/s

MW_a = molecular weight of air.

the biodegradation unit, and the concentrations and properties of the microorganisms.

4.4.2 Model Plant Parameters for Mechanically Aerated Impoundments

The dimensions of the treatment impoundment used as an example to estimate emissions were derived from the Westat data as described in Section 4.2.2 for storage impoundment. A median area of 1,500 m² and a depth of 1.8 m were chosen, which yields a total volume of 2,700 m³. The retention time in treatment impoundments is expected to be less than the retention time in storage impoundments. Two design manuals listed typical retention times for aerated (biologically active) ponds as 7 to 20 days⁴⁷ and 3 to 10 days.⁴⁸ For the example case, a retention time of 10 days was chosen from the design range of 3 to 20 days. The resulting flow rate is 3.1 L/s (0.0031 m³/s).

The correlations of Thibodeaux and Reinhardt given in Table 4-3 require values for the parameters that describe the mechanical aeration system. Metcalf and Eddy, Inc.,⁴⁹ suggest a range of 0.5 to 1.0 hp/1,000 ft³ for mixing in an impoundment. However, more power may be needed to supply additional oxygen or to mix certain treatment solutions. A review of trip reports showed power usage as high as 3.5 hp/1,000 ft³ at a specific TSDf impoundment.⁵⁰ For this analysis, a midrange value of 0.75 hp/1,000 ft³ from Metcalf and Eddy was used to generate an estimate of 75 hp required for mixing in the model unit.

Data from Reference 51 indicated that five aerators with 15-hp motors and 61-cm diameter propellers turning at 126 rad/s would agitate a volume of 441 m³ (15,590 ft³). Assuming a uniform depth in the impoundment of 1.8 m, the agitated surface area was estimated as 245 m² (441/1.8). The agitated surface is assumed to be turbulent and comprises 16 percent (245/1,500 x 100) of the total area. The balance of the surface area of the impoundment (84 percent) is assumed to be quiescent. As a comparison, Thibodeaux reported a turbulent area of 5.22 m²/hp and investigated a range of 0.11 to 20.2 m²/hp. The value of 5.22 m²/hp and a total of 75 hp yields an estimated turbulent area of 392 m² (26 percent), which is neater than the 16-percent turbulent area calculated by the above procedure.⁵² (Very

few data are available on the distribution of turbulent areas for aerated impoundments. The extent of turbulence depends in part on the number size, and placement of aerators. The example is based on typical aerator requirements to mix the contents of the impoundment.)

Typical values were chosen for the oxygen transfer rating of the aerator and the oxygen transfer correction factor. A value of 3.0 lb O₂/hp/h was chosen for oxygen transfer rating from a range of 2.9 to 3.0.⁵³ A value of 0.83 was used for the correction factor from a typical range of 0.80 to 0.85.⁵⁴ The transfer of power to the impeller was assumed to be 85 percent efficient, yielding an estimate of 64 hp for the impeller power.

The model for biodegradation requires the system's biomass concentration as an input parameter. The concentration of biomass in real systems can be highly variable depending upon the system's design and method of operation. For this analysis, the specified biomass is assumed to be actively degrading the constituent of interest. A value of 250 g/m³ (0.25 g/L) of biomass was chosen from the values presented in Table 4-5.

The example constituent (benzene) and the meteorological conditions chosen for the example calculation are the same as those chosen for storage impoundments. Input parameters for the mechanically aerated model unit are summarized in Table 4-7.

4.4.3 Example Calculation for Mechanically Aerated Treatment Impoundments

The example calculation for emissions from a mechanically aerated impoundment includes an estimate of the overall mass transfer coefficient for the turbulent zone. The overall mass transfer coefficient for the quiescent zone for storage impoundments is calculated as illustrated in Section 4.2.3 and will not be repeated here. Biodegradation is included as a competing removal mechanism.

- a. Calculate turbulent liquid-phase mass transfer coefficient, k_L . Use Thibodeaux (Table 4-6):

$$k_L(\text{m/s}) = [8.22 \times 10^{-9} J(\text{POWER})(1.024)^{T-20} 0.1 \times 10^6 \text{ MWL}/(V_a \nu \rho_L)] \left(\frac{D_w}{D_{O_2, w}} \right)^{0.5}$$

**TABLE 4-7. INPUT PARAMETERS--TREATMENT IMPOUNDMENTS
(MECHANICALLY AERATED)**

Area: 1,500 m ²	Number of impellers: 5
Depth: 1.8 m	Total power: 75 hp
Volume: 2,700 m ³	Power to impeller: 13 hp
Retention time: 10 days	Impeller speed: 126 rad/s
Flow: 0.0031 m ³ /s	Impeller diameter: 61 cm
Turbulent area: 240 m ² (16%)	O ₂ transfer: 3 lb/h/hp
Quiescent area: 1,260 m ²	O ₂ correction factor: 0.83
Temperature: 25 °C	
Windspeed: 4.47 m/s	
Viscosity of air: 1.8 x 10 ⁻⁴ g/cm•s	
Density of air: 1.2 x 10 ⁻³ g/cm ³	
Diffusivity of O ₂ in water: 2.4 x 10 ⁻⁵ cm ² /s	
Density of liquid: 1 g/cm ³	
Molecular weight of liquid: 18 g/g•mol	
Molecular weight of air: 29 g/g•mol	
Constituent: benzene with other biodegradable organics in water	
Concentration (benzene): 100 g/m ³ (100 ppm)	
Concentration (total organics): 250 g/m ³ (250 ppm)	
Henry's law constant (benzene): 5.5 x 10 ⁻³ atm•m ³ /g mol	

(continued)

TABLE 4-7 (continued)

Diffusivity in air (benzene): $0.088 \text{ cm}^2/\text{s}$

Diffusivity in water (benzene): $9.8 \times 10^{-6} \text{ cm}^2/\text{s}$

Maximum biorate (benzene and other organics): 19 mg/h/g of biomass =
 $5.28 \times 10^{-6} \text{ g/g biomass}\cdot\text{s}$

Limiting first-order biorate constant: $1.4 \text{ L/h/g} = 3.89 \times 10^{-7} \text{ m}^3/\text{s/g}$
biomass

Biomass concentration: $0.3 \text{ g/L} = 300 \text{ g/m}^3$

where

J = O_2 transfer rating, use 3.0 lb O_2 /h·hp

POWER = 75 hp

T = water temperature = 25 °C

O_t = O_2 transfer correction factor, use 0.83

MW_L = molecular wt of liquid (water) = 18 g/g mol

$$(V_{aV}) = \text{agitated area in ft}^2 = 240.0 \text{ m}^2 \left(\frac{\text{ft}^2}{0.0929 \text{ m}^2} \right) = 2,583 \text{ ft}^2$$

ρ_L = water density = 1 g/cm³

D_w = 9.8×10^{-6} cm²/s

$D_{O_2,w}$ = 2.4×10^{-5} cm²/s

$$k_L = (8.22 \times 10^{-9})(3)(75)(1.024)^5 \left[\frac{(0.83)(10^6)(18)}{(2,583)(1)} \right] \left[\frac{9.8 \times 10^{-6}}{2.4 \times 10^{-5}} \right]^{0.5}$$
$$= 7.7 \times 10^{-3} \text{ m/s}$$

- b. Calculate turbulent gas-phase mass transfer coefficient, k_G . Use Reinhardt (see Table 4-6):

$$k_G(\text{m/s}) = 1.35 \times 10^{-7} \text{ Re}^{1.42} p^{0.4} Sc_G^{0.5} Fr^{-0.21} D_a \text{ MW}_a/d$$

where

$$\text{Re} = \text{Reynold's number} = \frac{d^2 w \rho_a}{\mu_a}$$

d = impeller diameter = 61 cm

w = impeller speed = 126 rad/s

ρ_a = 1.2×10^{-3} g/cm³

μ_a = 1.81×10^{-4} g/cm·s

$$Re = \frac{(612) (126) (1.2 \times 10^{-3})}{1.81 \times 10^{-4}} = 3.1 \times 10^6$$

$$p = \text{power number} = \frac{P_I g_c}{\rho_L d^5 w^3}$$

$$P_I = 12.8 \text{ hp} \frac{550 \text{ ft lbf}}{\text{s} \cdot \text{hp}} = 7,040$$

$$g_c = 32.17 \frac{\text{lb} \cdot \text{ft}}{\text{s}^2 \text{ lbf}}$$

$$\rho_L = 62.37 \text{ lb/ft}^3$$

$$d^* = \text{impeller diameter in feet} = 2.0$$

$$w = 126 \text{ rad/s}$$

$$p = \frac{(7,040) (32.17)}{(62.37) (2)^5 (126)^3} = 5.6 \times 10^{-5}$$

$$Sc_G = 1.71 \text{ (from Section 4.2.3, part b)}$$

$$Fr = \text{Froude number} = \frac{d^* w^2}{g_c} = \frac{(2) (126)^2}{32.17} = 9.9 \times 10^2$$

$$D_a = 0.088 \text{ cm}^2/\text{s} \text{ (benzene)}$$

$$MW_a = 29 \text{ g/g mol}$$

$$d = \text{impeller diameter in cm} = 61 \text{ cm}$$

$$k_G = (1.35 \times 10^{-7}) (3.1 \times 10^6)^{1.42} (5.6 \times 10^{-5})^{0.4} (1.71)^{0.5} (9.9 \times 10^2)^{-0.21} (0.088) (29) / 61$$

$$k_G = 5.7 \times 10^{-2} \text{ m/s}$$

- c. Calculate overall mass transfer coefficient for turbulent area, K:

$$\frac{1}{K} = \frac{1}{k_L} + \frac{1}{K_{eq} k_G} = \frac{1}{7.7 \times 10^{-3}} + \frac{1}{(0.225)(5.7 \times 10^{-2})} = 2.4 \times 10^2$$

$$K = 7.71 \times 10^{-4} \text{ m/s}$$

- d. Calculate overall mass transfer coefficient for combined quiescent and turbulent areas, K:

From Section 4.2.3, K for quiescent area = $4.2 \times 10^{-6} \text{ m/s}$

From Part C, K for turbulent area = $7.7 \times 10^{-4} \text{ m/s}$

Turbulent area = 240 m^2

Quiescent area = $1,260 \text{ m}^2$

$$K \text{ (m/s)} = \frac{(4.2 \times 10^{-6})(1,260) + (7.7 \times 10^{-4})(240)}{(1,260 + 240)} = 7.7 \times 10^{-4} \text{ m/s}$$

(weighted by area)

- e. Calculate the effluent concentration for benzene for a well-mixed system from Equation (4-21):

$$C_L = [-b + (b^2 - 4ac)^{0.5}] / 2a$$

where

$$a = K' = (KA/V) (V/Q) + 1 = KA/Q + 1$$

$$b = K_S K' + (V/Q) K_{\max} b_i - C_0$$

$$c = -K_S C_0$$

$$K = 7.7 \times 10^{-4} \text{ m/s}$$

$$A = 1,500 \text{ m}^2$$

$$Q = 0.0031 \text{ m}^3/\text{s}$$

$$K_{\max} = 5.28 \times 10^{-6} \text{ g/s/g biomass}$$

$$b_i = 0.3 \text{ g/L} = 300 \text{ g/m}^3$$

$$V = 2,700 \text{ m}^3$$

$$C_o = 100 \text{ ppm} = 100 \text{ g/m}^3$$

$$K_s = K_{\max}/K_1 = (5.28 \times 10^{-6} \text{ g/s/g}) / (3.89 \times 10^{-7} \text{ m}^3/\text{s/g}) \\ = 13.6 \text{ g/m}^3$$

$$K_A = (7.7 \times 10^{-4} \text{ m/s})(1,500 \text{ m}^2) = 1.15 \text{ m}^3/\text{s}$$

$$a = K' = (1.15 \text{ m}^3/\text{s}) / (0.0031 \text{ m}^3/\text{s}) + 1 = 373$$

$$b = (13.6 \text{ g/m}^3)(373) + (2,700 \text{ m}^3 / 0.0031 \text{ m}^3/\text{s})(5.28 \times 10^{-6} \text{ g/s/g})(300 \text{ g/m}^3) - (100 \text{ g/m}^3) \\ = 7,965.6 \text{ g/m}^3$$

$$c = -(13.6 \text{ g/m}^3)(100 \text{ g/m}^3) = -1,360 \text{ g}^2/\text{m}^6$$

$$C_L = \{-[7,965.6 \text{ g/m}^3] + [(7,965.6 \text{ g/m}^3)^2 \\ - 4(373)(-136 \text{ g}^2/\text{m}^6)]^{0.5}\} / [2(373)] \\ = (-7,965.6 \text{ g/m}^3 + 7,982.2 \text{ g/m}^3) / 970 \\ = 0.22 \text{ g/m}^3$$

- f. Calculate the fraction emitted for a well-mixed system from Equation (4-22):

$$f_{\text{air}} = K A C_L / (Q C_o)$$

where

$$f_{\text{air}} = (1.15 \text{ m}^3/\text{s})(0.22 \text{ g/m}^3) / [(0.0031 \text{ m}^3/\text{s})(100 \text{ g/m}^3)] \\ f_{\text{air}} = 0.816$$

- g. Calculate benzene emissions for well-mixed system:

$$E(\text{g/s}) = f_{\text{air}} Q C_o \\ = (0.816)(0.0031 \text{ m}^3/\text{s})(100 \text{ g/m}^3) \\ = 2.5 \times 10^{-2} \text{ g/s} = 8.0 \text{ Mg/yr}$$

- h. For a plug-flow system, calculate the fraction removed with the effluent from Equation (4-27):

$$C_e/C_o = \exp(-K_1 b_i V/Q - K_A/Q) \\ K_1 = 3.89 \times 10^{-7} \text{ m}^3/\text{s/g biomass} \\ b_i = 0.3 \text{ g/L} = 300 \text{ g/m}^3 \\ V = 2,700 \text{ m}^3 \\ Q = 0.0031 \text{ m}^3/\text{s}$$

$$C_o = 100 \text{ ppm} = 100 \text{ g/m}^3$$

$$K = 1.0 \times 10^{-3} \text{ m/s}$$

$$A = 1,500 \text{ m}^2$$

$$K_1 b_1 V = (3.89 \times 10^{-7} \text{ m}^3/\text{s/g biomass})(300 \text{ g/m}^3)(2,700 \text{ m}^3) \\ = 0.315 \text{ m}^3/\text{s}$$

$$KA = (7.7 \times 10^{-4} \text{ m/s})(1,500 \text{ m}^2) = 1.15 \text{ m}^3/\text{s}$$

$$C_e/C_o = \exp \frac{-0.315 \text{ g/s}}{0.0031 \text{ m}^3/\text{s}} - \frac{1.15 \text{ m}^3/\text{s}}{0.0031 \text{ m}^3/\text{s}} = 0$$

- i. Calculate fraction emitted from Equation (4-28):

$$f_{air} = (1 - C_e/C_o)(KA)/(KA + K_1 b_1 V)$$

$$f_{air} = (1 - 0)(1.15 \text{ m}^3/\text{s})/(1.15 \text{ m}^3/\text{s} + 0.315 \text{ m}^3/\text{s})$$

$$f_{air} = 0.78$$

- j. Calculate benzene emissions for plug flow:

$$E(\text{g/s}) = f_{air} Q C_o \\ = (0.78)(0.0031 \text{ m}^3/\text{s})(100 \text{ g/m}^3) \\ = 0.24 \text{ g/s} = 7.7 \text{ Mg/yr}$$

4.4.4 Example Calculation for Activated Sludge Unit

As discussed in Section 4.2, an activated sludge unit usually consists of a concrete tank that is aerated and contains a relatively high concentration of active biomass. A model unit is defined in this section for this process, and the results of intermediate and final calculations are given. Detailed example calculations are not presented because the approach is exactly the same as that used for the mechanically aerated impoundment. The only significant difference in the method of operation is the recycle of solids back to the activated sludge unit, which results in a higher biomass concentration. For this model unit, a biomass concentration of 4 g/L (4,000 g/m³) was chosen from the range of 1.5 to 6 g/L in Table 4-3 and the recommended values in Table 4-5. Other differences between the aerated impoundment and activated sludge tank include, for the tank, a smaller surface area, a shorter retention time, a greater turbulent area, and a smaller F/D ratio. The aerated surface area was estimated as

described in Section 4.4.2. An aerator with a 7.5-hp motor will agitate a volume of 56.9 m^3 ($2,010 \text{ ft}^3$). For a uniform depth of 4 m, the agitated volume yields an agitated surface area of 14.2 m^2 ($56.9 \text{ m}^3/4 \text{ m}$). The input parameters are defined for this model unit in Table 4-8, and the results of the calculations are presented in Table 4-9.

4.5 DISPOSAL IMPOUNDMENTS WITH QUIESCENT SURFACES

4.5.1 Emission Model Equations

A disposal impoundment is defined as a unit that receives a waste for ultimate disposal rather than for storage or treatment. This type of impoundment differs from the storage and treatment impoundments in that there is no liquid flow out of the impoundment (seepage into the ground is neglected). For this case, the well-mixed system with a bulk concentration that is at equilibrium (i.e., the bulk concentration does not change with time) is not applicable. The quantity of a constituent in a disposal impoundment will decrease with time after the waste is placed in the impoundment because of the loss of volatiles to the air.

The calculation of the overall mass transfer coefficient is the same as that presented for impoundments with quiescent surfaces. If the disposal impoundment is aerated, K is calculated as described for aerated impoundments in Section 4.4. The emission estimating procedure differs in the calculation of the liquid-phase concentration that is the driving force for mass transfer to the air. For a disposal impoundment that is filled with a batch of waste, the rate of disappearance of a compound by biodegradation and air emissions is described by Equation (4-26).

Integrating Equation (4-26) from $C_t = C_0$ at $t = 0$ to $C_t = C_t$ at $t = t$ gives:

$$C_t/C_0 = \exp (-K_1 b_i t - KAt/V) \quad . \quad (4-31)$$

For an impoundment with a uniform depth, $V/A = D$. Substituting $V/A = D$ into Equation (4-31) yields:

$$C_t/C_0 = \exp (-K_1 b_i t - Kt/D) \quad . \quad (4-32)$$

TABLE 4-8. INPUT PARAMETERS--MECHANICALLY AERATED
ACTIVATED SLUDGE UNIT

Area: 27 m²
 Depth: 4 m
 Volume: 108 m³
 Retention time: 4 h
 Flow: 0.0075 m³/s
 Turbulent area: 19 m² (70%)
 Quiescent area: 8.0 m²

Total power: 7.5 hp
 Power to impeller: 6.4 hp
 Impeller speed: 126 rad/s
 Impeller diameter: 61 cm
 O₂ transfer: 3 lb/h/hp
 O₂ correction factor: 0.83

Temperature: 25 °C
 Windspeed: 4.47 m/s

Viscosity of air: 1.8×10^{-4} g/cm·s
 Viscosity of water: 9×10^{-3} g/cm·s
 Density of air: 1.2×10^{-3} g/cm³
 Diffusivity of O₂ in water: 2.4×10^{-5} cm²/s
 Density of liquid: 1 g/cm³
 Molecular weight of liquid: 18 g/g·mol
 Molecular weight of air: 29 g/g·mol

Constituent: benzene with other biodegradable organics in water
 Concentration (benzene): 10 g/m³ (10 ppm)
 Concentration (total organics): 250 g/m³ (250 ppm)

Henry's law constant (benzene): 5.5×10^{-3} atm·m³/g·mol
 Diffusivity in air (benzene): 0.088 cm²/s
 Diffusivity in water (benzene): 9.8×10^{-6} cm²/s
 Maximum biorate (benzene and other organics): 5.28×10^{-6} g/s/g biomass
 Limiting first-order biorate constant = 3.89×10^{-7} m³/s/g biomass
 Biomass concentration: 4.0 g/L = 4,000 g/m³

TABLE 4-9. INTERMEDIATE AND FINAL CALCULATION RESULTS
FOR ACTIVATED SLUDGE MODEL UNIT

Quiescent zone:

$$k_L = 6.5 \times 10^{-6} \text{ m/s}$$

$$k_g = 8.9 \times 10^{-3} \text{ m/s}$$

$$K = 6.5 \times 10^{-6} \text{ m/s}$$

Turbulent zone:

$$k_L = 9.7 \times 10^{-2} \text{ m/s}$$

$$k_g = 4.3 \times 10^{-2} \text{ m/s}$$

$$K = 4.88 \times 10^{-3} \text{ m/s}$$

$$\text{Overall mass transfer coefficient} = 3.4 \times 10^{-3} \text{ m/s}$$

For well-mixed system:

$$C_L = 3.17$$

$$f_{air} = 0.391$$

$$\text{Emissions} = 0.30 \text{ g/s} = 9.3 \text{ Mg/yr}$$

For plug-flow system:

$$f_{air} = 0.391$$

$$\text{Emissions} = 0.30 \text{ g/s} = 9.3 \text{ Mg/yr}$$

When Equation (4-32) is evaluated after some fixed time t , the ratio C_t/C_0 represents the fraction of the compound remaining in the impoundment; consequently, $1 - C_t/C_0$ represents the fraction that has been removed by biodegradation and air emissions. The fractions emitted to the air and biodegraded after some time (t) are calculated from their relative rates:

$$f_{\text{air}} = (1 - C_t/C_0)(KA)/(KA + K_1b_iV) \quad (4-33)$$

$$f_{\text{bio}} = (1 - C_t/C_0)(K_1b_iV)/(KA + K_1b_iV) \quad (4-34)$$

The quantity emitted after some time (t) is given by:

$$\text{Emitted quantity (g)} = f_{\text{air}} V C_0 \quad (4-35)$$

The average emission rate over the period of time = t is:

$$E \text{ (g/s)} = f_{\text{air}} V C_0/t \quad (4-36)$$

Alternatively, a simplifying assumption may be made that, because the impoundment is designed for disposal, all significantly volatile compounds are eventually emitted to the air. Emissions under this assumption would simply be QC_0 where Q equals the disposal rate in cubic meters/second. This assumption is probably valid for volatile compounds; however, compounds that are relatively nonvolatile may be removed slowly and the assumption may result in an overestimate of emissions.

4.5.2 Model Plant Parameters for Disposal Impoundments

The Westat data summary for impoundments indicated that disposal impoundments generally have higher surface areas and shallower depths than storage and treatment impoundments. The median surface area for disposal impoundments was approximately 9,000 m² (compared to 1,500 m² for storage impoundments), and the median depth was approximately 1.8 m. The disposal impoundment is assumed to be filled with waste every 6 mo (two turnovers per year).

The meteorological conditions and type of waste (water containing benzene and other organics for the example calculation are the same as those used for quiescent and aerated impoundments with biodegradation. The inputs for the example calculation of emissions from disposal impoundments are summarized in Table 4-10.

TABLE 4-10. INPUT PARAMETERS--DISPOSAL IMPOUNDMENTS

Area: 9,000 m²

Depth: 1.8 m

Volume: 16,200 m³

Turnovers per year: 2

Temperature: 25 °C

Windspeed: 4.47 m/s

Diffusivity in water (ether): 8.5 x 10⁻⁶ cm²/s

Viscosity of air: 1.81 x 10⁻⁴ g/cm•s

Density of air: 1.2 x 10⁻³ g/cm³

Constituent: benzene with other biodegradable organics in water

Concentration (benzene): 100 g/m³ (100 ppm)

Concentration (total organics): 250 g/m³ (250 ppm)

Henry's law constant (benzene): 5.5 x 10⁻³ atm•m³/g mol

Diffusivity in air (benzene): 0.088 cm²/s

Diffusivity in water (benzene): 9.8 x 10⁻⁶ cm²/s

Limiting first-order biorate constant: 3.89 x 10⁻⁶ m³/s/g biomass

Biomass concentration: 0.05 g/L = 50 g/m³

4.5.3 Example Calculations for Disposal Impoundments

Example calculations are presented below for the model unit defined to represent disposal impoundments.

- a. Calculate liquid-phase mass transfer coefficient, k_L . Use Springer's model (see Table 4-1):

$$\text{Effective diameter} = \left(\frac{\text{Area}}{\pi} \right)^{0.5} \times 2 = \left(\frac{9,000}{\pi} \right)^{0.5} \times 2 = 107 \text{ m}$$

$$F/D = \text{Effective diameter/depth} = \frac{107}{1.8} = 59.5$$

$$\text{Windspeed} = 4.47 \text{ m/s} \quad (U_{10} \geq 3.25 \text{ m/s})$$

$$F/D = 59.5$$

$$k_L = 2.611 \times 10^{-7} U_{10}^2 \left(\frac{D_w}{D_{\text{ether}}} \right)^{0.67} \text{ m/s}$$

where

$$U_{10} = \text{windspeed} = 4.47 \text{ m/s}$$

$$D_w = 9.8 \times 10^{-6} \text{ cm}^2/\text{s} \text{ (benzene)}$$

$$D_{\text{ether}} = 8.5 \times 10^{-6} \text{ cm}^2/\text{s} \text{ (ether)}$$

Then

$$k_L = 2.611 \times 10^{-7} (4.47)^2 \left(\frac{9.8 \times 10^{-6}}{8.5 \times 10^{-6}} \right)^{0.67}$$

$$k_L = 5.7 \times 10^{-6} \text{ m/s}$$

- b. Calculate gas-phase mass transfer coefficient, k_g . Use MacKay and Matasugu (see Table 4-1):

$$k_g = 4.82 \times 10^{-3} U^{0.78} Sc_G^{-0.67} d_e^{-0.11} \text{ (m/s)}$$

where

$$U = \text{windspeed} = 4.47 \text{ m/s}$$

$$Sc_G = \text{Schmidt No. for gas} = \frac{\text{viscosity of gas}}{(\text{gas density})(\text{diffusivity of } i \text{ in gas})}$$

$$\begin{aligned} \text{Gas} &= \text{air} \\ \text{Viscosity (air)} &= 1.81 \times 10^{-4} \text{ g/cm}\cdot\text{s} \\ \text{Density (air)} &= 1.2 \times 10^{-3} \text{ g/cm}^3 \\ \text{Diffusivity (benzene in air)} &= 0.088 \text{ cm}^2/\text{s} \end{aligned}$$

$$Sc_G = \frac{1.81 \times 10^{-4} \text{ g/cm}\cdot\text{s}}{(1.2 \times 10^{-3} \text{ g/cm}^3)(0.088 \text{ cm}^2/\text{s})} = 1.71$$

$$d_e = \text{effective diameter} = 107 \text{ m}$$

Then

$$\begin{aligned} k_G &= (4.82 \times 10^{-3}) (4.47)^{0.78} (1.71)^{-0.67} (107)^{-0.11} \\ &= 6.5 \times 10^{-3} \text{ m/s} \end{aligned}$$

c. Calculate overall mass transfer coefficient, K:

$$\frac{1}{K} = \frac{1}{k_L} + \frac{1}{K_{eq} k_G}$$

where

$$K_{eq} = \frac{H}{RT} = \frac{5.5 \times 10^{-3} \text{ m}^3 \cdot \text{atm/mol}}{(8.21 \times 10^{-5}) \left(\frac{\text{atm} \cdot \text{m}^3}{\text{mol K}} \right) (298 \text{ K})} = 0.225$$

Then

$$\frac{1}{K} = \frac{1}{5.7 \times 10^{-6}} + \frac{1}{(0.225)(6.5 \times 10^{-3})} = 1.76 \times 10^5$$

$$K = 5.7 \times 10^{-6} \text{ m/s}$$

d. Calculate the fraction remaining from Equation (4-32). The impoundment is filled with waste initially, and 6 mo later it will be filled again. Calculate the fraction remaining after the initial 6-mo period:

$$C_t/C_0 = \exp (-K_1 b_i t - Kt/D)$$

$$K_1 = 3.89 \times 10^{-7} \text{ m}^3/\text{s/g biomass}$$

$$b_i = 50 \text{ g/m}^3$$

$$t = 6 \text{ mo} = 1.58 \times 10^7 \text{ s}$$

$$C_0 = 100 \text{ g/m}^3$$

$$K = 5.7 \times 10^{-6} \text{ m/s}$$

$$D = 1.8 \text{ m}$$

$$K_1 b_i t = (3.89 \times 10^{-7} \text{ m}^3/\text{s/g biomass})(50 \text{ g/m}^3)(1.58 \times 10^7 \text{ s}) \\ = 307$$

$$Kt/D = (5.7 \times 10^{-6} \text{ m/s})(1.58 \times 10^7 \text{ s}) / 1.8 \text{ m} = 50.0$$

$$C_t/C_0 = \exp(-307 - 50) = 0$$

f. Calculate the fraction emitted from Equation (4-33):

$$f_{\text{air}} = (1 - C_t/C_0)(KA) / (KA + K_1 b_i V)$$

$$C_t/C_0 = 0$$

$$KA = (5.7 \times 10^{-6} \text{ m/s})(9,000 \text{ m}^2) = 0.051 \text{ m}^3/\text{s}$$

$$K_1 b_i V = (3.89 \times 10^{-7} \text{ m}^3/\text{s/g biomass})(50 \text{ g/m}^3)(16,200 \text{ m}^3) \\ = 0.315 \text{ m}^3/\text{s}$$

$$f_{\text{air}} = (1 - 0)(0.051 \text{ m}^3/\text{s}) / (0.051 \text{ m}^3/\text{s} + 0.315 \text{ m}^3/\text{s})$$

$$f_{\text{air}} = 0.14$$

g. Calculate the average emission rate over the 6-mo period from Equation (4-36):

$$E \text{ (g/s)} = f_{\text{air}} V C_0/t \\ = (0.14)(16,200 \text{ m}^3)(100 \text{ g/m}^3)/1.58 \times 10^7 \text{ s} \\ = 1.4 \times 10^{-2} \text{ g/s.}$$

4.6 DIFFUSED AIR SYSTEMS

4.6.1 Emission Model Equations

Some impoundments and open tanks (e.g., activated sludge units) are sparged with air to promote biodegradation or air stripping. To estimate emissions from diffused air systems, the model assumes that the air bubbling through the liquid phase reaches equilibrium with the liquid-phase concentration of the constituent. The emissions leaving with the diffused air are estimated by:

$$E = Q_a K_{eq} C_L \quad (4-37)$$

where

E = emissions, g/s

Q_a = air flow rate, m^3/s

K_{eq} = equilibrium constant

C_L = concentration in the liquid phase, g/m^3 .

Emissions can also occur from wind blowing across the surface. If the air sparging creates a very turbulent surface similar to the surface of mechanically aerated systems, then the emission rate should be based on values of K typical for mechanically aerated systems. If the air sparging rate does not result in a turbulent surface, then K can be estimated from the correlations given for quiescent surfaces in Section 4.2.

The approach to estimate total emissions for flowthrough tanks and impoundments sparged with diffused air is similar to that described for quiescent and aerated systems. Because the unit is sparged with air, the liquid phase is assumed to be well mixed and the plug-flow model is not used. A material balance around this well-mixed system is identical to Equation (4-18) in Section 4.3.2, but now K_{other} is:

$$K_{other} = (KA + Q_a K_{eq})/V \quad (4-38)$$

where all of the symbols have been previously defined. The steady-state liquid phase concentration (C_L) is then calculated using Equation (4-21). Air emissions are estimated as the sum from wind blowing across the surface and from the diffused air:

$$E = KAC_L + Q_a K_{eq} C_L \quad (4-39)$$

The fraction of the component feed emitted to the air (f_{air}) is:

$$f_{air} = (KC_L A + Q_a K_{eq} C_L)/Q_{Co} \quad (4-40)$$

For disposal impoundments with diffused air systems, the steady-state assumptions of the flowthrough models do not apply. Emissions are greatest

when the waste is first placed in the impoundment and gradually decrease with time. To incorporate the contribution to mass transfer from diffused air, an equivalent mass transfer coefficient is defined:

$$K_D = K_{eq}Q_a/A \quad (4-41)$$

where

K_D = equivalent mass transfer coefficient for diffused air, m/s

and all of the other symbols are as previously defined.

The mass transfer coefficient for wind blowing across the surface (K) is calculated as described previously for flowthrough systems. A combined overall mass transfer coefficient (K_C) is defined as:

$$K_C = K_D + K \quad (4-42)$$

The overall mass transfer coefficient (K_C) is used in the equations for disposal impoundments (Section 4.5.1) to estimate the fraction emitted (Equation 4-33) and the average emission rate (Equation 4-36). The combined overall mass transfer coefficient defined above includes the mass transfer effects from both removal mechanisms (wind and diffused air).

4.6.2 Model Unit Parameters for Activated Sludge Unit with Diffused Air

A model unit for the activated sludge process was defined in Section 4.4.4 and Table 4-8. The same dimensions are used here to define an activated sludge unit that uses diffused air instead of mechanical aeration. The only additional parameter that must be specified is the diffused air rate, which typically ranges from 0.3 to 0.5 m³/s per 1,000 m³ of volume (20 to 30 ft³/min per 1,000 ft³ of volume).⁵⁵ For the model unit with a volume of 108 m³, an estimate of 0.04 m³/s is recommended based on the midpoint of the design range. The model unit input parameters are summarized in Table 4-11.

4.6.3 Example Calculation for Diffused Air Activated Sludge Unit

An example calculation is presented below for the model unit defined in Table 4-11.

- a. Calculate the liquid-phase, gas-phase, and overall mass transfer coefficients. This procedure was illustrated for quiescent surfaces and the results for this model unit are given in Table 4-9:

TABLE 4-11. INPUT PARAMETERS--DIFFUSED AIR ACTIVATED SLUDGE UNIT

Area:	27 m ²
Depth:	4 m
Volume:	108 m ³
Retention time:	4 h
Flow:	0.0075 m ³ /s
Quiescent area:	8.0 m ²
Diffused air rate:	0.04 m ³ /s
Temperature:	25 °C
Windspeed:	4.47 m/s
Viscosity of air:	1.81 x 10 ⁻⁴ g/cm•s
Density of air:	1.2 x 10 ⁻³ g/cm ³
Diffusivity of O ₂ in water:	2.4 x 10 ⁻⁵ cm ² /s
Density of liquid:	1 g/cm ³
Molecular weight of liquid:	18 g/g•mol
Molecular weight of air:	29 g/g•mol
Constituent:	benzene with other biodegradable organics in water
Concentration (benzene):	100 g/m ³ (100 ppm)
Concentration (total organics):	250 g/m ³ (250 ppm)
Henry's law constant (benzene):	5.5 x 10 ⁻³ atm•m ³ /g•mol
Diffusivity in air (benzene):	0.088 cm ² /s
Diffusivity in water (benzene):	9.8 x 10 ⁻⁶ cm ² /s
Maximum biorate (benzene and other organics):	5.28 x 10 ⁻⁶ g/s/g biomass
Limiting first-order biorate constant:	3.89 x 10 ⁻⁷ m ³ /s/g biomass
Biomass concentration:	4.0 g/L = 4,000 g/m ³

$$k_L = 6.5 \times 10^{-6} \text{ m/s}$$

$$k_g = 8.7 \times 10^{-3} \text{ m/s}$$

$$K = 3.42 \times 10^{-3} \text{ m/s (weighted by area).}$$

- b. Calculate the equilibrium constant, K_{eq} . The compound is benzene in water, and K_{eq} has been presented as 0.225 in the previous sample calculations (from Equation 4-4).
- c. Calculate the equilibrium liquid concentration in the unit (C_L) from Equation (4-21):

$$Q = 0.0075 \text{ m}^3/\text{s}$$

$$C_0 = 100 \text{ g/m}^3$$

$$K = 3.42 \times 10^{-3} \text{ m/s}$$

$$A = 27 \text{ m}^2$$

$$Q_a = 0.04 \text{ m}^3/\text{s}$$

$$K_{eq} = 0.225$$

$$K_{max} = 5.28 \times 10^{-6} \text{ g/s/g biomass}$$

$$K_s = K_{max}/K_1 = (5.28 \times 10^{-6} \text{ g/s/g}) / (3.89 \times 10^{-7} \text{ m}^3/\text{s/g})$$

$$= 13.6 \text{ g/m}^3$$

$$b_i = 4,000 \text{ g/m}^3$$

$$V = 108 \text{ m}^3$$

$$Q_{C_0} = (0.0075 \text{ m}^3/\text{s})(100 \text{ g/m}^3) = 0.75 \text{ g/s}$$

$$KA = (3.42 \times 10^{-3} \text{ m/s})(27 \text{ m}^2) = 9.23 \times 10^{-2} \text{ m}^3/\text{s}$$

$$Q_a K_{eq} = (0.04 \text{ m}^3/\text{s})(0.225) = 9.0 \times 10^{-3} \text{ m}^3/\text{s}$$

$$K_{other} = (KA + Q_a K_{eq})/V \text{ (from Equation 4-38)}$$

$$= [(9.23 \times 10^{-2} \text{ m}^3/\text{s}) + (9.0 \times 10^{-3} \text{ m}^3/\text{s})] / (108 \text{ m}^3)$$

$$= 9.39 \times 10^{-4} \text{ 1/s}$$

$$V/Q = (108 \text{ m}^3) / (0.0075 \text{ m}^3/\text{s}) = 14,400 \text{ s}$$

$$a = K' = (9.39 \times 10^{-4} \text{ 1/s})(14,400 \text{ s}) + 1 = 14.5$$

$$b = (13.6 \text{ g/m}^3)(14.5) + (14,400 \text{ s})(5.28 \times 10^{-6} \text{ g/s/g})$$

$$(4,000 \text{ g/m}^3) - 100 \text{ g/m}^3$$

$$= 40.1 \text{ g/m}^3$$

$$c = -(13.6 \text{ g/m}^3)(100 \text{ g/m}^3) = -1,360 \text{ g}^2/\text{m}^6$$

$$C_L = \{ [-401 \text{ g/m}^3] + [(401 \text{ g/m}^3)^2 - 4(14.5)$$

$$(-1,360 \text{ g}^2/\text{m}^6)]^{0.5} \} / [2(14.5)]$$

$$= [(-401 \text{ g/m}^3) + (490 \text{ g/m}^3)] / 29$$

$$= 3.06 \text{ g/m}^3$$

d. Calculate air emissions from Equation (4-39).

$$\begin{aligned} E &= (9.23 \times 10^{-2} \text{ m}^3/\text{s})(3.06 \text{ g/m}^3) + (9.0 \times 10^{-3} \text{ m}^3/\text{s})(3.06 \text{ g/m}^3) \\ &= 0.31 \text{ g/s} = 9.7 \text{ Mg/yr} \end{aligned}$$

4.7 OIL FILM SURFACES

Some impoundments may have a floating film of oil on the surface. A rigorous approach to estimating emissions from this type of source would consider three resistances acting in series:

- From the aqueous phase to the oil
- Through the oil
- From the oil to the air.

Such an approach would require estimates of these three resistances and estimates of the equilibrium partitioning between both the aqueous and oil phases and the oil and air phases. Because these estimates are not generally available, a simplifying assumption is that the oil film is relatively thin and that mass transfer is controlled by the gas-phase resistance. For this case, Equation (4-2) reduces to:

$$K = k_g K_{eq} \quad (4-43)$$

where k_g is calculated from the correlation of MacKay and Matasugu (Table 4-1) and K_{eq} is calculated from Raoult's law by:

$$K_{eq} = P^* \rho_a MW_{oil} / (\rho_L MW_a P_o) \quad (4-44)$$

where

K_{eq} = dimensionless equilibrium constant

P^* = vapor pressure of the volatile compound of interest, atm

P_o = total pressure, 1 atm

ρ_a = density of air, g/cm³

ρ_L = density of oil, g/cm³

MW_{oil} = molecular weight of oil, g/g mol

MW_a = molecular weight of air, 28.8 g/g mol.

The value of K calculated above is substituted into the equations for flow-through systems to estimate emissions. For the well-mixed flow models, C_0 and C_L in Equations (4-1) and (4-6) represent the V_0 concentration in the oil phase (entering and leaving the impoundment, respectively), and the flowrate Q is the volumetric flow rate of oil. Biodegradation is neglected because the oil film inhibits the transfer of oxygen.

The procedure described above assumes that the oil layer in the impoundment is well mixed. For example, changes in wind direction in units with retention times on the order of days may tend to move the oil layer in different directions and result in mixing. However, some systems may be designed for or characterized by plug flow. This flow model assumes that the oil film moves across the impoundment's surface without backmixing. For plug flow of the oil film in flowthrough impoundments and tanks, the fraction of V_0 in the oil layer emitted to the air is given by Equation (4-11), and air emissions are estimated from Equation (4-12). In these equations, C_e is the V_0 concentration in the oily effluent, C_0 is the initial concentration in the oil layer entering the impoundment, τ is the residence time, D is the oil-film thickness, and Q is the volumetric flow-rate of oil.

For an oil film on a disposal impoundment, emissions are calculated as described in Section 4.5. However, biodegradation is neglected and Equation (4-32) reduces to:

$$C_t/C_0 = \exp (- Kt/D) \quad (4-45)$$

and the fraction emitted to the air is:

$$f_{air} = 1 - \exp (-Kt/D) \quad (4-46)$$

where

C_t = concentration in the oil film at time = t

C_0 = initial concentration in the oil film

D = oil-film thickness

and with the other symbols as previously defined. The average emission rate over the period of time equal to t is:

$$E \text{ (g/s)} = f_{air} V C_0/t$$

where

V = volume of oil in the impoundment, m^3

and with the other symbols as previously defined. An example calculation of this approach is given in Section 5.0 for applying an oil film to soil, which is analogous to an oil film on a disposal impoundment because there is no flow out in either case and emissions are a function of the time since disposal.

4.8 DISCUSSION OF ASSUMPTIONS AND SENSITIVITY ANALYSIS

4.8.1 Removal Mechanisms

The organic constituents present in wastes that are treated, stored, or disposed of in surface impoundments and open tanks may leave the unit by any of several mechanisms. Because of the large open surface area and relatively high volatility of many organic constituents, emissions to the air may be a primary removal mechanism for certain constituents. Other constituents may be destroyed in impoundments and tanks specifically designed for biodegradation. Aeration is often used to supply oxygen to biologically active systems. Unfortunately, aeration also greatly enhances the mass transfer of organic constituents to the air. Other removal mechanisms include adsorption on solids, seepage through the ground, or degradation (e.g., by photolysis or hydrolysis). For flowthrough systems, the organic constituents may leave the unit with the effluent that will subsequently be treated, stored, or disposed of.

Initial studies suggest that emission to air is a primary removal mechanism, especially for volatile constituents. Biodegradation in specific systems, particularly for semivolatiles, may also be significant. For flowthrough systems, the removal of semivolatiles with the effluent may also be a primary removal mechanism. Other forms of degradation, adsorption, and seepage are neglected in this analysis for several reasons. These mechanisms are not believed to be significant for most systems and most constituents; however, they may be removal routes in a specific system or for a specific constituent. For example, an open tank may be designed specifically for liquid-phase carbon adsorption. These mechanisms are also difficult to model in a manner that is generally applicable considering the relatively sparse data on such removal mechanisms, especially in hazardous

waste impoundments and tanks. Consequently, the modeling effort focuses on mass transfer to the air and some consideration of biodegradation.

Numerous studies have been conducted to assess mass transfer to the air; these include theoretical assessments, correlations based on laboratory and bench-scale measurements, and field measurements at actual sources. Additional data on specific wastes have been collected in air-stripping studies as more air-stripping columns have been used to remove VO constituents from water. The result is that the state of knowledge of mass transfer from the liquid to the gas phase (e.g., ambient air) is probably advanced compared to the state of knowledge of other removal mechanisms. The level of confidence in the air emission models is probably highest for the volatile constituents because of very high mass transfer rates. The level of confidence is somewhat lower for the relatively nonvolatile constituents because of potentially significant rates of removal by other mechanisms.

Much of the data on the performance of systems designed for biodegradation are reported as total removal from measurement of the influent and effluent concentrations. This total would include removal to the air and biodegradation. Some studies have been conducted in closed systems in which the biodegradation rate may be measured directly (loss to the air is deliberately prevented). These data are useful for comparing the relative rates of removal by biodegradation among constituents and make possible a ranking of these constituents with respect to biodegradability. In addition, the estimated rate of biodegradation may be compared to the estimated rate of air emissions to assess the relative extent of each.

The biodegradation model has not been validated and is used in this report as an approximate measure of the extent of biodegradation. For any specific treatment system, measurements of actual biodegradation rates should be used if available. Any user of the biodegradation model should be aware that the predicted rate is very sensitive to the choice of values for the biorate, biomass concentration, and the concentration of organic constituents in the waste. An environmentally conservative approach with respect to air emissions would be to neglect biodegradation (assume the rate is zero). This approach is probably valid for volatile constituents

in aerated systems; however, the approach may tend to overestimate emissions of relatively nonvolatile constituents that are destroyed in treatment systems specifically designed for biodegradation.

4.8.2 Major Assumptions

An inherent assumption in the emission estimating procedure is that the mass transfer correlations chosen earlier are generally applicable. A paper that compares several different models concludes that, in most cases, many different models yield comparable results for volatile constituents.⁵⁶ The choice of models may affect the estimated mass transfer coefficients for semivolatiles more than those for volatiles. The calculations indicate that emissions of volatiles are controlled by the liquid-phase resistance. Consequently, the value for the overall mass transfer coefficient (K) is primarily determined by the correlation used for the liquid-phase mass transfer coefficient (k_L). For constituents with decreasing volatility, both the liquid-phase and gas-phase resistance begin to contribute to the overall resistance to mass transfer. For these constituents, the choices of correlations for both k_g and k_L become important, and the choice of correlations may significantly affect the emission estimates.

The flow model chosen for storage and treatment impoundments assumes that the impoundment's contents are well mixed and that the system is operated at steady-state conditions. The flow for specific facilities may be better represented by plug flow or a model that accounts for axial dispersion. The choice of flow model does not make a significant difference in the estimated emissions. However, if the loading of the impoundment is cyclical or intermittent instead of continuous, the emissions from the impoundment are likely to be cyclical or intermittent. Estimates of short-term emission rates are very dependent upon the method of operation of the system. For disposal impoundments, peak emissions occur when the waste is first placed in the impoundment and then decrease with time. The approach used in this report estimates the average emission rate over a given period of time and does not provide an estimate of the initial peak emissions.

The calculation of Henry's law constant also contains inherent assumptions. The approach is valid for dilute solutions and has been applied successfully in the design of air-stripping columns. However, specific

mixtures may deviate from Henry's law because of component interactions or because of concentrations outside the range of applicability. Errors in applying Henry's law are generally environmentally conservative; i.e., the actual gas-phase concentration is not likely to be underestimated.

For concentrated mixtures of organics in a separate oil layer, the use of Raoult's law is recommended. This approach is valid for mixtures of constituents with similar properties, especially when the concentration of the component of interest is very high. A preferred approach would be to avoid the use of Henry's law or Raoult's law and actually measure the equilibrium partitioning between the liquid and gas phase of a waste. However, very few data are available for equilibrium partitioning that can be applied generally to hazardous waste mixtures.

4.8.3 Sensitivity Analysis

The emission correlations were evaluated for sensitivity to each of the input parameters.⁵⁷ In the analysis, each input parameter was varied individually over the entire range of reasonable values. The effect on emissions was noted, and the most sensitive parameters were identified.

Detention time is an important parameter that affects emissions from the impoundment. The emission estimates for volatile constituents are sensitive to short detention times, and the estimates for semivolatiles are sensitive to long detention times. Essentially all of the volatile constituents are emitted for longer detention times (several days), and very little of the semivolatiles are emitted for short detention times (a few days). However, significant emissions of the semivolatiles may occur for long detention times in storage impoundments or in disposal impoundments.

The value of Henry's law constant was not important for volatile constituents. The correlations indicated that these constituents are controlled by the liquid-phase resistance, which is not affected by Henry's law constant. The value of Henry's law constant has a direct effect on the emissions of semivolatiles (such as phenol), and the greatest effect is on those relatively nonvolatile compounds for which mass transfer is controlled by the gas-phase resistance.

Windspeed has a direct effect on the emission estimates for quiescent surfaces and has little effect on those from aerated systems. The results

showed that a standard windspeed of 4.5 m/s was reasonable compared with the results for windspeed distributions at actual sites.

Temperature did not affect the emission estimates for the volatile constituents. However, temperature did affect the emission estimates for nonvolatile constituents with mass transfer controlled by the gas phase. The temperature dependence of Henry's law constant accounts for this effect.

The diffusivity in air and water for a wide variety of constituents spans a relatively narrow range of values. The analysis showed that the emission estimates were not sensitive to the choice of values for diffusivity.

For mechanically aerated systems, the choice of values for impeller diameter, impeller speed, oxygen transfer rate, and oxygen correction factor did not affect the emission estimates significantly. The total horsepower and turbulent area had a direct effect on emissions of semi-volatiles (e.g., phenol). However, there was no significant effect on emissions of volatile constituents because the models predicted that they would be stripped almost completely from the water over the full range of aeration values.

The biodegradation model was very sensitive to all parameters investigated. The sensitive parameters include organic concentration, biomass concentration, and biorate.

Two meteorological parameters required in the models are temperature and windspeed. The emission estimates are based on a standard temperature of 25 °C and a windspeed of 4.47 m/s (10 mi/h). These standard values were evaluated by estimating emissions for windspeed/temperature combinations at actual sites based on their frequency of occurrence. Over a 1-yr period, the results from site-specific data on windspeed and temperature were not significantly different from the results using the standard values. Consequently, the standard values were judged adequate to estimate annual emissions. For short-term emissions, the actual temperature and windspeed over the short-term interval should be used to avoid underestimating emissions during high-windspeed/high-temperature conditions.

A sensitivity analysis was performed for three impoundment model units (storage, mechanically aerated, and disposal) presented in the example calculations in this section. Three compounds were chosen to represent relatively nonvolatile compounds (p-cresol), moderately volatile compounds (acetone), and relatively volatile compounds (benzene). Each of these compounds can be biodegraded. The results are given in Tables 4-12, 4-13, and 4-14. The key input parameters identified in the tables were increased by 50 percent from the base case to determine the effect on the percent of the compound in the waste that is emitted to the air.

For each of the different types of impoundments, the volatility appears to be important only for the low volatility category. As discussed previously, the windspeed (air turbulence) has a direct effect for each of the compounds in a storage impoundment and does not affect the mechanically aerated unit's results. The low volatility compounds are the most sensitive to changes in depth and biomass concentration for all three types of impoundments. An assumption of no biodegradation also has the most dramatic effect on the low volatility compound with smaller effects observed for the higher volatility compounds. The effects of retention time are small except for the results shown for the disposal impoundment after 5 days. The disposal impoundment results show that for short times, the time since disposal is an important parameter affecting emissions.

**TABLE 4-12. RESULTS OF SENSITIVITY ANALYSIS FOR QUIESCENT
STORAGE IMPOUNDMENT**

Key emission model inputs	Percent emitted for given Henry's law constant, atm•m ³ /mole		
	10 ⁻⁷	10 ⁻⁵	10 ⁻³
<u>Base case^a</u>	2.9	58	59
<u>50-percent increase from base case^b</u>			
Volatility	4.2 (45) ^c	61 (5)	59 (0)
Air turbulence	4.0 (38)	72 (24)	76 (29)
Retention time	3.2 (10)	62 (7)	62 (5)
Depth	2.1 (-28)	50 (-14)	49 (-17)
Biomass concentration	2.1 (-28)	52 (-10)	52 (-12)
<u>No biodegradation^d</u>	10 (245)	74 (28)	80 (36)

^aThis corresponds to the model unit for storage impoundments used in the example calculation.

^bEach parameter is increased individually by 50 percent from its base case value.

^cValues in parentheses are percent change from the base case.

^dBase case with no biodegradation.

**TABLE 4-13. RESULTS OF SENSITIVITY ANALYSIS FOR MECHANICALLY
AERATED IMPOUNDMENTS**

Key emission model inputs	Percent emitted for given Henry's law constant, atm•m ³ /mole		
	10 ⁻⁷	10 ⁻⁵	10 ⁻³
Base case^a	2.7	79	99
50-percent increase from base case^b			
Volatility	3.9 (44) ^c	85 (8)	99 (0)
Air turbulence	2.8 (4)	80 (1)	99 (0)
Water turbulence	3.6 (33)	85 (8)	99 (0)
Retention time	2.7 (0)	80 (1)	99 (0)
Depth	1.8 (-33)	73 (-8)	98 (-1)
Biomass concentration	1.8 (-33)	73 (-8)	98 (-1)
No biodegradation^d	20 (640)	94 (28)	100 (1)

^aThis corresponds to the model unit for mechanically aerated impoundments used in the example calculation.

^bEach parameter is increased individually by 50 percent from its base case value.

^cValues in parentheses are percent change from the base case.

^dBase case with no biodegradation.

TABLE 4-14. RESULTS OF SENSITIVITY ANALYSIS FOR DISPOSAL IMPOUNDMENTS

Key emission model inputs	Percent emitted for given Henry's law constant, atm·m ³ /mole		
	10 ⁻⁷	10 ⁻⁵	10 ⁻³
<u>Base case^a</u>	13	93	92
<u>50-percent increase from base case</u>			
Volatility	18 (38) ^b	94 (1)	92 (0)
Air turbulence	17 (31)	96 (3)	96 (4)
Retention time ^c	2.3 (-82)	55 (-41)	72 (-22)
Depth	9 (-31)	89 (-4)	88 (-4)
Biomass	9 (-31)	89 (-4)	89 (-3)
<u>No biodegradation^d</u>	84 (550)	100 (8)	100 (9)

^aBased on the dimensions given in the example calculation, 100 mg/L of the constituent in 1,000 mg/L total organics, and a time since disposal of 12 months.

^bValues in parentheses are percent change from base case.

^cA retention time of 5 days was selected here to show the sensitivity to retention time soon after disposal.

^dBase case with no biodegradation.

4.9 REFERENCES

1. Lunney, P. D. Characterization of Wind and Depth Effects Upon Liquid Phase Mass Transfer Coefficients: Simulation Studies. Master's thesis, University of Arkansas, Fayetteville, AR. January 1983. p. 119.
2. Springer, C., P. D. Lunney, and K. T. Valsaraj. Emission of Hazardous Chemicals from Surface and Near Surface Impoundments to Air. U.S. Environmental Protection Agency, Solid and Hazardous Waste Research Division. Cincinnati, OH. Project Number 808161-02. December 1984. p. 3-4 to 3-16.
3. Reference 2, p. 3-16 to 3-19.
4. Reference 2, p. 3-18.
5. Hwang, S. T. Toxic Emissions from Land Disposal Facilities. Environmental Progress. 1:46-52. February 1982.
6. Mackay, D., and A. Yeun. Mass Transfer Coefficient Correlations for Volatilization of Organic Solutes from Water. Environmental Science and Technology. 17:211-217. 1983.
7. Reference 6, p. 214.
8. GCA Corporation. Air Emissions for Quiescent Surface Impoundments--Emissions Data and Model Review. Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Contract No. 68-01-6871, Assignment 49. August 1985. p. 5-1 and 5-2.
9. Reference 8, p. 4-4.
10. Reference 5, p. 47.
11. Thibodeaux, L. J. Air Stripping of Organics from Wastewater. A Compendium. Air/Water. p. 373. (In publication.)
12. Westat Corporation. National Survey of Hazardous Waste Generators and TSD's Regulated Under RCRA in 1981. Prepared for the U.S. Environmental Protection Agency. Contract No. 68-01-6861. April 1984.
13. Metcalf and Eddy, Inc. Wastewater Engineering. New York, McGraw-Hill. 1972. p. 542-554.
14. Eckenfelder, W., M. Goronszy, and T. Quirk. The Activated Sludge Process: State of the Art. CRC Critical Reviews in Environmental Control. 15(2):148. 1984.
15. Beardsley, M., and J. Coffey. Bioaugmentation: Optimizing Biological Wastewater Treatment. Pollution Engineering. December 1985. p. 32.

16. Reference 13, p. 586.
17. Reference 15, p. 32.
18. Reference 13, p. 520-521.
19. U.S. Environmental Protection Agency. EPA Design Manual: Municipal Wastewater Stabilization Ponds. Publication No. EPA-625/1-83-015. October 1983. p. 3.
20. Reference 13, p. 557.
21. Englande, A. J. Performance Evaluation of the Aerated Lagoon System at North Gulfport, Mississippi. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-600/2-80-006. March 1980. p. 39-41.
22. Allen, C. Project Summary: Site Visits of Aerated and Nonaerated Surface Impoundments. Prepared for U.S. Environmental Protection Agency. Contract No. 68-03-3253. Assignment 2-8. June 1987. p. 2.
23. Petrasek, A., B. Austern, and T. Neiheisel. Removal and Partitioning of Volatile Organic Priority Pollutants in Wastewater Treatment. Presented at the Ninth U.S.-Japan Conference on Sewage Treatment Technology. Tokyo, Japan. September 1983. p. 16.
24. Bishop, D. The Role of Municipal Wastewater Treatment in Control of Toxics. Presented at the NATO/CCMS Meeting. Bari, Italy. September 1982. p. 18.
25. Hannah, S., B. Austern, A. Eralp, and R. Wise. Comparative Removal of Toxic Pollutants by Six Wastewater Treatment Processes. Journal WPCF. 58(1):30. 1986.
26. Kincannon, D., and E. Stover. Fate of Organic Compounds During Biological Treatment. Presented at ASCE Environmental Engineering Conference. 1981. p. 6.
27. Melcer, H. Biological Removal of Organic Priority Pollutants. Presented at Hazardous Substances in Wastewater Seminar. Toronto, Canada. November 1982. p. 20.
28. Reference 19, p. 75-146.
29. Reference 13, p. 481-573.
30. Reference 14, p. 119.
31. Bailey, J. E., and D. F. Ollis. Biochemical Engineering Fundamentals. New York, McGraw-Hill. 1977. p. 343-349.

32. Kincannon, D., and E. Stover. Determination of Activated Sludge Biokinetic Constants for Chemical and Plastic Industrial Wastewaters. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-600/2-83-073a. August 1983. p. 18-20.
33. Pitter, P. Determination of Biological Degradability of Organic Substances. Water Research. 10:231-235. 1976.
34. Kincannon, D., A. Weinert, R. Padorr, and E. L. Stover. Predicting Treatability of Multiple Organic Priority Pollutant Wastewater from Single Treatability Studies. Presented at the 37th Purdue Industrial Waste Conference, West Lafayette, IN. May 1982.
35. Reference 23, p. 4-16.
36. Reference 25, p. 27-34.
37. Coburn, J., C. Allen, D. Green, and K. Leese. Site Visits of Aerated and Nonaerated Impoundments. Revised Draft Summary Report. Prepared for U.S. Environmental Protection Agency. Contract No. 68-03-3253, Work Assignment No. 3-8. April 1988. p. A-1 to A-34.
38. Matter-Muller, C., W. Gujer, W. Giger, and W. Stumm. Non-Biological Elimination Mechanisms in a Biological Sewage Treatment Plant. Prog. Water Tech. 12:305. 1980.
39. Dobbs, R., M. Jelus, and K. Chang. Partitioning of Toxic Organic Compounds on Municipal Wastewater Treatment Plant Solids. Proceedings of the International Conference on Innovative Biological Treatment of Toxic Wastewaters. Scholze, R. J., Ed. Arlington, VA. June 1986. p. 585-601.
40. Reference 5, p. 46.
41. Reinhardt, J. R. Gas-Side Mass-Transfer Coefficient and Interfacial Phenomena of Flat-Bladed Surface Agitators. Ph.D. dissertation, University of Arkansas, Fayetteville, AR. 1977. 96 p.
42. GCA Corporation. Emissions Data and Model Review for Wastewater Treatment Operations. Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Contract No. 68-01-6871, Assignment 49. August 1985. p. 4-3.
43. Reference 42, p. 4-2.
44. Reference 5, p. 47.
45. Reference 42, p. 4-3.
46. Reference 41, p. 48.

47. Reference 19, p. 3.
48. Reference 13, p. 557.
49. Reference 13, p. 519.
50. GCA Corporation. Hazardous Waste TSD Waste Process Sampling. Prepared for U.S. Environmental Protection Agency. Report No. EMB/85-HNS-3. October 1985. p. 1-11.
51. GCA Corporation. Evaluation and Selection of Models for Estimating Air Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Prepared for U. S. Environmental Protection Agency. Publication No. EPA-450/3-84-020. December 1984. p. 69.
52. Thibodeaux, L., and D. Parker. Desorption Limits of Selected Gases and Liquids from Aerated Basins. AIChE Symposium Series. 72(156):424-434. 1976.
53. Reference 51, p. 67.
54. Reference 51, p. 67.
55. Reference 13, p. 519.
56. Allen, C. C. Prediction of Air Emissions from Surface Impoundments. Paper 31a. (Presented at 1986 Summer Meeting of AIChE. Boston, MA. August 1986.) 26 p.
57. Branscome, M., and A. Gitelman. Sensitivity Analysis: Emission Estimates for Surface Impoundments. Prepared for the U.S. Environmental Protection Agency. March 1986. 67 p.

5.0 LAND TREATMENT

This chapter presents the approach used to estimate air emissions from land treatment operations. Analytical models to estimate emissions, representative values of model input parameters, and example calculations are included.

5.1 NARRATIVE DESCRIPTION OF LAND TREATMENT AIR EMISSIONS

Land treatment is one of several land disposal methods used for final disposition of hazardous wastes. At land treatment facilities, wastes are either spread onto or injected into the soil, after which they are normally tilled into the soil. Other activities likely to occur at land treatment facilities include storage of wastes in tanks or surface impoundments, loading and unloading of wastes in vacuum trucks or dump trucks, and dewatering of wastes using filtration devices. All of these activities have emission points associated with them. The following paragraphs describe analytical models used to estimate emissions from the application, tilling, and final disposition of hazardous waste at a land treatment disposal site. Emissions from other land treatment activities, such as truck loading, storage tanks, and fugitive emissions from transfer and handling operations, are estimated using procedures described in Chapter 7.0 of this report.

Estimating emissions from land treatment may involve one to three independent steps depending on operating practices at a land treatment site. If waste is applied from a vacuum truck to the soil surface, allowed to remain on the surface for a period of time, and then tilled into the soil, emissions are estimated in three steps: (1) during application of waste onto the soil from a vacuum truck, (2) after waste application and before tilling, and (3) after tilling the waste into the soil. If waste is applied to the soil surface and immediately tilled into the soil, emissions

are estimated in only two steps: (1) during waste application, and (2) after tilling. If waste is applied by subsurface injection and immediately tilled, only one step is required to estimate emissions.

This section presents three separate analytical models that can be used to estimate air emissions from separate land treatment activities. Primary emphasis is given to the RTI land treatment model that is used to calculate emissions from waste that is mixed with the soil. This condition may exist when waste has been applied to the soil surface and has seeped into the soil or when waste has been injected beneath the soil surface or has been tilled into the soil. The RTI land treatment model is described below in Subsection 5.2.1, which includes separate discussions of the following topics:

<u>Subsection</u>	<u>Topic</u>
5.2.1.2	Biodegradation
5.2.1.4	Effective diffusivity
5.2.1.5	Waste partitioning
5.2.1.10	Model selection rationale

After waste is applied to the surface of the soil by spray application, it seeps into the soil. While the waste is on the surface, the concentrations at the surface can be approximated by the concentration in the waste. During this short period that the waste covers the surface of the soil, the maximum short-term emission rate is expected. For this special case, emissions are estimated as the product of an overall mass transfer coefficient, constituent concentration, and surface area of the land treatment site. The model for estimating the mass transfer coefficient from the surface to the wind was developed by McKay and Matsuga and is briefly discussed in Subsection 5.2.3.

Emissions from a waste stream as it is applied onto the soil surface from a vacuum truck, regardless of waste type, are also calculated as the product of an overall mass transfer coefficient, the surface area of the waste stream, and the concentration of a specific constituent. Preliminary calculations indicate that emissions from the spraying waste application are relatively small and can be ignored in most situations. Even so, a brief discussion of a model for estimating these emissions is presented in

Subsection 5.2.2, and the model can be used if desired. Also included in this section are Subsection 5.2.4, which discusses representative values of input parameters for the analytical models, and Subsection 5.2.6, which presents example calculations using each of the three models presented.

At many existing land treatment sites, waste is applied onto the soil from a vacuum truck and is allowed to remain for about 24 hours before being tilled into the soil. Under these conditions, three separate calculations may be needed to estimate air emissions. Emissions during waste application could be estimated using the waste application model described in Subsection 5.2.2; emissions after application but before tilling would be estimated using the RTI land treatment model as described in Subsection 5.2.1 (or, if a visible oil film exists on the soil surface, the oil film surface model as presented in Subsection 5.2.3); and emissions after tilling would be estimated using the RTI land treatment model. At other existing sites, waste is injected into the soil using subsurface injection and is immediately tilled. At these sites, only one calculation is needed to estimate emissions. In this situation, the RTI land treatment model would be used.

5.2 LAND TREATMENT

5.2.1 Land Treatment Emission Model Descriptions

5.2.1.1 Analytical Correlations. Emissions from land treatment after waste is applied to the soil are estimated using a model developed by Clark Allen of Research Triangle Institute (the RTI model). This model assumes that emissions from the surface of the soil/waste mixture are limited by the diffusion of vapors through the pore spaces in the soil/waste mixture and further assumes that an equilibrium concentration of organic vapors exists at all times within the pore spaces. The model is based on Fick's second law of diffusion applied to a flat slab as described by Crank¹ and includes a term to estimate biological decay assuming a decay rate that is first order with respect to waste loading in the soil.

The solution to the diffusion equation developed by Crank is for diffusion out of a slab that initially has a uniform concentration of diffusing material throughout and that has equal concentrations of diffusing material at each surface.

The general solution to the diffusion equation for those conditions, as presented by Crank, is:

$$F = \frac{M_t}{M_0} = 1 - \sum_{n=0}^{\infty} \frac{8}{(2n+1)^2 \pi^2} \exp\left(\frac{-D (2n+1)^2 \pi^2 t}{4 l^2}\right) \quad (5-1)$$

where

F = fraction of initially applied material that has diffused out of the slab at time t

M_t = mass of material that has diffused out of the slab at time t

M_0 = initial mass of material present

D = diffusion coefficient

l = distance from center to surface of slab

t = time after initial distribution of diffusing material into the slab.

This series solution converges very slowly for small values of time (i.e., $Dt/l^2 < 0.213$). Because of this slow convergence at short times (i.e., immediately after waste application or tilling), Crank presented an alternative solution that is valid during this short time. The following equation is obtained from the alternative solution:

$$F = \frac{M_t}{M_0} = \frac{2}{\sqrt{\pi}} \left(\frac{Dt}{l^2}\right)^{1/2} \quad \left[\text{for } Dt/l^2 < 0.213\right] \quad (5-2)$$

Equation (5-2) approximates the Crank solution but excludes a small error function correction used by Crank.

To verify the validity of Crank's solution for short times and to test the accuracy of an approximation for use over longer times, the values predicted by the solution for short times and the values obtained using the first term of the series solution [Equation (5-3)] are compared to the values obtained using the first three terms of the series solution. Table 5-1 presents the results for a range of values of the dimensionless parameter, Dt/l^2 :

$$F = 1 - \frac{8}{\pi^2} \exp \left[- \frac{0.213^2 t}{4l^2} \right] \quad (5-3)$$

Table 5-1 shows that, for values of the dimensionless parameter greater than 0.213, the first term of the series solution, Equation (5-3), can be used to estimate total emissions. The table also shows that the solution for short times, Equation (5-2), is valid for values of the dimensionless parameter below 0.213. Equations (5-2) and (5-3) give identical results for a parameter value of 0.213. This comparison indicates that sufficient accuracy can be attained under all conditions if the equation for short times is used for values of the dimensionless parameter below 0.213 and the first term of the general solution is used for values above 0.213. It is observed that the fraction of material that diffuses out of the slab is linear with respect to the square root of time up to the point where approximately 50 percent of the diffusing material is lost.²

The conditions defined for the above solutions by Crank are analogous to diffusion of volatile organics out of a surface layer of a soil/waste mixture as happens in land treatment operations. Because of the symmetry of conditions on which the above solutions are based, an impenetrable plane could, in theory, be inserted at the midpoint of the slab without changing the solution. One-half of the slab with an impenetrable boundary layer on the bottom would represent the surface layer of soil into which waste is mixed during land treatment.

In a land treatment operation, the applied material partitions into several phases including evaporation into a vapor phase, adsorption onto soil particles, and absorption into oil and water in the soil/waste mixture. Only the vapor phase is available for diffusion out of the soil/waste mixture. Therefore, to apply the above equations to land treatment, the amount of material in the vapor phase must be known. The amount of material that partitions into the vapor phase can be estimated by calculating equilibrium conditions within the soil/waste mixture. This equilibrium is estimated by defining K_{eq} , the ratio of the amount of organics in the vapor phase to the total amount of organics in the soil/waste mixture. The instantaneous emission rate, E , at any time, t , can be estimated by the following Equations (5-4) and (5-5), which are obtained by differentiating

TABLE 5-1. COMPARISON OF THE ESTIMATED FRACTION
EMITTED USING THREE DIFFERENT EQUATIONS
(INTEGRATED FLUX FROM SOIL)

Time parameter (Dt/l ²)	Short-term solution $2\left(\frac{Dt}{\pi l^2}\right)^{1/2}$	First term of series solution $1 - \frac{8}{\pi^2} \exp\left[-\frac{Dt\pi^2}{4l^2}\right]$	First three terms of series solution
0.000	0.000	0.189	0.067
0.025	0.178	0.238	0.179
0.050	0.252	0.284	0.252
0.075	0.309	0.326	0.309
0.100	0.357	0.367	0.357
0.125	0.399	0.405	0.399
0.150	0.437	0.440	0.437
0.175	0.472	0.474	0.472
0.200	0.505	0.505	0.504
0.213	0.521	0.521	0.520
0.250	0.564	0.562	0.562
0.275	0.592	0.589	0.589
0.300	0.618	0.613	0.613
0.325	0.643	0.636	0.636
0.350	0.668	0.658	0.658
0.375	0.691	0.679	0.679
0.400	0.714	0.698	0.698
0.425	0.736	0.716	0.716
0.450	0.757	0.733	0.733
0.475	0.778	0.749	0.749
0.500	0.798	0.764	0.764
0.525	0.818	0.778	0.778
0.550	0.837	0.791	0.791
0.575	0.856	0.804	0.803
0.600	0.874	0.816	0.816
0.625	0.892	0.827	0.827
0.650	0.910	0.837	0.837
0.675	0.927	0.847	0.847
0.700	0.944	0.856	0.856
0.725	0.961	0.864	0.865
0.750	0.977	0.873	0.873
0.775	0.993	0.880	0.880
0.800	1.009	0.887	0.887
0.825	1.025	0.894	0.894
0.850	1.040	0.900	0.900
0.875	1.056	0.906	0.906
0.900	1.070	0.912	0.912
0.925	1.085	0.917	0.917
0.950	1.100	0.922	0.922
0.975	1.114	0.927	0.927

Equations (5-2) and (5-3) and adding the equilibrium constant, K_{eq} , and a term to account for waste biodegradation, $e^{(-t/t_b)}$:

$$\text{(short times)} \quad E = \frac{M_o}{1} \left[\frac{K_{eq}D}{\pi t} \right]^{1/2} e^{-t/t_b} \quad , \quad (5-4)$$

and

$$\text{(longer times)} \quad E = M_o \left[\frac{2K_{eq}D}{1^2} \right] \exp \left(\frac{-K_{eq}D \pi^2 t}{4l^2} \right) e^{-t/t_b} \quad (5-5)$$

where

t_b = the biological degradation time constant.

The above equations account for the removal of organic material from the soil/waste mixture both biological degradation and air emissions. In a land treatment operation, the primary objective is to dispose of organic materials by biodegradation; thus, significant quantities of waste would be expected to be depleted from the soil by biological degradation. Other mechanisms of removal such as leaching and photolysis also are possible but are not accounted for in this model because of the estimated small amount of materials lost by these processes.

5.2.1.2 Biodegradation. Biodegradation at land treatment sites is generally considered to be a first-order process with respect to waste concentration in the soil up to the point where saturation is achieved.³ In addition to literature sources that make such statements, comments on a draft of this document provided by Chevron Research Company offer further evidence of the first-order nature of biodegradation at land treatment sites.⁴ A first-order decay process is defined in the literature as having the following form:⁵

$$\frac{dM}{dt} = - K_b M \quad ,$$

where

M = mass of organic material in the soil

K_b = biological decay constant.

Integrating and using the boundary conditions $M = M_0$ at $t = 0$ results in:

$$\frac{dM}{M} = -K_b dt$$

$$\log M = -K_b t + C_1$$

or

$$M = C_2 e^{-K_b t}$$

where C_1 and C_2 are constants of integration. Substituting the boundary conditions gives:

$$M = M_0 e^{-K_b t}$$

K_b has units of s^{-1} and can be expressed as the reciprocal of the biological decay time constant, $1/t_b$. The exponential was introduced directly into the rate relationship, Equations (5-4) and (5-5), to reduce the amount of material available for air emissions by the fraction of material removed by biooxidation.

5.2.1.3 Estimation of Equilibrium Coefficient, K_{eq} . Partitioning of volatile constituents in the waste is assumed to occur between the vapor space in the soil/waste mixture, adsorbent solids in the soil, and absorbent liquids in the soil and waste. Using 1 cm^3 of the soil/waste mixture as a basis for calculation, the total volume of gas (i.e., void space) in the cubic centimeter is described by the air porosity, ϵ_a . Using the ideal gas law, the number of moles of gas in 1 cm^3 of the soil/waste mixture is $P\epsilon_a/(RT)$, where P is the pressure of a constituent in the gas phase and is usually equal to XP^* (X is the mole fraction of the constituent in the liquid phase and P^* is the pure component vapor pressure). The moles of constituent in the gas phase in 1 cm^3 of the soil/waste mixture is thus $XP^*\epsilon_a/(RT)$. Oil loading in the soil/waste mixture in units of grams of oil per cubic centimeter of mixture is L ($\text{g}_{oil}/\text{cm}^3$ mixture), and the total moles of constituent per cubic centimeter of the mixture is XL/MW_{oil} . The equilibrium coefficient, K_{eq} , is defined as the moles of constituent in the gas phase per unit volume of the soil/waste mixture divided by the

total moles of constituent per unit volume of the soil/waste mixture. Therefore, the following equation can be written:

$$K_{eq} = \frac{XP^*\epsilon_a/(RT)}{X L/MW_{oil}} = \frac{P^*MW_{oil}\epsilon_a}{RTL}$$

This equation differs from the usual equation for equilibrium coefficient by the factor ϵ_a , which is included to account for the limited air space available within the soil/waste mixture. The ratio of moles per mole and grams per gram can be used interchangeably in this equation. The value of K_{eq} can be calculated from measurements, if available, of constituent concentrations in the pore space and in the soil/waste mixture.

In a similar manner, it can be shown that K_{eq} can be estimated for aqueous wastes with an assumed value of the Henry's law constant, H_c :

$$K_{eq} = \frac{H_c 10^6 \epsilon_a}{RT\epsilon_{waste}}$$

where

ϵ_{waste} = the volume fraction of the soil/waste mixture that is occupied by waste.

In the above equations, it is assumed that equilibrium is controlled by Raoult's law for oily wastes and by Henry's law for aqueous wastes. An aqueous waste is assumed to contain water and organic constituents that are dissolved in water. An example of an aqueous waste is a sludge containing 10 percent solids, 5 percent acetone, 1 percent methanol, 500 ppmw benzene, and the remainder water. If the waste contains oil mixed with the water, or the waste contains volatile constituents at concentrations greater than the solubility in water, it is modeled as an oily waste. It is important to specify the molecular weight of this separate organic phase for this Raoult's law calculation of K_{eq} .

5.2.1.4 Estimation of Effective Diffusivity. The diffusivity of specific compounds, as reported in the literature, assumes that the diffusion occurs in free air. In a land treatment operation, diffusion of vapors out of the soil must take place within the confines of the

air-filled voids within the soil. This characteristic of soil is referred to as the air porosity. The ratio of effective diffusivity of a constituent in the soil to its diffusivity in air can be described by the following equation:⁶

$$\frac{D_e}{D_a} = \frac{\epsilon_a^{10/3}}{\epsilon_T^2}$$

where

D_e = effective diffusivity of constituent in soil

D_a = diffusivity of constituent in air

ϵ_a = air porosity of soil

ϵ_T = total porosity of soil.

When air porosity and total porosity are the same (i.e., for dry soil), this equation reduces to:

$$\frac{D_e}{D_a} = \epsilon_a^{4/3} .$$

Total porosity refers to the fraction of the land treatment medium that is made up of nonsoil (or nonsolid) materials, i.e., the sum of the void space, water-filled space, and space occupied by the oil in the applied waste.

Soil air porosity undergoes substantial changes over time as soil dries out and when moisture is added by rainfall or by watering. As a result, accurately accounting for soil porosity in an analytical model is difficult. The use of average or typical values of soil porosity may be the most practical approach.

5.2.1.5 Waste Partitioning. A large percentage of wastes that are disposed of by land treatment are refinery sludges. These wastes are mostly sludge emulsions and consist of varying fractions of water, oil, and inorganic solids, where oil represents the total organic portion of the waste including volatile compounds. A much smaller amount of land-treated wastes are dilute aqueous solutions of water and organic compounds. When wastes are applied to a land treatment area, volatile materials in the soil

have the potential for partitioning into four different phases--a vapor phase, an oil phase where volatile material is dissolved in the oil, a water phase where volatile material is absorbed in the soil moisture, and a soil phase where volatile material is adsorbed by organic carbon within the soil. For oily wastes, VO compounds will preferentially dissolve in oil rather than water so that the fraction of volatile materials in the water phase is estimated to be very small. Partitioning of volatile materials into the soil phase by adsorption is a function of the amount of organic carbon in the soil. It is also estimated to account for only a small fraction of the applied organics because the surfaces in the soil are expected to contain oil from the application and tilling of waste materials that contain oil. This oil in the soil is expected to both absorb the volatile constituents and to interfere with the relatively lower adsorption rates on soil surfaces. For high molecular weight constituents present in aqueous wastes, adsorption may be more important. An equilibrium equation can be written that takes all four phases into account in the estimation of equilibrium vapor concentration in the soil. However, as presented here, the equilibrium equation in the RTI model includes only two phases. Calculations by one researcher looked at the difference in estimated emissions using two-phase partitioning of waste into an oil phase and vapor phase and using four-phase partitioning. The results of these comparisons are given in Table 5-2 and show for the conditions considered that, for soils having an organic carbon content of up to 10 percent, the estimated fraction of applied organics emitted using four-phase partitioning is only about 10 percent less than the estimated fraction emitted using two-phase partitioning. In a given situation, the amount of material adsorbed by organic carbon in the soil is relatively constant; thus, in soils with high organic carbon content, adsorption of materials in the soil may become more significant if low loading rates of oil or aqueous wastes are used. One of the products of biodegradation is organic carbon; thus, land treatment sites that have been active for an extended time may have elevated concentrations of organic carbon. Even so, with the normal oil loading used in land treatment, it is likely that a large fraction of the available adsorption sites would be occupied by the oil itself, thus limiting the effects of adsorption on emissions of the lighter constituents.

**TABLE 5-2. EMISSION ESTIMATES USING TWO DIFFERENT EQUATIONS
FOR THE VAPOR-SOIL PARTITION COEFFICIENT⁷**

Organic carbon content of soil (fraction)	Estimated emission fraction--two-phase partitioning	Estimated emission fraction--four-phase partitioning
0	0.622	0.622
0.001	0.622	0.621
0.010	0.622	0.614
0.100	0.622	0.559

For oily sludges, K_{eq} is calculated using vapor pressure and waste loading is calculated exclusive of water content. For dilute aqueous waste, partitioning is estimated to be in a water phase and a vapor phase, and the parameter K_{eq} is calculated using Henry's law constant; waste loading is calculated using the total waste applied. K_{eq} may be calculated from site-specific land treatment soil, vapor, and solids analyses if available. Table 5-3 summarizes the equations that make up the RTI land treatment model.

5.2.1.6 Emissions at Short Times. When a sludge containing volatile organics is applied onto or tilled into the soil at a land treatment site, the maximum rate of air emissions will occur immediately after application or tilling. Volatile organics will leave the surface and enter the environment through wind currents. Although the RTI model is based on the premise that emissions from land treatment are limited by vapor diffusion through the soil, the maximum rate of air emissions immediately after application or tilling will be limited by the gas-phase mass transfer coefficient, k_G . Within a few hours after application or tilling, the rate of air emissions from the volatile components will be substantially less than the maximum rate because the volatiles at the surface have been removed by the wind and the remaining volatiles must diffuse up through a layer of porous solids, a relatively slow process.

The land treatment model can be used for short-term emissions. The emissions from the short-term use of the land treatment model will be somewhat less than the oil-film model, although the initial rate from both of these models is equivalent. The oil film model is used to estimate maximum emission rates and the land treatment model is used to account for surface drying during short-term emission estimations. The equation for the emission rate immediately after application or tilling is:

$$E = \frac{M_0}{T} \left[\frac{1}{\frac{\epsilon_a}{k_G K_{eq}} + \sqrt{\frac{\pi t}{D_e K_{eq}}}} \right] e^{-t/t_b} \quad (5-6)$$

The basis of the above equation is a resistance in series model where the resistance (inverse of the mass transfer coefficient) is the sum of the

TABLE 5-3. RTI MODEL FOR LAND TREATMENT EMISSIONS

Emission rate equations

Short-term solution ($K_v t < 0.22$)

$$E = \frac{M_0}{T} \left[\frac{1}{\frac{\epsilon_a}{K_{eq} k_G} + \left(\frac{\pi t}{D_e K_{eq}} \right)^{1/2}} \right] e^{-t/t_b}$$

Long-term solution ($K_v t \geq 0.22$)

$$E = M_0 \left[\frac{2K_{eq} D_e}{l^2} \right] \exp \left(- \frac{K_{eq} D_e \pi^2 t}{4l^2} - \frac{t}{t_b} \right)$$

Fraction air emissions

Short-term solution [$K_v t < 0.22$]

$$F_{at} = \left[\frac{K_{eq} D_e t}{\pi l^2} \right]^{1/2} 2 \left[1 - 1/3 t/t_b \right]$$

Long-term solution ($K_v t \geq 0.22$)

$$F_{at} = \frac{8}{\pi^2} \left[1 + \frac{1}{K_d t_b} \right]^{-1} \left[1 - \exp \left[- K_d t - t/t_b \right] \right] + 0.1878$$

Long-term solution ($F_a < 0.33$ and $K_v t_b < 0.22$)

$$F_{at} = F_a [1 - \exp (-K_d t - t/t_b)]$$

Very long-term solution ($t \rightarrow \infty$) ($K_d t_b \geq 0.62$)

$$F_a = \frac{0.811 K_d t_b}{(K_d t_b + 1)} + 0.1878$$

Very long-term solution ($t \rightarrow \infty$) ($K_d t_b < 0.62$)

$$F_a = \left[\frac{K_{eq} D_e t_b}{l^2} \right]^{1/2}$$

(continued)

TABLE 5-3 (continued)

$$K_{eq} = \frac{P^* MW_{oil}}{RT} \frac{\epsilon_a}{L} \quad (\text{used for oily sludges})^a$$

$$K_{eq} = \frac{H_c}{RT} (10^6) \frac{\epsilon_a}{\epsilon_{waste}} \quad (\text{used for dilute aqueous waste})^a$$

$$k_G = 4.82 (10^{-3}) U^{0.78} Sc_G^{-0.67} de^{-0.11}$$

$$Sc_G = \frac{\mu_a}{\rho_a D_a}$$

$$M_o = L/C$$

$$K_v = \frac{K_{eq} D_e}{l^2} \quad (\text{volatilization constant})$$

$$K_d = \frac{\pi^2}{4} K_v$$

$$D_e = D_a \left(\frac{\epsilon_a^{10/3}}{\epsilon_T^2} \right) \quad (\text{if both air porosity and total porosity are known})$$

$$de = \left(\frac{4A}{\pi} \right)^{0.5}$$

$$D_e = D_a \epsilon_a^{4/3} \quad (\text{if only air porosity is known})$$

$$t_b = \frac{4.83 (10^7)}{B}$$

$$L = \frac{W f_{oil}}{A l} \quad (\text{for oily sludges}); L = \frac{W}{A l} \quad (\text{for dilute aqueous waste})$$

Variable	Definition	Data source
K_{eq}	Equilibrium coefficient of constituent in the soil (dimensionless)	Calculated

^aEquilibrium equations are adjusted to account for volume fractions of air and waste within the soil.

(continued)

TABLE 5-3 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
k_G	Gas-phase mass transfer coefficient (cm/sec)	Calculated
C	Concentration (weight fraction) of constituent in the oil phase or (for dilute aqueous waste) in water	Definition
D_a	Diffusion coefficient of constituent in air, cm^2/s	Data base
D_e	Effective diffusion coefficient of constituent in the soil, cm^2/s	Calculated
E	Emission rate of constituent, $\text{g}/\text{cm}^2/\text{s}$	Calculated
H_c	Henry's law constant for constituent, $\text{atm}\cdot\text{cm}^3/\text{g mol}$	Data base
l	Depth to which waste is mixed in the soil, cm	Facility operation
L	Oil or aqueous waste loading in the soil, g/cm^3	Calculated from annual throughput
M_t	Air emissions of constituent from the soil, g/cm^2	Calculated
M_o	Initial loading of constituent on the land treatment site, g/cm^2	Calculated
MW_{oil}	Average molecular weight of the oil, $\text{g}/\text{g mol}$	Estimated
MW	Molecular weight of constituent, $\text{g}/\text{g mol}$	Data base
P^*	Pure component vapor pressure of constituent, atm	Data base
R	Ideal gas constant, $82.1 \text{ atm}\cdot\text{cm}^3/\text{g mol}\cdot\text{K}$	Literature
t	Time after waste application to the land treatment site, s	Facility operation
t_b	Time constant for biological decay of constituent, s^b	Literature, or site specific

^bTime constant is the time required for biodegradation of 63.2 percent of a pollutant.

(continued)

TABLE 5-3 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
T	Temperature of vapor in soil, K	Assumed
ϵ_a	Volume fraction of air-filled voids in the soil (soil air-filled porosity) (dimensionless)	Estimated from literature data
ϵ_T	Total porosity of the soil (equivalent to dry basis bulk density divided by soil particle density) (dimensionless)	Estimated
ϵ_{waste}	Volume fraction of waste in the soil (dimensionless)	Calculated
B	Biorate of constituent, mg VO/g•h	Data base
K_v	Volatilization constant for constituent, s ⁻¹	Calculated
K_d	Modified volatilization constant, s ⁻¹	Calculated
K_b	Biodegradation constant for constituent, s ⁻¹	Calculated
F_a	Fraction of constituent emitted to the air after a long time	Calculated
F_{at}	Fraction of constituent emitted to the air at time t	Calculated
f_{oil}	Fraction by weight of applied waste that is oil (organic)	Definition
U	Windspeed, m/s	Estimated
W	Total waste applied to land treatment site, g	Definition
A	Area of land treatment site to which waste is applied, cm ² (m ² in calculation of de)	Definition
Sc _G	Schmidt number (gas phase)	Calculated
de	Effective diameter of land treatment area, m	Calculated
μ_a	Viscosity of air, g/cm•s	Literature
ρ_a	Density of air, g/cm ³	Literature

resistance of the soil and the resistance at the wind-porous solid interface. The mass transfer coefficient of the soil is defined in Equation (5-4) in the term $\left(\frac{K_{eq} D}{\pi t}\right)^{1/2}$. The resistance of the soil to mass transfer is the inverse of the above or $\left(\frac{K_{eq} D}{\pi t}\right)^{-1/2}$. The resistance at the air-soil interface is defined by $\frac{1}{K_{eq} k_G}$. Because K_{eq} has previously been defined as containing a factor to account for soil porosity, this factor (soil porosity) must be included in the above equation to maintain a consistent definition of K_{eq} throughout this discussion. The revised resistance is represented by $\frac{\epsilon_a}{K_{eq} k_G}$. Summing the two resistances and substituting into Equation (5-4) gives Equation (5-6). The gas-phase mass transfer coefficient, k_G , is calculated as described in Table 4-1 for a surface impoundment.

5.2.1.7 Estimating the Fraction Emitted at Short Times. The fraction of a constituent emitted to the air after some time, t , can be estimated by integrating the equation for air emissions from time 0 to time t :

$$F_{at} = \left[\frac{K_{eq} D_e}{\pi l^2} \right]^{1/2} \int_0^t t^{-1/2} e^{-t/t_b} dt$$

The exponential term can be replaced by a series,

$$e^{-t/t_b} = 1 - \frac{t}{t_b} + \frac{1}{2} \left(\frac{t}{t_b}\right)^2 - \frac{1}{6} \left(\frac{t}{t_b}\right)^3 + \frac{1}{24} \left(\frac{t}{t_b}\right)^4 - \dots$$

which can be substituted into the above integral, and each of the individual terms integrated. The results of these integrations are:

$$F_{at} = \left[\frac{K_{eq} D_e}{\pi l^2} \right]^{1/2} 2t^{1/2} \left[1 - \frac{1}{3} \left(\frac{t}{t_b}\right) + \frac{1}{10} \left(\frac{t}{t_b}\right)^2 - \frac{1}{42} \left(\frac{t}{t_b}\right)^3 + \dots \right]$$

This series solution converges with only a few terms for values of t/t_b less than 1. Therefore, the following simplification can be used to estimate the fraction emitted (i.e., integrated emissions) at short times:

$$F_{at} = \left[\frac{K_{eq} D_e}{\pi l^2} \right]^{1/2} 2t^{1/2} \left[1 - \frac{1}{3} \frac{t}{t_b} \right] \quad (5-7)$$

The resistance to emissions presented by gas-phase mass transfer at the soil surface is only considered important for the estimation of the emission rates immediately after application or tilling. This resistance is omitted in the above equation with little loss in accuracy.

The above equation is used to predict the fraction of a constituent emitted to the air when $K_v t$ is less than 0.22.

5.2.1.8 Estimating the Fraction Emitted for Longer Times. For longer times, when most of the constituent is not present in the soil, the short-term solution (Equations (5-4) and (5-7)) will overestimate air emissions. Under these conditions, Equation (5-5) can be integrated to estimate the fraction removed by volatilization. Equation (5-5) can be simplified by

defining the constant, K_d , as $\frac{K_{eq} D_e \pi^2}{4 l^2}$:

$$E = \frac{M_0 8 K_d}{\pi^2} \exp \left[- K_d t - t/t_b \right] \quad (5-8)$$

Integrating from time 0 to t gives:

$$F_{at} = \frac{8}{\pi^2} \left[1 + \frac{1}{K_d t_b} \right]^{-1} \left[1 - \exp \left[- K_d t - t/t_b \right] \right] + 0.1878 \quad (5-9)$$

In the above equation, terms after the first ($n > 0$) in the series solution are replaced by the constant 0.1878. This equation is used for estimating air emissions when $K_v t$ is greater than or equal to 0.22.

When $K_v t$ is less than 0.22, the following simplification can be used to estimate air emissions at longer times. An exponential decay factor is established to relate the fraction emitted at any time, t , to the fraction emitted at very long times (i.e., $t \rightarrow \infty$) as estimated using Equation (5-12), which follows. The resulting equation is:

$$F_{at} = F_a [1 - \exp (-K_d t - t/t_b)] \quad (5-10)$$

where

F_a = fraction of constituent emitted at very long times ($t \rightarrow \infty$).

For very long times (i.e., $t \rightarrow \infty$), the fraction emitted can be estimated using the following procedure. The integrated form of the general solution without dropping terms is:

$$F_{at} = \frac{8}{\pi^2} \sum_{n=0}^{\infty} \frac{1 - \exp\{-(2n+1)^2 K_d t - t/t_b\}}{(2n+1)^2 + \frac{1}{t_b K_d}}$$

This equation can be simplified using the following rationale: For large values of t , the exponential terms are negligibly small, and for large values of n , $1/(t_b K_d)$ becomes negligibly small compared to $(2n+1)^2$. If these conditions are true for all terms where $n > 0$, the simplified equation is:

$$F_a = \frac{8}{\pi^2} \left(\frac{K_d t_b}{K_d t_b + 1} + 0.2317 \right)$$

The value of 0.2317 was obtained by evaluating the first 125 terms of the series for $n > 0$ with negligibly small values of $1/(t_b K_d)$:

$$\sum_{n=1}^{125} \frac{1}{(2n+1)^2} = 0.2317$$

Combining terms and simplifying, the equation becomes:

$$F_a = \frac{0.81057 K_d t_b}{K_d t_b + 1} + 0.1878 \quad (5-11)$$

The assumptions used in developing Equation (5-11) are not valid for small values of $K_d t_b$ ($K_d t_b < 0.62$). The solution under these conditions is approximated by the following relationship:

$$F_a = \sqrt{\frac{K_{eq} D_e}{l^2} t_b} \quad (5-12)$$

This relationship was established by using multiple terms of the general solution to calculate values of F_a for a series of input values for the parameters $K_{eq} D_e / l^2$, which is identified as the volatilization constant,

K_v , and t_b and then using a curve-fitting routine to derive the relationship in Equation (5-12) for $d t_b < 0.62$.

Table 5-4 presents the results of calculations of the long-term fraction emitted (i.e., $t \rightarrow \infty$) using 100 terms of the general solution and inputting several values of the dimensionless ratio, $K_v t_b$, designated as I . This ratio is an indicator of the relative rates of volatilization and degradation. Table 5-4 also shows the results if the above approximating equations are used to calculate the long-term fraction emitted, and it shows good agreement between these results and the results obtained by the general solution.

Table 5-5 shows a comparison of the estimated emission fractions for a range of values of $K_v t$ and t/t_b using the first 100 terms of the general solution and using the approximations given in Equations (5-7) and (5-9). This table shows good agreement between the approximating equations and the rigorous solution.

To calculate the amount of waste remaining in the soil at any time, it is necessary to know both the amount emitted to the air and the amount biodegraded. At very long times (i.e., $t \rightarrow \infty$), all waste is assumed to disappear from the soil. Thus, the fraction of waste emitted plus the fraction biodegraded must be equal to 1 if other mechanisms of removal are ignored. Therefore, at very long times:

$$F_b = 1 - F_a, \quad (5-13)$$

where

F_b = fraction of constituent that is biodegraded after a long time (i.e., $t \rightarrow \infty$).

5.2.1.9 Tilling. To apply the model to a situation where the land treatment plot is retilled after the initial waste application and tilling, estimates of the amount of waste emitted to the air and the amount biodegraded are required. When retilling occurs, the amount of material remaining in the soil at the time of retilling is estimated using the following equation:

$$F_s = (1 - F'_{at}) e^{-t/t_b} \quad (5-14)$$

TABLE 5-4. ESTIMATED AIR EMISSION FRACTION AT LONG TIMES

Value of T $[T = K_v t_b]$	Estimated fraction (rigorous equation)	Estimated fraction ^a	Estimated fraction ^b
0.050	0.222	0.224	0.277
0.100	0.313	0.316	0.348
0.150	0.381	0.387	0.407
0.200	0.435	0.447	0.456
0.250	0.480	0.500	0.497
0.300	0.518	0.548	0.533
0.350	0.551	0.592	0.563
0.400	0.579	0.632	0.590
0.450	0.604	0.671	0.614
0.500	0.626	0.707	0.635
0.550	0.646	0.742	0.654
0.600	0.664	0.775	0.672
0.650	0.680	0.806	0.687
0.700	0.694	0.837	0.701
0.750	0.708	0.866	0.714
0.800	0.720	0.894	0.725
0.850	0.731	0.922	0.737
0.900	0.741	0.949	0.747
0.950	0.751	0.975	0.750
1.000	0.760	1.000	0.765
1.050	0.768	1.025	0.773
1.100	0.776	1.049	0.780
1.150	0.783	1.072	0.787
1.200	0.789	1.095	0.794
1.250	0.796	1.118	0.800
1.300	0.802	1.140	0.805
1.350	0.807	1.162	0.811
1.400	0.813	1.183	0.816
1.450	0.818	1.204	0.821
1.500	0.822	1.225	0.826
1.550	0.827	1.245	0.830
1.600	0.831	1.265	0.834
1.650	0.835	1.285	0.839
1.700	0.839	1.304	0.842
1.750	0.843	1.323	0.846
1.800	0.846	1.342	0.849
1.850	0.850	1.360	0.853
1.900	0.853	1.378	0.856
1.950	0.856	1.396	0.859
2.000	0.859	1.414	0.862

$$a_{Fa} = T^{0.5}$$

$$b_{Fa} = \frac{0.81057 K_d t_b}{K_d t_b + 1} + 0.1878$$

TABLE 5-5. RIGOROUS VS. APPROXIMATE ESTIMATES OF EMISSION FRACTIONS

t/t_b	$K_v t$	$K_v t_b$	Estimated fraction (rigorous)	Estimated fraction (approximated by) ^a	Estimated fraction (approximated by) ^b
0.01	0.05	5.00	0.25	0.25	0.28
0.01	0.10	10.00	0.35	0.36	0.36
0.01	0.15	15.00	0.43	0.44	0.44
0.01	0.20	20.00	0.50	0.50	0.50
0.01	0.25	25.00	0.56	0.56	0.56
0.01	0.30	30.00	0.61	0.62	0.61
0.01	0.35	35.00	0.65	0.67	0.65
0.01	0.40	40.00	0.69	0.71	0.69
0.01	0.45	45.00	0.73	0.75	0.73
0.01	0.50	50.00	0.76	0.79	0.76
0.01	0.55	55.00	0.79		0.79
0.01	0.60	60.00	0.81		0.81
0.01	0.65	65.00	0.83		0.83
0.01	0.70	70.00	0.85		0.85
0.01	0.75	75.00	0.87		0.87
0.01	0.80	80.00	0.88		0.88
0.01	0.85	85.00	0.90		0.90
0.01	0.90	90.00	0.91		0.91
0.01	0.95	95.00	0.92		0.92
0.01	1.00	100.00	0.93		0.93
0.10	0.05	0.50	0.24	0.24	0.28
0.10	0.10	1.00	0.34	0.34	0.36
0.10	0.15	1.50	0.42	0.42	0.43
0.10	0.20	2.00	0.49	0.49	0.49
0.10	0.25	2.50	0.54	0.55	0.54
0.10	0.30	3.00	0.59	0.60	0.59
0.10	0.35	3.50	0.64	0.65	0.64
0.10	0.40	4.00	0.67	0.69	0.68
0.10	0.45	4.50	0.71	0.73	0.71
0.10	0.50	5.00	0.74	0.77	0.74
0.10	0.55	5.50	0.77		0.77
0.10	0.60	6.00	0.79		0.79
0.10	0.65	6.50	0.81		0.81
0.10	0.70	7.00	0.83		0.83
0.10	0.75	7.50	0.85		0.85
0.10	0.80	8.00	0.86		0.86
0.10	0.85	8.50	0.87		0.88
0.10	0.90	9.00	0.89		0.89
0.10	0.95	9.50	0.90		0.90
0.10	1.00	10.00	0.91		0.91
0.30	0.05	0.17	0.23	0.23	0.27
0.30	0.10	0.33	0.32	0.32	0.34
0.30	0.15	0.50	0.39	0.39	0.41
0.30	0.20	0.67	0.46	0.45	0.46
0.30	0.25	0.83	0.51	0.51	0.52

See notes at end of table.

(continued)

TABLE 5-5 (continued)

t/t_b	$K_v t$	$K_v t_b$	Estimated fraction (rigorous)	Estimated fraction (approximated by) ^a	Estimated fraction (approximated by) ^b
0.30	0.30	1.00	0.56	0.56	0.56
0.30	0.35	1.17	0.60	0.60	0.60
0.30	0.40	1.33	0.63	0.64	0.64
0.30	0.45	1.50	0.67	0.68	0.67
0.30	0.50	1.67	0.70	0.72	0.70
0.30	0.55	1.83	0.72		0.73
0.30	0.60	2.00	0.75		0.75
0.30	0.65	2.17	0.77		0.77
0.30	0.70	2.33	0.79		0.79
0.30	0.75	2.50	0.80		0.80
0.30	0.80	2.67	0.82		0.82
0.30	0.85	2.83	0.83		0.83
0.30	0.90	3.00	0.84		0.84
0.30	0.95	3.17	0.85		0.86
0.30	1.00	3.33	0.86		0.87
1.00	0.05	0.05	0.19	0.17	0.25
1.00	0.10	0.10	0.26	0.24	0.30
1.00	0.15	0.15	0.32	0.29	0.35
1.00	0.20	0.20	0.37	0.34	0.40
1.00	0.25	0.25	0.42	0.38	0.44
1.00	0.30	0.30	0.46	0.41	0.47
1.00	0.35	0.35	0.49	0.44	0.51
1.00	0.40	0.40	0.52	0.48	0.54
1.00	0.45	0.45	0.55	0.50	0.56
1.00	0.50	0.50	0.58	0.53	0.59
1.00	0.55	0.55	0.60		0.61
1.00	0.60	0.60	0.62		0.63
1.00	0.65	0.65	0.64		0.65
1.00	0.70	0.70	0.66		0.67
1.00	0.75	0.75	0.68		0.68
1.00	0.80	0.80	0.69		0.70
1.00	0.85	0.85	0.71		0.71
1.00	0.90	0.90	0.72		0.72
1.00	0.95	0.95	0.73		0.74
1.00	1.00	1.00	0.74		0.75

^aApproximated by: $F_{at} = 1.128 \sqrt{K_v t} (1 - 1/3 t/t_b)$.

^bApproximated by: $F_{at} = \frac{8}{\pi^2} \left(1 + \frac{1}{K_d t_b} \right)^{-1} \left[1 - \exp \left[-K_d t - t/t_b \right] \right] + 0.1878$.

where

F_S = fraction of constituent remaining in the soil

F'_{at} = fraction of material emitted to the air at time t assuming no biodegradation (F'_{at} can be estimated by setting $t/t_b = 0$ in Equation (5-7) or (5-9), whichever is appropriate).

To continue modeling emissions after retilling occurs, M_o is set equal to $F_S M_o$ and t is reset to zero. If a reapplication of waste occurs, the total waste loading is the sum of the waste remaining in the soil and the newly applied waste:

$$M_o = F_S M_o + M_n \quad (5-15)$$

where

M_n = amount of constituent newly applied to the land treatment site.

To continue the modeling after waste reapplication and tilling, t is reset to zero.

5.2.1.10 Model Selection. The RTI model was selected for use in this regulatory effort after a review of three models of land treatment emissions. The models reviewed were the RTI model, the Thibodeaux-Hwang model, and a model developed by EPA's Office of Research and Development located in Ada, Oklahoma (the Ada model). The review considered three selection criteria: technical basis, representativeness, and availability of inputs.

The Ada model is the most ambitious of the three in attempting to account for mechanisms of pollutant removal other than air emissions and biodegradation. However, that model requires detailed site-specific model inputs that may not be available or reasonably estimated. Because of these characteristics of the Ada model, it was not considered appropriate for use in the current effort. Both the Thibodeaux-Hwang and the RTI models have input requirements that are reasonably available, both have been compared with available measured data, and both have shown reasonable agreement with the measurements.⁸ Apparently, either of these two models is satisfactory as a means of estimating emission rates at specific times for some organic compounds. However, if the Thibodeaux-Hwang model is used to estimate long-term, steady-state emissions, it would predict that all of the applied

volatile organics are emitted because it does not account for biodegradation. Such a prediction would contradict data obtained from laboratory and field studies that indicate biodegradation of some organic compounds in land treatment applications.⁹ The RTI model, in contrast, estimates biodegradation of individual compounds based on constituent-specific biodegradation rates. The RTI and the Thibodeaux Hwang models predict similar emission rates for initial volatile losses in the absence of biodegradation. Thus, the results of the RTI model show varying levels of biodegradation when used to evaluate the fate of different organic compounds.

In summary, the Ada model has had limited public review, accounts for multiple waste removal mechanisms, requires numerous detailed model inputs, and has no published comparisons of estimated and measured emissions. The Thibodeaux-Hwang model has been publicly reviewed, accounts for one major waste removal mechanism (volatilization), requires reasonably available model inputs, and there are published comparisons of measured and estimated emissions. The RTI model has had limited public review, accounts for the two major waste removal mechanisms (volatilization and biodegradation), requires reasonably available model inputs, and there are published comparisons of measured and estimated emissions. The peer review, emission comparisons, and data availability are compelling reasons for using the Thibodeaux-Hwang model. However, the absence of biodegradation in that model is considered a major shortcoming because of the importance attached to this removal mechanism by industry personnel and many independent researchers. Emission comparisons and data availability for the RTI model are roughly equivalent to those of the Thibodeaux-Hwang model, and the RTI model includes terms that account for biodegradation. The RTI and the Thibodeaux models predict similar emission rates for initial volatile losses in the absence of biodegradation. Thus, the RTI model was selected for use in the standards development process.

5.2.2 Waste Application Model

At land treatment facilities that do not use subsurface injection with immediate tilling, emissions may occur during the time that waste is being applied to the soil surface and while the waste lies on the soil before it is tilled into the soil. No existing models were identified that predict

emissions during application of an oily sludge to the soil surface. The approach selected for this case was to calculate an overall mass transfer coefficient of volatile material from the surface of the stream of sludge as it falls from the end of a hose to the soil surface. The mass transfer coefficients were calculated using an equation presented in Section 4.0 (Table 4-1). The constant in the equation for gas-phase resistance was increased by a factor of two in an attempt to account for an increase in mass transfer caused by the motion of the waste stream through the air. The equations for making this calculation are presented in Table 5-6 along with the definitions of the variables used and the sources of input data.

5.2.3 Oil Film Model

Emissions from waste lying on the soil surface are estimated in either of two ways depending on the form of the waste as it lays on the soil surface. In typical situations where the applied waste is spread over the surface of soil, the RTI land treatment model can be used to estimate emissions. The equation for short-term emissions given above as Equations (5-4) and (5-7) would be used for this situation. If the applied waste has a visible oil film on top, emissions immediately after spreading are estimated by calculating an overall mass transfer coefficient as described in Chapter 4.0 for an oil film on a surface impoundment. The mass transfer equation was developed by McKay and Matsuga and is based on data obtained from liquid hydrocarbon spills on land and water.¹⁰ The equations used to calculate emissions under this situation are given in Table 5-7 along with definitions of the variables used.

5.2.4 Model Inputs

Typical values of input parameters for the RTI model are based primarily on a data base developed by EPA¹¹ from site visits and contacts with State, regional, and industry sources supplemented by information from recent literature. These values were chosen as reasonably representative of average or typical practices currently used at land treatment operations. Oil loading in the soil is a model input that is calculated from several other parameters that might change independently. Varying the value of the oil loading rate, thus, has the same effect as varying any one or any combination of the other parameters. Oil loading is defined by waste throughput, the percent oil in the waste, area of the land treatment

TABLE 5-6. WASTE APPLICATION EMISSION MODEL

Emission equations

$$E = KCA$$

$$A = 2\pi r l$$

$$K = K_{eq} k_G \text{ (used for oily sludges); } \frac{1}{K} = \frac{1}{k_L} + \frac{1}{K_{eq} k_G} \text{ (used for dilute aqueous waste);}$$

$$k_G = 9.64(10^{-3})U^{0.78}Sc_G^{-0.67}de^{-0.11};$$

$$K_{eq} = \frac{H}{RT} \text{ (used for dilute aqueous waste)}$$

$$K_{eq} = \frac{p^*}{p_o} \frac{\rho_a}{\rho_L} \frac{MW_{oil}}{MW_{air}} \text{ (used for oily sludges)}$$

$$k_L = 1(10^{-6}) + 144(10^{-4})U^{*2.2}Sc_L^{-0.5}; Sc_L = \frac{\mu_L}{\rho_w D_w}$$

$$de = \left(\frac{4A}{\pi}\right)^{0.5}; Sc_G = \frac{\mu_a}{\rho_a Da}$$

$$U^* = 0.01U(6.1 + 0.63U)^{0.5}$$

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
E	Emission rate for constituent, g/s	Calculated
K	Overall mass transfer coefficient, m/s	Calculated
K _{eq}	Equilibrium coefficient, dimensionless	Calculated
H	Henry's law constant for constituent, atm cm ³ /g mol	Literature
R	Universal gas constant, atm cm ³ /g mol K	Literature
T	Temperature, K	Measured
p*	Vapor pressure of constituent, mm Hg	Literature
p _o	System pressure (atmospheric pressure), mm Hg	Definition

(continued)

TABLE 5-6 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
k_G	Gas-phase mass transfer coefficient, m/s	Calculated
U	Windspeed, m/s	Definition
Sc_G	Schmidt number on gas side	Calculated
Sc_L	Schmidt number on liquid side	Calculated
μ_a	Viscosity of air, g/cm \cdot s	Literature
ρ_w	Density of water, g/cm 3	Literature
ρ_a	Density of air, g/cm 3	Literature
ρ_L	Density of oil, g/cm 3	Estimated
D_a	Diffusivity of constituent in air, cm 2 /s	Literature
A	Surface area of cylindrical waste stream, m 2	Calculated
r	Radius of cylindrical waste stream, m	Measured
l	Length of cylindrical waste stream, m	Measured
C	Concentration of constituent in the waste, g/cm 3	Measured
d_e	Effective diameter of waste stream surface area, m	Calculated
MW_a	Molecular weight of air, g/g mol	Literature
MW_{oil}	Molecular weight of oil, g/g mol	Estimated
μ_L	Viscosity of water, g/cm \cdot s	Literature
D_w	Diffusivity of component in water, cm 2 /s	Data base
U^*	Friction velocity, m/s	Calculated
k_L	Liquid-phase mass transfer coefficient, m/s	Calculated

TABLE 5-7. OIL FILM SURFACE EMISSION MODEL

Emission rate equation

$$E = KC_t A$$

$$C_t = C_o [\exp (-Kt/D)]$$

$$K = k_G \text{ Keq (used for oily sludges)}$$

$$k_G = 4.82 (10^{-3}) U^{0.78} Sc_G^{-0.67} de^{-0.11}$$

$$Sc_G = \frac{\mu_a}{\rho_a D_a}$$

$$\text{Keq} = \frac{P^*}{P_o} \frac{\rho_a^{MW} \text{ oil}}{\rho_L^{MW} \text{ air}} \text{ (used for oily sludges);}$$

$$de = \left(\frac{4A}{\pi} \right)^{0.5}$$

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
E	Emission rate for constituent, g/s	Calculated
K	Overall mass transfer coefficient, m/s	Calculated
C _t	Concentration of constituent in the oil phase at time t	Calculated
C _o	Initial concentration of constituent in the waste	Definition
D	Oil film thickness, m	Measured
A	Area of land treatment, m ²	Measured
k _G	Gas-phase mass transfer coefficient, m/s	Calculated

(continued)

TABLE 5-7 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
U	Windspeed, m/s	Definition
Sc _G	Schmidt number--gas phase	Calculated
μ_a	Viscosity of air, g/cm•s	Literature
ρ_a	Density of air, g/cm ³	Literature
D _a	Diffusion coefficient of constituent in air, cm ² /s	Literature
d _e	Effective diameter of land treatment area, m	Calculated
K _{eq}	Equilibrium coefficient of constituent	Calculated
p*	Vapor pressure of constituent, mm Hg	Literature
P ₀	Atmospheric pressure, mm Hg	Definition
MW _{oil}	Molecular weight of the oil, g/g mol	Definition
MW _a	Molecular weight of air, g/g mol	Literature
ρ_L	Density of oil, g/cm ³	Estimated
R	Universal gas constant, atm cm ³ /g mol K	Literature
T	Temperature, K	Measured

site, and the depth to which the waste is mixed in the soil (tilling depth). Typical values of oil loading are defined from median values for those parameters by which it is defined. The data base shows annual throughput varying from about 2 Mg/yr to about 400,000 Mg/yr with a median value of about 1,800 Mg/yr. The area of land treatment sites ranges from less than 1 hectare (ha) to about 250 ha with a median value of 5 ha. The data base shows tilling depth varying from 15 cm to one case of 65 cm, with most being in the range of 15 to 30 cm. The single most frequently reported tilling depth is 20 cm, which is selected as a typical value. This value is in line with values of 15 to 30 cm reported in another study.¹² The data base shows oil content of the waste streams varying from about 2 to 50 percent, with a median value of about 12 percent and a mode value of 10 percent. The 10-percent figure is selected as typical.

Average molecular weight of the oil from which a particular constituent evaporates is one of the determining factors in the rate of evaporation and thus must be specified. Little data are available as guidance for selecting a value for this parameter. The distribution of constituents by molecular weight in land-treated wastes is not well known. In one field measurement study of land treatment emissions,¹³ a value of 282 g/g mol was used as the average molecular weight of the oil. This value was based on distillation of oil from a refinery sludge and identification of the constituent corresponding to the midpoint distillation temperature (i.e., the temperature at which 50 percent of the oil was distilled). The value 282 g/g mol is selected for use. A sensitivity analysis using the RTI model shows that emissions are not highly sensitive to this parameter.

Soil air porosity and total porosity impact the effective diffusivity of a constituent in the soil. Very little soil porosity information has been identified. One study reported measured values of soil porosity in a land treatment plot as ranging from 43.3 to 65.1 percent,¹⁴ with an average value of about 50 percent. The literature values are assumed to represent air porosity. Total soil porosity would include the air porosity and the space occupied by oil and water within the soil. One field study reported measured values of both total porosity and air-filled porosity.¹⁵ Measured values of total porosity ranged from 54.7 to 64.8 percent, with an average

value of 60.7 percent. Measured values of air-filled porosity ranged from 27.4 to 46.9 percent, with an average of 37.2 percent. The value of 61 percent for total porosity is assumed to be a representative value. A value of 0.5 is used in the model as a default for air porosity.

Biorate data used in the RTI model data base (CHEMDAT7) represent measured rates in aqueous systems. In order to use the aqueous biorate data in a land treatment process, a factor was established for converting aqueous data to land treatment values using measured data for benzene. A recent publication by the American Petroleum Institute (API) reported experimentally determined values of biological decay constants for land treatment studies using two different soil types.¹⁶ Decay constants were measured for six compounds including two compounds, benzene and toluene, that have aqueous biorates in the land treatment model data base (CHEMDAT7). For benzene, the ratio of the API data, measured in units of day^{-1} , and the aqueous data, measured in units of $\text{mg VO/g biomass} \cdot \text{hr}$, was calculated as 0.00179. This value is also a close approximation of the ratio of the two data points for toluene, the other compound for which data from both sources were available. The other compounds for which data were reported by API did not have referenced aqueous data in the data base. The above calculated relationship was used to calculate equivalent aqueous data values for those compounds. Reported and calculated values of aqueous biorates and land treatment biological decay constants are presented in Table 5-8. The ratio of 0.00179 is used for all compounds to convert from aqueous biorates to decay constants that can be used in the land treatment model. The input parameter for the land treatment model is a biological decay time constant, t_b , in units of seconds. The equation for calculating t_b from the aqueous biorate is derived as follows.

The biological decay time constant is, by definition, equal to the reciprocal of the biological decay constant, or

$$t_b = \frac{1}{K_b} \quad (5-16)$$

where K_b = biological decay constant. The ratio, r , of decay constant to aqueous biorate is:

TABLE 5-8. MEASURED AND ESTIMATED BIORATES AND DECAY CONSTANTS
FOR SELECTED ORGANIC CONSTITUENTS

Organic constituent	Aqueous biorate, mg VO/gbiomass•h	Calculated decay constant, day ⁻¹	Measured decay constant, day ^{-1a}	
			Nunn ^b	Kidman ^c
Benzene	19.0	0.034	0.034	0.013
Ethylbenzene	46.4 ^d	0.083	0.083	0.076
Xylene(-o)	40.8 ^d	0.073	0.073	0.026
Naphthalene	42.5 ^d	0.076	0.076	0.050
Toluene	73.5	0.132	0.106	0.119
Methyl naphthalene	24.0 ^d	0.043	0.043	0.059

^aReference 17.

^bData obtained using a clay loam soil (Nunn soil).

^cData obtained using a sandy loam soil (Kidman soil).

^dValues calculated from API experimental data.

$$r = \frac{K_b}{B} \frac{g_{bio} h}{mg \cdot VO \cdot day} \text{ or } K_b = rB \text{ day}^{-1} .$$

Substituting into Equation (5-16) gives:

$$t_b = \frac{1}{rB} \text{ day}$$

To obtain a result in seconds, this equation must be multiplied by 86,400 s/day. Making this conversion and inserting the value of r (i.e., 0.00179) gives:

$$t_b = \frac{86,400}{0.00179 B} = \frac{4.83 (10^7)}{B} .$$

For situations in which petroleum wastes are landfarmed and no information is known about the nature of the volatile materials, it is possible to estimate a default biorate from the average decay constant values reported in the API investigation, 0.07 day^{-1} , which corresponds to a biorate, B , of 40 mg/g-h . This value is between the values for benzene and toluene in the data base. The average value of the biological rate constant in the two soils investigated by API was not significantly different.

In summary, parameters and selected typical values for use in the RTI model are as follows:

Annual waste throughput	= 1,800 Mg
Area of land treatment	= 5 ha
Oil content of waste	= 10 percent
Average molecular weight of the oil	= 282 g/g mol
Soil air porosity	= 0.5
Soil total porosity	= 0.61
Tilling depth	= 20 cm
Temperature	= 25 °C.

5.2.5 Estimation of Total VO Emissions

The preceding discussion has been limited to estimating emissions of a single constituent in a hazardous waste. Using the models presented here to estimate total VO emissions can be accomplished using any of several approaches. The most obvious approach, and the one that should give the most accurate results, would be to obtain a detailed analysis of the constituents in the waste being land treated. The emission equations could be used to calculate emissions of each constituent, and total emissions could be calculated by summing the emissions of individual constituents. In many cases, a detailed analysis of the applied waste may not be available, and other, less accurate methods may be needed to estimate total VO emissions. An alternative to the constituent approach could make use of a boiling curve or steam stripping test of a sample of the waste. Experimental data developed by Chevron Research Company¹⁸ indicate that a large fraction of the constituents that boil at temperatures of 400 °F or lower will be emitted to the atmosphere and that those constituents with higher boiling points will tend to remain in the soil for a sufficient time to undergo biodegradation. Similar results are obtained by applying the RTI land treatment model to the constituents in the CHEMDAT7 data base. If a sample of waste were subjected to a laboratory boiling test or other equivalent test at a temperature of 400 °F, the fraction of oil evaporated would approximate the fraction that potentially would be emitted to the air in a land treatment operation.

A third approach to estimating total VO emissions would again make use of the experimental results generated by Chevron Research. The test results showed that approximately 25 percent of the applied oil in the land treatment test was emitted to the air. In the absence of a detailed constituent analysis and with no boiling or steam stripping test of the waste, a crude estimate of total VO emissions could be made by assuming that emissions are equal to 25 percent of the applied oil. This approximating alternative would only apply to raw oily refinery wastes that have not undergone any pretreatment to remove VO.

5.2.6 Example Calculations

5.2.6.1 Emissions from Land Treatment Soil. The following calculation demonstrates the use of the RTI model to calculate the long-term

fraction of applied material emitted to the atmosphere and to calculate the short-term and long-term emission rates and emission fractions. The calculations are made for benzene at a concentration of 2,000 ppm by weight in a waste stream that is 10 percent oil.

Input values are:

Land area	= 2.5 ha (half of total area of 5 ha assumed active)
Annual throughput	= 1,800 Mg
Oil content of waste	= 10% (by weight)
Benzene concentration in oil	= 2,000 ppm (by weight) (2 mg/g oil)
Tilling depth	= 20 cm
Soil air porosity	= 0.5
Soil total porosity	= 0.61
Average molecular weight of oil	= 282 g/g mol

- a. Calculate oil loading (Equation from Table 5-3):

$$L = \frac{(1,800 \times 10^6 \text{ g}_{\text{waste}})(0.1 \text{ g}_{\text{oil}}/\text{g}_{\text{waste}})}{(2.5 \times 10^8 \text{ cm}^2)(20 \text{ cm})} = 0.036 \text{ g}_{\text{oil}}/\text{cm}^3$$

- b. Identify constituent properties:

Benzene properties:

$$B = 19.00 \text{ mg VO/gbiomass}\cdot\text{h}$$

$$D_a = 0.088 \text{ cm}^2/\text{s}$$

$$P^* = 95.2 \text{ mm Hg} = 0.125 \text{ atm.}$$

- c. Calculate the equilibrium coefficient (Equation from Table 5-3):

$$K_{eq} = \frac{P^* \text{ MW}_{\text{oil}} \epsilon_a}{RT L} = \frac{(0.125 \text{ atm})(282 \text{ g/g mol})(0.5 \text{ cm}^3/\text{cm}^3)}{(82.05 \text{ atm}\cdot\text{cm}^3/\text{g mol}\cdot\text{K})(298 \text{ K})(0.036 \text{ g/cm}^3)}$$

$$K_{eq} = 0.02002$$

- d. Calculate the biological degradation time constant (Equation from Table 5-3):

$$t_b = \frac{4.83(10^7)}{8} = \frac{4.83(10^7)}{19} = 2.54(10^6) \text{ s} .$$

- e. Calculate the effective diffusivity of constituent in the soil (Equation from Table 5-3):

$$D_e = D_a \frac{\epsilon_a^{10/3}}{\epsilon_T^2} = 0.088 \text{ cm}^2/\text{s} \frac{(0.5)^{10/3}}{0.61^2} = 0.0235 \text{ cm}^2/\text{s} .$$

- f. Calculate the value of $K_d = \frac{\pi^2 K_{eq} D_e}{4 l^2}$:

$$K_d = \frac{(9.87)(0.0235) \text{ cm}^2/\text{s} (0.02002)}{4 (400) \text{ cm}^2} = 2.9(10^{-6}) \text{ s}^{-1} .$$

- g. Calculate the fraction of constituent emitted to the air after a long time (Equation (5-11)):

$$F_a = \frac{0.81 K_d t_b}{K_d t_b + 1} + 0.1878 = 0.90 .$$

- h. Calculate the long-term emission rate after 60 h (216,000 s).

$$K_{eq} \frac{D_e t}{l^2} = \frac{0.02002 \times 0.0235 \times 216,000}{400} = 0.254 .$$

Use Equation (5-5) (long-term equation):

$$E = M_o \frac{2 K_{eq} D_e}{l^2} e \left[\frac{\pi^2 K_{eq} D_e t}{4 l^2} \right] e^{-t/t_b}$$

$$\begin{aligned} E &= \frac{2 \times 0.72 \times 2 \times 0.02002 \times 0.0235}{400} \times e \left[\frac{-0.02002 \times 0.0235 \times 9.87 \times 216,000}{1,600} \right] \\ &\quad \times e[-216,000/2.54(10^6)] \\ &= 3.38(10^{-6}) e(-0.627) e(-0.085) = 3.38(10^{-6}) e(-0.712) \\ E &= 3.38(10^{-6}) (0.491) = 1.7(10^{-6}) \frac{\text{mg}}{\text{cm}^2 \text{ s}} \end{aligned}$$

c. Calculate the short-term emission rate after 15 min (900 s):

$$K_{eq} \frac{D_e t}{l^2} = \frac{0.02002 \times 0.0235 (900)}{400} = 0.0010$$

$$k_G = 4.82 (10^{-3}) U^{0.78} Sc_G^{-0.67} de^{-0.11} ,$$

where

$$U = \text{windspeed} = 4.47 \text{ m/s}$$

de = effective diameter of land treatment area

$$de = \left(\frac{4A}{\pi} \right)^{0.5} = 178 \text{ m}$$

$$Sc_G = \frac{\mu_g}{\rho_a D_a}$$

where

$$\mu_g = \text{viscosity of air} = 1.81(10^{-4}) \text{ g/cm}\cdot\text{s}$$

$$\rho_a = \text{density of air} = 1.2 (10^{-3}) \text{ g/cm}^3$$

$$D_a = 0.088 \text{ cm}^2/\text{s}$$

$$Sc_G = \frac{1.81 (10^{-4})}{1.2 (10^{-3}) (0.088)} = 1.71$$

$$k_G = 4.82(10^{-3}) (4.47)^{0.78} (1.71)^{-0.67} (178)^{-0.11} = 0.0061 \text{ m/s} = 0.61 \text{ cm/s}$$

$$E = \frac{M_0}{l} \left[\frac{1}{\frac{\epsilon_a}{k_G K_{eq}} + \sqrt{\frac{\pi t}{D_e K_{eq}}}} \right] e^{-t/t_b}$$

$$E = \frac{0.72 \text{ g} \cdot 2 \text{ mg}}{1 \text{ cm}^2 \cdot 20 \text{ cm} \cdot \text{g}} \left[\frac{1}{\frac{0.5}{0.61 \times 0.02002} + \sqrt{\frac{3.14 \times 900}{0.0235 \times 0.02002}}} \right] e^{(-900/2.54(10^6))}$$

$$= 0.072 (0.0004) e^{(-0.0004)} = 2.87 (10^{-5}) \frac{\text{mg}}{\text{cm}^2 \cdot \text{s}} .$$

Table 5-9 shows estimated emission rates and emission fractions for various times up to 40 days (960 hours).

5.2.6.2 Emissions from Waste Application. The following is an example calculation for the application of an oily waste to a land treatment plot using the equations in Table 5-6. For benzene in waste oil, the calculation is:

Input values:

$$\begin{aligned}
 r &= 0.038 \text{ m} \\
 L &= 0.46 \text{ m} \\
 \mu_a &= 1.81 (10^{-4}) \text{ g/cm}\cdot\text{s} \\
 \rho_a &= 1.2 (10^{-3}) \text{ g/cm}^3 \\
 U &= 4.47 \text{ m/s} \\
 R &= 82.05 \text{ atm}\cdot\text{cm}^3/\text{K}\cdot\text{g mol} \\
 T &= 298 \text{ K} \\
 C &= 200 \text{ ppm} = 200 \mu\text{g/g} = 0.0002 \text{ g/cm}^3 = 200 \text{ g/m}^3 \\
 &\quad (\text{assuming a density of } 1 \text{ g/cm}^3) \\
 A &= 2 \pi r L = 2(3.14)(0.038 \text{ m})(0.457 \text{ m}) = 0.11 \text{ m}^2 \\
 \rho_L &= 1 \text{ g/cm}^3 \\
 MW_a &= 29 \text{ g/g mol} \\
 MW_{oil} &= 282 \text{ g/g mol}.
 \end{aligned}$$

- a. Calculate the effective diameter of the waste stream surface (Equation from Table 5-6):

$$d_e = \left(\frac{4A}{\pi} \right)^{0.5} = 0.37 \text{ m}.$$

- b. Calculate the Schmidt number (Equation from Table 5-6):

$$Sc_G = \frac{\mu_a}{\rho_a D_a} = \frac{1.81 (10^{-4}) \text{ g/cm}\cdot\text{s}}{[1.2 (10^{-3}) \text{ g/cm}^3] (0.088 \text{ cm}^2/\text{s})} = 1.71.$$

- c. Calculate the equilibrium coefficient (Equation from Table 5-6):

$$K_{eq} = \frac{P^*}{P_o} \frac{\rho_a MW_{oil}}{\rho_L MW_a} = \frac{(95.2 \text{ mm Hg}) [1.2 (10^{-3}) \text{ g/cm}^3] (282 \text{ g/g mol})}{(760 \text{ mm Hg}) (1 \text{ g/cm}^3) (29 \text{ g/g mol})} = 0.0015$$

TABLE 5-9. ESTIMATED EMISSION RATES AND FRACTIONS EMITTED
VERSUS TIME FOR EXAMPLE LAND TREATMENT CALCULATION

Time after application/tilling, h	Emission rate, 10^{-6} mg/cm ² •s	Equation used	Fraction emitted, fraction	Equation used
1	14.4	Short term	0.073	Short term
2	10.3	Short term	0.104	Short term
4	7.30	Short term	0.147	Short term
8	5.12	Short term	0.207	Short term
24	2.90	Short term	0.356	Short term
48	1.98	Short term	0.497	Short term
96	1.08	Long term	0.674	Long term
192	0.348	Long term	0.827	Long term
480	0.011	Long term	0.899	Long term
960	0.00004	Long term	0.901	Long term

- d. Calculate the gas-phase mass transfer coefficient (Equation from Table 5-6):

$$\begin{aligned} k_G &= 9.64(10^{-3}) U^{0.78} Sc_G^{-0.67} de^{-0.11} \\ &= 9.64(10^{-3})(4.47)^{0.78}(1.71)^{-0.67}(0.37)^{-0.11} \\ &= 0.024 \text{ m/s} \end{aligned}$$

- e. Calculate the overall mass transfer coefficient (Equation from Table 5-6):

$$K = k_G K_{eq} = (0.0015)(0.024) \text{ m/s} = 0.000036 \text{ m/s}$$

- f. Calculate emissions from a unit volume of waste (Equation from Table 5-6):

$$\begin{aligned} E &= KCA = 0.000036 \text{ m/s} (200 \text{ g/m}^3) (0.11) \text{ m}^2 \\ &= 7.92 (10^{-4}) \text{ g/s} \end{aligned}$$

Using a calculated fall time of 0.305 s:

$$E = (7.92 \cdot 10^{-4} \text{ g/s})(0.30 \text{ s}) = 2.38 (10^{-4}) \text{ g}$$

$$\begin{aligned} \text{Stream volume} &= \pi r^2 L = (3.14)(0.038 \text{ m})^2(0.46 \text{ m}) = 0.002 \text{ m}^3 \\ &= 2,073 \text{ cm}^3 \end{aligned}$$

$$\text{Mass of constituent} = (0.002 \text{ m}^3) (200 \text{ g/m}^3) = 0.4 \text{ g}$$

- g. Calculate the fraction of constituent emitted to the air:

$$\text{Fraction emitted} = \frac{2.38(10^{-4}) \text{ g}}{0.4 \text{ g}} = 0.00059 = 0.06\%$$

5.2.6.3 Emissions from an Oil Layer on Soil Prior to Tilling. An example calculation for butanol-1 in an oil layer on the soil surface of a land treatment site is given below using the equation from Table 5-7.

Input values:

$$\begin{aligned} \mu_a &= 1.81 (10^{-4}) \text{ g/cm} \cdot \text{s} \\ &= 1.2 (10^{-3}) \text{ g/cm}^3 \\ &= 4.47 \text{ m/s} \end{aligned}$$

$$\begin{aligned}
 MW_{oil} &= 282 \text{ g/g mol} \\
 \rho_L &= 1 \text{ g/cm}^3 \\
 MW_a &= 29 \text{ g/g mol} \\
 C &= 0.0002 \text{ g/cm}^3 = 200 \text{ g/m}^3 \\
 A &= 25,000 \text{ m}^2 \\
 t &= 24 \text{ h} = 86,400 \text{ s} \\
 d &= 0.072 \text{ m}
 \end{aligned}$$

- a. Calculate the effective diameter of the soil surface (Equation from Table 5-7):

$$d_e = \left(\frac{4A}{\pi} \right)^{0.5} = \left(\frac{4 \times 25,000}{\pi} \right)^{0.5} = 178 \text{ m}$$

- b. Calculate the Schmidt number (Equation from Table 5-7):

$$Sc_G = \frac{\mu_a}{\rho_a D_a} = \frac{1.81 (10^{-4}) \text{ g/cm}\cdot\text{s}}{[1.2 (10^{-3}) \text{ g/cm}^3] (0.080 \text{ cm}^2/\text{s})} = 1.885$$

- c. Calculate the equilibrium coefficient (Equation from Table 5-7):

$$\begin{aligned}
 K_{eq} &= \frac{P^* \rho_g MW_{oil}}{P_o \rho_L MW_a} = \frac{(6.5 \text{ mm Hg}) [1.2 (10^{-3}) \text{ g/cm}^3] (282 \text{ g/g mol})}{(760 \text{ mm Hg}) (1 \text{ g/cm}^3) (29 \text{ g/g mol})} \\
 &= 1.0 (10^{-4})
 \end{aligned}$$

- d. Calculate the gas-phase mass transfer coefficient (Equation from Table 5-7):

$$\begin{aligned}
 k_G &= 4.82 (10^{-3}) u^{0.78} Sc_G^{-0.67} d_e^{-0.11} = 4.82 (10^{-3}) (4.47)^{0.78} (1.89)^{-0.67} \\
 &\quad (178)^{-0.11} \\
 &= 5.7 (10^{-3})
 \end{aligned}$$

- e. Calculate the overall mass transfer coefficient (Equation from Table 5-7):

$$K = k_G K_{eq} = [5.7 (10^{-3}) \text{ m/s}] [1.0 (10^{-4})] = 5.70 (10^{-7}) \text{ m/s}$$

- f. Calculate the fraction of constituent emitted to the air at time t (Equation from Table 5-7):

$$f = 1 - \frac{C_t}{C_o} = 1 - e^{-Kt/D} = 1 - e^{-5.70 (10^{-7})(86,400)/0.072}$$

$$= 1 - 0.50 = 0.50$$

5.2.6.4 Examples of the Use of the Land Treatment Model for Specific Cases. To illustrate how the land treatment model is used to estimate long-term and short-term emissions from various waste management options, the following case studies are presented. Each of these examples represents the land treatment of API separator sludge/DAF float, with the exception of example 5. The waste contains 10 percent organics and is applied at a rate of 2,500 Mg of waste per year. The loading, benzene concentration, and porosity are identical to the example presented in Section 5.2.6.1. The land area is 35,000 square meters.

Example 1. Waste is applied monthly for 9 months of the year. No waste is applied during December, January, or February because the soil is frozen or saturated with water during those months. The waste is applied from a vacuum truck by spraying onto the soil surface with a nozzle. The soil is tilled 24 hours after application and is tilled again (no waste application) after 2 weeks.

Two calculations of air emissions are required: after application and after tilling. The first time period is 1 day and the second time period is 14 days. The amount of waste applied is 2,520 Mg/9 or 280 Mg/application. It is assumed that the oil content in the soil is 0.036 grams of oil/cm³. The amount of oil applied each application is 0.036/9 or 0.004 g/cm³. The concentration of benzene in the waste is 2,000 ppmw and the concentration of benzene added to the oil in the soil is 2,000/9 or 222 ppmw each application. The land area is 3,500 square meters. After application, the liquid is assumed to seep into the soil to a depth of 5 cm, and the oil loading in the waste on the soil surface was assumed to be the same as the oil content of the soil.

To estimate the amount of air emissions between application and tilling, the following parameters are used in CHEMDAT7:

- Concentration of benzene: 2,000 ppmw
- Tilling depth: 5 cm
- Time of calculation: 1 day
- Loading (10 percent oil): 0.036 g/cm³.

The fraction lost during the first day is 98 percent with 0.007 percent lost to biological decay.

The fraction lost after the first tilling is estimated by the use of the following parameters:

- Concentration of benzene: 222 ppmw
- Tilling depth: 20 cm
- Time of calculation: 14 days
- Loading: 0.036 g oil/cm³.

The calculated fraction lost during the first tilling period is 0.89 to air emissions and 0.095 to biodegradation. This fraction is independent of the concentration of benzene and is expected to also apply to the second tilling period.

The loss of benzene during the month is 97.8 percent during the spreading period and $(1 - 0.978 - 0.0067) \times 0.89 \times 100$ or 1.4 percent during the tilling periods, for a total air emission loss of 99.2 percent.

Example 2. Waste is applied weekly except when the ground is saturated with water. Waste is applied from a vacuum truck, and the waste is spread over the surface of the soil. The soil is tilled on the day following the application and weekly between applications. The waste is applied monthly to the land treatment plot throughout the year.

As in example 1, two calculations of emissions are required: after application and after tilling. The first time period is 1 day and the second time period is 7 days. The amount of waste applied is 2,520 Mg/12 or 210 Mg/application. It is assumed that the oil content in the soil is 0.036 grams of oil/cm³. The amount of oil applied each application is 0.036/12 or 0.003 g/cm³. The concentration of benzene in the waste is 2,000 ppmw and the concentration of benzene added to the oil in the soil is 2,000/12 or 167 ppmw each application. The land area is 3,500 square meters. After application, the liquid is assumed to seep into the soil to

a depth of 5 cm, and the oil loading in the waste on the soil surface was assumed to be the same as the oil content of the soil.

To estimate the amount of air emissions between application and tilling, the following parameters are used in CHEMDAT7:

- Concentration of benzene: 2,000 ppmw
- Tilling depth: 5 cm
- Time of calculation: 1 day
- Loading (10 percent oil): 0.036 g/cm³.

The fraction lost during the first day is 98 percent with 0.007 percent lost to biological decay.

The fraction lost after the first tilling is estimated by the use of the following parameters:

- Concentration of benzene: 167 ppmw
- Tilling depth: 20 cm
- Time of calculation: 7 days
- Loading: 0.036 g oil/cm³.

The calculated fraction lost during the first tilling period is 0.80 to air emissions and 0.083 to biodegradation. This fraction is independent of the concentration of benzene and is expected to also apply to the second tilling period.

The fraction loss of benzene during four tilling periods is $(0.805) + (0.0995)(0.805) + (0.805)(0.0995)^2 + 0.805(0.0995)^3$ or 0.89.

The loss of benzene during the month is 97.8 percent during the spreading period and $(1 - 0.978 - 0.0067) \times 0.89 \times 100$ or 1.4 percent during the tilling periods, for a total air emission loss of 99.2 percent.

Example 3. The waste is dewatered prior to land treatment. The parameters are the same as those used in example 1, except the waste is dewatered and the filter cake is land-treated. The oil content of the filter cake is 20 percent. The waste is applied from a dump truck and is spread by a bulldozer. The waste is tilled into the soil on the day following spreading. It is assumed that the dewatering process removes 60 percent of the oil from the waste.

As in the preceeding examples, two calculations of air emissions are required: after spreading and after tilling. The first time period is 1 day and the second time period is 14 days. The amount of waste applied is 2,520 Mg/9 or 280 Mg/application. It is assumed that the oil content in the soil is 0.036 grams of oil/cm³. The amount of oil applied each application is 0.036/9 or 0.004 g/cm³. The concentration of benzene in the waste is 2,000 ppmw and the concentration of benzene added to the oil in the soil is 2,000/9 or 222 ppmw each application. The land area is 3,500 square meters. After application, the liquid is assumed to be retained in the waste, and the oil loading in the waste on the soil surface is assumed to be the same as the oil content of the waste.

To estimate the amount of air emissions between application and tilling, the following parameters are used in CHEMDAT7:

- Concentration of benzene: 2,000 ppmw
- Tilling depth: 2 cm
- Time of calculation: 1 day
- Loading (20 percent oil): 0.2 g/cm³.

The fraction lost during the first day is 98 percent with 0.006 percent lost to biological decay.

The fraction lost after the first tilling is estimated by the use of the following parameters:

- Concentration of benzene: 222 ppmw
- Tilling depth: 20 cm
- Time of calculation: 14 days
- Loading: 0.036 g oil/cm³.

The calculated fraction lost during the first tilling period is 0.89 to air emissions and 0.095 to biodegradation. This fraction is independent of the concentration of benzene and is expected to also apply to the second tilling period.

The air emission loss of benzene during the application period is 98.3 percent during the spreading period and $(1-0.983-0.006) \times 0.89 \times 100$ or 1.2 percent during the tilling periods, for a total air emission loss of 99.5 percent. The air emissions on the basis of the untreated waste would

depend on the recovery of oil in the dewatering process and the air emissions from the dewatering process.

Example 4. The waste is tilled as it is applied to the soil. The tilling depth is 20 cm. The period between tillings is 3 days. Waste is applied monthly.

Only one calculation of air emissions is required: after tilling. The amount of waste applied is 2,520 Mg/12 or 210 Mg/application. It is assumed that the oil content in the soil is 0.036 grams of oil/cm³. The amount of oil applied each application is 0.036/12 or 0.003 g/cm³. The concentration of benzene in the waste is 2,000 ppmw, and the concentration of benzene added to the oil in the soil is 2,000/12 or 167 ppmw each application. The land area is 3,500 square meters.

The fraction lost after the first tilling is estimated by the use of the following parameters:

- Concentration of benzene: 167 ppmw
- Tilling depth: 20 cm
- Time of calculation: 3 days
- Loading: 0.036 g oil/cm³.

The calculated fraction lost during the first tilling period is 0.60 to air emissions and 0.055 to biodegradation. This fraction is independent of the concentration of benzene and is expected to also apply to the following tilling periods.

The loss of benzene during the first tilling period is 60 percent with (100-60-5.5) or 34.5 percent remaining. The loss of benzene during the second tilling period is 0.60 (34.5) or 20.7 percent, with a fraction of (0.345)² or 0.119 of benzene remaining. The total loss of benzene for the month is 92 percent.

Example 5. Waste is applied monthly for 9 months of the year. No waste is applied during December, January, or February because the soil is frozen or saturated with water during those months. The waste is applied from a vacuum truck by spraying onto the soil surface with a nozzle. The soil is tilled 24 hours after application and is tilled again (no waste application) after 2 weeks. The waste is aqueous containing 10 percent organics and 2,000 ppmw benzene. The waste contains 10 percent solids by weight.

Two calculations of air emissions are required: after application and after tilling. The first time period is 1 day and the second time period is 14 days. The amount of waste applied is 2,520 Mg/9 or 280 Mg/application. It is assumed that the water content in the soil is 0.10 grams/cm³. The concentration of benzene in the waste is 2,000 ppmw and the concentration of benzene added to the water in the soil is 300 ppmw each application. The land area is 3,500 square meters. After application, the liquid is assumed to seep into the soil to a depth of 5 cm, and the water loading in the waste on the soil surface was assumed to be the same as the water content of the soil.

To estimate the amount of air emissions between application and tilling, the following parameters are used in CHEMDAT7:

- Concentration of benzene: 300 ppmw
- Tilling depth: 5 cm
- Time of calculation: 1 day
- Loading (10 percent): 0.10 g/cm³.

The fraction lost to air emissions during the first day is 99.8 percent with less than 0.01 percent lost to biological decay.

5.2.7 Assumptions and Sensitivity Analyses

The RTI model incorporates the following assumptions to simplify development and use of the model:

- Volatilization and biodegradation are the predominant waste removal mechanisms (i.e., other mechanisms can be ignored).
- Waste is mixed uniformly within a surface layer of the soil.
- Waste does not flow as a liquid within the soil.
- The adsorption isotherm of a constituent is linear within the application surface layer and does not change with time.
- No bulk flow of gas is induced within the soil.
- The rate of biological decay/chemical reaction is a first-order process.
- The diffusion coefficient does not vary with either concentration or time.

- The concentration of a constituent in the gas phase at the surface of the soil is much lower than the concentration of that constituent in the gas phase within the soil.
- There is no diffusion of waste into the soil beneath the zone of incorporation.
- Liquid-vapor equilibrium is established at all times within the soil.

The RTI model was evaluated for sensitivity to each of the input parameters. In the analysis, each input parameter was varied over the entire range of reasonable values. The effect on emissions of parameter variations was noted, and the parameters showing the highest sensitivity were identified.

Individual constituent properties were found to have the most significant impact on land treatment emissions. These properties include vapor pressure, diffusivity in air, and biodegradation rate. The more volatile compounds are mostly emitted to the atmosphere unless a volatile compound also has a high biodegradation rate or low diffusivity. Compounds with low vapor pressures tend also to have lower diffusivities; thus, even if such compounds also have moderate or low biorates, they are mostly biodegraded rather than emitted to the air.

Operating and field parameters also have an impact on emissions but to a lesser extent than compound properties. Tilling depth, soil porosity, and waste loading all have an impact on air emissions, with the largest impact on the more volatile compounds. Tilling depth can have a substantial impact on air emissions of volatile compounds, especially if a compound also has a relatively high biorate. As tilling depth increases, materials remain in the soil for a longer time and thus have a greater chance of being biodegraded.

Waste loading can be varied in two ways: by increasing the concentration of a compound in the waste or by increasing the amount of material applied to the soil per unit area. If the concentration of a compound is changed, air emissions change in direct proportion to the change in concentration (i.e., the fraction of the compound that is emitted to the air remains constant). If total waste loading is changed, air emissions change in the same direction as the change in loading but not in

direct proportion (i.e., the fraction emitted is lower for higher loading rates). These results assume that a treatment site is not overloaded to the point where biodegradation ceases to be a first-order process.

Average molecular weight of the oil has an effect on air emissions, but the magnitude of the effect is less than that of the other parameters studied. As average molecular weight goes up, the fraction emitted for a specific constituent increases; the fraction emitted decreases if the average molecular weight is reduced.

5.3 REFERENCES

1. Crank, J. The Mathematics of Diffusion. London, Oxford University Press. 1970. p. 45-47.
2. Crank, J. Diffusion in Polymers. New York, Academic Press. 1968. p. 16.
3. Huddleston, R. L., C. A. Bleckman, and J. R. Wolfe. Land Treatment Biological Decay Processes. In: Land Treatment, a Hazardous Waste Management Alternative. Water Resources Symposium Number Thirteen. Austin, The University of Texas at Austin. 1985. p. 44.
4. Letter and attachment from Klett, R. J., Chevron Corporation, to Wyatt, Susan R., U.S. Environmental Protection Agency, July 8, 1987, p. 1-2. Comments on Draft Emission Models Report.
5. Levenspiel, O., Chemical Reaction Engineering. John Wiley and Sons, Inc., New York, New York. May 1967. p. 47.
6. Millington, R. J, and J. P. Quirk. Permeability of Porous Solids. Trans. Faraday Society. 57:1200-1207. 1961.
7. Letter and attachments from Sonenville, G. F., Chevron Research Company, to Thorneloe, S. A., U.S. EPA. May 22, 1986. p. 19. Comments on preliminary draft BID for land treatment.
8. GCA Corporation. Air Emissions from Land Treatment--Emissions Data and Model Review. Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. September 1985. Chapter 4.
9. Radiation Technologies, Inc. Treatability Data in Support of a Treatment Zone Model for Petroleum Refining Land Treatment Facilities. Prepared for American Petroleum Institute. Washington, DC. March 1986. p. 154.
10. McKay, Donald, and Ronald S. Matsuga. Evaporation Rates of Liquid Hydrocarbon Spills on Land and Water. The Canadian Journal of Chemical Engineering. 51:434-439. August 1973.

11. Memorandum from Thorneloe, S., EPA/OAQPS, to Durham, J., EPA/OAQPS, January 31, 1986. Land treatment data base.
12. Environmental Research and Technology. Land Treatment Practices in the Petroleum Industry. Prepared for American Petroleum Institute. Washington, DC. June 1983. p. 1-2.
13. Radian Corporation. Field Assessment of Air Emissions and Their Control at a Refinery Land Treatment Facility. Volume I. Prepared for U.S. Environmental Protection Agency. Cincinnati, OH. September 1986. p. 154 and 179.
14. Reference 13, p. 43.
15. Utah Water Research Laboratory. Evaluation of Volatilization of Hazardous Constituents at Hazardous Waste Land Treatment Sites. For U.S. Environmental Protection Agency, Office of Research and Development, Ada, Oklahoma. Undated, p. 55.
16. American Petroleum Institute. Land Treatability of Appendix VIII Constituents Present in Petroleum Refinery Waste: Laboratory and Modeling Studies. API Publication No. 4455. American Petroleum Institute, Washington, DC. April 1987. P. 3-10 and 3-12.
17. Reference 16.
18. Ricciardelli, A. J., et al. 1986. Landfarm Simulator Program. Summary Report. Chevron Corporation, Richmond, California. July 1987. p. 18-24.

6.0 LANDFILLS AND WASTEPILES

6.1 INTRODUCTION

The main facilities in this category that constitute the model plants are waste fixation units, open landfills, closed landfills, and wastepiles.

All wastes that contain free liquids and that are destined for disposal in a landfill must be treated to eliminate the free liquids. This is often accomplished by adding a "fixative" to the waste, such as portland cement, cement kiln dust, or lime flue dust. These materials react with water in the waste and set up to form a dry material that encapsulates or binds the organic constituents of the waste. This fixation process is most often conducted in lined open pits or open tanks into which the liquid waste is poured. The fixative then is added and the materials are thoroughly mixed, most often with a backhoe. Alternatively, mechanical mixers such as pugmills can be used to blend the waste and fixative. Emissions are generated for as long as the waste remains in the pit. Emissions from this process may be estimated by using the open dump model.

A landfill is a facility, usually an excavated, lined pit, into which wastes are placed for permanent disposal. Emissions from open landfills, those still receiving wastes, can be estimated by applying the Research Triangle Institute (RTI) land treatment model.¹ Emissions from closed landfills, those filled to design capacity and with a cap (final cover) installed, can be estimated with the RTI closed landfill model.

Wastepiles are temporary accumulations of waste. They serve a storage function and have a limited life span. Emissions from wastepiles can be estimated by applying the RTI land treatment model.²

6.2 CLOSED LANDFILLS

6.2.1 Emission Model Equations

The RTI closed landfill model is used to estimate the time-dependent behavior of emissions from waste placed in a closed (capped) landfill that

is vented to the atmosphere and (as a special case) open-landfill waste covered with daily earth covers. This model accounts for escape of the constituent of interest via two primary, independent mechanisms: diffusion through the cap and convective loss from barometric pumping through the vent(s). It is the purpose of this section to describe the model and its evolution in a general way and to present all model equations and major assumptions.

The model is based primarily upon the work of Farmer et al.,³ who applied Fick's first law for steady-state diffusion to closed landfills. Farmer's equation utilizes an effective diffusion coefficient for the soil cap based on the work of Millington and Quirk.⁴ A previous EPA study⁵ was dedicated to the evaluation of available models for estimating emissions from hazardous waste treatment, storage, and disposal facilities (TSDF), including closed landfills. This study endorsed the models of Farmer et al.⁶ and Thibodeaux⁷ for closed landfills, apparently because of their treatment of soil-pore diffusion. Of the two, the Farmer et al.⁸ model alone has received experimental verification (although to a limited degree) via a laboratory experiment using hexachlorobenzene-containing waste in a simulated landfill.

The diffusion model of Farmer et al.⁹ was subsequently modified by RTI to allow for convective losses of the constituent of interest from the landfill, which can occur from barometric pumping. Furthermore, the decline in the emission rate from closed landfills over the long term was accounted for via the incorporation of a time-dependent decay function.

The barometric pumping emission mechanism results from changes in atmospheric pressure--as the pressure is lowered, gas flows out of the landfill through the vent(s) to equalize internal pressure. The contribution to total emissions resulting from barometric pumping equals the concentration of the constituent of interest in the gas within the landfill multiplied by the total flow of gas from the landfill. It is recognized that under certain conditions (e.g., the presence of significant biomass) biogas could be generated in a landfill. Biogas consists of methane and carbon dioxide, which is produced from the action of bacteria on organic material. Because of the convective or purging action of biogas in removing the constituent of interest in vapor form, biological decay (if it

occurs) results in a net increase in the emission rate. However, it should be noted that there is no evidence that there is significant biomass (necessary for biogas generation) in any chemical waste landfill. Therefore, in this analysis it is assumed, as suggested in the literature, that the toxic property of the waste will inhibit biological processes and thus prevent biogas generation.¹⁰ Hence, closed landfill model equations presented in this document account for diffusion through the cap and barometric pumping only.

The equations inherent in the RTI closed landfill model are as follows: Fick's first law for steady-state diffusion, based on the work of Farmer et al.,¹¹ for a landfill is given as:

$$J_i = -D_{ei} (C_{2i} - C_{si})/l \quad (6-1)$$

where

J_i = vapor flux of the constituent through the soil surface,
g/cm²·s

D_{ei} = effective diffusion coefficient, cm²/s

C_{2i} = concentration of constituent in the air above the cap,
g/cm³ air

C_{si} = concentration of the constituent in the vapor space beneath
the cap, g/cm³

l = cap thickness, cm.

(Because the concentration of the constituent at the surface is negligible, $C_{2i} \approx 0$.)

Emissions associated with diffusion alone (E_{1i} , g/s) are obtained from the above equation by multiplying by the landfill surface area (A) in cm²:

$$E_{1i} = J_i \times A \quad (6-2)$$

The effective diffusion coefficient of the constituent in soil, D_{ei} , is computed (using the expression developed by Millington and Quirk¹² and applied by Farmer et al.¹³) from the diffusion coefficient of the constituent in air, D_{ai} , as:

$$D_{ei} = D_{ai} (\epsilon_a^{3.33} / \epsilon_T^2) \quad (6-3)$$

where

D_{ai} = vapor diffusion coefficient in air, cm^2/s

ϵ_a = soil cap air-filled porosity, cm^3/cm^3 (the actual air-filled porosity of the moist soil)

ϵ_T = total porosity of the soil cap.

The concentration of the constituent of interest in the vapor space beneath the cap is computed using the ideal gas law as follows:

$$C_{si} = P_i MW_i / RT' = P_i MW_i / R(T + 273) \quad (6-4)$$

where

P_i = equilibrium partial pressure of constituent, atm

MW_i = molecular weight of constituent, g/g mol

R = gas constant, $82.05 \frac{\text{cm}^3 \cdot \text{atm}}{\text{K} \cdot \text{g mol}}$

T' = absolute temperature in the landfill, K

T = temperature in the landfill, °C.

Calculation of the equilibrium partial pressure, P_i , depends on the type of waste liquid as follows:

- a. For dilute aqueous solutions (where Henry's law applies), the equilibrium partial pressure of constituent within the landfill (P_i , atm) is computed as:

$$P_i = \frac{H_{ci} \rho_{\text{liquid}} \cdot X_i}{MW_{\text{liquid}}} \times 10^6 \frac{\text{cm}^3}{\text{m}^3} \quad (6-5)$$

where

H_{ci} = Henry's law constant, $\text{m}^3 \cdot \text{atm}/\text{mol}$

ρ_{liquid} = density of waste liquid, g/cm^3 (1 g/cm^3 is generally a good estimate for this parameter)

X_i = mole fraction of constituent i in waste liquid

where

$$X_i = (C_i/MW_i) / [C_{H_2O}/18 + C_i/MW_i]$$

where

C_i = weight fraction of constituent i in the original waste liquid

C_{H_2O} = weight fraction of water in the original waste liquid

MW_{liquid} = average molecular weight of waste liquid, $\frac{g}{g \text{ mol}}$.

- b. For two-phase (water + organic liquid) or organic liquid waste, the equilibrium partial pressure of the constituent of interest within the landfill (P_i , atm) is computed using Raoult's law:

$$P_i = X_i P_i^* \quad (6-6)$$

where

X_i = mole fraction of constituent in the organic liquid phase

where

$$X_i = (C_i/MW_i) / [C_i/MW_i + C_{oil}/MW_{oil}]$$

where

C_i = weight fraction of constituent i in the original waste liquid

C_{oil} = weight fraction of oil carrier-liquid in the original waste liquid

MW_{oil} = molecular weight of oil carrier-liquid, $g/g \text{ mol}$

P_i^* = pure component vapor pressure of the constituent of interest, atm.

Emissions from barometric pumping are computed as:

$$E_{2i} = Q \times C_{Si} \times A \quad (6-7)$$

where

E_{2i} = emissions from barometric pumping, g/s

Q = flow rate of gas through the vent, expressed as a flux, $cm^3/cm^2 \text{ landfill area} \cdot s$

C_{sf} = concentration of constituent in the gas within the landfill,
g/cm³ gas (see Equation (6-4))

A = surface area of the landfill, cm².

The gas flow rate, Q , is estimated using the following procedure.

- a. Compute volume of gas available for barometric pumping, assuming the entire void-volume of the waste is available:

$$V_C = D \times A \times \epsilon_{fw} \quad (6-8)$$

where

V_C = volume of void space, cm³

D = thickness of waste bed within landfill, cm

A = surface area of the landfill, cm².

ϵ_{fw} = air porosity fraction of fixed waste (dimensionless).

- b. Compute the total volume of gas (cm³) exiting the vent of the landfill due to changes in barometric pressure and/or temperature within the landfill:

$$V_B = V_C \left[\left(\frac{P_{ref}}{P_1} \right) \left(\frac{T_1 + 273}{T_{ref} + 273} \right) - 1 \right] \quad (6-9)$$

where

V_B = total volume of gas exiting landfill, cm³

P_{ref} = initial (reference) barometric pressure, mm Hg

P_1 = final barometric pressure, mm Hg

T_1 = final landfill temperature, °C

T_{ref} = initial (reference) landfill temperature, °C.

For cases in which $P_1 > P_{ref}$ and/or $T_1 < T_{ref}$, the computed value of V_B may be negative (indicating a condition of gas flow into the landfill). Because this condition results in no emissions associated with barometric pumping, V_B should be set equal to zero to avoid calculational errors in the following steps.

- c. Compute the average flow rate of gas from the landfill over the time interval of interest:

$$Q = \frac{V_B}{\Delta t A} \quad (6-10)$$

where

Q = average flow rate of gas from the vent due to barometric pumping, cm^3/cm^2 landfill area \cdot s

Δt = time interval over which the change in pressure and/or temperature occurred, s

A = landfill area, cm^2 .

In an average day, barometric pressure drops 4 mbar from a typical value of 1,013 mbar. Landfill temperature is assumed to remain constant. Hence, under these conditions, $P_{\text{ref}} = 1,013$ mbar, $P_1 = 1,009$ mbar, $T_{\text{ref}} = T_1 = 15^\circ\text{C}$, and $\Delta t = 8.64 \times 10^4$ s.

Having computed the instantaneous emissions associated with diffusion through the cap and barometric pumping, the total initial emission rate at the time of landfill closure, E_i^* (g/s), is computed as the sum:

$$E_i^* = E_{1i} + E_{2i} \quad (6-11)$$

The total instantaneous emission rate at any time then is computed via an exponential decay function:

$$E_i(t) = \frac{(3,600 \text{ s/h})(24 \text{ h/d})(365.25 \text{ d/yr})E_i^*}{10^6} \exp(-\lambda t)$$

$$E_i(t) = 31.56 E_i^* \exp(-\lambda t) \quad (6-12)$$

where

$E_i(t)$ = total time-dependent emission rate, Mg/yr

E_i^* = initial emission rate, at time of landfill closure, g/s

t = time since landfill closure, mo

λ = "decay" constant, mo⁻¹.

The "decay" constant, λ , is computed as follows:

$$\lambda = \frac{(3,600 \text{ s/h}) \times (24 \text{ h/d}) \times 365.25 \text{ d/yr} \times E_i^*}{12 \text{ mo/yr} \times M_{oi}}$$

$$\lambda = 2.63 \times 10^6 E_i^*/M_{oi} \quad (6-13)$$

where M_{oi} is the total mass of the constituent of interest in the landfill (g). This parameter can be computed from the weight fraction of the constituent in the original waste liquid (C_i), the mass of original waste liquid in a unit volume of fixed waste (W), the landfill surface area (A), and the thickness of the waste layer within the landfill (D):

$$M_{oi} = C_i W A D \quad (6-14)$$

The average emission rate from a closed, vented landfill over the time since landfill closure is equal to the integral of the emission rate equation over the time period divided by the time period, which yields the following expression:

$$E_{Ai}(t) = \frac{(3,600 \text{ s/h})(24 \text{ h/d})(365.25 \text{ d/yr}) E_i^*}{(10^6 \text{ g/Mg})\lambda t} [1 - e^{-\lambda t}]$$

$$E_{Ai}(t) = \frac{31.56 E_i^*}{\lambda t} [1 - e^{-\lambda t}] \quad (6-15)$$

where

$E_{Ai}(t)$ = average emission rate over the time since landfill closure, Mg/yr

t = time since landfill closure, mo.

Table 6-1 summarizes the equations necessary to apply the RTI closed landfill model.

The model is highly sensitive to the air porosity of the clay cap (ϵ_a), which largely determines the diffusion rate through the cap. The model is sensitive to the properties of the constituent of interest, particularly the vapor pressure (P_i^*), Henry's law constant (H_{Ci}), and mole fraction in the waste liquid (X_i). Because temperature affects volatility, the model is sensitive to temperature. Other parameters to which the model is sensitive include the depth of the fixed waste (D), the air porosity of the fixed waste (ϵ_{fw}), the landfill surface area (A), and the barometric pressure change ($P_{ref} - P_1$). This latter group of parameters is significant in that it impacts the barometric pumping rate or the volume of gas available for pumping. In contrast to these parameters, the model exhibits relatively low sensitivity to the diffusivity of the constituent in air (D_{ai}), the cap thickness (l), and the total mass of the constituent in the landfill (M_{0i}).

The major assumptions associated with the RTI closed landfill model are as follows:

- The liquid waste containing the constituent of interest is assumed to be bound in the fixed waste within the landfill.
- The constituent of interest in the gas within the landfill is assumed to be in equilibrium with the liquid in the waste.
- Adsorption of the constituent of interest onto the soil of the cap is assumed to be negligible.
- The fraction of air-filled space in the landfill cap (air porosity) is assumed to remain relatively constant over the long term.
- The effective diffusion coefficient of the cap is assumed not to vary with either the concentration of the constituent of interest or time.
- The concentration of the constituent of interest in air at the top of the landfill cap is assumed to be effectively 0.
- No biodegradation (with concurrent production of biogas) is assumed to occur due to the suppression of biological processes by the toxic waste.

TABLE 6-1. RTI CLOSED LANDFILL MODEL

$$E_i(t) = 31.56 E_i^* \exp(-\lambda t)$$

$$E_{Ai}(t) = \frac{31.56 E_i^*}{\lambda t} \left(1 - e^{-\lambda t} \right)$$

$$\lambda = 2.63 \times 10^6 E_i^* / M_{oi}$$

$$M_{oi} = C_i W A D$$

$$E_i^* = E_{1i} + E_{2i}$$

$$E_{1i} = J_i \times A$$

$$J_i = -D_{ei} (C_{2i} - C_{si}) / l$$

$$D_{ei} = D_{ai} (\epsilon_a^{3.33} / \epsilon_T^2)$$

$$C_{2i} = 0$$

$$C_{si} = P_i MW_i / R (T + 273)$$

$$P_i = \frac{H_{ci} \rho_{liquid} X_i}{MW_{liquid}} \times 10^6 \frac{cm^3}{m^3} \quad (\text{dilute aqueous waste liquids})$$

$$X_i = (C_i / MW_i) / (C_{H_2O} / 18 + C_i / MW_i) \quad (\text{dilute aqueous waste liquids})$$

$$P_i = X_i P_i^* \quad (\text{two-phase liquid or organic liquid waste})$$

$$X_i = (C_i / MW_i) / (C_i / MW_i + C_{oil} / MW_{oil}) \quad (\text{two-phase liquid or organic liquid waste})$$

$$E_{2i} = Q C_{si} A$$

$$Q = V_B / (\Delta t A)$$

$$V_B = V_C \left[\left(\frac{P_{ref}}{P_1} \right) \left(\frac{T_1 + 273}{T_{ref} + 273} \right) - 1 \right]$$

$$V_C = D A \epsilon_{fw}$$

(continued)

TABLE 6-1 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
A	Landfill surface area, cm^2	Westat survey ^a
C_{si}	Concentration of constituent i in the gas within the landfill, g/cm^3 gas	Calculated
C_{2i}	Concentration of constituent i in air above the cap, g/cm^3	Assumed
C_i	Weight fraction of constituent i in the original waste liquid (dimensionless)	Definition
$C_{\text{H}_2\text{O}}$	Weight fraction of water in the original waste liquid (dimensionless)	Definition
C_{oil}	Weight fraction of oil carrier-liquid in the original waste liquid (dimensionless)	Definition
D	Thickness of waste bed within landfill, cm	Westat survey ^b
D_{ai}	Diffusivity of constituent i in air, cm^2/s	Literature
D_{ei}	Effective diffusion coefficient of constituent i in clay cap, cm^2/s	Calculated
$E_{Ai}(t)$	Average emission rate of constituent i over time t since landfill closure, Mg/yr	Calculated
$E_i(t)$	Total instantaneous emission rate of constituent i at time t since landfill closure, Mg/yr	Calculated
E_i^*	Total initial emission rate of constituent i at time of landfill closure, g/s	Calculated
E_{1i}	Initial emission rate of constituent i at landfill closure due to diffusion alone, g/s	Calculated

^aReference 14.^bReference 15.

(continued)

TABLE 6-1 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
E_{2i}	Initial emission rate of constituent i at landfill closure due to barometric pumping alone, g/s	Calculated
H_{ci}	Henry's law constant for constituent i , $m^3 \cdot \text{atm} / \text{mol}$	Literature
J_i	Initial diffusion flux of constituent i , $\text{g} / \text{cm}^2 \cdot \text{s}$	Calculated
l	Landfill cap thickness, cm	Literature
M_{oi}	Initial mass of constituent i in the landfill, g	Definition or calculated
MW_i	Molecular weight of constituent i , g/g mol	Literature
MW_{liquid}	Average molecular weight of the dilute aqueous waste liquid, g/g mol (assumed to be 18 g/g mol)	Estimated
MW_{oil}	Molecular weight of the oil carrier-liquid, g/g mol	Definition or estimated
P_i^*	Pure component vapor pressure of constituent i , atm	Literature
P_i	Equilibrium partial pressure of constituent i in the vapor space, atm	Calculated
P_{ref}	Initial (reference) barometric pressure, mm Hg	Meteorological information
P_1	Final barometric pressure after Δt , mm Hg	Meteorological information
Q	Average flow rate of gas from landfill vent(s) due to barometric pumping, $\text{cm}^3 / \text{cm}^2$ landfill area $\cdot \text{s}$	Calculated
R	Ideal gas constant, $82.05 \text{ cm}^3 \cdot \text{atm} / \text{g mol} \cdot \text{K}$	Literature
t	Time since landfill closure, mo	Definition

(continued)

TABLE 6-1 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
Δt	Time interval used to determine average barometric pumping rate, s	Definition
T	Temperature within landfill, °C	Estimated from literature data
T_{ref}	Initial (reference) landfill temperature, °C	Estimated from literature data
T_1	Final landfill temperature after Δt , °C	Estimated
V_B	Total volume of gas exiting landfill in Δt , cm ³	Calculated
V_C	Total volume of void space within waste, cm ³	Calculated
W	Mass of original waste liquid in a unit volume of fixed waste, g/cm ³	Definition or estimated
X_i	Mole fraction of constituent i in the aqueous liquid (for dilute aqueous waste) or in the organic phase (for two-phase or organic liquid waste) (dimensionless)	Definition
ϵ_a	Air porosity of the clay cap (dimensionless)	Estimated from clay property data
ϵ_T	Total porosity of the clay cap (dimensionless)	Estimated from clay property data
ϵ_{fw}	Air porosity of the fixed waste (dimensionless)	Estimated from fixed waste property data
ρ_{liquid}	Density of dilute aqueous waste liquid, g/cm ³ (generally equals 1 g/cm ³)	Definition or estimated
λ	Exponential decay constant, mo ⁻¹	Calculated

- The landfill is assumed to be vented to the atmosphere. The volume of gas available for barometric pumping is assumed to consist of the total void-volume of the waste bed.
- No transport of the constituent of interest in moving water is assumed to occur.

6.2.2 Model Plant Parameters for Closed Landfills

The characteristics of a model closed landfill facility are discussed here. This model facility is used as the basis for an example calculation in Section 6.2.3.

The model facility for closed landfills has an area of $1.417 \times 10^8 \text{ cm}^2$ (3.5 acres): This value represents an approximately midrange value from the Westat survey.¹⁶ A reasonable value of landfill depth, also selected from the Westat survey,¹⁷ is 458 cm (15 ft). The landfill cap is assumed to be composed of compacted clay. The cap thickness value of 107 cm (3.5 ft) represents the average of extremes in thickness of clay caps reported in site studies (2 ft to 6 ft).¹⁸ The value used for air porosity of the clay cap is 0.08 (8 percent), while the total porosity is 0.41 (41 percent). These values were computed based on reasonable physical properties and level of compaction for compacted clay.¹⁹ The landfill is assumed to be vented to the atmosphere. The temperature beneath the landfill cap is estimated at 15 °C, which represents the temperature of shallow ground water at a midlatitude U.S. location.²⁰ This temperature is assumed to remain constant. The landfill is assumed to be exposed to a nominal barometric pressure of 1,013 mbar, which represents an estimate of the annual average atmospheric pressure in the United States.²¹ Barometric pumping is estimated for the landfill using a daily pressure drop from the nominal value of 4 mbar. The 4 mbar value represents an estimate of the annual average diurnal pressure drop.²²

The model closed landfill facility is assumed to contain fixed waste. The waste liquid (before fixation) selected for the facility is assumed to be a two-phase aqueous/organic containing 20 percent chloroform, 20 percent low-volatility organic,* and 60 percent water (by weight). This liquid has an average density of 1.16 g/cm^3 . The fixation industry indicates that

*For modeling purposes, this component of the waste liquid represents the oil carrier-liquid.

waste liquid, when combined with fixative, may in actuality increase in volume by as much as 50 percent.^{23,24} The volume change, which is a function of the specific waste being fixed and the specific formulation of the fixative, can only be determined experimentally. In view of the inherent variability in the fixation process and the lack of real data, for the purpose of this calculation the assumption is made that the fixation process does not change the waste volume. This assumption is environmentally conservative and may result in an overestimation of the landfill emissions. Actual volume changes that may take place as a result of fixation can easily be accounted for because the change in the calculated emissions is inversely proportional to the change in waste volume. One industry contact indicated that, for the purposes of estimating emissions, the assumption of no volume change during fixation was reasonable.²⁵ Based on the waste liquid density and the assumption of no volume increase from fixation, the mass of waste liquid in a unit volume of fixed waste is 1.16 g/cm^3 . The air porosity of the fixed waste (used to estimate the total volume of gas available for barometric pumping) is taken to be 0.25 (25 percent). This value was inferred from measurements of total porosity and moisture content of various fixed wastes,²⁶ and, for the purposes of this analysis, is assumed to pertain to waste within the landfill as opposed to waste immediately following fixation. As discussed previously, there is no evidence for significant biomass in any chemical waste landfill. Therefore, in this analysis it is assumed, as suggested in the literature, that the toxic property of the waste will inhibit biological processes and thus prevent biogas generation.²⁷ Hence, the waste biomass concentration is taken to be 0 g/cm^3 .

The properties of chloroform that are pertinent to this analysis include the molecular weight (119.4 g/g mol), pure component vapor pressure at 15°C (0.162 atm), diffusivity in air at 15°C ($0.10 \text{ cm}^2/\text{s}$) and density (1.49 g/cm^3). The low-volatility organic liquid present in the waste has a molecular weight of 147 g/g mol and a density of 1.31 g/cm^3 .

Table 6-2 summarizes the model facility parameters for closed landfills used in the example calculation in Section 6.2.3. For facilities that accept more than one type of waste, the weighted average constituent concentrations may be used.

TABLE 6-2. INPUT PARAMETERS--CLOSED LANDFILL

Parameter	Value
Area	$1.417 \times 10^8 \text{ cm}^2$ (3.5 acres)
Waste bed thickness	457 cm (15 ft)
Cap thickness	107 cm (3.5 ft)
Cap air porosity	0.08 (8%)
Cap total porosity	0.41 (41%)
Type landfill	Vented
Temperature beneath cap	15 °C
Typical barometric pressure	1,013 mbar
Daily barometric pressure drop	4 mbar
Waste liquid (before fixation)	Two-phase aqueous/organic
Liquid composition	20% chloroform, 20% low-volatility organic (oil), 60% water (by weight)
Liquid/fixative	1 unit volume liquid + dry fixative = 1 unit volume fixed waste
Liquid in fixed waste	1.16 g/cm^3
Air porosity fixed waste	0.25 (25%)
Biomass concentration	0 g/cm^3
<u>Chloroform properties</u>	
Molecular weight	119.4 g/g mol
Vapor pressure (15 °C)	0.162 atm (123 mm Hg)
Diffusivity in air (15 °C)	$0.10 \text{ cm}^2/\text{s}$
Density	1.49 g/cm^3
<u>Low-volatility organic^a properties</u>	
Molecular weight	147 g/g mol
Density	1.31 g/cm^3

^aAlso referred to as oil "carrier-liquid."

6.2.3 Example Calculation for Closed Landfill

This section presents a step-by-step calculation of emissions from a closed landfill that is vented to the atmosphere. The equations discussed in Section 6.2.1 and summarized in Table 6-1 are used with the model unit parameters in Section 6.2.2 to estimate emissions from a fixed, two-phase aqueous/organic waste containing chloroform:

- Waste liquid (before fixation): 20 percent chloroform, 20 percent low-volatility organic liquid, 60 percent water (by weight)
 - Liquid/fixative: 1 unit volume liquid + dry fixative = 1 unit volume fixed waste
 - Waste biomass concentration: 0 g/cm³
 - Landfill area: 1.417 x 10⁸ cm² (3.5 acres)
 - Waste bed thickness: 457 cm (15 ft)
 - Cap thickness: 107 cm (3.5 ft)
 - Type landfill: vented
 - Temperature beneath cap: 15 °C
 - Time period for emission calculation: 1 yr.
- a. Compute the effective diffusion coefficient, D_{ei} (cm²/s) (Equation (6-3)):

$$D_{ei} = D_{ai} (\epsilon_a^{3.33} / \epsilon_T^2)$$

$$D_{ei} = (0.10 \text{ cm}^2/\text{s}) (0.08)^{3.33} / (0.41)^2$$

$$D_{ei} = 1.32 \times 10^{-4} \text{ cm}^2/\text{s}$$

- b. Compute the equilibrium partial pressure of chloroform in the vapor space, P_i (atm):

The waste before fixation was a two-phase liquid. Hence, Raoult's law applies. The mole fraction for this case is computed as:

$$X_i = (C_i / MW_i) / (C_i / MW_i + C_{oil} / MW_{oil})$$

$$X_i = (0.20/119.4 \text{ g/g mol}) + [0.20/119.4 \text{ g/g mol} + 0.20/147 \text{ g/g mol}]$$

$$X_i = 0.55$$

From Raoult's law (Equation (6-6)):

$$P_i = X_i P_i^*$$

$$P_i = (0.55)(0.162 \text{ atm})$$

$$P_i = 8.91 \times 10^{-2} \text{ atm}$$

- c. Compute the concentration of chloroform in the vapor space beneath the cap, C_{si} (g/cm³ void space) (Equation (6-4)):

$$C_{si} = P_i MW_i / R(T + 273)$$

$$C_{si} = \frac{(8.91 \times 10^{-2} \text{ atm})(119.4 \text{ g/g mol})}{(82.05 \text{ cm}^3 \cdot \text{atm/g mol} \cdot \text{K})(15 + 273)}$$

$$C_{si} = 4.50 \times 10^{-4} \text{ g/cm}^3$$

- d. Compute initial chloroform emission flux resulting from diffusion through the cap only, J_i (g/cm²·s) (Equation (6-1)):

$$J_i = -D_{ei}(C_{2i} - C_{si})/l$$

$$J_i = -(1.32 \times 10^{-4} \text{ cm}^2/\text{s})(0 \text{ g/cm}^3 - 4.50 \times 10^{-4} \text{ g/cm}^3)/107 \text{ cm}$$

$$J_i = 5.55 \times 10^{-10} \text{ g/cm}^2 \cdot \text{s}$$

- e. Compute initial chloroform emissions resulting from diffusion through the cap only, E_{1i} (g/s) (Equation (6-2)):

$$E_{1i} = J_i \times A$$

$$E_{1i} = (5.55 \times 10^{-10} \text{ g/cm}^2 \cdot \text{s})(1.417 \times 10^8 \text{ cm}^2)$$

$$E_{1i} = 7.86 \times 10^{-2} \text{ g/s}$$

- f. Estimate the barometric pumping-induced gas flow rate through the vent(s):

1. Compute the volume of gas available for barometric pumping, V_C (cm^3) (Equation (6-8)):

$$V_C = D \times A \times \epsilon_{fw}$$

$$V_C = (457 \text{ cm})(1.417 \times 10^8 \text{ cm}^2)(0.25)$$

$$V_C = 1.62 \times 10^{10} \text{ cm}^3$$

2. Compute volume of gas exiting the vent due to barometric pressure change, V_B (cm^3) (Equation (6-9)):

$$V_B = V_C \left[\left(\frac{P_{\text{ref}}}{P_1} \right) \left(\frac{T_1 + 273}{T_{\text{ref}} + 273} \right) - 1 \right]$$

For this case, $T_1 = T_{\text{ref}} = 15^\circ\text{C}$, and barometric pressure drops by 4 mbar from the nominal value of 1,013 mbar:

$$V_B = 1.62 \times 10^{10} \text{ cm}^3 \left[\left(\frac{1,013 \text{ mbar}}{1,009 \text{ mbar}} \right) \left(\frac{15^\circ\text{C} + 273 \text{ K}}{15^\circ\text{C} + 273 \text{ K}} \right) - 1 \right]$$

$$V_B = 6.42 \times 10^7 \text{ cm}^3$$

3. Compute the average flow rate of gas over the time interval, Q ($\text{cm}^3/\text{cm}^2 \text{ landfill area} \cdot \text{s}$) (Equation (6-10)):

The average diurnal pressure drop of 4 mbar occurs within a 24-h period. For convenience, the gas flow from this pressure change is averaged over 24 h (equals $8.64 \times 10^4 \text{ s}$).

$$Q = \frac{V_B}{\Delta t A}$$

$$Q = \frac{6.42 \times 10^7 \text{ cm}^3}{(8.64 \times 10^4 \text{ s})(1.417 \times 10^8 \text{ cm}^2)}$$

$$Q = 5.25 \times 10^{-6} \text{ cm}^3/\text{cm}^2 \cdot \text{s}$$

4. Compute the barometric pumping-induced emission rate, E_{2i} (g/s) (Equation (6-7)):

$$E_{2i} = Q \times C_{si} \times A$$

$$E_{2i} = (5.25 \times 10^{-6} \text{ cm}^3/\text{cm}^2 \cdot \text{s})(4.50 \times 10^{-4} \text{ g/cm}^3)(1.417 \times 10^8 \text{ cm}^2)$$

$$E_{2i} = 0.335 \text{ g/s} .$$

- g. Compute the total initial emission rate, E_i^* (g/s) (Equation (6-11)):

$$E_i^* = E_{1i} + E_{2i}$$

$$E_i^* = 7.86 \times 10^{-2} + 0.335$$

$$E_i^* = 0.413 \text{ g/s} .$$

- h. Compute the time-dependent instantaneous emission rate:

1. Compute total mass of constituent i in landfill, M_{oi} :

First compute W , the mass of original waste liquid in a unit volume of fixed waste. Assuming one unit volume of waste liquid results in one unit volume of fixed waste, this parameter can be computed using the densities of the waste liquid components and their weight fractions as follows:

$$\begin{aligned} W &= [(1.49 \text{ g/cm}^3)(0.2) + (1.31 \text{ g/cm}^3)(0.2) + (1 \text{ g/cm}^3)(0.6)] \\ &\quad \times 1 \text{ cm}^3 \text{ liquid/cm}^3 \text{ fixed waste} \\ &= 1.16 \text{ g/cm}^3 . \end{aligned}$$

M_{oi} is then computed as:

$$M_{oi} = C_i W A D$$

$$\begin{aligned} M_{oi} &= \frac{20 \text{ g chloroform}}{100 \text{ g liquid}} \times \frac{1.16 \text{ g liquid}}{\text{cm}^3 \text{ fixed waste}} \times 1.417 \times 10^8 \text{ cm}^2 \\ &\quad \times 457 \text{ cm} = 1.50 \times 10^{10} \text{ g chloroform} \end{aligned}$$

2. Compute the decay constant, λ (mo^{-1}) (Equation (6-13)):

$$\lambda = 2.63 \times 10^6 E_i^*/M_{oi}$$

$$\lambda = (2.63 \times 10^6)(0.413 \text{ g/s})/1.50 \times 10^{10} \text{ g}$$

$$\lambda = 7.25 \times 10^{-5} \text{ mo}^{-1}$$

3. Compute the instantaneous emission rate, E_i , in Mg/yr, after 1 yr (Equation (6-12)):

$$E_i(t) = 31.56 E_i^* \exp(-\lambda t)$$

$$E_i = (31.56)(0.413 \text{ g/s}) \exp(-7.25 \times 10^{-5} \text{ mo}^{-1} \times 12 \text{ mo})$$

$$E_i = 13.0 \text{ Mg/yr}$$

i. Compute the average emission rate in the first year, E_{Ai} , in Mg/yr (Equation (6-15)):

$$E_{Ai}(t) = \frac{31.56 E_i^*}{\lambda t} [1 - e^{-\lambda t}]$$

$$E_{Ai} = \frac{(31.56)(0.413 \text{ g/s})}{(7.25 \times 10^{-5} \text{ mo}^{-1})(12 \text{ mo})} [1 - \exp\{-12 \text{ mo} \times 7.25 \times 10^{-5} \text{ mo}^{-1}\}]$$

$$E_{Ai} = 13.0 \text{ Mg/yr}$$

6.3 FIXATION PITS

This section is currently under review
for potential revisions.

6.4 OPEN LANDFILLS AND WASTEPILES

6.4.1 Emission Model Equations

The RTI land treatment model (also known as the Allen model,³⁴ discussed in Section 5.2) is used to estimate the air emission rate of the constituent of interest from open (active) landfills and wastepiles. This

model is based on the theory of diffusion out of an infinite flat slab and was intended originally for use in estimating emissions from land treatment operations. The intent of this section is to discuss use of the model with regard to the estimation of emissions from open landfills and wastepiles; a detailed description of the model relevant to land treatment operations and the theoretical basis for the model are presented in Section 5.0 of this report and will not be repeated here.

A land-treatment-type model was selected for estimating emissions from open landfills and wastepiles because (1) no adequate models exist for these sources, and (2) there are a number of similarities in physical characteristics of open landfills, wastepiles, and land treatment operations. A previous EPA study³⁵ dedicated to the evaluation of models for estimating emissions from hazardous waste TSDF identified only one model for open waste dumps such as landfills and wastepiles--the open dump model. A serious limitation of the model for this application, however, is that it does not account for depletion of the volatilizing chemical from the waste surface. Hence, the open dump model is judged unsuitable for the estimation of emissions from landfills and wastepiles over the time period of interest (months or longer).

The similarity in physical characteristics among open landfills, wastepiles, and land treatment operations is apparent upon close examination--in all three, the waste liquid is ultimately mixed homogeneously with a "carrier" matrix (soil in the case of land treatment; dry fixative in the case of active landfills; and soil, fixative, or some other solid matrix in the case of wastepiles). In all cases, the matrix is porous and permeable, allowing the diffusion of the constituent of interest through the matrix and into the air. Hence, in all cases, diffusion theory can be used to model the emission rate. The notable difference between land treatment operations and open landfills/wastepiles is the presence of an additional mechanism affecting emissions in the case of land treatment--biological decay of the constituent. Because biodegradation is not thought to occur*

*There is no evidence that there is significant biomass (necessary for biological decay) in any chemical waste landfill. It is assumed that the toxic property of the waste will inhibit biological processes.³⁶

in open landfills/wastepiles, however, its effect is not accounted for in the modeling of air emissions.

The RTI land treatment model, which was selected for estimating emissions from open landfills and wastepiles, has the following characteristics: a sound basis in scientific theory, limited validation against measured emissions from land treatment operations, and reasonably available input data.³⁷ The model considers effects such as evaporation of the constituent of interest from interstitial surfaces of the carrier matrix and diffusion of material through air-filled pore spaces.

The equations necessary to apply the land treatment model to open landfills and wastepiles are summarized in Table 6-6. These equations, extracted from Chapter 5.0, can be used to estimate the fraction of the constituent emitted (F_t) and the instantaneous emission rate (E). It should be noted that the absence of biodegradation represents a special case that allows some simplification of several of the Chapter 5.0 equations, e.g., Equations (5-4) and (5-5). (The absence of biomass implies that biomass concentration equals 0. Hence, t_b , the time constant for biological decay, equals infinity. Consequently, the exponential term e^{-t/t_b} becomes unity.) Also, the absence of biodegradation implies that the fraction of the constituent emitted after a long time, F_a , would equal unity.

Because the land treatment model was derived originally for land treatment operations, model input parameters are not necessarily in the most convenient units and terminology for open landfills and wastepiles. Hence, several points should be noted:

- Fixed waste is analogous (for modeling purposes) to the waste-laden soil in land treatment.
- M_0 , the area-loading of the constituent in g/cm^2 , is geared toward land treatment operations. For open landfills and wastepiles, it should be computed as indicated in Table 6-6.
- No "tilling" (as discussed in Chapter 5.0) is performed in open landfills or wastepiles.

- Waste liquid is "applied" or mixed with fixative only once. Hence, waste "reapplication" (used in the sense discussed in Chapter 5.0) does not occur in open landfills and waste-piles.
- The waste bed depth in open landfills and wastepiles is analogous to the "depth to which waste is mixed" in land treatment, as discussed in Chapter 5.0.

The approach required to estimate emissions from open landfills or waste-piles is as follows, based on equations in Table 6-6:

1. Compute the loading of waste liquid (L) in the fixative or soil, using the known waste composition. (For two-phase aqueous organics or organic liquid wastes, L should be computed as grams organic phase per cubic centimeter solid material. For dilute aqueous waste liquids, L equals grams aqueous liquid per cubic centimeter solid material.)
2. Compute the effective diffusion coefficient (D_e).
3. Compute the partition coefficient (K_{eq}).
4. Use the appropriate emission equation to compute the fraction of constituent emitted (F_t) and/or the instantaneous emission rate (E). For wastepile calculations, the time input to these equations should be no greater than the life of the wastepile (retention time).

The sensitivity of the land treatment model to some parameters differs in its application to open landfills and wastepiles from that in land treatment operations because of the difference (in some cases) in the expected range of the parameters. In general, it can be stated that, for application to open landfills and wastepiles, the model is sensitive to the air porosity of the solid waste, the liquid loading in the solid waste, the waste depth, the concentration of the constituent in the waste, and the volatility of the constituent under consideration. In contrast, the model exhibits a relatively low sensitivity to the diffusion coefficient of the constituent in air.

The following major assumptions are associated with the RTI land treatment model and its application to open landfills and wastepiles:

- The waste liquid is mixed uniformly with the carrier matrix. (either fixative, soil, or some other granular solid material) before placement in an open landfill or wastepile.

TABLE 6-6. RTI LAND TREATMENT MODEL APPLIED TO OPEN
LANDFILLS AND WASTEPILES (NO BIODEGRADATION)

Emission fraction^a

$$F_{at} = 0.72 (K_d t)^{1/2} \quad \text{for } \frac{K_{eq} D_e t}{l^2} < 0.25 \quad (\text{valid for no biodegradation only})$$

$$F_{at} = \frac{8}{\pi^2} \left[1 - \exp(-K_d t) \right] + 0.1878 \quad (\text{for } \frac{K_{eq} D_e t}{l^2} \geq 0.25 - \text{no biodegradation})$$

$$F_a = 1 \quad (t \rightarrow \infty - \text{no biodegradation}).$$

$$K_d = \frac{K_{eq} D_e \pi^2}{4l^2}$$

$$D_e = D_a \epsilon_a^{3.33/\epsilon_T^2} \quad \left[D_e/D_a = \epsilon_a^{3.33/\epsilon_T^2} \right]$$

$$K_{eq} = \frac{H_c (10^6 \text{ cm}^3/\text{m}^3)}{RT} \times \frac{\epsilon_a}{\epsilon_{\text{waste}}} \quad (\text{for aqueous waste})$$

$$K_{eq} = \frac{P^* MW_{\text{oil}}}{RT} \frac{\epsilon_a}{L} \quad (\text{for two-phase aqueous/organic or organic liquid waste})$$

Emission rate

$$E = \frac{2 M_o}{l^2} \frac{K_{eq} D_e}{\pi^2} \left[\exp(-\tau) \right] \quad \text{for } K_{eq} D_e t/l^2 \geq 0.213$$

$$E = \frac{M_o}{l} \left[\frac{1}{\frac{\epsilon_a}{k_G K_{eq}} + \left(\frac{\pi t}{K_{eq} D_e} \right)^{1/2}} \right] \quad (\text{for all other times})$$

$$\tau = \frac{D_e K_{eq} \pi^2 t}{4l^2}$$

$$M_o = l L C$$

$$d_e = (4A/\pi)^{1/2}$$

$$k_G = 4.82 (10^{-3}) U^{0.78} Sc_G^{-0.67} de^{-0.11}$$

See notes at end of table.

(continued)

TABLE 6-6 (continued)

Variable	Definition	Data source
C	Weight fraction of constituent in the oil (organic) phase (for two-phase or organic liquid waste), or weight fraction of constituent in the water (for aqueous waste)	Definition
D _a	Diffusion coefficient of constituent in air, cm ² /s	Data base
D _e	Effective diffusion coefficient of constituent in the solid waste, cm ² /s	Calculated
E	Emission rate of constituent, g/cm ² /s	Calculated
F _a	Fraction of constituent emitted to the atmosphere at infinity (equals unity for no biodegradation)	Definition
F _t	Fraction of constituent emitted to the atmosphere after time t	Calculated
H _C	Henry's law constant for constituent, atm•m ³ /g mol	Data base
K _d	Volatilization constant for constituent, s ⁻¹	Calculated
k _G	Gas-phase mass transfer coefficient, m/s	Calculated
K _{eq}	Ratio of gas-phase constituent to total constituent in the solid waste	Calculated
l	Depth of waste in open landfill or wastepile, cm	Literature
L	Waste loading in fixative or soil. For two-phase aqueous/organics or organic liquids, L = g organic (oil) phase/cm ³ solid material. For dilute aqueous waste liquids, L = g aqueous liquid/cm ³ solid material	Definition
A	Area of open landfill, m ²	Definition
M ₀	Area loading of constituent, g/cm ²	Calculated
See notes at end of table.		(continued)

TABLE 6-6 (continued)

<u>Variable</u>	<u>Definition</u>	<u>Data source</u>
MW_{oil}	Average molecular weight of the oil (less constituent), g/g mol	Estimated
p^*	Pure component vapor pressure of constituent, atm	Data base
R	Ideal gas constant, $82.05 \text{ atm}\cdot\text{cm}^3/\text{g mol}\cdot\text{K}$	Literature
T	Temperature of vapor in solid waste, K	Assumed
U	Windspeed (m/s)	Assumed
t	Time variable for emission calculation, s (represents time lapse from initial waste composition)	Definition
τ	Dimensionless parameter used in the instantaneous emission rate expression	Calculated
ϵ_a	Void fraction (air porosity) of solid waste (dimensionless)	Estimated from fixed waste property data
d_e	Effective diameter of land treatment area, m	Calculated
Sc_G	Schmidt number	Calculated
ϵ_T	Total porosity of solid waste (dimensionless)	Industry personnel
ϵ_{waste}	Volume fraction of waste liquid in solid waste (dimensionless) (can be computed as $L/[\text{density aqueous liquid in g/cm}^3]$)	Calculated

^aThe first equation presented represents a special case of Equation (5-7) for no biodegradation.

^bThis equation represents the first term of the series in Equation (5-5), for the special case of no biodegradation. The exponential terms are expressed, for convenience, in terms of the dimensionless parameter " τ ".

- The liquid waste containing the constituent of interest is assumed to be bound in the waste after fixation and placement in the open landfill or wastepile.
- The waste liquid does not flow within the carrier matrix.
- The adsorption isotherm of the constituent of interest is linear within the depth of the waste and does not change with time.
- No bulk flow of gas is induced within the waste matrix.
- The diffusion coefficient does not vary with either concentration or time.
- The concentration of the constituent of interest in the gas phase at the surface of the open landfill/wastepile is much lower than the concentration of the constituent of interest in the gas phase within the waste matrix.
- No diffusion of the waste liquid into depths below the waste layer is assumed.
- Liquid-vapor equilibrium is established at all times within the waste matrix.
- For the case of fixed waste in the landfill or wastepile, the fixed waste mixture behaves as a soil with regard to diffusion of the constituent of interest.
- No biodegradation of the constituent of interest occurs in open landfills or wastepiles.

6.4.2 Model Plant Parameters for Open Landfills and Wastepiles

The characteristics of model facilities for open landfills and wastepiles are discussed here. The model open landfill facility is used as the basis for an example calculation using the model.

6.4.2.1 Parameters for Open Landfills. The model facility for open landfills has a surface area of $1.42 \times 10^8 \text{ cm}^2$ (3.5 acres). This value represents an approximately midrange value from the Westat survey.³⁸ A reasonable value of landfill depth from the Westat survey³⁹ was 458 cm (15 ft). The model open landfill is assumed to be half full, and hence has a waste depth of 229 cm (7.5 ft). The landfill is assumed to contain fixed waste. A standard temperature of 25 °C is assumed to apply.

The waste liquid (before fixation) selected for this model facility is assumed to be a two-phase aqueous/organic containing 20 percent chloroform,

20 percent low-volatility organic, and 60 percent water (by weight). This liquid has an average density of 1.16 g/cm^3 . The fixation industry indicates that waste liquid, when combined with fixative, may increase in volume by up to 50 percent,⁴⁰ depending on the specific combination of waste and fixative. In view of the inherent variability in the fixation process and the lack of real data on volume changes, for purposes of this report, the assumption is made that the waste volume does not change during fixation. Measurements⁴¹ performed on various types of fixed waste yielded a broad range of total porosities. Fifty percent,* as used in this study, is a reasonable estimate of this parameter. A 25-percent* air porosity appears to be a reasonable value; this value was inferred from measurements of total porosity and moisture content.⁴² As discussed previously, there is no evidence of significant biomass in any chemical waste landfill. Therefore, in this analysis it is assumed, as suggested in the literature, that the toxic property of the waste will inhibit biological processes and thus prevent biogas generation.⁴³ Hence, the waste biomass concentration is taken to be 0 g/cm^3 .

The properties of chloroform that are pertinent to this analysis include the molecular weight (119.4 g/g mol), pure component vapor pressure (208 mm Hg), and diffusivity in air ($0.104 \text{ cm}^2/\text{s}$). The low-volatility organic liquid present in the waste has a molecular weight of 147 g/g mol .

Table 6-7 summarizes the model facility parameters for open landfills used in the example calculation in Section 6.4.3.

6.4.2.2 Parameters for Wastepiles. A review of information in the Westat survey⁴⁴ led to the selection of an approximately midrange value for basal area of $4.65 \times 10^6 \text{ cm}^2$. For modeling purposes, the pile is assumed to be flat. A uniform height of 100 cm was inferred, using the Westat information and engineering judgment. All waste ultimately disposed of in the landfill is assumed to be stored initially in the wastepile. The open landfill described previously (Section 6.4.2.1) is assumed to be filled to capacity in 1 yr. Based on the filled landfill volume ($1.42 \times 10^8 \text{ cm}^2 \times 458 \text{ cm depth} = 6.50 \times 10^{10} \text{ cm}^3$), the wastepile volume ($4.65 \times 10^6 \text{ cm}^2 \times$

*These porosity values are assumed to be representative of waste in an open landfill, rather than waste that has recently undergone fixation.

TABLE 6-7. INPUT PARAMETERS--OPEN LANDFILL

Area	$1.42 \times 10^8 \text{ cm}^2$ (3.5 acres)
Waste depth	229 cm (7.5 ft) ^a
Volume	$3.25 \times 10^{10} \text{ cm}^3$
Temperature	25°C
Waste liquid (before fixation)	Two-phase aqueous/organic
Liquid composition	20% chloroform, 20% low-volatility organic, 60% water (by weight)
Liquid density (average)	1.16 g/cm^3
Liquid/fixative	1 unit volume liquid + dry fixative = 1 unit volume fixed waste
Air porosity fixed waste	0.25 (25%)
Total porosity fixed waste	0.50 (50%)
Biomass concentration	0 g/cm^3
<u>Chloroform properties</u>	
Molecular weight	119.4 g/g mol
Vapor pressure	208 mm Hg
Diffusivity in air	$0.104 \text{ cm}^2/\text{s}$
<u>Low-volatility organic properties</u>	
Molecular weight	147 g/g mol

^aRepresents half full.

100 cm = $4.65 \times 10^8 \text{ cm}^3$), and the filling time of 1 yr, it can be concluded that the wastepile undergoes a turnover rate of 140 turnovers/yr. Hence, the wastepile retention time is 2.6 d/turnover. The properties of the waste liquid and the resulting fixed waste accommodated by the model wastepile are identical to those for the open landfill (Section 6.4.2.1) and will not be repeated here. Table 6-8 summarizes the model facility parameters used for wastepiles.

6.4.3 Example Calculation for Open Landfill

This section presents a step-by-step calculation of emissions from an open landfill. The equations identified in Table 6-6 are used with the model unit parameters in Section 6.4.2 to estimate emissions from a fixed, two-phase aqueous/organic waste containing chloroform; the same equations would be applied to the estimation of emissions from wastepiles:

- Waste liquid (before fixation): 20 percent chloroform,
20 percent low-volatility
organic liquid, 60 percent
water

• Liquid/fixative: 1 unit volume liquid + dry fixative = 1 unit
volume fixed waste

- Waste biomass concentration: 0 g/cm^3
- Landfill area: $1.42 \times 10^8 \text{ cm}^2$ (3.5 acres)
- Landfill depth: 229 cm (7.5 ft)
- Temperature: 25 °C
- Time period for emission calculation: $3.15 \times 10^7 \text{ s}$ (1 yr).

a. Compute waste loading, L:

Liquid density before fixation = 1.16 g/cm^3

1 cm^3 liquid waste + fixative = 1 cm^3 fixed waste

$$L = \text{g organic phase/cm}^3 \text{ fixed waste} \\ = (0.20 + 0.20) \times 1.16 \text{ g/cm}^3 = 0.46 \text{ g/cm}^3$$

(Note that weight fraction of chloroform in the oil phase [C] = $0.2/(0.2 + 0.2) = 0.50$.)

TABLE 6-8. INPUT PARAMETERS--WASTEPILES

Surface area	$4.65 \times 10^6 \text{ cm}^2$
Average height	100 cm
Turnover rate	139/yr
Retention time	2.6 d
Temperature	25 °C
Windspeed	4.92 m/s
Waste type	Fixed waste
Waste liquid (before fixation)	Two-phase aqueous/organic
Liquid composition	20% chloroform, 20% low-volatility organic, 60% water (by weight)
Liquid density (average)	1.16 g/cm^3
Liquid/fixative	1 unit volume liquid + dry fixative = 1 unit volume fixed waste
Air porosity fixed waste	0.25 (25%)
Total porosity fixed waste	0.50 (50%)
Biomass concentration	0 g/cm^3
<u>Chloroform properties</u>	
Molecular weight	119.4 g/g mol
Vapor pressure (25 °C)	208 mm Hg
Diffusivity in air (25 °C)	$0.104 \text{ cm}^2/\text{s}$
<u>Low-volatility organic properties</u>	
Molecular weight	147 g/g mol

b. Compute effective diffusion coefficient for fixed waste:

$$D_e = D_a \frac{\epsilon_a^{3.33}}{\epsilon_T^2}$$

where

ϵ_a = air porosity fixed waste = 0.25

ϵ_T = total porosity fixed waste = 0.50.

Then

D_a = diffusivity of chloroform in air = 0.104 cm²/s

$$D_e = (0.104 \text{ cm}^2/\text{s}) \frac{(0.25)^{3.33}}{(0.50)^2}$$

$D_e = 4.11 \times 10^{-3} \text{ cm}^2/\text{s}$. (Note: $D_e/D_a = 3.96 \times 10^{-2}$.)

-c.- Compute "partition" coefficient, K_{eq} (ratio of gas-phase chloroform to total chloroform in the waste):

For oily waste,

$$K_{eq} = \frac{P^* MW_{oil}}{R T} \frac{\epsilon_a}{L}$$

where

P^* = pure component vapor pressure of chloroform = (208 mm Hg) / (760 mm Hg/atm) = 0.274 atm

MW_{oil} = molecular weight low-volatility organic = 147 g/g mol

R = ideal gas constant = 82.05 cm³•atm/g mol•K

T = temperature within solid waste, K

$T = 273 \text{ K} + 25 \text{ }^\circ\text{C} = 298 \text{ K}$

$$K_{eq} = \frac{(0.274 \text{ atm})(147 \text{ g/g mol})(0.25)}{(82.05 \text{ cm}^3 \cdot \text{atm/g mol} \cdot \text{K})(298 \text{ K})(0.46 \text{ g/cm}^3)}$$

$K_{eq} = 8.95 \times 10^{-4}$.

d. Compute fraction of total chloroform emitted, F_t , after 1 year:

First, determine which solution applies by computing $\frac{K_{eq} D_e t}{l^2}$ (Table 6-6):

$$\begin{aligned}\frac{K_{eq} D_e t}{l^2} &= \frac{8.95 \times 10^{-4} \times 4.11 \times 10^{-3} \text{ cm}^2/\text{s}}{(229 \text{ cm})^2} \\ &= 7.01 \times 10^{-11} \text{ s}^{-1}\end{aligned}$$

Therefore,

$$\begin{aligned}\frac{K_{eq} D_e t}{l^2} &= 7.01 \times 10^{-11} \text{ s}^{-1} \times 3.15 \times 10^7 \text{ s} \\ &= 2.21 \times 10^{-3}\end{aligned}$$

$$K_d t = \frac{K_{eq} D_e t}{l^2} \frac{\pi^2}{4} = 5.45 (10^{-3})$$

Because $K_{eq} D_e t / l^2$ is less than 0.25, the first equation of Table 6-6 applies, and

$$F_t = 0.72 (K_d t)^{1/2}$$

$$F_t = 0.72 (5.45 \times 10^{-3})^{1/2}$$

$$F_t = 0.053$$

e. Compute instantaneous emission rate, E , after 1 yr:

1. Compute initial mass of chloroform in landfill:

$$M_{oi} = l L C$$

where

$$l = \text{waste depth} = 229 \text{ cm}$$

$$L = \text{g organic/cm}^3 \text{ fixed waste} = 0.46 \text{ g/cm}^3$$

$$C = \text{weight fraction chloroform in oil} = 0.50.$$

Then

$$M_0 = (229 \text{ cm})(0.46 \text{ g/cm}^3)(0.50)$$

$$M_0 = 52.7 \text{ g/cm}^2$$

3. Compute instantaneous emission rate, E_i . Because $Keq D_e t/l^2 < 0.213$, use the following equation to compute the emission rate:

$$E = \frac{M_0}{t} \left[\frac{1}{\frac{\epsilon_a}{k_G Keq} + \frac{\pi t}{Keq D_e}} \right]$$

$$k_G = 4.82 (10^{-3}) U^{0.78} Sc_G^{-0.67} de^{-0.11}$$

$$U = \text{windspeed} = 4.92 \text{ m/s}$$

$$de = \text{effective diameter of landfill area} = \left(\frac{4A}{\pi} \right)^{0.5} = 134 \text{ m}$$

$$Sc_G = \frac{\mu g}{\rho_a D_a}$$

$$\text{where: } \mu g = \text{viscosity of air} = 1.8 (10^{-4}) \text{ g/cm/s.}$$

$$\rho_a = \text{Density of air} = 1.2 (10^{-3}) \text{ g/cm}^3.$$

$$D_a = 0.104 \text{ cm}^2/\text{s}.$$

$$Sc_G = \frac{1.81 (10^{-4})}{1.2 (10^{-3}) (0.104)} = 1.45$$

$$k_G = 4.82 (10^{-3}) (4.92)^{0.78} (1.45)^{-0.67} (134)^{-0.11}$$

$$= 0.0076 \text{ m/s} = 0.76 \text{ cm/s}$$

$$E = \frac{52.7}{229} \left[\frac{1}{\frac{0.25}{0.76 \times 8.95(10^{-4})} + \frac{3.14 \times 3.15(10^7)}{8.95(10^{-4}) \times 4.11(10^{-3})}} \right] = 0.230 \left[\frac{1}{368 + 5.18(10^6)} \right]$$

$$E = 4.43 (10^{-8}) \text{ g/cm}^2/\text{s}.$$

6.5 REFERENCES

1. GCA Corporation. Air Emissions from Land Treatment-Emissions Data and Model Review. Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. August 1985. 120 pp.

2. Reference 1.
3. Farmer, W. J., M. S. Yang, J. Letey, W. F. Spencer, and M. H. Roulier. Land Disposal of Hexachlorobenzene Wastes: Controlling Vapor Movement in Soils. Fourth Annual Research Symposium. U.S. Environmental Protection Agency. Publication No. EPA-600/9-78-016. August 1978.
4. Millington, R. J., and J. P. Quirk. Permeability of Porous Solids. Trans. Faraday Society. 57:1200-1207. 1961.
5. U.S. Environmental Protection Agency. Evaluation and Selection of Models for Estimating Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Office of Air Quality Planning and Standards, Research Triangle Park, NC. Publication No. EPA-450/3-84-020. December 1984.
6. Reference 3.
7. Thibodeaux, L. J. Estimating the Air Emissions of Chemicals from Hazardous Waste Landfills. Journal of Hazardous Materials. 4:235-244. 1981.
8. Reference 3.
9. Reference 3.
10. Shen, T. T. Estimating Hazardous Air Emissions from Disposal Sites. Pollution Engineering. 31-34. August 1981.
11. Reference 3.
12. Reference 4.
13. Reference 3.
14. Westat, Inc. National Survey of Hazardous Waste Generators and TSD Facilities Regulated Under RCRA in 1981. Prepared for U.S. Environmental Protection Agency. Contract No. 68-61-6861. 1981.
15. Reference 14.
16. Reference 14.
17. Reference 14.
18. Ely, R. L., G. L. Kingsbury, M. R. Branscome, L. J. Goldman, C. M. Northeim, J. H. Turner, and F. O. Mixon, Jr. Performance of Clay Caps and Liners for Disposal Facilities. Research Triangle Institute, Research Triangle Park, NC. Prepared for U.S. Environmental Protection Agency, Cincinnati, OH. EPA Contract No. 68-03-3149. March 1983.

19. Telecon. Goldman, Leonard, Research Triangle Institute, with Borden, Roy, Department of Civil Engineering, North Carolina State University, Raleigh, NC. August 13, 1986.
20. Gerachty, J. J., D. W. Miller, F. Vander Leeden, and F. L. Troise. Water Atlas of the United States. Water Information Center, Inc., Port Washington, NY. 1973. Plate 30.
21. Telecon. Goldman, Leonard, Research Triangle Institute, with Hughes, John, National Climatic Center, Asheville, NC. May 15, 1986.
22. Reference 21.
23. Telecon. Goldman, Leonard, Research Triangle Institute, with Boyenga, Dave, MBI Corporation, Dayton, OH, November 20, 1985.
24. Telecon. Goldman, Leonard, Research Triangle Institute, with Webber, Emlyn, VFL Technology Corporation, Malvern, PA. November 12, 1985.
25. Telecon. Massoglia, Martin, Research Triangle Institute, with Webber, Emlyn, VFL Technology Corporation, Malvern, PA. January 13, 1987.
26. Telecon. Goldman, Leonard, Research Triangle Institute, with Hannak, Peter, Alberta Environmental Center. April 4, 1986.
27. ~~Reference~~ 10.
28. Reference 5.
29. Arnold, J. H. Studies in Diffusion: III. Unsteady-State Vaporization and Absorption. Transactions of the American Institute of Chemical Engineers. 40:361-379. 1944.
30. Reference 29.
31. Reference 29.
32. Letter and attachment from Shen, T., New York State Department of Environmental Conservation, to McDonald, R., EPA/OAQPS. Modification of Arnold's equation. February 16, 1986. (See also Reference 10.)
33. Trip Report. Goldman, Leonard, Research Triangle Institute, with Chemical Waste Management, Sulphur, Louisiana. February 25, 1986.
34. Reference 1.
35. Reference 5.
36. Reference 10.

37. Memorandum and attachment from M. Wright, Research Triangle Institute, to S. Thorneloe, EPA/OAQPS. May 30, 1986. Selection of an emissions model for land treatment.
38. Reference 14.
39. Reference 14.
40. References 23, 24, and 25.
41. Reference 26.
42. Reference 26.
43. Reference 10.
44. Reference 14.

7.0 TRANSFER, STORAGE, AND HANDLING OPERATIONS

7.1 NARRATIVE DESCRIPTION OF MODEL PLANTS AND EMISSIONS

This chapter presents models for estimating VO emissions of hazardous wastes from container loading, storage, and cleaning; stationary tank loading and storage; spills; fugitive sources; and vacuum truck loading.

7.2 CONTAINER LOADING

This section addresses emission-estimating practices for hazardous waste loading into tank trucks, railroad tank cars, marine vessels, and 55-gal (0.208-m³) drums.

7.2.1 Emission Model for Container Loading

The process of loading containers with volatile hazardous wastes generates emissions. If containers were assumed to be clean when loaded, only those vapors generated by the loaded waste could be emitted. If containers hold residues of previous volatile wastes, additional emissions from the residues will also be generated.

To calculate container loading losses, the AP-42¹ equation for loading petroleum liquids is applied. This equation was derived for tanks, cars, and marine vessels. It is also applied to 55-gal drums in this case because the loading principles are similar and because no equation has been developed exclusively for small containers such as drums. The loading equation is as follows:

$$L_L = \frac{12.46 \text{ SMP}^*}{T} \quad (7-1)$$

where

L_L = loading loss, lb/10³ gal of liquid loaded

M = molecular weight of vapors, lb/lb mol

P^* = true vapor pressure of liquid loaded, psia

T = bulk temperature of liquid loaded, °R (°F + 460).

S = saturation factor, dimensionless (see Table 7-1).

Equation (7-1) for estimating emissions from containers is not applicable to open dumpsters because they are designed with no tops, unlike drums that have limited venting through bungs.

7.2.2 Model Parameters

Containers are considered to be splash-loaded (as opposed to submerged-loaded) for emission-estimating purposes. This loading method creates larger quantities of V₀ vapors and increases the saturation factor, S , of each volatile compound within the container. A saturation factor is a dimensionless quantity that "represents the expelled vapors' fractional approach to saturation and accounts for the variations observed in emission rates from the different unloading and loading methods."² A saturation factor of 1.45 was selected for these emission estimates, based on previous documentation of splash-loading petroleum liquids.^{3,4}

Typical capacities for containers are assumed to be as follows:

- Drums: 55 gal (0.208 m³)
- Tank trucks: 7,000 gal (26.5 m³)
- Railroad tank cars: 30,000 gal (114 m³)
- Marine vessels: 20,000 tons.

It is assumed that 55-gal drums and tank trucks are the predominant containers used in the waste management industry. Bulk liquid hazardous waste is shipped predominantly by highway; therefore, hazardous waste tank truck models are used for estimating emissions.

Molecular weight and vapor pressure are functions of the waste loaded, and 25 °C is considered an annual average ambient operating temperature.

7.2.3 Sample Calculation for Tank Loading

The following sample calculation may be used to estimate V₀ emissions from tank truck loading of an organic liquid. Waste stream compounds and properties for the sample calculation are as follows. The same waste stream is employed in each sample calculation in this section; only the type of emission source is varied.

TABLE 7-1. S FACTORS FOR CALCULATING PETROLEUM
LOADING LOSSES

Cargo carrier	Mode of operation	S factor
Tank trucks and tank cars	Submerged loading of a clean cargo tank	0.50
	Splash loading of a clean cargo tank	1.45
	Submerged loading: normal dedicated service	0.60
	Splash loading: normal dedicated service	1.45
	Submerged loading: dedicated vapor balance service	1.00
	Splash loading: dedicated vapor balance service	1.00
Marine vessels ^a	Submerged loading: ships	0.2
	Submerged loading: barges	0.5

^aTo be used for products other than gasoline.

Constituent	Weight fraction	Molecular weight (lb/mol)	Vapor pressure (psia)	Mole fraction
Benzene	0.3	78	1.84	0.368
Naphthalene	0.3	128	0.0044	0.224
Phenol	0.4	94	0.0066	0.408

The input parameters for the truck loading model are as follows:

- True vapor pressure of loading liquid, psia: 0.68 (calculated in a., below)
- Molecular weight of vapor, lb/mol: 78.23 (calculated in b., below)
- Saturation factor, dimensionless: 1.45 (splash loading)
- Bulk temperature of liquid loaded, °R: 537
- Annual throughput, gal/yr: 28,000

a. Calculate P^* , true vapor pressure of liquid, by Raoult's Law:

$$P^* = (P_1 \cdot X_1) + (P_2 \cdot X_2) + (P_3 \cdot X_3)$$

where

P^* = true vapor pressure, psia

P_1 , P_2 , and P_3 = vapor pressures of pure components

X_1 , X_2 , and X_3 = mole fractions of VO components in liquid

$$\begin{aligned}
 P^* &= (1.84 \text{ psia} \times 0.368) + (0.0044 \text{ psia} \times 0.224) + (0.0066 \\
 &\quad \text{psia} \times 0.408) \\
 &= 0.68 \text{ (psia)}.
 \end{aligned}$$

b. Calculate M , molecular weight of vapors:

$$M = \frac{(P_1 \cdot X_1)}{P^*} \cdot M_1 + \frac{(P_2 \cdot X_2)}{P^*} \cdot M_2 + \frac{(P_3 \cdot X_3)}{P^*} \cdot M_3$$

where

M = molecular weight of vapor

M₁, M₂, and M₃ = molecular weight of each component

$$\begin{aligned} M &= \left(\frac{1.84 \times 0.368}{0.68} \right) \times 78 + \left(\frac{0.0044 \times 0.224}{0.68} \right) \\ &\quad \times 128 + \left(\frac{0.0066 \times 0.408}{0.68} \right) \times 94 \\ &= 78.23 \text{ (lb/mol)}. \end{aligned}$$

c. Calculate emissions from truck loading:

$$\begin{aligned} L_L &= \frac{12.46 \text{ SMP}^*}{T} \\ &= \frac{12.46 \times 1.45 \times 78.23 \times 0.68}{537 \text{ }^\circ\text{R}} \\ &= 1.79 \text{ lb/1,000 gal} \end{aligned}$$

$$\begin{aligned} \text{Annual emissions, } L_L &= \frac{1.79 \times 10^{-3} \text{ lb/gal} \times 28,000 \text{ gal/yr}}{2,205 \text{ lb/Mg}} \\ &= 0.023 \text{ Mg/yr} \end{aligned}$$

7.3 CONTAINER STORAGE

This section addresses storage emissions from tank trucks, railroad tank cars, 55-gal drums, marine vessels, and open dumpsters.

7.3.1 Emission Model for 55-Gal Drums, Tank Trucks, and Railroad Tank Cars

With regard to 55-gal drums, container storage is considered a location where multiple drums are most likely to accumulate and be stored for more than 90 days. Because drums are designed to be stored with a sealed lid and bung, the potential for breathing losses is minimal. Therefore, breathing loss is assumed to be negligible. However, the potential does exist for a drum to rupture or become damaged and leak during storage. Thus, the emissions from drum storage may be estimated using the same spill fraction used for drum handling-- 10^{-4} (to be discussed in more detail in Section 7.7, Spills). The following equation is used to estimate emissions from drum storage:⁵

$$E = 10^{-4} \times I \times W_i \times V_i \quad (7-2)$$

where

E = emission from drum storage, Mg/yr

I = throughput, Mg/yr

W_i = V0 weight fraction

V_i = volatilization fraction.

Emission-estimating methodologies have not been developed for storage in tank trucks and railroad tank cars. Only loading information was available in the literature for these containers. The assumed same emission estimates principle for drum storage is applied with an emission factor of 10^{-5} (to be discussed in more detail in Section 7.7, Spills).⁶

7.3.2 Model Parameters for Drum Storage

It is assumed that 50 percent of the V0 storage loss from drum spill or rupture will be volatilized. The remaining volatiles will be captured with RCRA spill response measures taken at the facility.

7.3.3 Sample Calculations for Drum Storage

Input parameters:

Waste stream: organic liquid

(See Section 7.2.3 for constituents.)

Waste density: 1.04 Mg/m³

Drum storage capacity: 182 drums (0.208 m³/drum)

Turnovers per year: 12

Spill fraction: 10^{-4}

Weight fraction: 1

Volatilization fraction: 0.5.

a. Calculate annual throughput, I , Mg/yr:

$$\begin{aligned} I &= 182 \times 0.208 \text{ m}^3 \times 12 \times 1.04 \text{ Mg/m}^3 \\ &= 472 \text{ Mg/yr} \end{aligned}$$

b. Calculate air emissions:

$$\begin{aligned} E &= 10^{-4} \times I \times W_i \times V_i \\ &= 10^{-4} \times 472 \text{ Mg/yr} \times 1 \times 0.5 \\ &= 0.024 \text{ Mg/yr} \end{aligned}$$

7.3.4 Emission Model for Open Dumpsters

No information was found in readily available literature to estimate VO emissions from the storage of hazardous wastes in open-top dumpsters. The wastes held in dumpsters may range from sludges to contaminated filters. An emission algorithm^{7,8} was developed for losses from an open dump. Because an open dumpster is similar to an open dump, this algorithm was used to calculate storage emissions. (See Section 6.3 for a detailed derivation of the open dump model.)

$$E_i = \frac{2 P_o MW_i Y_i^* w}{RT} \sqrt{\frac{D_i l U}{\pi F_v}} \quad (7-3)$$

The open dump model is valid for short-term emission estimates only. For long-term emission estimates, to avoid overestimates, it is important to subtract the emissions from the waste's VO content on a daily, weekly, or monthly basis depending on the waste volatility; this is done to identify the point in time when no VO remain in the waste.

7.3.5 Model Parameters for Open Dumpster Storage

This section is currently under review for potential revisions.

P_o = atmospheric pressure (mm Hg)

MW_i = molecular weight of compound i , g/g-mole

$Y_i^* = \frac{X_i P^*}{P_t}$ where X_i = mole fraction of compound (i) in liquid phase

P^* = vapor pressure of compound (i), mm Hg

P_t = total system operating pressure, mm Hg

E_i = Air Emissions of compound (i), g/dumpster

T = temperature ($^{\circ}\text{K}$)

R = ideal gas constant = $82.05 \frac{\text{cm}^3 \cdot \text{atm}}{\text{g mole} \cdot ^{\circ}\text{K}}$

D_i = diffusivity of component (i) in air.

U = wind speed (m/sec)

7.4 CONTAINER CLEANING

7.4.1 Emission Model for Container Cleaning

An AP-42 document⁹ on tank truck cleaning is used as the primary source for container-cleaning emission estimates. AP-42 states that tank truck cleaning typically involves washing the truck interior with agents such as water, steam, detergents, or other chemicals. The document also

provides emission factors that are a function of vapor pressure and viscosity. These factors have been applied to emission estimates for cleaning all types of containers, as follows:

Tank truck residue to be removed	Emission factor g/truck (lb/truck)
High vapor pressure, low viscosity	215 (0.474)
Medium vapor pressure, medium viscosity	32.4 (0.071)
Low vapor pressure, low viscosity	5.5 (0.012)

The following equation is used to estimate emissions for container cleaning:

$$E = F_C \times N \times W_i \times 10^{-6} \quad (7-3)$$

where

E = cleaning loss, Mg/yr

F_C = emission factor for cleaning, g/container

N = number of cleanings per year

W_i = VO weight fraction.

7.4.2 Model Parameters

In all containers, the primary input parameter for estimating cleaning emissions is the cleaning emission factor, which is determined from (1) residue vapor pressure and viscosity (functions of waste handled), and (2) container volume.

Based on AP-42,¹⁰ a typical tank truck volume of 26.5 m³ (7,000 gal) is assumed.

Because no data are currently available for drum cleaning, the emission factors for tank trucks were used to calculate cleaning emissions from drums by comparing the proportion of drum volume (55 gal) to that of the tank truck (7,000 gal).

55-Gal drum residue to be removed	Emission factor	
	g/drum	(lb/drum)
High vapor pressure, low viscosity	1.69	(0.0037)
Medium vapor pressure, medium viscosity	0.25	(0.0006)
Low vapor pressure, low viscosity	0.04	(0.00009)

Emissions from marine vessels have not been addressed because of the low usage of such vessels in the waste management industry.

7.4.3 Sample Calculation for Tank Truck Cleaning

The general assumptions for truck cleaning are as follows:

- Residue: pure organic liquid (benzene)
- Number of truck cleanings per year: 4
- Truck capacity: typical truck
- Weight fraction: 1.

- a. Determine the cleaning emission factor, F_c :

(215 g/truck was used because of high vapor pressure and low viscosity of pure benzene residue).

- b. Calculate cleaning emissions:

$$\begin{aligned}
 E &= F_c \times N \times W_i \times 10^{-6} \\
 &= 215 \text{ g} \times 4 \times 1 \times 10^{-6} \text{ Mg/g} \\
 &= 8.6 \times 10^{-4} \text{ Mg/yr}
 \end{aligned}$$

7.5 STATIONARY TANK LOADING

7.5.1 Emission Model for Stationary Tank Model

Stationary tank working losses are those emissions from waste loading and unloading operations. AP-42's "Storage of Organic Liquids"¹¹ provides an equation to estimate loading and unloading emissions from storage tanks. The equation was developed for handling VO liquid in the following industries:

- Petroleum producing/refining
- Petrochemical and chemical manufacturing
- Bulk storage and transfer operations
- Other industries consuming or producing organic liquids.

Because hazardous wastes have the potential to contain VO compounds, as do organic liquids, and because they are most commonly stored in the same fashion as these liquid products, the following equation was selected from AP-42:

$$L_w = 2.40 \times 10^{-5} M_v \cdot P^* \cdot V \cdot N \cdot K_n \cdot K_c \quad (7-4)$$

where

L_w = working losses, lb/yr

M_v = molecular weight of vapor in storage tank, lb/lb mol

P^* = true vapor pressure at bulk liquid conditions, psia

N = number of turnovers per year (dimensionless)

$N = \frac{\text{total throughput per year (gal)}}{\text{tank capacity, } V \text{ (gal)}}$

V = tank capacity, gal

K_n = turnover factor, dimensionless (for turnovers ≤ 36 , $K_n = 1$;
for turnovers > 36 , $K_n = \frac{180 + N}{6N}$)

K_c = product factor, dimensionless (for crude oil, $K_c = 0.84$; for all other organic liquids, $K_c = 1$).

7.5.2 Model Parameters

It is assumed that all stationary tanks are fixed-roof. According to responses to the 1982 Westat Mail Survey,¹² which were examined by the GCA Corporation,¹³ there are four sizes of tanks that best represent the waste management industry:

- 5.3 m³ (1,500 gal)
- 30.3 m³ (8,000 gal)
- 75.0 m³ (20,000 gal)
- 795 m³ (210,000 gal).

Table 7-2 lists typical input parameters for these model tanks. Turnovers per year were selected based on volume of waste processed in waste management scenarios recorded in various documents. Molecular weight and vapor pressure are a function of the waste loaded.

7.5.3 Sample Calculation for Tank Loading Emission Model

Input parameters:

Waste stream: organic liquid (see Section 7.2.3 for constituents)

M_v, molecular weight of vapor, lb/lb mol: 78.23

P*, true vapor pressure of loading liquid, psia: 0.68

K_c, product factor for working loss: 1

V, fixed-roof tank capacity, gal: 20,000

N, turnovers per year: 44

K_n, turnover factor, dimensionless: 0.848.

a. Calculate M_v, molecular weight of vapor:

(see Section 7.2.3 for calculation).

b. Calculate P*, true vapor pressure of loading liquid:

(see Section 7.2.3 for calculation).

c. Calculate K_n, turnover factor: because N = 44, $K_n = \frac{180 + N}{6N} = 0.848$

d. Calculate air emissions:

$$\begin{aligned}
 L_w &= 2.40 \times 10^{-5} \times M_v \cdot P^* \cdot V \cdot N \cdot K_n \cdot K_c \\
 &= 2.40 \times 10^{-5} \times 78.23 \times 0.68 \times 20,000 \times 44 \times 0.848 \times 1 \\
 &= 953 \text{ lb/yr} \\
 &= 0.43 \text{ Mg/yr}
 \end{aligned}$$

TABLE 7-2. PERTINENT FIXED-ROOF TANK SPECIFICATIONS^{14,15,16}

Specifications	Model A	Model B	Model C	Model D
Capacity, m ³ (gal)	5.3 (1,500)	30.3 (8,000)	75.7 (20,000)	795 (210,000)
Tank height, m	2.4	2.4	2.7	12.2
Tank diameter, m	1.7	4	5.8	9.1
Average vapor space height, m	1.2	1.2	1.4	6.1
Adjustment for small diameter (dimensionless)	0.26	0.65	0.86	1
Average diurnal temp. change, °C (°F)	11 (20)	11 (20)	11 (20)	11 (20)
Paint factor (dimensionless)	1	1	1	1
Relation of tank to ground	Above	Above	Above	Above
Product factor	1	1	1	1

7.6 STATIONARY TANK STORAGE

7.6.1 Model Description

Fixed-roof tank storage of hazardous wastes results in V0 "breathing" emissions through vents as ambient temperature and barometric pressure fluctuate. Emissions occur in the absence of any liquid level change in the tank. An existing AP-42¹⁷ equation was used to estimate V0 breathing losses from hazardous waste storage tanks as follows:

$$L_b = 2.26 \times 10^{-2} M_v \left(\frac{P^*}{P_A - P^*} \right)^{0.68} \cdot D^{1.73} \cdot H^{0.51} \cdot \Delta T^{0.5} \cdot F_p \cdot C \cdot K_C \quad (7-5)$$

where

L_b = fixed-roof breathing loss, lb/yr

M_v = molecular weight of vapor in tank, lb/lb mol

P^* = true vapor pressure at bulk liquid conditions, psia

P_A = average atmospheric pressure at tank location, psia

D = tank diameter, ft

H = average vapor space height, ft (assumed to be one-half of tank height)

ΔT = average ambient diurnal temperature change, °F (20 °F assumed as a typical value)

F_p = paint factor, dimensionless (see Table 7.3)

C = adjustment factor for small diameter tanks, dimensionless
(for diameter ≥ 30 ft, $c = 1$; for diameter < 30 ft,
 $c = 0.0771 D - 0.0013 D^2 - 0.1334$)

K_C = product factor, dimensionless (for crude oil, $K_C = 0.65$, for all other organic liquids, $K_C = 1.0$).

The above equation requires an estimation of the true vapor pressure using the liquid concentration. For very volatile constituents, the liquid concentration depends on the amount lost as air emissions. To correct for the loss to the air in estimating the liquid concentration, the following equation may be used:

$$\text{fraction lost to air} = \frac{L_b}{L_b + L_t}$$

where L_t is the tank input of the volatile constituent in pounds per year.

TABLE 7-3. PAINT FACTORS FOR FIXED-ROOF TANKS¹⁸

Tank color		Paint factors (F_p)	
		Paint condition	
Roof	Shell	Good	Poor
White	White	1.00	1.15
Aluminum (specular)	White	1.04	1.18
White	Aluminum (specular)	1.16	1.24
Aluminum (specular)	Aluminum (specular)	1.20	1.29
White	Aluminum (diffuse)	1.30	1.38
Aluminum (diffuse)	Aluminum (diffuse)	1.39	1.46
White	Gray	1.30	1.38
Light gray	Light gray	1.33	1.44 ^a
Medium gray	Medium gray	1.40	1.58 ^a

^aEstimated from the ratios of the seven preceding paint factors.

7.6.2 Model Parameters

Table 7-3 identifies the model parameters for estimating tank breathing losses. Molecular weight and vapor pressure are functions of the waste stored.

7.6.3 Sample Calculation for Tank Storage Emission Model

Input parameters:

Waste stream, organic liquid (see Section 7.2.3 for constituents)

M_v , molecular weight of vapor, lb/lb mol: 78.23

P^* , true vapor pressure of loading liquid, psia: 0.68

K_C , product factor for breathing loss: 1

v , fixed-roof tank capacity, gal: 20,000

D , tank diameter, ft: 19

H , average vapor space height, ft: 4.5

ΔT , diurnal temperature change, °F: 20

F_p , paint factor, dimensionless: 1

C , adjustment factor for small tanks: 0.86 (calculate in c., below)

a. Calculate molecular weight of vapor:

(see Section 7.2.3 for calculation).

b. Calculate true vapor pressure of loading liquid:

(see Section 7.2.3 for calculation).

c. Calculate adjustment factor for small tanks:

$$C = 0.0771 \times 19 - 0.0013(19)^2 - 0.1334 \\ = 0.86$$

d. Calculate air emissions:

$$L_b = 2.26 \times 10^{-2} M_v \left[\frac{P^*}{P_A - P^*} \right]^{0.68} \cdot D^{1.73} \cdot H^{0.51} \cdot \Delta T^{0.5} \cdot F_p \cdot C \cdot K_C \\ = 2.26 \times 10^{-2} \times 78.23 \times \left[\frac{0.68}{14.7 - 0.68} \right]^{0.68} \times (19)^{1.73} \times (4.5)^{0.51} \\ \times (20)^{0.5} \times 1 \times 0.86 \times 1 \\ = 300 \text{ lb/yr} \\ = 0.14 \text{ Mg/yr}$$

7.7 SPILLS

7.7.1 Model Description

An ICF study¹⁹ of truck transport to and from TSDF and truck emissions at TSDF terminals provided the background information necessary to estimate spillage losses during TSDF and trucking operations. As a result of this study, spill fractions of 10^{-4} and 10^{-5} were assumed for drum movement of wastes and all other remaining waste movement, respectively. Thus, for every 10,000 Mg of drummed hazardous waste moved, 1 Mg is assumed to be spilled. The following equation is used to estimate spill emissions:

$$E = F_s \times I \times W_i \times V_i \quad (7-6)$$

where

E = spill emissions, Mg/yr

F_s = emission fraction, 10^{-4} or 10^{-5}

I = annual throughput, Mg/yr

W_i = VO weight fraction

V_i = fraction for volatilization.

7.7.2 Model Parameters

In both cases of spills, it is assumed that 50 percent of the volatiles in the waste are lost. The remaining 50 percent are recovered by RCRA spill plan response. Therefore, most spills would be mitigated before 100 percent of VO is lost to the atmosphere.

It is assumed that spills do not occur during the transfer of waste into a stationary tank if loading is automated through fixed piping.

7.7.3 Sample Calculation for Drum Storage Model

Input parameters:

Waste stream: organic liquid (see Section 7.2.3 for constituents)

Waste density: 1.04 Mg/m^3

Emission fraction: 10^{-4}

Weight fraction: 1

Volatilization fraction: 0.5

Number of drums handled: 2,184 ($0.208 \text{ m}^3/\text{drum}$).

a. Calculate annual throughput, I, Mg/yr:

$$I = 2,184 \times 0.208 \text{ m}^3 \times 1.04 \text{ Mg/m}^3 \\ = 472 \text{ Mg/yr} .$$

b. Calculate air emissions:

$$E = 10^{-4} \times 472 \text{ Mg/yr} \times 1 \times 0.5 \\ = 0.024 \text{ Mg/yr} .$$

7.8 FUGITIVE EMISSIONS

7.8.1 Emission Model for Fugitives

Waste transfer operations often involve pumping wastes through pipelines into a variety of containment units. Such pumping creates the potential for fugitive emission losses from pumps, valves, and flanges. Table 7-4 presents the Synthetic Organic Chemical Manufacturing Industries (SOCMI) emission factors²⁰ that had been developed to estimate VO that leak from pump seals, valves, and flanges. These factors are independent of the throughput, type, or size of the process unit.

TABLE 7-4. SOCMI EMISSION FACTORS FOR FUGITIVE LOSSES

Equipment	Type of service	Emission factor (kg/h-source)
Pump seals	Light liquid	4.94 E-2
Valves	Light liquid	7.10 E-3
Flanges	--	8.30 E-4

The following equation is used to estimate fugitive emissions:

$$E = \Sigma (F_f \times N_i) \times h \times 10^{-3} \quad (7-7)$$

where

E = fugitive emissions, Mg/yr

F_f = emission factor per source-type, kg/h-source (see Table 7-4)

N_i = number of sources per source-type

h = residence time in the equipment (assume = 8,760 h/yr).

7.8.2 Model Parameters

The major input parameters required for the emission model are emission factor, number of sources, and residence time. It is assumed that waste remains in the transfer equipment 24 h/d, 365 d/yr; therefore, V0 are continuously being leaked to the atmosphere.

Minimal information has been compiled on typical quantities of pumps, valves, and flanges at waste management facilities. Therefore, previous contractors have turned to data collected from the petroleum refining industry and SOCFI. GCA recommended that "for any hazardous waste filling operation, transfer operation, or handling operation involving pumps, the estimate of two pumps, 35 valves, and 80 flanges be used. This includes tank filling, tank truck or car filling, and drum filling."²¹ Because the relationship 2:35:80 appears to be too high for pumping waste into a single drum, one pump, three valves, and eight flanges are used for estimating emissions. GCA recommended that smaller quantities of pumps, valves, and flanges identified by SOCFI be applied for transfer operations to injection wells and incinerators, i.e., 1 pump, 18 valves, and 40 flanges.²²

7.8.3 Sample Calculation for Fugitive Emission Model

Estimate the annual fugitive emissions from a set of piping lines that connect to a storage tank, given the following information.

Input parameters:

Assume 2 pumps, 35 valves, and 80 flanges associated with the piping equipment.

Assume the waste stream is organic liquid.

Assume waste remains in piping line 24 h/d, 365 d/yr.

a. The emission factor for light liquids was used because of the high V0 content.

b. Calculate fugitive emissions:

$$E = (0.0494 \text{ kg/h} \times 2 + 0.0071 \text{ kg/h} \times 35 + 8.3 \times 10^{-4} \text{ kg/h} \times 8) \\ \times 8,760 \text{ h/yr} \times 10^{-3} \text{ Mg/kg} = 3.62 \text{ Mg/yr}$$

7.9 VACUUM TRUCK LOADING

7.9.1 Emission Model for Vacuum Truck Loading

Emissions from vacuum truck loading are estimated by calculating an equilibrium concentration of organic vapors in the vacuum truck at its

operating conditions and assuming that a total volume of gas equal to the vacuum truck volume is emitted to the atmosphere for each loading episode. Equations for making the calculations are presented as follows:

$$E_i = N_v \times Y_i \times MW_i$$

$$Y_i = \frac{X_i P^*}{P_t} \quad (\text{for oily waste})$$

$$N_v = \frac{V}{[P_o V_G (T/273)]/P_t}$$

where

E_i = air emissions of compound i , g/truckload

N_v = total moles of vapor discharged, g mol

Y_i = mole fraction of compound i in vapor phase

X_i = mole fraction of compound i in liquid phase

MW_i = molecular weight of compound i , g/g mol

P^* = vapor pressure of compound i , mm Hg

P_t = total system operating pressure, mm Hg

P_o = atmospheric pressure, mm Hg

V = vacuum truck volume, m^3

V_G = volume of 1 g mol of gas at STP, $0.0224 m^3/g \text{ mol}$

T = operating temperature, K.

7.9.2 Model Parameters

Based on information obtained during site visits to refineries using land treatment, vacuum trucks have a capacity of about $21 m^3$ (5,500 gal) and operate at a pressure of approximately 303 mm Hg. These values are used in estimating vacuum truck emissions.

Molecular weight and vapor pressure are functions of waste loaded, and 25 °C is considered a standard operating temperature.

7.9.3 Sample Calculation

The following is a sample calculation of benzene emissions during loading of a vacuum truck with organic liquid.

Input parameters:

Waste stream: organic liquid (see Section 7.2.3 for constituents)

V0 constituent: benzene

MW_i, molecular weight, g/g mol: 78

P*, pure compound vapor pressure: 95.2

P_t, system operating pressure, mm Hg: 303

P₀, atmospheric pressure, mm Hg: 760

X_i, mole fraction in liquid: 0.368

V, vacuum truck volume, m³: 21

V_G, volume of 1 g mol of gas at STP, m³/g mol: 0.0224

T, operating temperature, K: 298

N, turnovers per year, truckload/yr: 10.

- a. Calculate total moles of vapor discharged, g mol:

$$\begin{aligned} N_v &= \frac{V}{[P_a V_G (T/273)]/P_t} \\ &= \frac{21 \text{ m}^3}{(760 \text{ mm Hg} \times 0.0224 \text{ m}^3/\text{g mol} \times 298 \text{ K}/273 \text{ K})/303 \text{ mm Hg}} \\ &= 342.41 \text{ g mol/truckload} \end{aligned}$$

- b. Calculate mole fraction of benzene in vapor phase, Y_i:

$$Y_i = \frac{P^* X_i}{P_t} = \frac{95.2}{303} \times 0.368 = 0.1156$$

- c. Calculate air emissions per truckload, g/truckload:

$$\begin{aligned} E_i &= N_v \times Y_i \times \text{MW}_i \\ &= (342.41 \text{ g mol/truckload}) (0.1156) (78 \text{ g/g mol}) \\ &= 3,087 \text{ g/truckload} \end{aligned}$$

- d. Calculate annual emissions for benzene, Mg/yr:

$$\begin{aligned} \text{Annual emission} &= E_i \times N \\ &= 3,087 \text{ g/truckload} \times 10 \text{ truckload/yr} \\ &= 30,870 \text{ g/yr} \\ &= 0.031 \text{ Mg/yr} \end{aligned}$$

- e. Repeat the above procedures, a through d., to compute emissions for each constituent as follows:

Constituent	E _i , g/truckload	Annual emissions, Mg/yr
Benzene	3,087	0.031
Naphthalene	7	0.00007
Phenol	14	0.00014
Total emissions	3,108	0.0312

7.10 REFERENCES

1. U.S. Environmental Protection Agency. Transportation and Marketing of Petroleum Liquids. In: AP-42. Compilation of Air Pollutant Emission Factors. Third Edition, Supplement 12, Section 4.4. Research Triangle Park, NC. Office of Air Quality Planning and Standards. July 1979. 13 pp.
2. GCA Corporation. Air Emission Estimation Methods for Transfer, Storage, and Handling Operations. Draft Technical Note. Prepared for U.S. Environmental Protection Agency, Office of Air Quality Planning and Standards. Research Triangle Park, NC. Contract No. 68-01-6871. August 1985.
3. Reference 1.
4. Reference 2.
5. U.S. Environmental Protection Agency. Assessing the Release and Costs Associated with Truck Transport of Hazardous Wastes. PB 84-224-468 (Contract No. 68-01-0021). Washington, DC. January 1984. 151 p.
6. Reference 5.
7. Shen, T. T. Estimating Hazardous Air Emissions from Disposal Sites. Pollution Engineering. 31-34. August 1981.
8. GCA Corporation. Air Emissions of VOC from Waste Piles at Hazardous Waste Treatment, Storage, and Disposal Facilities. Prepared for U.S. Environmental Protection Agency, Office of Air Quality Planning and Standards. Research Triangle Park, NC. Contract No. 68-01-6871. August 1985.
9. U.S. Environmental Protection Agency. Tank and Drum Cleaning. In: AP-42. Compilation of Air Pollutant Emission Factors. Fourth Edition, Section 4.8. Research Triangle Park, NC. Office of Air Quality Planning and Standards. September 1985. 4 pp.
10. Reference 9.

11. U.S. Environmental Protection Agency. Storage of Organic Liquids. In: AP-42. Compilation of Air Pollutant Emission Factors. Fourth Edition, Section 4.3. Research Triangle Park, NC. Office of Air Quality Planning and Standards. September 1985. 25 pp.
12. Westat, Inc. National Survey of Hazardous Waste Generators and Treatment, Storage, and Disposal Facilities Regulated Under RCRA in 1981. Prepared for U.S. Environmental Protection Agency. Office of Solid Waste, Washington, DC. April 1984.
13. Addendum to Memorandum dated September 6, 1985, from Eichinger, Jeanne, GCA Corporation, to Hustvedt, K. C., EPA/OAQPS. September 12, 1985. TSDF model source parameters and operating practices data base.
14. Reference 11.
15. Reference 13.
16. Graver Standard Cone-Roof, Flat-Bottom Tanks. Sizes and Capacities.
17. Reference 11.
18. TRW-Environmental, Inc. Background Documentation for Storage of Organic Liquids. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. Contract No. 68-02-3174. May 1981.
19. Reference 5.
20. U.S. Environmental Protection Agency. Control of Volatile Organic Compound Leaks from Synthetic Organic Chemical and Polymer Manufacturing Equipment. Research Triangle Park, NC. Publication No. EPA-450/3-83-006. March 1984.
21. Reference 2.
22. Reference 2.

8.0 COMPARISON OF MODEL RESULTS WITH FIELD TEST DATA

8.1 INTRODUCTION

Predictions from TSDf emission models are compared with field test data in this chapter. In general, considering the uncertainty of field emission measurements, agreement between measured and predicted values is considered reasonable. Measured and predicted emissions generally agree within an order of magnitude.

The following caveats must be considered in any evaluation of the comparison results presented in the following sections:

1. The field test data did not always include all of the input parameters required to use the emission models. In such cases, parameter values representative of field operations were used as defaults.
2. The emission models use average influent and effluent concentrations to estimate annual emissions. Variations in concentrations and constituents are not reflected.
3. Field test data provide information on a limited number of hazardous constituents. Extrapolation of comparisons on limited constituents to all constituents of interest may not always be possible.
4. The method of measuring emissions (e.g., flux chambers and other enclosure systems) could alter the real-world system being tested and affect the representativeness of the measured emissions.

8.2 SURFACE IMPOUNDMENTS AND OPEN TANKS

8.2.1 Summary

Emission test data were available from tests of five quiescent surface impoundments. The overall mass transfer coefficients determined in these tests agreed generally within an order of magnitude with the overall coefficient predicted by the mass transfer correlations described in

Chapter 4.0. Predicted values were higher than measured values in some cases and lower in others.

The emission models used for impoundments also were applied to open tanks. The comparison of measured and predicted values for the overall mass transfer coefficient for open wastewater treatment tanks yielded mixed results. For tanks with quiescent surfaces (e.g., clarifiers and equalization basins), the model predictions were generally lower than measured values but agreed within an order of magnitude. For the aerated systems, the model predictions agreed well with material balance and ambient air measurements for an open aerated system.

8.2.2 Details of Comparisons

The approach to the comparison of predicted and measured values is to estimate the overall mass transfer coefficient from the correlations given in Chapter 4.0 and to compare this value to the overall mass transfer coefficient from the test data. The overall mass transfer coefficient from the test data is calculated from a measured emission rate and a measured or estimated bulk concentration in the liquid phase. Note that errors in either the measured emission rate or liquid-phase concentration have a direct effect on the errors in the calculated mass transfer coefficient.

Most of the measured emission data were obtained by flux chamber measurements. At a few sources, ambient air monitoring and material balances were used to determine the emission rate for calculation of the overall mass transfer coefficient.

GCA Corporation performed an analysis of data from impoundments. The results are given in Tables 8-1 through 8-4 for four ponds at two different sites. Site 5 is a commercial hazardous waste facility with a wastewater treatment system onsite. The reducing lagoon receives wastes classified as reducing agents from tank trucks. The lagoon is operated on a batch basis and was observed to contain a zone of solids and a surface with a floating oil film. The holding pond receives aqueous wastes from the water treatment system and is filled (and discharged) on a monthly basis. The oxidizing lagoon receives oxidizing agents including halogens and other organic compounds. The accumulation of solids and oil film also was observed on this lagoon. Site 4 also is a commercial hazardous waste facility, and its

TABLE 8-1. COMPARISON OF RESULTS FOR REDUCING LAGOON 1
AT SITE 5^{1,2}

Constituent	Mass transfer coefficient ($\times 10^6$ m/s)	
	Average flux chamber measurement ^a	Model predictions (for 5 to 10 m/s windspeed) ^b
Benzene	4.9	4.2-17
Toluene	5.0	3.9-15
Ethylbenzene	5.5	3.6-14
Naphthalene	2.6	3.5-14
Methylene chloride	12	4.7-19
Chloroform	5.7	4.3-17
1,1,1-Trichloroethane	7.6	3.9-15
Carbon tetrachloride	11	3.9-16
p-Dichlorobenzene	2.6	3.6-14
Styrene	5.7	3.7-15

^aCalculated from reported emission rate and corresponding liquid-phase concentration.

^bWindspeed during the test ranged from 5 to 10 m/s.

TABLE 8-2. COMPARISON OF RESULTS FOR HOLDING POND 6
AT SITE 5^{3,4}

Constituent	Mass transfer coefficient ($\times 10^6$ m/s)	
	Average flux chamber measurement ^a	Model predictions (for 5 to 10 m/s windspeed) ^b
Benzene	2.7	5.3-21
Toluene	2.3	4.9-19
Ethylbenzene	2.6	4.6-18
Naphthalene	1.6	4.4-18
Methylene chloride	3.1	6.0-24
Chloroform	2.2	5.4-21
1,1,1-Trichloroethane	3.9	4.9-19
Chlorobenzene	<0.039	4.9-20
p-Dichlorobenzene	4.3	4.6-18
Acetaldehyde	3.4	5.7-19

^aCalculated from reported emission rate and corresponding liquid-phase concentration.

^bWindspeed during the test ranged from 5 to 10 m/s.

TABLE 8-3. COMPARISON OF RESULTS FOR OXIDIZING LAGOON 2
AT SITE 55,6

Constituent	Mass transfer coefficient ($\times 10^6$ m/s)	
	Average flux chamber measurement ^a	Model predictions (for 5 to 10 m/s windspeed) ^b
Toluene	0.38	3.8-15
Ethylbenzene	0.037	3.6-14
1,1,1-Trichloroethane	35	3.9-15

^aCalculated from reported emission rate and corresponding liquid-phase concentration.

^bWindspeed during the test ranged from 5 to 10 m/s.

TABLE 8-4. COMPARISON OF RESULTS FOR SURFACE IMPOUNDMENT AT SITE 47,8

Constituent	Mass transfer coefficient ($\times 10^6$ m/s)		
	Flux chamber measurement ^a		Model predictions (for 5 to 10 m/s windspeed) ^b
	Average	Range	
Toluene	2.4	1.9-2.7	6.3-25.1
Ethylbenzene	1.0	0.46-1.4	5.9-23.5
Methylene chloride	8.4	5.6-10.0	7.7-30.5
1,1,1-Trichloroethane	2.6	1.1-3.6	6.3-24.7
Chloroform	12.0	5.4-15.0	7.0-27.6
p-Dichlorobenzene	0.44	0.079-0.75	5.9-23.1

^aResults for June 22, 1984.

^bWindspeed during the test ranged from 5 to 10 m/s.

impoundment is used to contain aqueous wastes. Table 8-5 presents a comparison of results for Site 3, which is a chemical manufacturing plant that produces primarily nitrated aromatics and aromatic amines. This impoundment is a wastewater holding pond for the wastewater treatment system at the plant. Two wastewater streams that enter the treatment system are distillation bottoms from aniline production (K083) and the nitrobenzene production wastewater (K104).

The results in Tables 8-1 through 8-5 show a reasonable agreement between measured and predicted values of the overall mass transfer coefficient. The measured results for the impoundment in Table 8-3 may have been affected by an oil film observed on the surface or from poor mixing in the impoundment, which can complicate representative sampling of the liquid-phase concentration. Table 8-5 shows good agreement of results for toluene and benzene, which were present in the liquid phase at 2.6 and 17 mg/L, respectively. The liquid-phase concentrations of the other four compounds in Table 8-5 ranged from 0.029 to 0.15 mg/L. The differences in measured and predicted values for these four compounds may have been affected by the accuracy of sampling and analysis of the liquid. The compounds listed in Tables 8-1 through 8-5 are controlled by the liquid-phase mass transfer. Consequently, the results are most dependent on Springer's correlation for k_L (the liquid-phase mass transfer coefficient) and suggest that Springer's model is probably accurate within an order of magnitude.

GCA, in a separate document, examined measured and predicted mass transfer coefficients for open tanks that are part of wastewater treatment systems.¹⁰ The results for Site 8, which is an industrial wastewater treatment operation, included a primary clarifier, an equalization basin, and aerated stabilization basins. The various influent and effluent liquid streams were analyzed, and air emissions around the unit were monitored. Overall mass transfer coefficients were calculated from material balance data and from ambient air monitoring. These values are listed in Tables 8-6 through 8-8 along with the predicted values from the mass transfer correlations given in Chapter 4.0. The primary clarifier, equalization basin, and the quiescent portion of the stabilization basin were modeled as quiescent surfaces, and the correlations of Springer and MacKay/Matasugu

TABLE 8-5. COMPARISON OF RESULTS FOR WASTEWATER HOLDING LAGOON
AT SITE 3⁹

Constituent	Mass transfer coefficient ($\times 10^6$ m/s)	
	Flux chamber measurement	Predicted ^a
Cyclohexane	0.39	3.8
Tetrachloroethylene	0.10	3.7
Toluene	9.0	3.8
Benzene	3.7	4.1
n-Undecane	0.21	2.8
Methylchloride	35.0	3.1

^aBased on an average measured windspeed of 3.7 m/s and an average temperature of 22 °C.

TABLE 8-6. COMPARISON OF RESULTS FOR PRIMARY CLARIFIERS
AT SITE 8¹¹

Constituent	Mass transfer coefficient ($\times 10^6$ m/s)		
	Material balance	Ambient monitors	Model prediction
Tetralin	--	227.0	
2-Ethyl hexanol	96.0	42.0	2.0
2-Ethyl hexyl acrylate	--	123.0	2.7
Naphthalene	179.0	92.0	3.4
1,2-Dichloroethane	58.0	2.9	4.0
Benzene	5.4	18.0	4.1
Toluene	35.0	50.0	3.8
Ethyl benzene	156.0	39.0	3.5

TABLE 8-7. COMPARISON OF RESULTS FOR EQUALIZATION BASIN
AT SITE 812

Constituent	Mass transfer coefficient ($\times 10^6$ m/s)		
	Material balance	Ambient monitors	Model prediction
1,2-Dichloroethane	20	19.0	5.0
Benzene	20	8.9	5.1
Toluene	25	42.0	4.7
Ethyl benzene	25	5.4	4.4

TABLE 8-8. COMPARISON OF RESULTS FOR AERATED STABILIZATION BASINS
AT SITE 813

Constituent	Mass transfer coefficient ($\times 10^4$ m/s)		
	Material balance	Ambient monitors	Model prediction
2-Ethyl hexanol	0.05	0.01	0.17
2-Ethyl hexyl acrylate	4.8	8.3	2.9
1,2-Dichloroethane	2.0	0.52	5.7
Benzene	12.4	1.1	10.6
Toluene	5.0	5.8	10.1
Ethyl benzene	2.9	0.55	9.9

were used. The turbulent portion of the stabilization basins was modeled using the correlations of Thibodeaux and Reinhardt.

Useful conclusions from the comparison of measured and predicted values are difficult because of the lack of consistent results from air monitoring, probably due to very short sampling periods, changes in the windspeed and direction, and the contribution to emissions from sources near the mentioned source. In addition, material balance calculations are subject to error from changes in influent/effluent flow rates and concentrations of specific compounds. In general, the model predictions for the primary clarifier and equalization basin are lower than the measured values. For the aerated stabilization basin, most of the predicted mass transfer coefficients are higher than the measured values; however, the agreement is within an order of magnitude. The measured values for the primary clarifier may have been higher than the predicted values because of the contribution from nearby sources to measured air concentrations or because of the contribution from the falling film at the clarifier.

GCA also performed an analysis on an aerated lagoon at Site 7.¹⁴ This lagoon was covered and was purged with air at a rate of $1.4 \text{ m}^3/\text{s}$ ($3,000 \text{ ft}^3/\text{min}$). Predicted and calculated values for the mass transfer coefficients are given in Table 8-9 and show that predicted values are 1 to 2 orders of magnitude higher. The basis of the predicted values includes Thibodeaux and Reinhardt's correlations for aerated systems. No strong conclusions on the model's validity can be drawn from these data because the dome enclosure over the system and its evacuation rate probably have a direct effect on emissions. In addition, difficulties with air measurements and determination of the appropriate driving-force concentration (assumed to be the bulk liquid concentration) can lead to errors in the calculated values of the overall mass transfer coefficient.

The results of the biodegradation model were also compared to available data from biodegradation measurements. The most desirable comparison would be for a system in which the air emission rate and biodegradation rate were measured directly. However, the extent of biodegradation from studies of real systems has usually been determined by difference from a material balance (what enters the system minus what leaves with the effluent and with air emissions).

TABLE 8-9. COMPARISON OF RESULTS FOR COVERED AERATED LAGOON
AT SITE 715,16

Constituent	Mass transfer coefficient ($\times 10^4$ m/s)	
	Vent rate measurement	Predicted ^a
1,2-Dichloroethane	0.05	7.2
Benzene	0.30	8.9
Toluene	0.95	8.8

^aBased on an estimated windspeed (not measured) of 5 m/s¹⁷ and an estimated turbulent area of about 50 percent.¹⁸

Petrasek et al. performed such a study on a large pilot-scale activated sludge system with diffused air aeration.¹⁹ The activated sludge unit was enclosed, and the diffused air that was removed was sampled (for flow rate and concentration) to determine air emissions. This system was designed for a flow rate of 2.2 L/s (35 gal/min) with an air purge rate of 57 L/s. A summary of the operating parameters is given in Table 8-10. The study used a synthetic wastewater that contained individual volatile compounds at levels of 32 to 300 ppb. The biomass concentration was 2 g/L, and the resultant food-to-microorganism (F/M) ratio of 0.5 is well within the recommended design range of 0.2 to 0.6.

Petrasek reported the percent of each compound entering the activated sludge unit that was emitted with the diffused air; the fraction biodegraded could be determined by difference assuming all unrecovered material was biodegraded. The results are summarized in Table 8-11 and show a range of measured values from 5 percent for chlorobenzene to 62 percent for 1,1,1-trichloroethane. The predictions of the biodegradation model discussed in Chapter 4.0 are also presented in Table 8-11 for comparison. The comparison shows that the model predictions agree well with the Petrasek measurements for nearly every compound.

Another type of comparison between measurements and predictions involves effluent concentrations for well-defined wastewater treatment systems. Namkung and Rittman²² reported influent and effluent concentrations of volatile organics for two Chicago wastewater treatment plants that receive large shares of industrial discharges. The measurements were made for two large activated sludge units aerated by diffused air. In addition, the system's operational parameters were defined (Table 8-12) and provided the necessary inputs for the mathematical model that includes air emissions (diffused air system) and biodegradation.

The results of measured and predicted effluent concentrations are summarized in Table 8-13. The most convincing comparison is the close match for both plants for tetrachloroethylene, which the authors stated was not biodegradable in these systems. Therefore, a biodegradation rate equal to zero was used in the model for this compound. The close agreement between measured and predicted effluent concentrations suggests that this compound is almost

TABLE 8-10. DESCRIPTION OF PETRASEK'S ACTIVATED
SLUDGE SYSTEM²⁰

Parameter	Value
Flow rate (L/s)	2.2
Volume (m ³)	59.8
Residence time (h)	7.5
Air rate (L/s)	57
Fraction of surface agitated	0
Biomass concentration (g/L)	2.0
Concentration range for organics (ppm)	0.032 - 0.30
F/M ^a	0.5

^aF/M^a = Food to microorganism ratio (lb/lb biomass • day)
^a based on chemical oxygen demand.

TABLE 8-11. COMPARISON OF PETRASEK'S MEASUREMENTS AND MODEL PREDICTIONS

Compound	Reported fraction emitted ^a	Predicted fraction emitted ^b	Fraction assumed biodegraded ^a	Predicted fraction biodegraded ^b
Benzene	0.15	0.19	0.80	0.78
Carbon tetrachloride	0.59	0.54	0.41	0.44
Chlorobenzene	0.05	0.02	0.95	0.97
Chloroform	0.34	0.20	0.66	0.75
Dichloropropane (1,2)	0.32	0.09	0.68	0.88
Ethyl benzene	0.21	0.15	0.79	0.82
Tertachloroethane and -ethene	0.27	0.37 ^c	0.73	0.58 ^c
Toluene	0.20	0.15	0.80	0.84
Trichloroethane (1,1,1)	0.62	0.57	0.38	0.40
Trichloroethane (1,1,2)	0.25	0.06 ^d	0.75	0.87 ^d
Trichloroethene	0.41	0.37	0.59	0.59

^aData from Petrasek et al.;²¹ the fraction biodegraded is assumed to be the fraction unaccounted for based on the analyses of the sludge, the air, and the effluent.

^bModel predictions based on the equations presented in Chapter 4.0 assuming influent VO concentrations of 0.10 mg/L and operating parameters as provided in Table 8-10.

^cArithmetic average for the removal fractions calculated for 1,1,2,2-tetrachloroethane and tetrachloroethene.

^dEmployed 1,1,1-trichloroethane's biodegradation rate constants.

TABLE 8-12. DESCRIPTION OF TWO CHICAGO ACTIVATED SLUDGE UNITS²³

	Calumet	West-southwest
<u>Operating parameters</u>		
Volume (m ³)	184,500	802,300
Depth (m)	1.8	1.8
Wastewater flow (m ³ /s)	10.0	36.6
Air rate (m ³ /s)	55	193
Residence time (h)	5.1	6.1
Total organics (mg/L)	115	180
Biomass (g/L)	2.2	2.0
Fraction surface agitated	0	0
<u>Concentrations (ppb)</u>		
Chloroform, in	4.0	4.4
out	7.1	2.4
Ethylbenzene, in	18	10
out	0.5	BDL
Methylene chloride, in	9.8	48
out	11	11
Tetrachloroethylene, in	16	12
out	2.1	1.6
Toluene, in	85	22
out	6.2	BDL
1,1,1-Trichloroethane, in	13	15
out	2.9	2.2
Trichloroethylene, in	9.7	22
out	0.5	2.1

BDL = Below detection limit.

TABLE 8-13. COMPARISON OF MEASURED AND PREDICTED EFFLUENT CONCENTRATIONS FOR CHICAGO WASTEWATER TREATMENT PLANTS²⁴

Compound	Calumet effluent concentrations, ppb		West-southwest effluent concentrations, ppb	
	Measured	Predicted ^a	Measured	Predicted ^a
Chloroform	b	b	2.4	2.3
Ethyl benzene	0.5	0.68	c	0.16
Methylene chloride	b	b	11	7.1
Tetrachloroethylene	2.1	1.0	1.6	0.77
Toluene	6.2	2.9	c	0.69
1,1,1-Trichloroethane	2.9	1.0	2.2	1.1
Trichloroethylene	0.5	0.75	2.1	1.6

^aBased on the equations presented in Chapter 4.0.

^bNo comparison possible because measured concentration in effluent was greater than measured concentration in influent.

^cMeasured effluent concentration was below detection limit.

entirely removed by air stripping, and the quantity predicted to be air stripped by the model is reasonably accurate.

The results in Table 8-13 also indicate that 1,1,1-trichloroethane and trichloroethylene are biodegraded. The model predictions used a biorate for these two compounds that was derived from Petrasek's data in Table 8-11. Both Petrasek's data and the comparison in Table 8-13 indicate that these compounds are biodegraded to some extent; otherwise, the measured effluent concentrations in Table 8-13 would have been higher than those predicted by the model with biodegradation included.

Tabak et al.²⁵ conducted an extensive study of the biodegradability of numerous toxic compounds. They found that, when the microbial culture is properly acclimated, almost all nonpesticide compounds could be, at least partially, biodegraded. Although biodegradation rate constants were not determined, the percent of compound biodegraded was shown to be dependent on the acclimation of the culture, and (although to a lesser extent) dependent on the concentration of the compound used. For every compound tested, the percent biodegraded by the third subculture (presumably the most acclimated) always decreased when the concentration was doubled (unless both cultures were either 100 percent or 0 percent biodegraded), and this decrease was rarely a decrease of a factor of two or more.²⁶ If biodegradation were strictly a first-order process, the percent biodegraded would be independent of the concentration. If biodegradation were strictly a zero-order process, the percent biodegraded would decrease by a factor of two (for those compounds not biodegraded, 100 percent) when the concentration was doubled. Because an intermediate effect was generally realized, Tabak's results suggest Monod-type biodegradation rate kinetics are appropriate.

Another comparison that can be made is based on a series of field studies, as reported by Coburn et al.,²⁷ in which batch, biodegradation rate studies were performed while controlling air emissions. The experimental first-order biodegradation rate constant and the predicted, apparent first order rate constant based on the Monod model can be compared in the last two columns of Table 8-14. Note for compounds whose log mean concentrations are near or are greater than the appropriate half-saturation

constant (e.g., formaldehyde or methanol), the predicted first-order rate constant according to the Monod model provides a better estimate of the observed biodegradation rate than would be provided assuming simple first-order kinetics (i.e., using K_1 straight from the data base as provided, for comparison, in Table 8-14). Additionally, using the recommended biodegradation rate constants and modeling approach, the predicted biodegradation rates presented in Table 8-14 agree well with the reported biodegradation rates for nearly every experimental run.

A separate study was conducted for EPA to evaluate measured and predicted emissions for aerated waste treatment systems.²⁹ The correlations of Thibodeaux and Reinhardt were used (as recommended in Chapter 4.0) to estimate the mass transfer coefficients of the turbulent zone. The results showed an agreement between measured and predicted values that were within an order of magnitude. The report concluded that, when adequate descriptions of plant operating parameters are available, reliable emission estimates can be obtained from the models (within the accuracy that results from variations in sampling and chemical analysis). When plant operating parameters are known, the major limitations in the models result from a lack of accurate biooxidation rates and vapor/liquid equilibrium data for specific compounds.³⁰

8.2.3 Recommendations for Additional Data

The estimate of emissions from open liquid surfaces is provided by the product of the mass transfer coefficient, driving-force concentration, and surface area. Surface area can be determined with reasonable accuracy. The previous comparison of mass transfer coefficients indicated that they can be estimated within an order of magnitude. Probably the greatest source of uncertainty is in the estimate of the appropriate driving force for mass transfer. The concentration is likely to vary with time and location in the impoundment. The type of flow system and extent of mixing in the liquid also will affect this concentration.

For the less volatile compounds that may be controlled by gas-phase mass transfer, the collection of equilibrium data may be useful to compare with the estimated values used in the models. The comparisons presented in this section primarily address compounds with high volatility in water

TABLE 8-14. COMPARISON OF MEASURED AND PREDICTED BIODEGRADATION RATES

Compound	Log mean conc., ^b mg/L	Rate constants ^a			First-order rate constant, L/g/h	
		K _{max} , mg/g/h	K _s , mg/L	K ₁ , L/g/h	Experiment ^b	Predicted ^c
Acetone	1.35	1.3	1.1	1.15	1.15	0.53
	2.56				0.34	0.36
Benzene	0.005	19	13.6	1.4	0.36	1.4
	0.10				2.1	1.4
Chloroform	0.008	2.94	3.7	0.79	0.36	0.79
	0.002				0.29	0.79
Dimethylphthalate	4.2	2.2	0.71	3.1	0.36	0.45
Ethanol	4.9	8.8	9.8	0.90	0.70	0.60
Ethylbenzene	0.005	6.8	3.2	2.1	0.36	2.1
Ethylene oxide (oxirane)	1.7	4.2	4.6	0.91	0.81	0.67
	3.2				0.75	0.54
	3.9				0.37	0.49
Formaldehyde	8.0	5.0	20	0.25	0.13	0.17
	62				0.077	0.057
Methanol	250	18	90	0.20	0.067	0.053
	480				0.018	0.032
	490				0.040	0.031
	495				0.023	0.031
Methyl ethyl ketone	0.10	2.0	10	0.20	0.24	0.20
	0.27				0.18	0.19
	0.37				0.19	0.19
	0.80				0.16	0.18
Methylene chloride	0.028	22	55	0.40	0.11	0.40
	0.031				0.11	0.40
	0.053				0.36	0.40
	0.15				0.20	0.40
	0.23				0.57	0.40

(continued)

TABLE 8-14 (continued)

Compound	Log mean conc., ^b mg/L	Rate constants ^a			First-order rate constant, L/g/h	
		K _{max} , mg/g/h	K _s , mg/L	K ₁ , L/g/h	Experiment ^b	Predicted ^c
2-Propanol	2.9	15	200	0.75	0.069	0.074
	6.2				0.085	0.073
Thiobismethane	1.07	0.16	0.17	0.93	0.13	0.11
Toluene	0.014	73.5	30.6	2.4	0.28	2.4
	0.016				0.34	2.4
	0.081				0.63	2.4
	0.14				1.9	2.4
1,1,1-Trichloroethane	0.040	3.5	4.73	0.74	0.38	0.73
	0.16				0.88	0.72
Trichloroethene	0.004	3.9	4.43	0.88	0.41	0.88
Total xylenes	0.097	40.8	22.7	1.8	>2.2 ^d	1.8

^aRecommended rate constants from Appendix D, Table D-1.

^bFrom data reported by Coburn et al.;²⁸ the log mean concentration was calculated as follows:

$$C_{LM} = (C_i - C_f) / \ln(C_i/C_f)$$

where

C_i = initial concentration, mg/L
C_f = final concentration, mg/L.

^cCalculated as the apparent first-order rate constant using the Monod model (Equation 4-13) based upon the log mean concentration as follows:

$$K_1 = K_{max} / (K_s + C_{LM})$$

^dFinal concentration was below detection limit. The final concentration was assumed to be at the detection limit to calculate the first-order rate constant. The actual rate constant should be greater than the reported value.

(high Henry's law constant). Because semivolatile compounds also can be emitted to a significant extent, air emission measurements for these less volatile compounds would be useful for comparison with model predictions.

8.3 LAND TREATMENT

Field data from four test sites and one laboratory simulation were used as a basis for comparing measured emissions with estimated emissions using the RTI land treatment model. Two other laboratory simulations of land treatment are under way or near completion, but the documentation on those tests is not yet sufficiently complete so that comparisons of measured and estimated emissions can be made. These include an EPA-sponsored study being conducted by Radian Corporation and a simulation study by Chevron Research Company in Richmond, California. Table 8-15 summarizes the tests evaluated. Generally, estimated emissions are within an order of magnitude of measured values. Values of estimated emissions varied both above and below measured values.

Comparisons of estimated and measured emission flux rates are presented graphically in this section. Comparisons of estimated and measured emissions by weight percent of applied material are presented in the next section.

Considering the potential for error in measuring or estimating values for parameters that are input to the model and the potential for error in measuring emissions, differences in the range of an order of magnitude are not unexpected. In making the comparisons, values for all model inputs sometimes were not available in the emission test reports. In these cases, values were estimated using averages of field data or values identified previously as typical or representative of actual land treatment practices.

In the 1985 test at a Midwest petroleum refinery (Case 1),³¹ emission measurements were made at sample locations in six test plots. For each plot, emission measurements were made after waste application but before the plot was tilled, again after the waste was tilled, and for another period after a second tilling. All measurements were made using a flux chamber and tenax traps. Emission rates were measured for six specific organic constituents: benzene, toluene, p-xylene, o-xylene, m-xylene, and naphthalene. Benzene and toluene were selected as a basis for comparing

TABLE 8-15. SUMMARY OF LAND TREATMENT TESTING AND TEST RESULTS

Site No.	Test site location	Test description	Test year	Test sponsor	Test procedures	Test duration	Test results	
							Waste constituent	Emissions, wt. %
12	West Coast corporate research facility	Laboratory simulation	1986 - 1987	Private corporation	Run 1 (raw waste)	2.5 months	Oil	35
					Run 2 (raw waste)	22 days	Oil	11
					Run 2 (treated waste)	22 days	Oil	1
13	Southwest research facility	Laboratory simulation	1986	EPA	Run 1 ^a (API separator sludge)	31 days		
					Box #1		Oil	5.2
					Box #2		Oil	NA
					Box #3		Oil	6.5
					Box #4		Oil	6.7
					Run 2 ^a (IAF float)	31 days		
					Box #1		Oil	15
					Box #2		Oil	NA
					Box #3		Oil	18
					Box #4		Oil	19
14	Midwestern refinery	Flux chamber	1986	ORD	Plot A	8 days	Benzene	81
							Toluene	41
							Ethylbenzene	195
							p-Xylene	18
							m-Xylene	39
							o-Xylene	28
							Naphthalene	1
					Plot B	8 days	Benzene	110
							Toluene	66
							Ethylbenzene	402
							p-Xylene	21
							m-Xylene	83
							o-Xylene	38
							Naphthalene	2

See notes at end of table.

(continued)

TABLE 8-15 (continued)

Site No.	Test site location	Test description	Test year	Test sponsor	Test procedures	Test duration	Test results	
							Waste constituent	Emissions, wt. %
14 (con.)								
					Plot C	8 days	Benzene	39
							Toluene	17
							Ethylbenzene	140
							p-Xylene	25
							m-Xylene	25
							o-Xylene	17
							Naphthalene	
					Plot D	8 days	Benzene	142
							Toluene	86
							Ethylbenzene	353
							p-Xylene	55
							m-Xylene	79
							o-Xylene	52
							Naphthalene	2
					Plot E	8 days	Benzene	187
							Toluene	63
							Ethylbenzene	345
							p-Xylene	43
							m-Xylene	52
							o-Xylene	39
							Naphthalene	1
					Plot F	8 days	Benzene	84
							Toluene	47
							Ethylbenzene	288
							p-Xylene	13
							m-Xylene	28
							o-Xylene	24
							Naphthalene	1
15	West Coast refinery	Flux chamber	1984	ORD	Surface application	5 weeks	n-Heptane	68
							Methylcyclohexane	61
							3-Methyl-heptane	52
							n-Nonane	56
							1-Methylcyclohexene	49
							1-Octene	58
							β -Pinene	17
							Limonene	22

See notes at end of table.

(continued)

TABLE 8-15 (continued)

Site No.	Test site location	Test description	Test year	Test sponsor	Test procedures	Test duration	Test results			
							Waste constituent	Emissions, wt. %		
15 (con.)							Toluene	37		
							p-,m-Xylene	35		
							1,3,5-Trimethylbenzene	21		
							o-Ethyl-toluene	32		
							Total VO	38		
							Total oil	1.2		
							Subsurface	5 weeks	n-Heptane	94
									Methylcyclohexane	88
									3-Methyl-heptane	77
									n-Nonane	88
						1-Methylcyclohexane			78	
						1-Octene			74	
						β -Pinene			21	
						Limonene			26	
						Toluene			58	
						p-,m-Xylene			48	
						1,3,5-Trimethylbenzene	27			
						o-Ethyl-toluene	42			
						Total VO	38			
						Total oil	1.4			
16	Southwest research facility	Laboratory simulation	1983	API/EPA	Run no. 18	8 hours ^b	Oil	9.1		
							Run no. 21	Oil	4.4	
							Run no. 24	Oil	8.82	
							Run no. 27	Oil	8.6	
							Run no. 28	Oil	9.1	
							Run no. 32	Oil	3.6	
							Run no. 33	Oil	2.6	
							Run no. 34	Oil	8.01	
							Run no. 35	Oil	8.9	
							Run no. 38	Oil	78.8	
							Run no. 37	Oil	9.9	
							Run no. 40	Oil	8.7	
							Run no. 41	Oil	2.8	
							Run no. 44	Oil	4.9	
							Run no. 45	Oil	49.9	
							Run no. 46	Oil	7.7	
							Run no. 47	Oil	6.9	
							Run no. 48	Oil	5.8	
							Run no. 49	Oil	9.7	
							Run no. 50	Oil	1.1	
Run no. 51	Oil	8.47								

See notes at end of table.

(continued)

TABLE 8-15 (continued)

Site No.	Test site location	Test description	Test year	Test sponsor	Test procedures	Test duration	Test results	
							Waste constituent	Emissions, wt. %
16	Gulf Coast commercial TSDf	Flux chamber	1983	ORD	Single test ^c	69 hours	Total VO	6.77
						56 hours	Benzene	3.91
17	Midwestern refinery	Flux chamber	1979	API	Centrifuge sludge			
					Test no. 5	19.9 hours	Oil	6.1
					Test no. 6	367 hours	Oil	2.6
					API separator sludge ^d			
					Test no. 7	619 hours	Oil	13.6
					Test no. 8	122 hours	Oil	1.1
					Test no. 9	528 hours	Oil	13.6

API = American Petroleum Institute.

IAF = Induced air flotation.

ORD = Office of Research and Development.

^aSludge applied to Box #1 and Box #3 as duplicate tests; sludge treated with mercuric chloride to eliminate (or reduce) bioactivity applied to Box #4 and no sludge applied to Box #2, which served as a control.

^bEach run for which results are reported was 8 hours.

^cTest was conducted using aged wastes.

^dAllowed to weather for 14 days in open 5-gal buckets in an outdoor open shelter prior to application.

measured and estimated emissions in this test. The comparison was made for test plot A after the waste was tilled for the first time. Estimated emissions for each compound are higher than the measured values but generally are within a factor of 10. Estimated and measured values are shown graphically for benzene and toluene in Figures 8-1 and 8-2, respectively.

At the West Coast refinery (Case 2),³² emission tests were made using three adjacent plots marked off in the land treatment site. The center plot was used as a control and had no waste applied while waste was applied to the other two plots. One plot had waste applied to the soil surface and the other had waste applied by subsurface injection. Flux chambers were situated on each test plot and emission measurements were made during three different test periods each lasting 4 days. Canister air samples, sludge samples, and liquid samples were analyzed by gas chromatography (GC). Emissions of both total V₀ and selected specific constituents were measured during the test. For comparing measured and estimated emissions, total V₀ and toluene emissions from the surface application plot were used. Estimated emission rates for both toluene and total V₀ agree reasonably well with measured rates but range from higher to lower than measured rates at different times. Estimated cumulative emissions over the entire test period agree reasonably well with the measured values. For both toluene and total V₀, estimates covered a 4-day period with a till occurring after 2 days. Estimated and measured values over the 4-day period for which the comparison is made are generally within an order of magnitude, as can be seen in Figures 8-3 and 8-4. Measured values were reported as half-day average emission rates.

For the test at the commercial hazardous waste site in 1983, (Case 3),³³ waste was applied to a single test plot and tilled into the soil. Air emission measurements were made over a 3-day period using a flux chamber and gas canisters. Sampling locations were selected randomly, with a control point used to provide a common sampling position each day. Sample analyses were made by GC. Emission comparisons of measured and estimated emissions were made for total nonmethane hydrocarbon (NMHC) emissions using data generated by GCA in a separate study of the data from this test.³⁴ As with previous tests, estimated emission flux rates were greater

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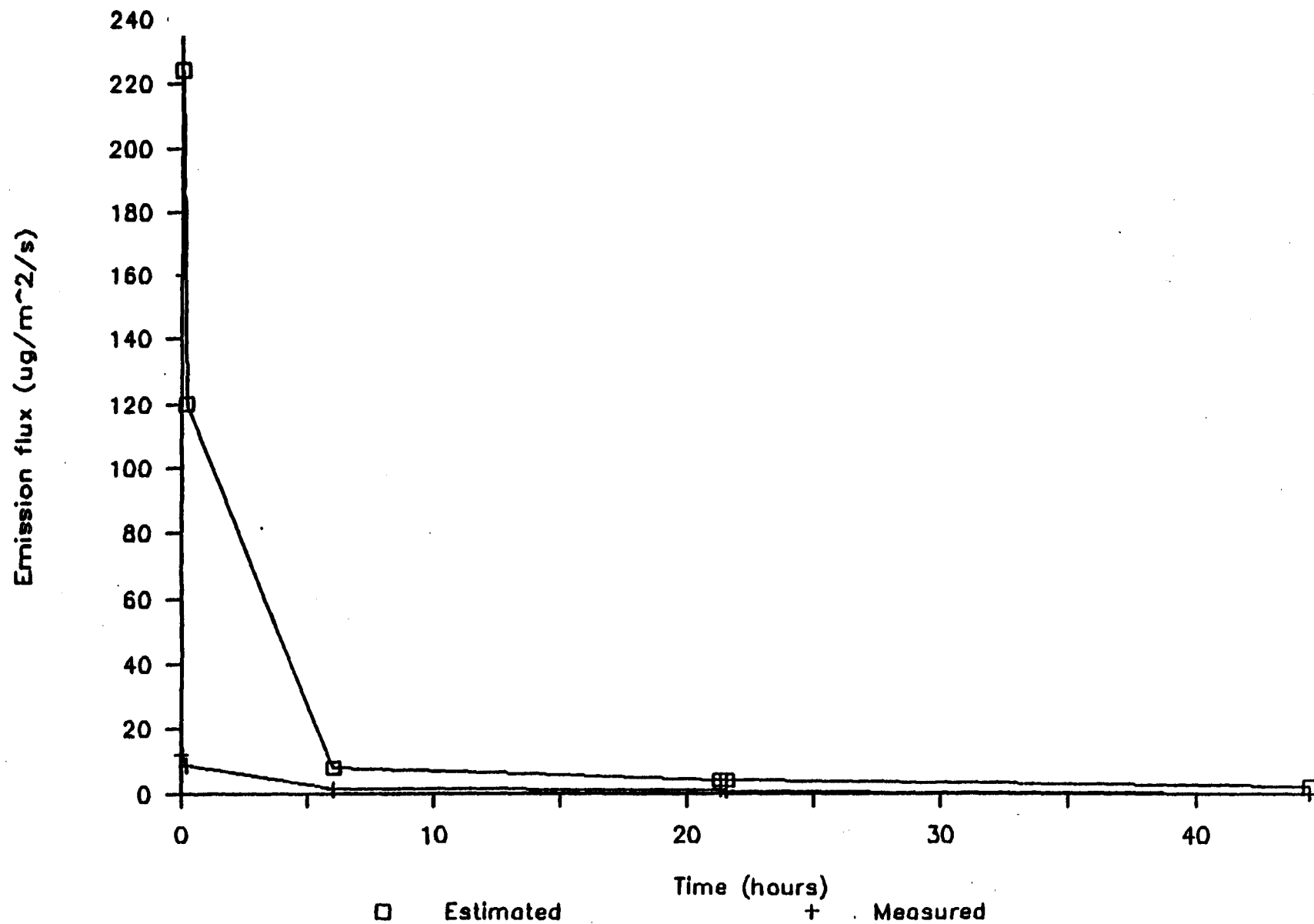


Figure 8-1. Estimated vs. measured benzene emission flux rates—Case 1.

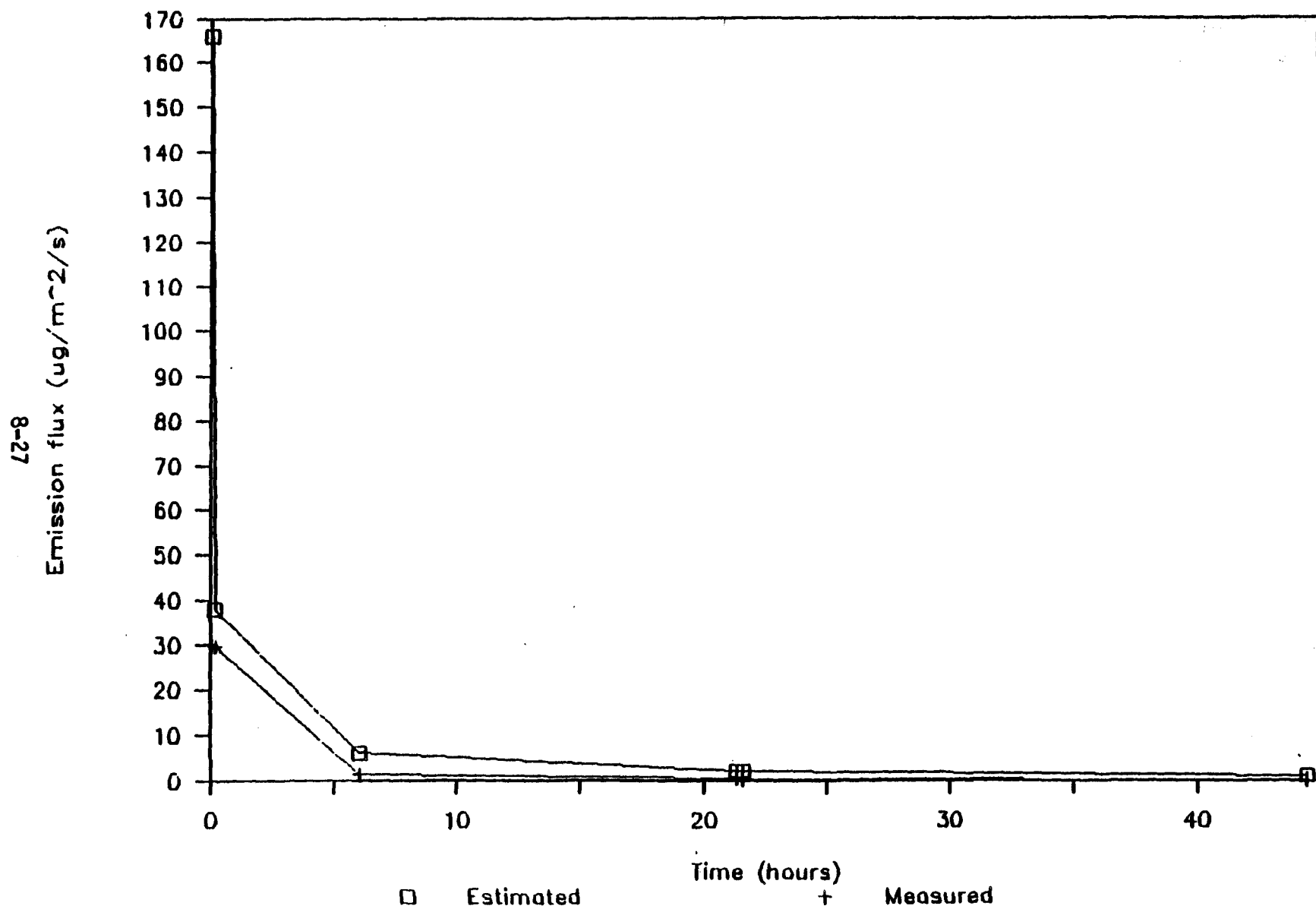


Figure 8-2. Estimated vs. measured toluene emission flux rates—Case 1.

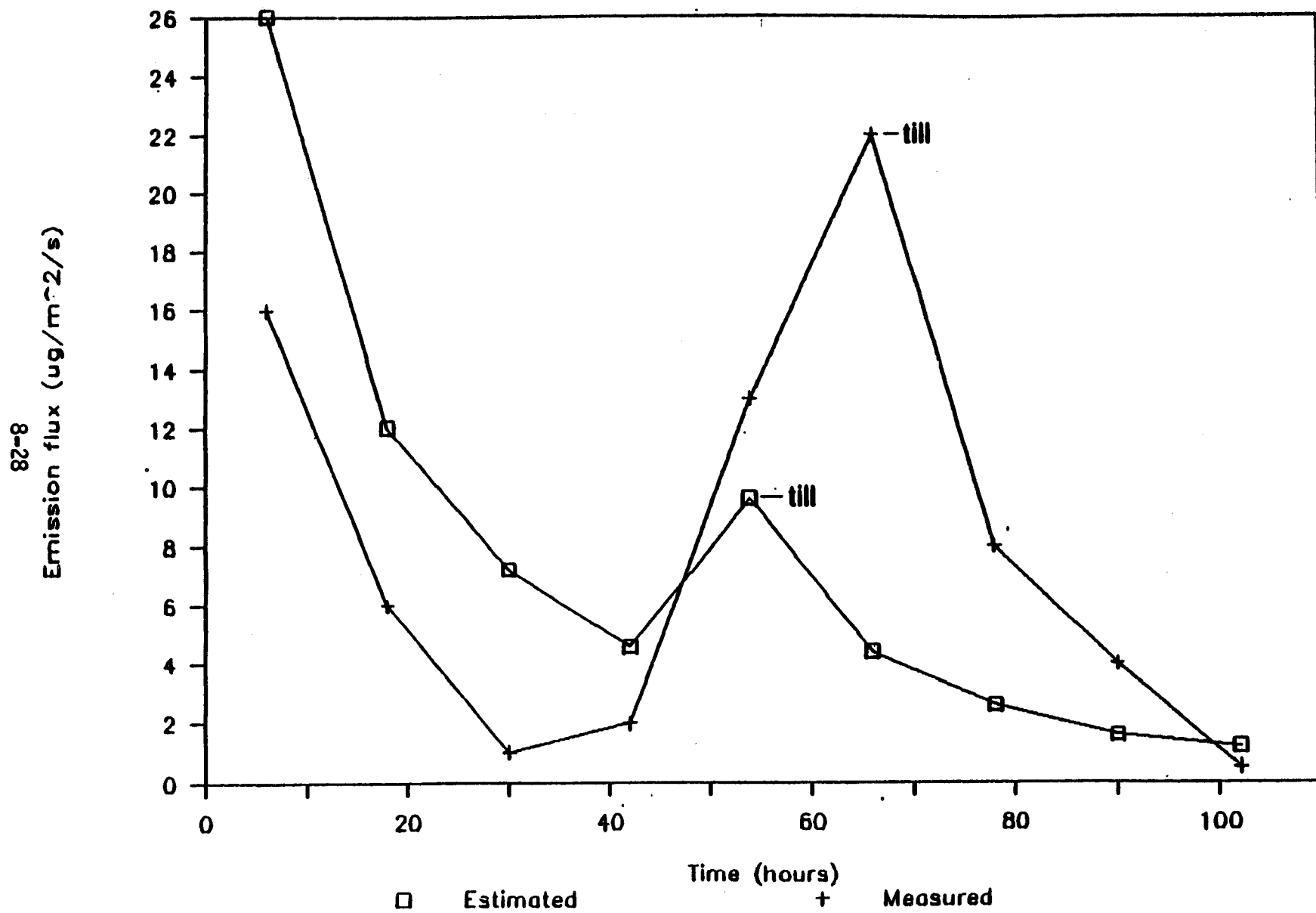


Figure 8-3. Estimated vs. measured toluene emission flux rates—Case 2 (data for 4 days only).

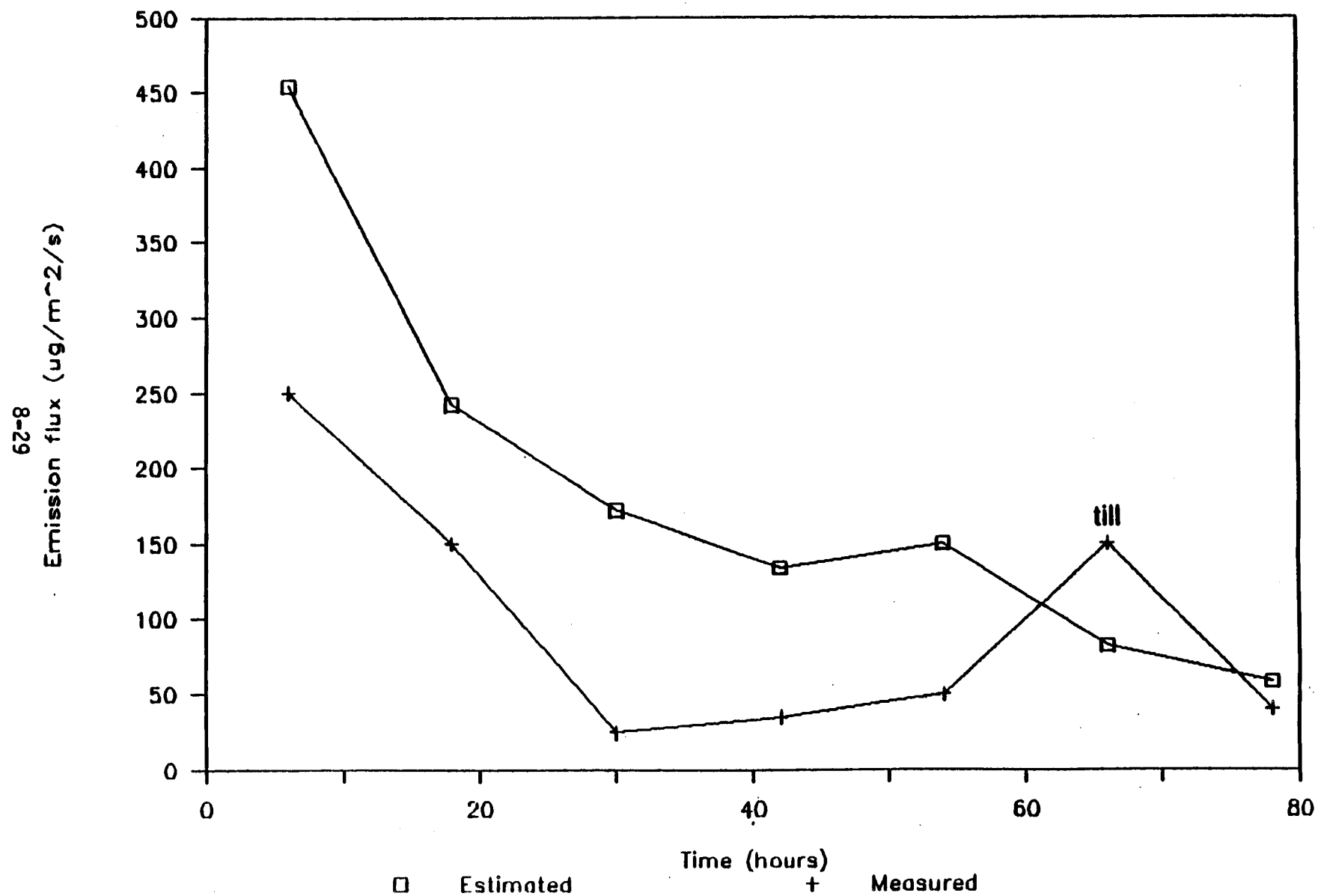


Figure 8-4. Estimated vs. measured total VO emission flux rates—Case 2.

than measured values but mostly were within a factor of 10 or less of the measurements. Estimated cumulative emissions also were substantially higher than measured values. Estimated and measured values of instantaneous emission flux rates are shown in Figure 8-5.

In the 1979 test at the Midwest petroleum refinery (Case 4),³⁵ three test plots were laid out. One plot was used as a control and had no waste applied, one plot had an API separator sludge applied, and the other plot had a centrifuge sludge applied. A 1-ft² collector box was placed on the test plot and continuously purged with fresh air. The outlet from the box was analyzed for total V₀ (as methane and NMHC) using a continuous hydrocarbon analyzer. For one test run, total V₀ emissions were estimated with the land treatment model for comparison with the measured values. Measured and estimated values are shown graphically in Figure 8-6. As can be seen, the estimated and measured values agree quite well for this test. Total cumulative emissions for each test were also estimated using the model and compared with measured values. The estimated values were generally higher than measured values for these emissions.

8.3.1 Midwest Refinery--1985 (Case 1)

Table 8-16 presents the model input values used to compare estimated and measured emissions for plot A of the Case 1 test data. The information in Table 8-16 represents data for plot A as reported in the test report. Similar information was reported for plots B through F and those data were used as appropriate for input to the model. Table 8-17 shows measured emissions of six constituents made during the test. In this test, the waste was allowed to stay on top of the soil for 24 hours before it was tilled into the soil. Measured emissions during the first 24 hours were combined with measured emissions after tilling to get total emissions. Table 8-17 shows variations in measured emissions among the different test plots and shows emissions greater than applied material for some plots and some waste constituents. In Table 8-17, weight fraction represents the fraction of applied material that is emitted to the air. For ethylbenzene, all plots have measured emissions in excess of the amount applied. To compare measured and estimated emissions, the RTI land treatment model can be used for estimating emissions both before and after tilling. Estimated

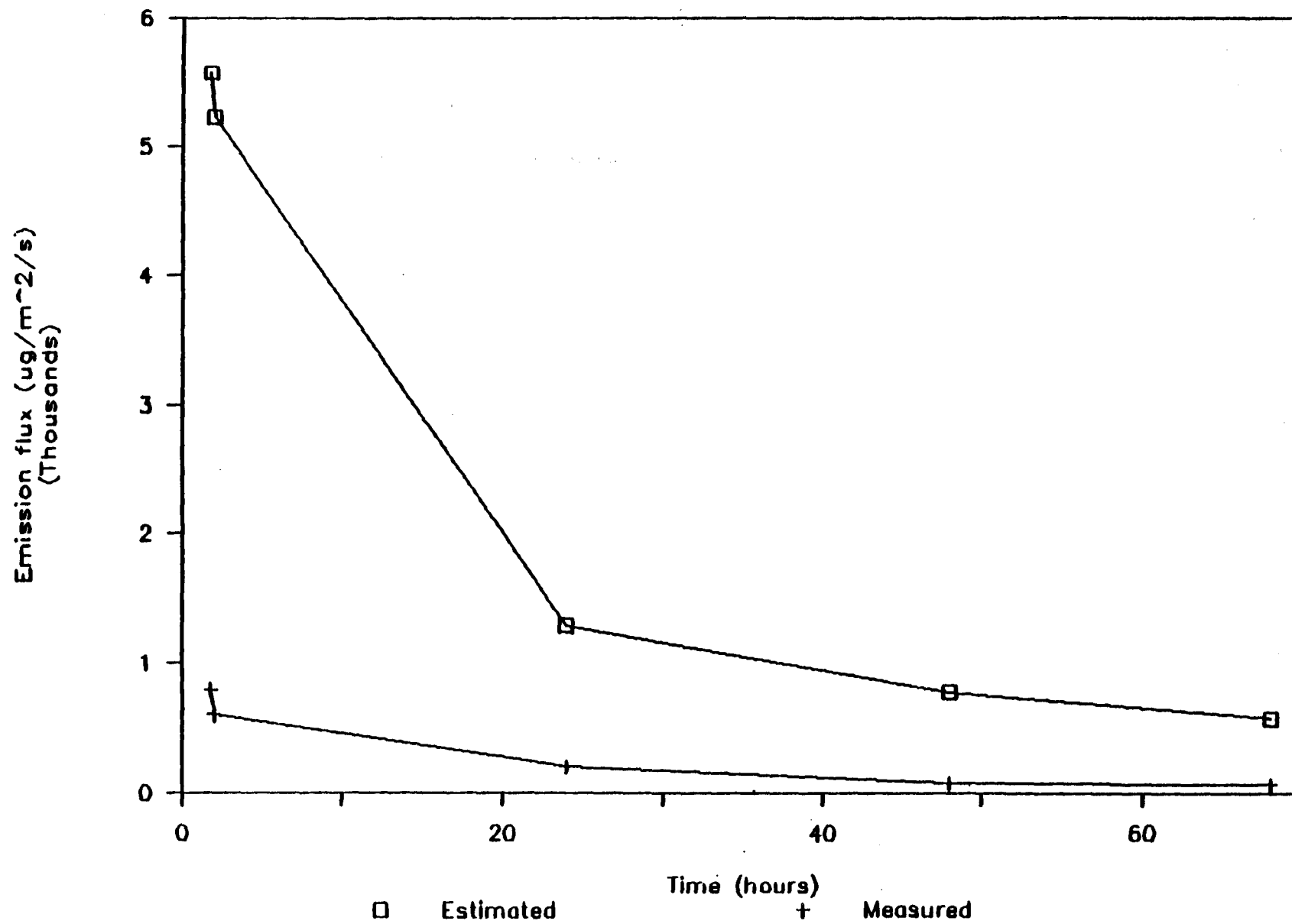


Figure 8-5. Estimated vs. measured VO emission flux rates—Case 3.

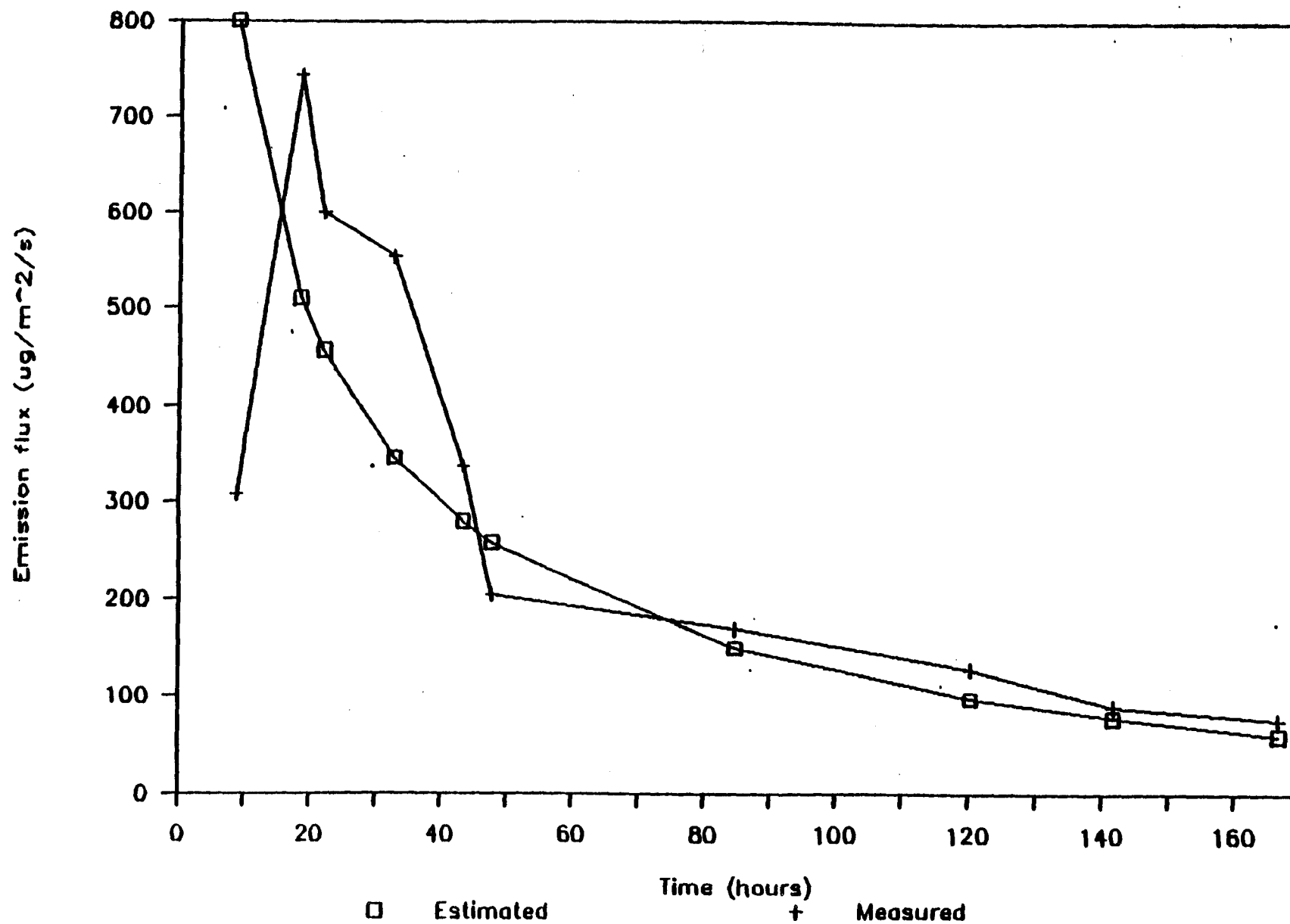


Figure 8-6. Estimated vs. measured emission flux rates—Case 4.

TABLE 8-16. INPUT PARAMETERS FOR RTI LAND TREATMENT MODEL^a

Parameter	Value	Source
Organic loading	0.0236 g/cm ³	Calculated from field data
Tilling depth	20 cm	Field data
Soil air porosity	0.40	Field data
Soil total porosity	0.61	Field data
Benzene concentration ^b	0.000249	Calculated from field data
Toluene concentration ^b	0.000632	Calculated from field data
Benzene diffusivity	8.80 E-02 cm ² /s	Data base
Toluene diffusivity	8.70 E-02 cm ² /s	Data base
Benzene vapor pressure	95.2 mm Hg	Data base
Toluene vapor pressure	30.0 mm Hg	Data base
Benzene biorate	19.0 mg V0/g•h	Data base
Toluene biorate	73.0 mg V0/g•h	Data base
Molecular weight of oil	282 g/g mol	Assumed

^aSource of field data: Reference 36. Data represent conditions in plot A.

^bWeight fraction of oil.

TABLE 8-17. MEASURED AND ESTIMATED EMISSIONS--CASE 1

Measured cumulative emissions ^a														
Test location	Benzene		Toluene		Ethylbenzene		p-Xylene		m-Xylene		o-Xylene		Naphthalene	
	$\mu\text{g}/\text{cm}^2$	wt. frac.	$\mu\text{g}/\text{cm}^2$	wt. frac.	$\mu\text{g}/\text{cm}^2$	wt. frac.	$\mu\text{g}/\text{cm}^2$	wt. frac.	$\mu\text{g}/\text{cm}^2$	wt. frac.	$\mu\text{g}/\text{cm}^2$	wt. frac.	$\mu\text{g}/\text{cm}^2$	wt. frac.
A	271.01	0.03	340.71	0.41	57.97	1.06	7.39	0.10	96.40	0.30	21.11	0.20	2.15	0.01
B	209.00	1.10	454.20	0.00	96.40	4.02	7.50	0.21	103.04	0.03	23.10	0.30	2.31	0.02
C	100.35	0.30	209.00	0.17	59.27	1.40	16.03	0.26	67.17	0.26	10.76	0.17	3.00	0.01
D	459.42	1.42	703.00	0.00	101.05	3.53	23.92	0.55	105.32	0.70	30.02	0.52	3.35	0.02
E	302.23	1.07	576.10	0.03	109.31	3.45	20.74	0.43	130.30	0.52	31.50	0.30	2.40	0.01
F	324.00	0.04	404.07	0.47	71.55	2.00	6.07	0.13	76.04	0.20	21.30	0.24	2.44	0.01
Estimated cumulative emissions														
Test location	Benzene		Toluene											
	$\mu\text{g}/\text{cm}^2$	wt. frac.	$\mu\text{g}/\text{cm}^2$	wt. frac.										
All	--	0.03	--	0.53										

^aTest duration was approximately 8 days.

cumulative emissions for benzene and toluene for all plots are shown in Table 8-17 and show reasonable agreement with measured values.

8.3.2 West Coast Refinery (Case 2)

The data in Table 8-18 were used to estimate emissions of toluene and total V0 from the surface application plot at the Case 2 land treatment facility. Estimated and measured cumulative emissions are compared in Table 8-19. The comparisons were made for total V0 (as determined by purge and trap) and for toluene.

8.3.3 Commercial Waste Disposal Test (Case 3)

Table 8-20 shows the inputs used to estimate emissions from the Case 3 land treatment operation. No specific constituent data were available so emissions were estimated using average characteristics of the total organic phase. Results are shown in Table 8-21. The comparison is made for the estimated versus measured cumulative weight percent of applied oil that is emitted after 24 hours and after 68 hours, which is the duration of the entire test.

8.3.4 Midwest Refinery--1979 (Case 4)

The information in Table 8-22 was used to estimate emissions from the Case 4 facility test. No specific constituent data were available; emissions were estimated for total organics using average parameter values. Results are presented in Table 8-23. The comparisons are for the cumulative weight percent of applied oil that was emitted over the entire period of each test.

8.4 LANDFILLS AND WASTEPILES

Emission testing has been performed on at least one active (open) landfill at each of five sites. Only three of these sites have closed or inactive landfills at which emission measurements were performed. No emission test data are available for wastepiles.

Meaningful comparisons can be performed of emission test data with mathematical model predictions provided that all key model input parameters are available from the tests. A review of documentation from the emission tests indicates that generally more than half of the needed model input parameters (other than chemical property data) are unknown, despite the fact that several emission tests were performed with the stated intention

TABLE 8-18. INPUT PARAMETERS FOR RTI LAND TREATMENT MODEL^a

Parameter	Value	Source
Organic (oil) loading	0.0328 g/cm ³	Estimated from field data
Tilling depth	20 cm	Field data
Soil porosity	0.5	Field data
Molecular weight of oil	282 g/g mol	Field data
Toluene concentration	0.00157 (wt. fraction of oil)	Calculated from field data
Toluene diffusivity	8.70 E-02 cm ² /s	Data base
Toluene vapor pressure	30.0 mm Hg	Data base
Toluene biorate	73.0 mg VO/g•h	Data base
VO concentration	0.04 (wt. fraction of oil)	Calculated from field data
VO diffusivity	6.60E-02 cm ² /s	Average from field data
VO vapor pressure	14.6 mm Hg	Average from field data
VO biorate	23.68 mg VO/g•h	Average from field data

^aSource of field data: Reference 37.

TABLE 8-19. ESTIMATED VS. MEASURED EMISSIONS--CASE 2

	Time after tilling, day/h	Estimated emissions, wt. %	Measured emissions, wt. %
Toluene	33/793	31	37
Total VO	33/793	32	30
Total oil	33/793	1.3	1.2

TABLE 8-20. INPUT PARAMETERS FOR RTI LAND TREATMENT MODEL^a

Parameter	Value	Source
Organic loading	0.0406 g/cm ³	Calculated from field data
Tilling depth	19.6 cm	Field data
Soil porosity	0.5	Assumed
Molecular weight of oil	282 g/g mol	Assumed
Vapor pressure	0.57 mm Hg	Calculated by GCAB
Diffusivity in air	2.70 E-02 cm ² /s	Average from field data
Biorate	23.68 mg VO/g•h	Average from data base

^aSource of field data: Reference 38.^bReference 39.

TABLE 8-21. ESTIMATED VS. MEASURED TOTAL VO EMISSIONS--CASE 3

Time after tilling, h	Estimated emissions, wt. % total applied oil	Measured emissions, wt. % total applied oil
68.00	4.5	0.77

TABLE 8-22. INPUT PARAMETERS FOR RTI LAND TREATMENT MODEL^a

Parameter	Value	Source
Organic loading	0.002125 g/cm ³	Estimated from field data
Tilling depth	20 cm	Assumed
Soil porosity	0.5	Assumed
Molecular weight of oil	282 g/g mol	Assumed
Diffusivity in air	9.12 E-02	Average from data base
Vapor pressure	0.76 mm Hg	Calculated by GCAB ^b
Biorate	23.68 mg VO/g•h	Average from data base

^aSource of field data: Reference 40.

^bReference 41.

TABLE 8-23. ESTIMATED VS. MEASURED EMISSIONS--CASE 4

Test	Elapsed time, day/h	Estimated emissions, wt. % total applied oil	Measured emissions, wt. % total applied oil
5	1/20	5.0	0.14
6	13/307	14.0	2.5
7	26/619	16.0	13.5
8	5/122	14.0	1.1
9	22/520	28.0	13.4

of validating emission models. Examples of key model input parameters that are generally unknown or poorly defined include waste porosities (air and total), average waste bed temperature (for active and closed landfills), waste composition at depths greater than the surface layer, barometric pressures, clay cap porosities (air and total), clay cap thickness, waste bed depth, and (for active landfills particularly) time between core sampling and air emissions determination. To apply the models, representative default values have been used where necessary. Because of the necessity to estimate key input parameters, the comparisons that follow are of extremely limited value for model validation. To achieve validation of emission models, additional field tests or laboratory experiments are needed for active and closed landfills and wastepiles.

Field data from two sites were used for comparison with the land treatment model as applied to active landfills. These sites (5 and 8) were chosen because of similarity in constituency of selected chemicals and relative availability of model input parameters. However, it should be noted that at each of the sites more than half of the needed model input parameters were not available from the tests and thus required estimation.

Information on the waste composition within closed landfills was insufficient to allow use of the closed landfill model. At two of the three closed/inactive landfill sites (4 and 5), no solid samples of waste were taken; at the remaining site (Site 9), a single soil core was apparently extracted from the 3-ft clay cover, providing no information about the composition of the waste below the cover. However, it should be noted that Farmer et al.⁴² (who developed the precursor* to the RTI closed landfill model) mentioned that their model has received experimental verification via a laboratory experiment using hexachlorobenzene-containing waste in a simulated landfill.

Following are the results of the comparison for active landfills at Sites 5^{43,44} and 8⁴⁵. Table 8-24 presents model input parameters used in

*The Farmer et al. model accounts for diffusion through the clay cap only (not barometric pumping).

TABLE 8-24. MODEL INPUT PARAMETERS USED IN APPLICATION OF THE RTI LAND TREATMENT MODEL TO AN ACTIVE LANDFILL AT SITE 5^a

Parameter	Value	Data source
L, total organic loading in soil	$2.65 \times 10^{-3} \text{ g/cm}^3$	Inferred from field data (solid sample analysis) assuming soil density = 2.3 g/cm^3
C _i , weight fraction of constituent i in organic phase	Xylene: 0.178 Methylene chloride: 8.48×10^{-4} Tetrachloroethylene: 1.37×10^{-3}	Computed from field data (solid sample analysis)
T, temperature of constituent vapor in soil	25 °C	Default value
l, depth of waste in landfill	229 cm (7.5 ft)	Default value
ε _T , total porosity of waste	0.50 (50%)	Default value
ε _a , air porosity of waste	0.25 (25%)	Default value
S _b , soil biomass concentration	0 g/cm ³	Default value
MW _{oil} , molecular weight of organic carrier liquid	150 g/g mol	Default value
t, time between soil sampling and air emission measurement	900 s (15 min)	Engineering judgment

^aLandfill 10, General Organic Cell.46,47

the application of the RTI land treatment model to an active landfill at Site 5. Table 8-25 presents a comparison of measured and predicted emission rates for the Site 5 landfill.

Model predictions for the chemicals assessed here are higher than field data values by a factor ranging from 13 to 441. This discrepancy may be largely a result of the presence of daily earth covers (6 in. thickness) and layers of drums within the waste bed--neither of which are accounted for by the model. Other contributing factors may include the estimation of key model input parameters (e.g., air porosity of waste, temperature of the constituent within soil) and the nonrepresentative nature of the waste sample (obtained at the surface) for describing the waste composition at depth.

Table 8-26 presents model input parameters used in the application of the RTI land treatment model to an active landfill at Site 8. Table 8-27 presents a comparison of measured and predicted emission rates for the Site 8 landfill.

Model predictions of the emissions at Site 8 are, in general, closer to field data than were the predictions made for Site 5. Better overall agreement may be due to the absence of drums and daily earth covers in this landfill.

8.5 TRANSFER, STORAGE, AND HANDLING OPERATIONS

8.5.1 Models Documented in AP-42

Emission methods for the following models were taken from AP-42; they have been developed from the field data in the petroleum industry and should be applicable to TSDF:

- Container loading (from AP-42, Section 4.4)⁵²
- Container cleaning (from AP-42, Section 4.8)⁵³
- Stationary tank loading (from AP-42, Section 4.3)⁵⁴
- Stationary tank storage (from AP-42, Section 4.3).⁵⁵

8.5.2 Fugitive Emissions

Fugitive emission sources have been studied extensively for the petroleum and Synthetic Organic Chemical Manufacturing Industries

TABLE 8-25. COMPARISON OF MEASURED AND PREDICTED EMISSION RATES
FOR SITE 5 ACTIVE LANDFILL^a

Chemical	Field data result, $\mu\text{g}/\text{m}^2\cdot\text{s}$	Model prediction, $\mu\text{g}/\text{m}^2\cdot\text{s}$
Xylene	32.8	440.0
Methylene chloride	0.734	14.0
Tetrachloroethylene	0.0111	4.9

^aLandfill 10, General Organic Cell. 48,49

TABLE 8-26. MODEL INPUT PARAMETERS USED IN APPLICATION OF THE RTI LAND TREATMENT MODEL TO AN ACTIVE LANDFILL AT SITE 850

Parameter	Value	Source
L, total organic loading in soil	$1.71 \times 10^{-6} \text{ g/cm}^3$	Field data
C _i , weight fraction of VO _i in organic phase	Xylene: 0.012 1,1,1-TCE: 0.19 Tetrachloroethylene: 0.096	Computed from field data (solid sample analysis)
T, temperature of VO vapor in soil	25 °C ^a	Default value
l, depth of waste in landfill	229 cm (7.5 ft)	Default value
ε _T , total porosity of waste	0.50 (50%) ^b	Default value
ε _a , air porosity of waste	0.25 (25%) ^b	Default value
S _b , soil biomass concentration	0 g/cm ³	Default value
MW _{oil} , molecular weight of organic carrier liquid	150 g/g mol	Default value
t, time between soil sampling and air emission measurement	900 s (15 min)	Engineering judgment

^aSoil surface temperatures at this site were reported at 26 to 36 °C. The model unit default value of 25 °C is applied to the constituent within the soil in this analysis.

^bA single "porosity" value of 31.7 percent was reported for this site, based on one soil sample. Because this value is not defined explicitly, and may not be representative of typical waste in the landfill, the model unit default values of air porosity and total porosity were applied in this analysis.

TABLE 8-27. COMPARISON OF MEASURED AND PREDICTED EMISSION RATES
FOR THE SITE 8 ACTIVE LANDFILL⁵¹

Chemical	Field data result, $\mu\text{g}/\text{m}^2\cdot\text{s}$	Model prediction, $\mu\text{g}/\text{m}^2\cdot\text{s}$
Total xylene	6.21	0.23
1,1,1-Trichloroethane	3.57	3.8
Tetrachloroethylene	6.31	1.9

(SOCMI) facilities.⁵⁶ These SOCMI emission factors are assumed to be applicable to similar operations at TSDF.

8.5.3 Spillage

An ICF⁵⁷ study of truck transport to and from TSDF and truck emissions at TSDF terminals provided the information necessary to estimate spillage losses during TSDF and trucking operation. However, none of the reports listed in Appendix F of the BID contains field test data for comparison.

8.5.4 Open Dumpster Storage Emissions

No field data were available for comparison.

8.6 REFERENCES

1. GCA Corporation. Air Emissions for Quiescent Surface Impoundments--Emissions Data and Model Review, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Contract No. 68-01-6871, Assignment 49. August 1985.
2. Radian Corporation. Hazardous Waste Treatment, Storage, and Disposal Facility Area Sources: VOC Air Emissions. Prepared for U.S. Environmental Protection Agency, Emissions Standards and Engineering Division. Research Triangle Park, NC. Contract No. 68-02-3850. January 25, 1985.
3. Reference 1.
4. Reference 2.
5. Reference 1.
6. Reference 2.
7. Reference 1.
8. Radian Corporation. Hazardous Waste Treatment, Storage, and Disposal Facility Area Sources: VOC Air Emissions at Chemical Waste Management, Inc., Kettleman Hills Facility. Volume 1. Prepared for U.S. Environmental Protection Agency, Emissions Standards and Engineering Division. Research Triangle Park, NC. EMB Report 85-HW5-2. December 1984.
9. GCA Corporation. First Chemical Corporation Wastewater Holding Lagoon Field Study. Prepared for U.S. Environmental Protection Agency. Contract No. 68-02-3851 (WA10). 143 p. August 1986.
10. GCA Corporation. Emissions Data and Model Review for Wastewater Treatment Operations. Draft Technical Note. Prepared for U.S.

Environmental Protection Agency. Contract No. 68-01-6871, Assignment 49. August 1985.

11. Reference 10.
12. Reference 10.
13. Reference 10.
14. Reference 10.
15. Reference 10.
16. Nelson, Thomas P., et al. (Radian). Field Assessment of Air Emissions and Their Control at Hazardous Waste Facilities (Draft). Prepared for U.S. Environmental Protection Agency, Industrial Environmental Research Laboratory. Cincinnati, Ohio. December 1984. 77 p.
17. Reference 10.
18. Reference 16.
19. Petrasek, A., B. Austern, and T. Meiheisel. Removal and Partitioning of Volatile Organic Priority Pollutants in Wastewater Treatment. Presented at the Ninth U.S.-Japan Conference on Sewage Treatment Technology. Tokyo, Japan. September 1983. 31 p.
20. Reference 19, p. 2-4.
21. Reference 19, p. 16.
22. Namkung, E., and B. Rittman. Estimating Volatile Organic Compound Emissions from Publicly Owned Treatment Works. Journal WPCF. 59(7):677.
23. Reference 22, p. 671-672.
24. Reference 22, p. 672.
25. Tabak, H., S. Quave, C. Mashni, and E. Barth. Biodegradability Studies with Priority Pollutant Organic Compounds. Staff Report. Wastewater Research Division. U.S. Environmental Protection Agency. Cincinnati, Ohio. 1980.
26. Reference 25.
27. Coburn, J., C. Allen, D. Green and K. Leese. Site Visits of Aerated and Non-aerated Impoundments Revised Draft Summary Report. Prepared for U.S. Environmental Protection Agency. Contract No. 68-03-3253, Work Assignment No. 3-8. April 1988. p. A-1 to A-34.

28. Reference 27.
29. Allen, C. C., et al. Preliminary Assessment of Aerated Waste Treatment Systems at Hazardous Waste TSDFs (Draft). Prepared for U.S. Environmental Protection Agency. Contract No. 68-02-3992. December 1985. 108 p.
30. Reference 29.
31. Utah State University. Evaluation of Volatilization of Hazardous Constituents at Hazardous Waste Land Treatment Sites. Prepared for U.S. Environmental Protection Agency. Ada, Oklahoma. Undated. 157 p.
32. Radian Corporation. Field Assessment of Air Emissions and Their Control at a Refinery Land Treatment Facility. Volumes I and II. Prepared for U.S. Environmental Protection Agency. Cincinnati, OH. September 1986. 699 p.
33. Radian Corporation. Hazardous Waste Treatment, Storage, and Disposal Facility Area Sources--VOC Air Emissions. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. January 1985. 141 p.
34. GCA Corporation. Air Emissions from Land Treatment--Emissions Data and Model Review. Draft Technical Note. Prepared for U.S. Environmental Protection Agency. August 1985. p. 4-36.
35. Suntech, Inc. Atmospheric Emissions from Oily Waste Land Spreading. Final Report SWM-8(563). Prepared for American Petroleum Institute. Washington, DC. Undated. 63 p.
36. Reference 31.
37. Reference 32.
38. Reference 33.
39. Reference 34.
40. Reference 35.
41. Reference 34.
42. Farmer, W. J., M. S. Yang, J. Letey, W. F. Spencer, and M. H. Roulter. Land Disposal of hexachlorobenzene Wastes: Controlling Vapor Movement in Soils. Fourth Annual Research Symposium. U.S. Environmental Protection Agency. Publication No. EPA-600/9-78-016. August 1978.
43. Radian Corporation. Hazardous Waste Treatment, Storage and Disposal Facility Area Sources: VOC Air Emissions. Prepared for U.S. Environ-

mental Protection Agency. Research Triangle Park, NC. DCN 85-222-078-17-09. January 25, 1985. 141 p.

44. Radian Corporation. Evaluation of Air Emissions from Hazardous Waste Treatment, Storage and Disposal Facilities in Support of the RCRA Air Emission Regulatory Impact Analysis (RIA): Data Volume for Site 4 and Site 5. Prepared for U.S. Environmental Protection Agency. Cincinnati, OH. DCN 83-203-001-63-19. January 11, 1984. 454 p.
45. Radian Corporation. Hazardous Waste Treatment, Storage and Disposal Facility Area Sources--VOC Air Emissions. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. EMB Report 85-HWS-1. May 1985. 54 p.
46. Reference 43.
47. Reference 44.
48. Reference 43.
49. Reference 44.
50. Reference 45.
51. Reference 45.
52. U.S. Environmental Protection Agency. Transportation and Marketing of Petroleum Liquids. In: AP-42. Compilation of Air Pollutant Emission Factors. Third Edition, Supplement 12, Section 4.4. Office of Air Quality Planning and Standards. Research Triangle Park, NC. July 1979. 13 p.
53. U.S. Environmental Protection Agency. Tank and Drum Cleaning. In: AP-42. Compilation of Air Pollutant Emission Factors. Third Edition, Supplement 12, Section 4.8. Office of Air Quality Planning and Standards. Research Triangle Park, NC. February 1980. 4 p.
54. U.S. Environmental Protection Agency. Storage of Organic Liquids. In: AP-42. Compilation of Air Pollutant Emission Factors. Third Edition, Supplement 12, Section 4.3. Office of Air Quality Planning and Standards. Research Triangle Park, NC. April 1981. 25 p.
55. Reference 47.
56. U.S. Environmental Protection Agency. Control of Volatile Organic Compound Leaks from Synthetic Organic Chemical and Polymer Manufacturing Equipment. Research Triangle Park, NC. Publication No. EPA-450/3-86-006. March 1984.
57. U.S. Environmental Protection Agency. Assessing the Release and Costs Associated with Truck Transport of Hazardous Wastes. PB 84-224-468 (Contract No. 68-01-0021). Washington, DC. January 1984. 151 p.

APPENDIX A
CHEMDAT7 USER'S GUIDE

APPENDIX A

CHEMDAT7 USER'S GUIDE

A.1 INTRODUCTION

CHEMDAT7 is a Lotus 1,2,3 spreadsheet that includes analytical models for estimating volatile organic (VO) compound emissions from treatment, storage, and disposal facility (TSDF) processes under user-specified input parameters. The available models include disposal impoundments, closed landfills, land treatment facilities, and aeration and nonaeration impoundment processes. Predicted emissions can be viewed on the screen or printed. A graphical presentation of the relationships between emission prediction and vapor pressure and between emission prediction and the partition coefficient is also available to the user. The resulting scatter diagrams can be printed via PrintGraph, another Lotus procedure.

VO emission models from hazardous waste TSDF were described in Chapters 4.0, 5.0, 6.0, and 7.0 of this report. The emission rates from some TSDF can be estimated via CHEMDAT7. In this regard, Exhibit A-1 specifies the appropriate CHEMDAT7 model to estimate emissions from particular TSDF. For example, the nonaerated impoundment model in CHEMDAT7 can estimate emissions from storage impoundments. The CHEMDAT7 model for predicting emissions from treatment impoundments is the aerated impoundment model. Furthermore, the land treatment model in CHEMDAT7 can estimate emissions from land treatment soil, open landfills, and wastepiles. Emissions from an oil film surface in a land treatment facility or an oil film on a surface impoundment are predicted via the oil film model in CHEMDAT7. When a CHEMDAT7 model is not available to predict emissions, the reader must resort to the prediction equations in the previous sections of this report to estimate VO emissions from TSDF.

CHEMDAT7 calculates the fractions of waste constituents of interest that are distributed among the pathways (partition fractions) applicable to the facility under analysis. Estimated annual emissions from many of the

TSDF are calculated by this spreadsheet. Otherwise, estimated annual emissions can be calculated offline by multiplying the throughput of the constituent of interest by the partition fraction.

The seventh version of the CHEMDAT spreadsheet contains several major operational modifications. In CHEMDAT7, the user can select a subset of target compounds for investigation. The user can also specify which TSDF processes are to be considered during a session. These two selections improve the efficiency of CHEMDAT7 relative to some of the earlier versions by minimizing storage requirements as well as the actual loading and execution time.

Default input parameters in the accompanying CHEMDAT7 diskette demonstrate example calculations in Chapters 4.0, 5.0, and 6.0 of this report. However, the user can readily change the input parameters to reflect different hazardous waste TSDF characteristics and then recalculate emissions under these modified conditions. The diskette with this user's guide is write-protected. It is suggested that a copy be used in estimating emissions. Furthermore, the list of compounds in CHEMDAT7 can be augmented by any of the 700 chemicals in Appendix D. Procedures for introducing additional compounds into CHEMDAT7 are described in Appendix D.

Instructions on the use of CHEMDAT7 appear in this appendix. (We have assumed throughout this appendix that the user has some minimal knowledge of Lotus 1,2,3.) CHEMDAT7 contains a data base of component-specific properties used to generate internally the inputs for the environmental fate models of waste disposal practices. Section A.2 specifies how to get started in Lotus 1,2,3. Section A.3 details proper utilization and modification of CHEMDAT7 via the alternative command menu. The data base and model parameters are described in Section A.4. The steps required to obtain a printout of the graphs are described in Section A.5. The referenced exhibits complete this appendix.

A.2 GETTING STARTED IN LOTUS 1,2,3

The applications program, Lotus 1,2,3, Version 2.0, should be installed on the computer system for use. Place the diskette containing CHEMDAT7 in a low density drive.

Access the 1,2,3 mode of operation in Lotus. A skeleton of a spreadsheet will appear on the screen, and the message READY will appear in

the upper right corner of the screen. (Consider this and other messages as road signs that guide you through CHEMDAT7.) Either copy the contents of the CHEMDAT7 diskette to your default directory or set the directory in Lotus 1,2,3 to the disk drive containing the CHEMDAT7 diskette. To retrieve CHEMDAT7, press the following sequence of keys, i.e., those keys that appear between brackets { }:

{/} {F} {R} .

Note that {/} calls the active command mode. (The active command mode is indicated by the message MENU in the upper right corner of the screen.) The F represents file and the R represents retrieve. In other words, these are the commands to retrieve a file. We indicate the file to retrieve by typing the drive label, followed by a colon, followed by CHEMDAT7,

{B:CHEMDAT7} ,

and then pressing {RETURN}.

An equivalent approach is to position the cursor over CHEMDAT7, which appears below the command line, and then press {RETURN}. The cursor controls=(keys with arrows on top, i.e., ↑, ↓, →, and ←) position the cursor. After CHEMDAT7 loads for a minute or two (note the message WAIT flashing at the bottom of the screen), the user will be in the help mode of the alternative command menu developed specifically for CHEMDAT7. The message CMD at the bottom of the screen indicates that the alternative command menu is active. After reviewing the appropriate help screens (see Section A.6 for further details), press {Q} for quit to return to the main alternative command menu. A discussion of this menu follows.

A.3 ALTERNATIVE COMMAND MENU

The alternative command menu is a menu-driven set of instructions specific to CHEMDAT7. Choose an option (DATA-FORMS, VIEW, SORT, PRINT, SELECT, HELP, or QUIT) by positioning the cursor over the selection and pressing {RETURN}. Alternatively, the first letter or set of letters that uniquely identify the selection can be entered: e.g., {P} can be pressed to choose the PRINT option. Alternative command menu options and suboptions are listed in Exhibit A-2; please study this guide carefully. If you get lost after accessing the alternative command of CHEMDAT7, press {ESC} several times to completely exit the macro. Then the first level of

options (DATA-FORMS, VIEW, etc.) can be accessed via {ALT M}. "ALT M" stands for the alternative command menu.

A.3.1 DATA-FORMS Option

The DATA-FORMS option in the alternative command menu is the crux of this software package. It is the vehicle by which the user can change model input parameters in CHEMDAT7.

Once the user has specified {DATA-FORMS} and has selected the desired model, the corresponding input parameters will appear highlighted on the computer screen. Move the cursor to the cell location containing the model input parameter to be modified. After the proper cell has been located in the spreadsheet with the cursor, type the new input parameter, then press {RETURN}. The new input parameter will appear on the screen. At this point, the message CALC will appear in the lower right side of the screen indicating that the model predictions must be recalculated to reflect the new input parameter. Once all the input parameters have been modified, press {F9} to recalculate the emission predictions under the new input parameters. The WAIT message will flash as the model predictions are recalculated. The CALC message will disappear once the model predictions have been revised. The user must perform this latter step so that the emission results will reflect the modified input parameters. Furthermore, this revised spreadsheet must be saved if updated graphs are desired.

To save the revised spreadsheet, access the active command mode (press {ESC}). Then type

B: / F S {ENTER} R ,

where F represents file, S represents save, the file name automatically appears, and R represents replace. Consequently, the old version of the CHEMDAT7 spreadsheet is replaced by the new, modified version.

The DEFAULT suboption for DATA-FORMS is used to replace model input parameters with the default values, i.e., those values corresponding to example calculations described in Section A.4.3. The model default values for impoundments (non-aerated, diffused, oil film, or disposal), aerated wastewater treatment, land treatment soil, open landfills, and closed landfills appear in Exhibits A-3, A-4, A-5, A-6, A-7, and A-8, respectively. After input parameters are replaced with default values, remember to recalculate model predictions.

The CONC suboption for DATA-FORMS is used when individual concentrations are to be specified for each VO in the data set of interest. If you select this option, you may specify the individual concentrations for each of the units that support this feature, but you must enter a zero as the concentration in the standard data forms. Failure to do this will result in the concentration used for each of the compounds defaulting to the concentration on the standard data form.

The IMPORT suboption is used with a specially designed program, CHEM7, that complements CHEMDAT7. One of the uses of the {IMPORT} {RESTORE} option is to replace the data in CHEMDAT7 with the original data supplied with the diskette. The {IMPORT} {GO} option loads a set of compounds into the spreadsheet under the compounds selected with the {SORT} {SELECT} option.

A.3.2 VIEW Option

A.3.2.1 Viewing Model Emission Results. The VIEW option enables the user to view on the computer screen emission results from disposal impoundments, land treatment facilities, open landfills/wastepiles, closed landfills, and nonaerated and aerated models. Exhibit A-2 describes the options and suboptions available. To exit the model results, press {ALT M}.

A.3.2.2 Viewing Graphs. The user is able to view graphs on the computer screen by pressing {VIEW} {GRAPH} and then selecting the desired model. The user must exit from this procedure via {QUIT} in order to reformat the spreadsheet.

A.3.2.3 Viewing Mass Transfer Coefficients. The user is able to view mass transfer coefficients on the computer screen by pressing {VIEW} {K} and selecting the desired facility. A list of the facilities that use mass transfer coefficients is presented on screen, preceded by a number. After the user presses the desired number, the mass transfer coefficients and Henry's law constants appear on the screen for each compound. The units of the mass transfer coefficients are M/sec.

A.3.3 SORT Option

This option in the alternative command menu enables the user to rearrange the order of the compounds in the CHEMDAT7 spreadsheet. Suboptions include sorting by biorate, selected compound, alphabetically, by

compound type (class), and by vapor pressure. Model predictions must be recalculated after sorting.

The {SELECT} suboption under {SORT} is a very powerful feature of CHEMDAT7. Analogous to a "cut and paste" procedure, {SORT} {SELECT} enables the user to select a subset of the 61 compounds for entry into the CHEMDAT7 spreadsheet. Model calculations are only performed for the selected compounds, which improves the performance of the spreadsheet. Consequently, emission results are limited to the selected compounds. This {SORT} {SELECT} option was employed to choose the compound of interest, benzene, in previous exhibits.

Two approaches are available for selecting compounds. To select a set of chemicals, use a "1" to flag those compounds of interest. Those compounds preceded with a "0" will be ignored. The user may also order the selected compounds so that compound-specific emission results are printed in a desired sequence. If there are n compounds of interest, place an integer from 1 to n in front of each compound. Once {ALT} {Z} is typed, CHEMDAT7 will rearrange the compounds in descending order according to this integer. As before, emission results will not be estimated for those compounds preceded by a "0".

{SORT} {DATA} is an option used to add additional compounds to the {SORT} {SELECT} compounds that are selected. An extended data base of approximately 300 compounds is added to the spreadsheet, starting at Cell {A 255}. The compounds in this extended data base are selected in an identical fashion to the compound selection process under {SORT} {SELECT}. After the compounds of interest are selected, press {ALT A} to add the selected compound to your CHEMDAT7 compound list. CAUTION: This option will destroy data for compounds that are not selected in the {SORT} {SELECT} option, so perform the {SORT} {SELECT} step first. If you change your mind later, you can always restore the original data set by {DATA-FROM} {IMPORT} {RESTORE} and start over.

A.3.4 PRINT Option

The PRINT option allows the user to obtain a hard copy of model-specific emission results (via the suboptions LAND-TREAT, DISPOSAL, AERATED, LANDFILL, and NON-AERATED) and the corresponding set of input parameters, as well as mass transfer coefficients (via the K suboption) and the data summaries (via the DATA suboption). The data summaries include

biorate data (refer to columns N through P in Exhibit A-9), land treatment data (see columns AM and AN in Exhibit A-9), and selected data tables (see columns F, G, H, and I in Exhibit A-9) that are printed via the commands {PRINT} {DATA} {BIORATE}, {PRINT} {DATA} {LAND-TREAT}, or {PRINT} {DATA} {DATA}, respectively. Examples of printed model input parameters and emission results appear in the exhibits.

The PRINT option has another important function: it enables the user to store the graph file on the diskette. Enter the commands {PRINT} {GRAPH} followed by the selected model. This procedure is required to obtain a hard copy of the model-specific graph. Further details are provided in Section A.5.

A.3.5 SELECT Option

The SELECT option in the main menu, another "cut and paste" feature of CHEMDAT7, allows the user to activate only a subset of the TSDf models for entry into the spreadsheet. Models are selected through the 0-1 (off-on) flag system described previously. Note that, if the user attempts to use a model that is not active, error messages will be printed instead of model emission results.

A.3.6 HELP Option

The HELP option provides the user with information regarding the proper application and various assumptions of the specific models in CHEMDAT7. A general help screen is also available. The help screens are presented in Section A.6.

A.3.7 QUIT Option

The active command mode can be accessed from the alternative command menu by locating the cursor over {QUIT} and pressing {RETURN}. In this mode, all the features of Lotus 1,2,3 are available by pressing {/}.

A.3.8 Adding Additional Compounds

There are three ways to add new compounds to the CHEMDAT data base: overwriting existing data, selecting compounds from the supplemental list, and using a separate compound data program (CHEM7).

A.3.8.1 Overwriting Existing Data. To overwrite existing data, use the cursor to locate a compound that you do not need and wish to overwrite:

- Use the {F5} key to locate the cursor in the cell containing the name of the compound to be overwritten. The data for this compound appears in the corresponding row.

- Enter the data for the new compound in that row. Data sources include:
 - Reference books
 - CHEM7
 - Default values.
- Use the alternative command menu {ALT}{M} option of {Sort}{Select} to select the compound and reform the worksheet by pressing {ALT}{Z}.

A.3.8.2 Selecting from a Supplemental List. To select compounds from a supplemental list, use the command menu {ALT}{M} option of {Sort}{Data}. Select the compounds in the supplemental list by tagging each selected compound with a number greater than 1 in column A of the row containing the compound name. After the selection is complete, press {ALT}{A} to reform the spreadsheet.

The compounds selected from the supplemental list will be sorted from the list and transferred to the bottom of the working list in rows 14 through 82. The spreadsheet will then be sorted with all selected compounds (both from the working list and the supplemental list) transferred to the top of the list, in descending order of the tag number in column A.

There is a limit of 20 compounds when individual concentrations are specified for each compound. Row (DP43..DY43) can be copied in lower rows to expand the compound number limit for individual concentration specifications.

A.3.8.3 Loading Compounds from a Master List. To load a compound from a master list, a separate program (CHEM7) is available. To use CHEM7, you must quit CHEMDAT7 by the {/}{Q}{Y} option. Change the directory to the same directory that you are using for the CHEMDAT7 program, then type CHEM7 and press {ENTER}.

For the users of a hard disk system; copy the files on the CHEM7 diskette to your hard disk directory used by CHEMDAT7. Be sure to save a backup copy of all your original diskettes.

Users of a floppy disk system will need to use the CHEM7 diskette. Again, be sure to make a backup copy before using your diskette. When you load the compound data, you need to put a diskette containing the compound data (MASTERCL.CH7) in the disk drive. When you print your data to diskette, you need to use a diskette that you will use with the CHEMDAT7 program. You may place the CHEMDAT7 diskette (CHEMDAT7.WK1) in the disk drive.

A separate manual is available to describe the program CHEM7. The user must load compound data from a master data base, select compounds, and print the data to disk. The user can optionally append compound data to the master data base. If the user only wants to select additional compounds from an extended list and let the computer automatically fill in missing data, the command CHEM7 1 can be entered, with a space between the 7 and the 1.

After using CHEM7, return to CHEMDAT7 and load the files you have created. In CHEMDAT7 you will use the command menu {ALT}{M} option of {Data-form}{Import}{Go}.

A.4 ANATOMY OF THE CHEMDAT7 SPREADSHEET

This section describes the structure of the CHEMDAT7 spreadsheet; a generalized layout of the spreadsheet appears in Attachment A. In general, rows 1 through 5 of this spreadsheet contain column labels or names. The compound-specific data base and calculation results appear in rows 14 through 83. The locations of model-specific input parameters are specified throughout Section A.4.3.

A.4.1 Data Base

The data base in the CHEMDAT7 spreadsheet is a matrix of component-specific properties or calculations. Sixty-one chemicals or compounds appear in rows 14 through 83 of the spreadsheet. These compounds are listed in Exhibit A-10.

Compound properties and model computational results appear in columns B through BU of the spreadsheet. For example, molecular weight appears in column D. Thus, each compound-specific data item has a unique cell address "cr" where c (alphabetic) represents the appropriate column and r (numeric) the row. Suppose data on toluene appear in row 16 of the CHEMDAT7 spreadsheet. Then, the molecular weight of toluene would appear in cell D16.

A.4.2 Column Labels

In general, column labels or names appear in rows 1 through 5 of the spreadsheet. Exhibit A-9 lists the column labels in CHEMDAT7.

A.4.3 Model Input Parameters and Predicted Emissions

Brief descriptions and locations of the CHEMDAT7 model-specific parameters are presented in the following subsections. Additionally, use of CHEMDAT7 is demonstrated via example calculations presented previously in this report. VO emissions estimated from CHEMDAT7 models are compared to the example calculation results.

Missing data in the CHEMDAT7 spreadsheet (e.g., vapor pressure for dioxin) frequently preclude estimating emission rates for the affected compound. In such cases, "ERR" is printed in place of the estimated emission. The user is encouraged to insert these missing data as values become available. The procedure for modifying input parameters is described in Section A.3.1.

A.4.3.1 Nonaerated Model. CHEMDAT7 nonaerated model input parameters are located in cells C06 through C015 of the spreadsheet and are illustrated in Exhibit A-11. (This and subsequent exhibits depict what the user will see on the computer screen.) The windspeed in meters/second is placed in C06. C07 contains the depth of the nonaerated impoundment (in meters). The surface area of the impoundment (in meters squared) and the flow rate (in cubic meters/second) appear in C08 and C09, respectively. The V0 inlet concentration expressed in milligrams/liter is placed in C010. Cell C011 contains the sum of the organics (in milligrams/liter) entering the facility. The conversion of these organics to biomass is used to estimate the rate of adsorption as a pathway for the removal of organics from the system. Note that the V0 inlet concentration in C010 should be less than the overall concentration of organics in C011. Because biodegradation is presumed to be nonexistent in a storage impoundment, the amount of active biomass (C012) has been set at zero. If the uniqueness of the TSDf permits biodegradation as a pathway and the user desires to consider its impact on emissions, C012 can be changed to reflect active biomass using standard Lotus procedures (see Section A.3.1). The input parameter, biomass solids in (C013), is appropriate for municipal facilities only. An approximation of the solids input (C014) for adsorption in municipal facilities is 200 ppmw. Of these solids, approximately 50 percent are removed in the primary clarifier, leaving 0.1 g/L solids input to the treatment basin. The ambient air temperature for the facility in degrees Celsius is placed in C015.

The nonaerated model input parameters in Exhibit A-11 have the same values as those used in the example calculation for storage impoundments in Section 4.2.3 where the compound of interest is benzene. The predicted emissions for benzene from CHEMDAT7 (see Exhibit A-11) are to those model results presented in the latter portion of Section 4.2.3. The predicted fraction of benzene that will be emitted to the air is calculated as 0.801.

The estimated annual air emissions of benzene from the example nonaerated surface impoundment total 0.39 Mg/yr. CHEMDAT7 model input parameters and results are printed using the PRINT option as discussed below in Section A.3.4.

In contrast, the example calculation in Section 4.3.3 considers biodegradation as a pathway. The amount of active biomass is 0.05 g/L; see Exhibit A-3. The remaining model input parameters are identical to those in Section 4.2.3. Almost 11 percent of the benzene is emitted to the air so that the annual emission rate is 0.052 Mg/yr.

A.4.3.2 Aerated Model. CHEMDAT7 parameters for the aerated impoundment model are located in column C0, rows 78-88, and column CS, rows 79-85 (see Exhibit A-4). The equations for estimating relative pathways include those for the quiescent surface of nonaerated systems, but these must be supplemented to account for a turbulent zone.

Because biodegradation is assumed to occur in aerated impoundments, the amount of active biomass (grams/liter) must be specified in C082. Cell C087 contains the fraction of the impoundment's surface area that is agitated. The submerged air flow (m^3/s) in C088 accommodates a mechanical air source located below the surface of the impoundment. Consequently, emissions from a diffused air flow-through system, i.e., one that emphasizes biodegradation, are estimated by placing an appropriate value in C088. The remaining input parameters in column C0 are the same as those explained in the previous section.

The number of impellers is specified in CS79. The oxygen transfer rating of the aerator (pounds O_2 /horsepower/hour) appears in CS80. The total power of the mechanical aeration system expressed in horsepower is placed in cell CS81. Cell CS82 contains the power efficiency factor (no units). The remaining parameter inputs include the water temperature (CS83) in degrees Celsius, the impeller diameter (CS84) in centimeters, and the impeller speed (CS85) in rads/second. If the impoundment's surface is not agitated (C087=0), the final seven model input parameters (CS79-CS85) are ignored in the emission calculations.

The aerated model input parameter values in Exhibit A-4 are the same as those used in the example for a mechanically aerated treatment impoundment discussed in Section 4.4.3. The compound of interest is benzene. CHEMDAT7 reproduces the materials balance results presented in

Section 4.4.3; see Exhibit A-4. The predicted fraction of benzene that will be emitted to the air is 0.83. The estimated annual air emissions of benzene from such aerated impoundments are 0.81 Mg.

Emission results for the activated sludge unit in Section 4.4.4 and the diffused air activated sludge unit in Section 4.6.3 are also available from the CHEMDAT7 aerated model. See Exhibits A-5 and A-12, respectively, for the model input parameters, the materials balance results, and the annual air emission rate.

Diffused aerated systems retain the input parameters for mechanical aeration. It is possible to have both in an activated sludge system. When not physically present, the mechanical aeration parameters can be used to estimate the mass transfer coefficients of exposed surfaces agitated by diffusing air bubbles.

The diffused air option in the aerated impoundment model can be used with or without surface aeration. If the surface is not agitated by the churning action of submerged aeration, the fraction agitated can be specified as zero. If the surface is visibly agitated, the fraction agitated can be specified as the fraction of the surface that is estimated to be agitated by visual inspection or calculations. The mass transfer coefficients of the agitated zone are estimated the same as for surface aeration, so the default parameters of the surface-agitated units are used for the diffused air model.

A.4.3.3 Land Treatment Model. The CHEMDAT7 land treatment model can predict emissions from land treatment soil, open landfills, and wastepiles. A general description of the land treatment model parameters is followed by specific guidelines for each application.

The land treatment model parameters for CHEMDAT7 are located in cells CV7 to CV18 as illustrated in Exhibit A-6. The oil loading must be obtained from the site operator or manually calculated offline as indicated in the report. It then is entered in CV7. The weight concentration of the VO (ppm by weight) (see CV8) is in the oil phase (for a two-phased liquid), in the water (for a dilute aqueous liquid), or in the liquid (for an organic liquid waste). The depth (CV9) is the depth of tilling in the land treatment facility in centimeters. The total porosity and the air porosity appear in CV10 and CV11, respectively. CV12 contains the average molecular weight of oil. In cell CV13, the value "1" is entered if the waste liquid

is a dilute aqueous solution or a "0" is entered for an (Raoult's law) organic waste or a two-phase (water + organic liquid) waste model. An intermediate time period over which emissions are to be calculated is specified in CV14 in units of days. Cell CV15 is an indicator for biological activity in the TSDF. When the user selects the land treatment model to predict emissions from land treatment soil, this flag is automatically set to 1. When the land treatment model is used to predict emissions from open landfills or wastepiles, biodegradation is assumed not to occur, and this indicator is automatically set to zero. The user may change this value if biodegradation is to be considered in the open landfill or waste-pile. The remaining three parameters have been described previously.

The values of these parameters in Exhibit A-6 estimate relative emissions from land treatment soil for benzene under the scenario detailed in Section 5.2.6.1. The estimated emissions for benzene presented in this section correspond to the CHEMDAT7 predictions in Exhibit A-6. Because an intermediate time period of 365.25 days was specified in CV14, the intermediate time emission fractions reflect annual estimates. The predicted annual fraction of benzene that will be emitted to the air is calculated as 0.90. This is equivalent to the example calculation result for land treatment facilities in Section 5.2.6.1.g. The long-term emission fraction for this example is equal to the intermediate result. Annual air emissions of benzene from the land treatment soil can be estimated from these results as follows:

Calculation

From Section 5.2.6.1:

Annual throughput: 1,800 Mg

Oil content of waste: 10 percent

Benzene concentration in oil: 2,000 ppm

$$E = 0.903 \times 1,800 \text{ Mg} \times 0.10 \times 0.002 = 0.33 \text{ Mg/yr}$$

Benzene emission from the land treatment facility is 0.33 Mg/yr.

Short-term emission results for the land treatment facility are also available from CHEMDAT7. Exhibit A-7 contains the estimated emission rates for benzene at 1/4, 1, 4, 12 and 24 hours after application to the land treatment soil.

Different tilling frequencies can also be accommodated by using the CHEMDAT7 land treatment model to predict emissions for each time period

defined by the tilling frequencies. This approach additionally requires that initial weight concentration of the VO (CV8) for each time period be adjusted for the estimated mass of the VO emitted or biodegraded during the previous time period. For each run of the model, the time period (in days) over which emissions are to be calculated is placed in CV14.

As indicated previously, the CHEMDAT7 land treatment model can be used to estimate emissions from open landfills. Exhibit A-13 contains the CHEMDAT7 land treatment model parameters and emission results that correspond to the example calculation in Section 6.4.3 for predicting the emission fraction of chloroform from open landfills. Because biodegradation is presumed not to occur in an open landfill, CV15 has been set to zero.

One year post-application, the fraction of chloroform lost to the atmosphere from open landfills is 0.053 from both CHEMDAT7 and the example calculation in Section 6.4.3.d. The following calculation demonstrates how to convert this emission fraction into an annual emission rate for an open landfill.

Calculation

From Section 6.4.3:

Landfill depth: 229 cm

Landfill area: $1.42 \times 10^8 \text{ cm}^2$

Loading: 0.46 g/cm^3

Weight fraction of chloroform in oil: 0.5

$$\begin{aligned} E &= 229 \text{ cm} \times 1.42 \times 10^8 \text{ cm}^2 \times 0.46 \text{ g/cm}^3 \times 0.5 \times 1 \text{ Mg}/10^6 \text{ g} \times 0.053 \\ &= 4.0 \times 10^2 \text{ Mg/yr.} \end{aligned}$$

Estimates of long-term emission and biological fractions from open landfills for chloroform are 1.0 and 0.0, respectively.

Finally, emissions from wastepiles also can be estimated via the CHEMDAT7 land treatment model under the restriction that the time parameter (CV14) must be less than or equal to the life of the wastepile. In the wastepile scenario discussed in Section 6.4.2.2, the retention time is 2.6 days. The user can estimate the emission fraction for any time period less than or equal to 2.6 days via CHEMDAT7 by placing the selected time period (expressed in days) in CV14. The average height of the wastepile, 100 cm, is placed in CV9. The remaining CHEMDAT7 input parameters would be identical to those in Exhibit A-13. The emission rate for the time period specified in CV14 is calculated from the resultant emission fraction as for

the open landfill. Multiply this value by 140, the estimated number of turnovers per year, to obtain the annual emission rate of chloroform for the wastepile. The user is reminded that, for open landfills and wastepiles, the biomass concentration in the land treatment model defaults to zero (CV15).

A.4.3.4 Disposal Impoundment Model. CHEMDAT7 disposal impoundment model input parameters are located in C046 through C056 of the spreadsheet. Please refer to Exhibit A-14. The concentration of V0 (C051) is the initial concentration in the waste expressed in milligrams/liter. The adsorbing biomass concentration (in C050) is the concentration available to remove the V0. This and the active biomass concentration (grams/liter in C049) are set to zero when adsorption and biodegradation are assumed to be nonexistent in a disposal impoundment. The remaining input parameters were previously defined.

The parameters specified in Exhibit A-14 reflect the example calculation for benzene emissions from a disposal impoundment described in Section 4.5.3. The CHEMDAT7 emission results also shown in this exhibit suggest that 14 percent of the benzene in the disposal impoundment will have been emitted to the air in the specified period of 6 months. Section 4.5.3 contains the same fractional result. With two turnovers per year, the corresponding emission rate is 0.04 Mg/yr.

The CHEMDAT7 disposal impoundment model can also estimate emissions from a diffused air system. This modification is achieved by defining a positive (nonzero) submerged air flow (m^3/second) in C055. If the user wishes to consider biodegradation as a pathway in the diffused air system, the concentration of active biomass (g/L) should be placed in C049.

A.4.3.5 Closed Landfill Model. Cells CV30 through CV52 in the CHEMDAT7 spreadsheet contain the closed landfill model input parameters; see Exhibit A-8. The user must first select the appropriate model. The options include the aqueous model (CV50=1) and the (Raoult's law) two-phased or organic liquid model (CV50=0). Additional CHEMDAT7 specifications are required for each of these models.

With the dilute aqueous model:

- The weight fraction oil (CV41) is set at zero (consequently, the sum of weight fraction water (CV42) and weight fraction V0 (CV43) must be one).

- MW-liquid (CV52) and rho-liquid (CV51) are used in the estimation procedure. Default values of 18 g/g mol and 1 g/cm³, respectively, appear in the software as information to the user. These values cannot be changed.
- mwt oil (CV45) does not contribute to the emission prediction.
- Liquid in waste (CV44) is set at 1 g/cm³ in accordance with dilute aqueous waste.

With the (Raoult's law) two-phased model:

- The values of MW-liquid (CV52) and rho-liquid (CV51) are ignored in the emission estimation process.

Finally, the closed landfill model can accommodate biodegradation. The input parameter CC/GVOC CONV (CV48) is the amount of gas produced in cubic centimeters per gram of VO biodegraded and is only applicable when biodegradation is considered a pathway. The user may wish to change the given representative value of CC/GVOC CONV to investigate other scenarios. The amount of active biomass (g/cc) is placed in CV49. The emissions then are augmented by those resulting from this additional pathway. Please note that the gas conversion input is ignored if biodegradation is presumed not to occur (i.e., when biomass in CV49 is set to zero). The remaining closed landfill model parameters are self-explanatory.

The values of the closed landfill model parameters in Exhibit A-8 correspond to those in the example calculations of chloroform emissions from a closed landfill (see Section 6.2.3). A two-phased model was used, and both the instantaneous emission rate after 1 year and the average emission rate in the first year were equivalent (13.0 Mg/yr of chloroform is the value for both rates from the sample calculations) to the estimated emission values from the CHEMDAT7 landfill model as shown in Exhibit A-8.

A.4.3.6 Oil Film Model. The CHEMDAT7 oil film model predicts emissions from an oil film surface impoundment or from an oil layer on land treatment soil. Input parameters for the oil film model are located in cells CV69 to CV78; see Exhibit A-15. Depth in meters (CV70) refers to the oil film thickness measured in meters. Area (CV71) is the surface area of the facility in square meters. Flow (CV72) is the rate of flow through a surface impoundment and, consequently, is not applicable when predicting emissions from an oil layer on land treatment soil. When there is no flow,

set CV72 to zero and enter the number of months for disposal in CV77. This latter input is needed to calculate residence time.

The input parameters specified in Exhibit A-15 reflect the example calculation for butanol-1 emissions from an oil layer on the soil surface of a land treatment site as described in Section 5.2.3.3. This exhibit shows the equivalent CHEMDAT7 emission result.

A.5 PRINTING A GRAPH

A.5.1 Graph Selection

Two types of graphs are available in CHEMDAT7. The first is a plot of emission prediction versus the partition coefficient; the second is a plot of emission prediction versus vapor pressure. Both graphs are available for all models except the oil film. Select a graph in the active command mode via

{/} {G} {N} {U} ,

where G represents graph, N represents name, and U represents use. (Recall that you can also position the cursor over the desired option and press {RETURN}.) Now select the desired graph. GRAPH-PC is the plot of emission prediction versus the partition coefficient, and GRAPH-VP is the plot of emission prediction versus vapor pressure. You can ignore the plot that appears on your screen. To return to the alternative command menu, press {ESC}, {QUIT}, {ALT M}.

A.5.2 Storing the Graph File

The sequence {PRINT} {GRAPH} followed by selection of the desired model will create the plot chosen via Section A.5.1 for the selected model and simultaneously store it in a specific file on the diskette (see Section A.3.1). CHEMDAT7 will store only one plot for each model. Furthermore, it stores the most recently created plot. Consequently, the user must first create the plot for which he wants a hard copy using the procedures described in this and the previous section.

A.5.3 The Hard Copy

To obtain a hard copy of a plot, access the active command mode; follow the commands to exit Lotus 1,2,3; and then access Lotus {PRINTGRAPH}. Once in this mode, {SELECT} plots for printing using the space bar to flag (#) the desired plots. The most recently created plot for closed landfill emissions is Q1; for land treatment emissions, Q2; for

nonaerated process emissions, Q3; for aerated process emissions, Q4; and for disposal impoundment emissions, Q5. Press {RETURN} and then {Go} to print the selected plots.

A.6 HELP SCREENS

Exhibits A-16 to A-22 present help screens that are available in CHEMDAT7. These help screens are provided to assist the user when other documentation may not be available.

TSDF (Section)

CHEMDAT7 Model

SURFACE IMPOUNDMENTS AND OPEN TANKS

Storage impoundments (4.2)
Treatment impoundments, mechanically
aerated systems (4.3)
Disposal impoundments (4.4)
Diffused air systems (4.5)

Oil film surfaces (4.6)
Diffused air-activated sludge

Nonaerated impoundment
Aerated impoundment
Disposal impoundment
Disposal impoundment
Disposal or aerated
impoundments
Oil film
Aerated impoundment

LAND TREATMENT

Land treatment soil (5.2)
Waste application (5.2)
Oil film surface (5.2)

Land treatment
NA
Oil film

LANDFILLS AND WASTEPILES

Closed landfills (6.2)
Fixation pits (6.3)
Open landfills (6.4)
Wastepiles (6.4)

Closed landfill
NA
Land treatment (modified)
Land treatment (modified)

TRANSFER, STORAGE, AND HANDLING OPERATIONS

Container loading (7.2)
Container storage (7.3)
Container cleaning (7.4)
Stationary tank loading (7.5)
Stationary tank storage (7.6)
Spills (7.7)
Fugitive emissions (7.8)
Vacuum truck loading (7.9)

NA
NA
NA
NA
NA
NA
NA
NA

NA = Not available in CHEMDAT7.

EXHIBIT A-1. RELATIONSHIP BETWEEN HAZARDOUS WASTE TSDF AND CHEMDAT7 MODELS

Selection

Options (explanation)

DATA-FORMS

(Go to data-entry forms)

IMPOUND (Go to the data entry form MENU for impoundments)

NON-AERATED (Go to the data entry form for flowthrough impoundments)

DIFFUSED (Go to the data entry form for diffused-air disposal)

FILM (Go to the data entry form for oil films)

DISPOSAL IMPOUNDMENTS (Go to the data entry form for disposal impoundments)

QUIT

AERATED (Go to the data entry form for activated sludge processes)

CLOSED-LANDFILL (Go to the data entry form for capped landfill)

OPEN-LF/WP (Go to the data entry form for open landfill/wastepile)

LAND-TREAT (Go to the data entry form for land application of wastes)

CONC (Enter concentration for each compound, permits different concentration for different compounds)

A IMPORT (Import compound data selected from the Master Data Set)

DEFAULT (Replace parameters with default parameters)

Municipal (municipal waste aerated impoundment parameters)

Aerated (default aerated model parameters)

Non-aerated (default non-aerated model parameters)

Landfill (default landfill model parameters)

Land-tr. (default land treatment model parameters)

VIEW

Open lf (set open landfill/wastepile to default)
Disposal (default disposal impoundment model parameters)
Quit (Return to data forms menu)
QUIT (Return to main menu)
(Go to a portion of the worksheet)
IMPOUND. (Go to the MENU for impoundments)
NON-AERATED (Go to the calculations for flowthrough impoundments)
DIFFUSED (Go to the calculations for diffused air disposal impoundments)
FILM (Go to the calculations for oil film impoundments)
DISPOSAL (Go to the calculations for disposal impoundments)
QUIT
AERATED (View results)
OPEN LF/WP (Use land treatment model to simulate an open landfill)
CLOSED-LF (View calculations for a closed landfill)
LAND-TREAT (View results)
TOTAL-EMISSIONS (View the results for long-term integrated emissions)
EMISSION-RATE (View the results for initial emission rates)
RETURN (Main menu)
GRAPH (VIEW GRAPH)
LANDFILL (View a graph)
LAND-TREAT (View a graph)

EXHIBIT A-2 (continued)

	NON-AERATED (View a graph)
	AERATED (View a graph)
	DISPOSAL (View a graph)
	QUIT
	K (View the mass transfer coefficients)
SORT	(Rearrange the order of the compound listings)
	SELECT (Sort by selected compounds)
	DATA (Select additional compounds from an extended data base)
	ALPHABETIC (Sort alphabetically)
	BIOLOGICAL (Sort in descending biological rates)
	CLASS (Sort by compound type)
	V-PRESSURE (SORT BY VAPOR PRESSURE)
	QUIT (RETURN TO MENU)
PRINT	(Print a portion of the worksheet)
	LAND-TREAT (Print land treatment, open landfill, or wastepile results)
	LONG-TERM (Print long-term environmental fate)
	SHORT-TERM (Print short-term emission estimates)
	DISPOSAL (Print disposal lagoon results)
	AERATED (Print aerated process results)
	LANDFILL (Print closed landfill results)
	NON-AER. (Print non-aerated impoundment results)
	K (Print mass transfer coefficients for selected facilities)
	DATA (Print data summaries)
	BIORATE (Print biorate data)
	LAND-TREAT (Print land treatment data)

	DATA (Print chemdat information)
	GRAPH (Put a graph file on the disk for later printing)
	FILM (Print thin film lagoon results)
SELECT	(Select which models are to be used)
HELP	(Look at an instructional screen)
	GENERAL (View general information)
	MODEL (Help in selecting the data form for your unit)
	DATA (Discussion of data entry)
	LANDTREAT (Information for the use of the land treatment models)
	LANDFILL (Information for the use of the landfill model)
	BIO-RATE (Information for the use of the biological reaction rates)
	IMPOUND (Information for the use of the impoundment models)
	QUIT (Return to the main menu)
QUIT	(Return to spreadsheet command mode)

EXHIBIT A-2. (continued)

NON-AERATED WASTEWATER TREATMENT

WINDSPEED	4.47 m/s
depth	1.8 m
AREA	1500 m ²
FLOW	0.00156 m ³ /s
VO inlet conc.	100 mg/l
TOTAL ORGANICS IN	250 mg/l
ACTIVE BIOMASS	0.05 g/l
BIOMASS SOLIDS IN	0 g/l
TEMPERATURE	25 deg. C

TOTAL AIR EMISSIONS 0.61 Mg/yr

COMPOUND NAME	EFFLUENT EMISS. BIOL.			PHOTOL.ADSORB. air & HYDRO.	emiss. (Mg/yr)	
BENZENE	0.031	0.124	0.845	0.000	0.001	0.6101

EXHIBIT A-3. NONAERATED MODEL INPUT PARAMETERS AND EMISSION RESULTS

AERATED WASTEWATER TREATMENT

WINDSPEED	4.47 m/s
DEPTH	1.8 m
AREA	1500 m2
FLOW	0.0031 m3/s
ACTIVE BIOMASS	0.25 g/l
BIOMASS SOLIDS IN	0 g/l
VO INLET CONC.	100 mg/l
TOTAL ORGANICS IN	250 mg/l
TOTAL BIORATE	19 mg/g bio-hr
FRACT. AGITATED	0.16
SUBMERGED AIR FLOW	0 m3/s
Number impellers	5
Oxygen trans. rat.	3 lbO2/h-hp
POWR (total)	75 HP
Power efficiency	0.83
Temperature	25 deg C
impeller dia	61 cm
impeller speed	126 rad/s

TOTAL AIR EMISSIONS 7.984 Mg/yr

COMPOUND NAME	RELATIVE AERATED WASTEWATER VOC PATHWAYS					emiss. (Mg/yr)
	EFFLUENT EMISS.	BIOL.	PHOTOL.	ADSORB. air & HYDRO.		
BENZENE	0.002	0.816	0.182	0.000	0.000	7.9842

EXHIBIT A-4. AERATED MODEL INPUT PARAMETERS AND EMISSION RESULTS
FOR MECHANICALLY AERATED TREATMENT IMPOUNDMENT

AERATED WASTEWATER TREATMENT

WINDSPEED	4.47 m/s
DEPTH	4 m
AREA	27 m2
FLOW	0.0075 m3/s
ACTIVE BIOMASS	4 g/l
BIOMASS SOLIDS IN	0 g/l
VO INLET CONC.	100 mg/l
TOTAL ORGANICS IN	250 mg/l
TOTAL BIORATE	19 mg/g bio-hr
FRACT. AGITATED	0.7
SUBMERGED AIR FLOW	0 m3/s
Number impellers	1
Oxygen trans. rat.	3 lbO2/h-hp
POWR (total)	7.5 HP
Power efficiency	0.83
Temperature	25 deg C
impeller dia	61 cm
impeller speed	126 rad/s

TOTAL AIR EMISSIONS 9.257 Mg/yr

COMPOUND NAME	RELATIVE AERATED WASTEWATER VOC PATHWAYS					emiss. (Mg/yr)
	EFFLUENT EMISS.	BIOL.	PHOTOL.	ADSORB. air & HYDRO.		
BENZENE	0.032	0.391	0.576	0.000	0.001	9.2572

EXHIBIT A-5. AERATED MODEL INPUT PARAMETERS AND EMISSION RESULTS FOR ACTIVATED SLUDGE UNIT

LAND TREATMENT MODEL DATA
 (land treatment)
 L>Loading (g oil/cc soil) 0.036
 Concentration in oil(ppmw) 2000
 l,Depth of tilling (cm) 20
 Total porosity 0.61
 Air Porosity(0 if unknown) 0.5
 MW oil 282
 For aqueous waste, enter 1 0
 Time of calc. (days) 365.25
 For biodegradation,enter 1 1
 Temperature (Deg. C) 25
 Wind Speed (m/s) 4.47
 Area (m2) 25000

COMPOUND NAME	LANDTREATMENT FRACTION LOST		INTERMEDIATE TIME 365.25 days	
	AIR	BIOL.	AIR	BIOL.
BENZENE	0.903	0.097	0.903	0.097

EXHIBIT A-6. LAND TREATMENT MODEL INPUT PARAMETERS AND EMISSION
 RESULTS FOR LAND TREATMENT SOIL

LAND TREATMENT MODEL DATA

(land treatment)
 L, Loading (g oil/cc soil) 0.036
 Concentration in oil (ppmw) 2000
 l, Depth of tilling (cm) 20
 Total porosity 0.61
 Air Porosity (0 if unknown) 0.5
 MW oil 282
 For aqueous waste, enter 1 0
 Time of calc. (days) 365.25
 For biodegradation, enter 1 1
 Temperature (Deg. C) 25
 Wind Speed (m/s) 4.47
 Area (m2) 25000

COMPOUND NAME	LAND TREATMENT EMISSION RATES (g/cm2-s)				
	TIME (hours)				
	0.25	1	4	12	48
BENZENE	2.89E-08	1.46E-08	7.28E-09	4.16E-09	1.98E-09

EXHIBIT A-7. LAND TREATMENT MODEL INPUT PARAMETER FOR OPEN LANDFILLS AND WASTEPILES

CLOSED LANDFILL

A, LF area (cm2)	1.4170E+08
l, cap thickness (cm)	107
ea, cap air porosity	0.08
et, cap total porosity	0.41
T, avg. LF temp. (C)	15
efw, waste porosity	0.25
Pref, ref. pressure (mbar)	1013
Pl, new pressure (mbar)	1009
Tref, reference temp. (C)	15
Tl, new temp. (C)	15
dt, bar.pump time int.(s)	86400
weight fraction oil	0.2
weight fraction water	0.6
weight fraction VO	0.2
W, liquid in waste (g/cm3)	1.16
mwt, oil	147
t, time of calc. (mo.)	12
D, depth of waste (cm)	457
CC/GVOC CONV	1750
active biomass (g/cc)	0.00E+00
VO diss. in water, enter 1	0
rho-liquid, density (g/cm3)	1
MW-liquid, (g/g mol)	18

CLOSED LANDFILL AIR EMISSIONS		EMISSION RATES	
COMPOUND NAME	EMISS. FRACTION	BIOL. FRACTION	12 months AVERAGE INST.
CHLOROFORM	1.000	0.000	12.99431 12.98869

EXHIBIT A-8. CLOSED LANDFILL MODEL INPUT PARAMETERS AND EMISSION RESULTS

<u>Column</u>	<u>Column label [explanation]</u>
B	COMPOUND NAME
C	COMPOUND TYPE
D	M.W. [molecular weight]
E	DENSITY (g/cc)
F	VAP.PRESS. (mmHg) [vapor pressure at 25 °C]
G	H LAW CONST (atm•m ³ /mol) [Henry's law constant]
H	DIFF. WAT (cm ² /s) [diffusion coefficient in water]
I	DIFF. AIR (cm ² /s) [diffusion coefficient in air]
J	BOILING POINT (°C)
K	VAPOR PRESSURE COEFFICIENT - A
L	VAPOR PRESSURE COEFFICIENT - B
M	VAPOR PRESSURE COEFFICIENT - C
V	K(Y/X) [Henry's law coefficient, mol fraction] (calculated)
W	LAND-TREATMENT BIORATE (DAY ⁻¹) (calculated)
N	LOG OCT/WATER PARTITION
O	BIORATE (mg/g-hr)

EXHIBIT A-9. LIST OF COLUMN LABELS IN CHEMDAT7

Column	Column label [explanation]	
P	K1 L/g-hr.	
Q	HYDROL (s-1) [rate const. photolysis] + [rate const. hydrolysis]	
X	ADS. (m ³ /kg solids) [adsorption]	
Y	DISPOSAL IMPOUNDMENT - total fraction removed	
Z	OIL FILM - emissions fraction	
AA	VP in landfill (mm Hg	
AB	ln(K)	
AC	temp correction factor	
AD	K _S (g/cm ³ per g/cm ³) [partition factor, V _O into sludge]	
AE	Biorate (hr ⁻¹)	NON-AERATED
AF	K _O (g mol/cm ² -s) = [overall mass transfer coefficient (m/s)] x 5.56	↓
AG	Co (Mg/yr)	AERATED
AH	Temperature correction CS83	↓
AI	biorate (s-1)	↓
AJ	K _O (g mol/cm ² -s) = [overall mass transfer coefficient (m/s)] x 5.56	↓
AK	Co(mg/L)	↓
AL	ADS. COEF. (VAP/SOL) [adsorption coefficient]	↓
AM	BIO RATES (1/tb s-1) [biological rates]	↓
AN	DECAY PARAMETER	↓
AO	EMISS. FRACTION [emission fraction for specified period]	CLOSED LANDFILL
AP	BIOL. FRACTION [biological removal fraction for specified period]	↓
AQ	AVERAGE EMISSION RATE (Mg/specified time period)	↓

EXHIBIT A-9 (continued)

<u>Column</u>	<u>Column label [explanation]</u>	
AR	INSTANTANEOUS EMISSION RATE (Mg/specified time period)	
AS	EMISS. [long-term emission fraction]	LAND TREATMENT
AT	BIOL. [long-term biological removal fraction]	↓
AU	EMISS. [intermediate emission fraction]	
AV	BIOL. [intermediate biological removal fraction]	
AW	EFFLUENT [annual effluent fraction]	NON-AERATED
AX	EMISS. [annual emission fraction]	↓
AY	BIOL. [annual biological removal fraction]	
AZ	PHOTOL. & HYDROL. [annual fraction removed via photolysis and hydrolysis]	
BA	ADSORB. [annual fraction removed via adsorption]	
BB	AIR EMISSIONS (Mg/year)	
BC	EFFLUENT [annual effluent fraction]	AERATED
BD	EMISS. [annual emission fraction]	↓
BE	BIOL. [annual biological removal fraction]	
BF	PHOTOL. & HYDROL. [annual fraction removed via photolysis and hydrolysis]	
BG	ADSORB. [annual fraction removed via adsorption]	
BH	AIR EMISSIONS (Mg/year)	
BI	AQUEOUS [fraction remaining]	DISPOSAL IMPOUNDMENT
BJ	EMISS. [emission fraction for specified period]	↓
BK	BIOL. [biological removal fraction for specified period]	

EXHIBIT A-9 (continued)


<u>Column</u>	<u>Column label [explanation]</u>	
BL	PHOTOL. & HYDROL. [fraction removed via photolysis and hydrolysis for specified period]	
BM	ADSORB [fraction removed via adsorption for specified period]	
BN	Air emission (Mg/yr)	
BO	VP aqueous landfill	
BP	K' aerated	
BQ	K_0 (g mol/cm ² -s) = [overall mass transfer coefficient (m/s)] x 5.56	
BR	SUM RATES DISPOSAL [sum of the various rate processes]	DISPOSAL IMPOUNDMENT
BS	lambda	
BT	CONC (g/cm ³)	
BU	K' non-aerated	

EXHIBIT A-9 (continued)

COMPOUND NAME	M.W.	DENSITY (g/cc)	VAP.PRESS (mmHg)	H LAW CONST. (atm-m3/mol)	DIFF. MAT (cm2/sec)
ACETALDEHYDE	44.0	0.79	768	9.50E-05	1.41E-05
ACETONE	58.0	0.79	286	2.50E-05	1.14E-05
ACROLEIN	56.1	0.84	244.2	5.66E-05	1.22E-05
ACRYLONITRILE	53.1	0.81	114	8.80E-05	1.34E-05
ALLYL ALCOHOL	58.1	0.85	23.3	1.80E-05	1.14E-05
BENZENE	78.1	0.87	95.2	5.50E-03	9.80E-06
BENZYL CHLORIDE	126.6	1.10	1.21	4.33E-04	7.80E-06
BUTANOL-1	74.1	0.81	6.5	8.90E-06	9.30E-06
CARBON DISULFIDE	76.1	1.26	366	1.68E-02	1.00E-05
CARBON TETRACHLORIDE	153.8	1.59	113	3.00E-02	8.80E-06
CHLOROBENZENE	112.6	1.11	11.8	3.93E-03	8.70E-06
CHLOROFORM	119.4	1.49	208	3.39E-03	1.00E-05
CHLOROPRENE	88.5	0.958	273	0.331	1.00E-05
CRESOL(-m)	108.1	1.03	0.08	4.43E-07	1.00E-05
CRESOL(-o)	108.1	1.05	0.24	2.60E-06	8.30E-06
CRESOL(-p)	108.1	1.03	0.11	4.43E-07	1.00E-05
CRESYLIC ACID	108.0	1.05	0.3	1.70E-06	8.30E-06
CUMENE (isopropylbenzene)	120.2	0.86	4.6	1.46E-02	7.10E-06
CYCLOHEXANONE	98.2	0.95	4.8	4.13E-06	8.62E-06
DICHLOROBENZENE(1,2) (-o)	147.0	1.31	1.5	1.94E-03	7.90E-06
DICHLOROBENZENE(1,4) (-p)	147.0	1.29	1.2	1.60E-03	7.90E-06
DICHLOROETHANE(1,2)	99.0	1.26	80	1.20E-03	9.90E-06
DICHLOROPROPANE(1,2)	113.0	1.16	40	2.30E-03	8.73E-06
DIMETHYL NITROSAMINE	74.08	1.005			1.00E-05
DIOXIN	322.0	1.83	1.52E-09	8.12E-05	5.60E-06
EPICHLOROHYDRIN	92.5	1.18	17	3.23E-05	9.80E-06
ETHYLACETATE	88.1	0.90	100	1.28E-04	9.66E-06
ETHYLBENZENE	106.2	0.87	10	6.44E-03	7.80E-06
ETHYLENEOXIDE	44.0	0.87	1250	1.42E-04	1.45E-05
ETHYLETHER	74.1	0.71	520	6.80E-04	9.30E-06
FORMALDEHYDE	30.0	0.00	3500	9.76E-05	1.98E-05
FREONS	120.92	1.486	5000	4.01E-01	1.00E-05
HEXACHLOROBUTADIENE	260.8	1.67	0.15	2.56E-02	6.16E-06
HEXACHLOROCYCLOPENTADIENE	272.8	1.71	0.081	1.60E-02	6.16E-06
HNHS	98.4		1	5.38E-03	9.40E-06
HNLS	144.0		1	3.00E-02	9.39E-06
HNMS	117		1	1.18E-03	8.24E-06
ISOBUTYL ALCOHOL	74.0	0.79	10	2.20E-06	9.30E-06
LHNS	97.3		1	1.58E-07	9.64E-06
LHMS	78.4		1	1.58E-07	1.13E-05
MALEIC ANHYDRIDE	98.1	0.93	0.0001	4.00E-08	1.11E-05
METHANOL	32.0	0.79	114	2.70E-06	1.64E-05
METHYL ACETATE	74.1	0.92	235	1.02E-04	1.00E-05
METHYL ETHYL KETONE	72.1	0.82	100	4.35E-05	9.80E-06
METHYL ISOBUTYL KETONE	100.2	0.80	15.7	4.95E-05	7.80E-06
METHYLENE CHLORIDE	85.0	1.34	438	3.19E-03	1.17E-05
METHYLNAPHTHALENE (1)	142.2				
HNHS	69.9		1	6.80E-05	1.16E-05
HNLS	112.0		1	2.22E-05	8.60E-06
HNMS	57		1	4.08E-05	1.18E-05
NAPHTHALENE	128.2	1.14	0.23	1.18E-03	7.50E-06
NITROBENZENE	123.1	1.20	0.3	1.31E-05	8.60E-06
NITROSOMORPHOLINE	116.14				1.00E-05
PHENOL	94.1	1.07	0.341	4.54E-07	9.10E-06
PHTHALIC ANHYDRIDE	148.1	1.33	0.0015	9.00E-07	8.60E-06
POLYCHLORINATED BIPHENYLS	290	1.45		2.94E-04	1.00E-05
PROPYLENE OXIDE	58.1	0.83	524.5	1.34E-03	1.00E-05
PYRIDINE	79.1	0.98	20	2.36E-05	7.60E-06
TETRACHLOROETHANE(1,1,2,2)	168.0	1.59	6.5	3.80E-04	7.90E-06
TETRACHLOROETHYLENE	165.83	1.624	19	2.90E-02	8.20E-06
TOLUENE	92.4	0.87	30	6.68E-03	8.60E-06
TRICHLOROETHANE(1,1,1)	133.4	1.33	123	1.72E-02	8.90E-06
TRICHLOROETHANE(1,1,2)	133.4	1.3	25	7.40E-04	8.80E-06
TRICHLOROETHYLENE	131.4	1.40	75	9.10E-03	9.10E-06
TRICHLOROFLUOROMETHANE	137.4	1.49	796	5.83E-02	9.70E-06
TRICHLORO(1,1,2)TRIFLUORO	187.38	1.58	300	4.35E-01	8.20E-06
VINYL CHLORIDE	62.5	0.91	2660	8.60E-02	1.23E-05
VINYLDIENE CHLORIDE	97.0	1.21	591	1.50E-02	1.04E-05
XYLENE(-o)	106.2	0.88	7	5.27E-03	1.00E-05

EXHIBIT A-10. ALPHABETICAL LIST OF COMPOUNDS IN CHEMDAT7

COMPOUND NAME	DIFF. AIR (cm ² /sec)	BOILING POINT (deg.C)	VAPOR PRESSURE COEFFICIENTS		
			A	B	C
ACETALDEHYDE	1.24E-01	20.8	8.005	1600.017	291.809
ACETONE	1.24E-01	56.2	7.117	1210.595	229.664
ACROLEIN	1.05E-01	53.0	2.387833		
ACRYLONITRILE	1.22E-01	77.4	7.038	1232.53	222.47
ALLYL ALCOHOL	1.14E-01	97.0	1.367406		
BENZENE	8.80E-02	80.1	6.905	1211.033	220.79
BENZYL CHLORIDE	7.50E-02	179.4	0.082788		
BUTANOL-1	8.00E-02	117.7	7.476	1362.39	178.77
CARBON DISULFIDE	1.04E-01	46.3	6.942	1169.11	241.59
CARBON TETRACHLORIDE	7.80E-02	76.8	6.9339	1242.43	230
CHLOROBENZENE	7.30E-02	132.0	6.978	1431.05	217.55
CHLOROFORM	1.04E-01	61.2	6.493	929.44	196.03
CHLOROPRENE	1.04E-01	59.4	6.161	783.45	179.7
CRESOL (-m)	7.40E-02	202.0	7.508	1856.36	199.07
CRESOL (-o)	7.40E-02	190.8	6.911	1435.5	165.16
CRESOL (-p)	7.40E-02	203.0	7.035	1511.08	161.85
CRESYLIC ACID	7.40E-02	235.0	-0.52289		
CUMENE (isopropylbenzene)	6.50E-02	153.0	6.963	1460.793	207.78
CYCLOHEXANONE	7.84E-02	157.0	0.681266		
DICHLOROBENZENE(1,2) (-o)	6.90E-02	179.0	0.176097		
DICHLOROBENZENE(1,4) (-p)	6.90E-02	173.4	0.079184		
DICHLOROETHANE(1,2)	1.04E-01	83.4	7.025	1272.3	222.9
DICHLOROPROPANE(1,2)	7.82E-02	96.8	6.98	1380.1	22.8
DIMETHYL NITROSAMINE	1.04E-01	153	ERR		
DIOXIN	1.04E-01	421.4	12.88	6465.5	273
EPICHLOROMYDRIN	8.60E-02	117.0	1.230494		
ETHYLACETATE	7.32E-02	77.0	7.101	1244.95	217.88
ETHYLBENZENE	7.50E-02	136.2	6.975	1424.255	213.21
ETHYLENEOXIDE	1.04E-01	10.7	7.128	1054.54	237.76
ETHYLETHER	7.40E-02	34.5	6.92	1064.07	228.8
FORMALDEHYDE	1.78E-01	-14.0	7.195	970.6	244.1
FREONS	1.04E-01	-29.8	3.699106		
HEXACHLOROBUTADIENE	5.61E-02	215.0	-0.82393		
HEXACHLOROCYCLOPENTADIENE	5.61E-02	234.0	-1.09155		
HHMB	8.73E-02				
HHLB	8.76E-02				
HHMB	7.40E-02				
ISOBUTYL ALCOHOL	8.60E-02	107.9	1.000036		
LHMB	8.27E-02				
LHMB	1.00E-01				
MALEIC ANHYDRIDE	9.50E-02	200.0	-4.00014		
METHANOL	1.50E-01	65.0	7.897	1474.08	229.13
METHYL ACETATE	1.04E-01	54.0	7.065	1157.63	219.73
METHYL ETHYL KETONE	8.08E-02	79.6	6.97421	1209.6	216
METHYL ISOBUTYL KETONE	7.50E-02	115.8	6.672	1168.4	191.9
METHYLENE CHLORIDE	1.01E-01	39.8	7.409	1325.9	252.6
METHYLNAPHTHALENE (1)					
HHMB	9.50E-02				
HHLB	7.60E-02				
HHMB	1.15E-01				
NAPHTHALENE	5.90E-02	218.0	7.01	1733.71	201.86
NITROBENZENE	7.60E-02	210.8	7.115	1746.6	201.8
NITROSONORPHOLINE	5.90E-02	225	ERR		
PHENOL	8.20E-02	182.0	7.133	1516.79	174.95
PHTHALIC ANHYDRIDE	7.10E-02	284.0	8.022	2868.5	0
POLYCHLORINATED BIPHENYLS	1.04E-01		ERR		
PROPYLENE OXIDE	1.04E-01	33.9	2.719846		
PYRIDINE	9.10E-02	115.5	7.041	1373.8	214.98
TETRACHLOROETHANE(1,1,2,2)	7.10E-02	146.2	6.631	1228.1	179.9
TETRACHLOROETHYLENE	7.20E-02	121.4	6.976	1386.92	217.53
TOLUENE	8.70E-02	110.7	6.954	1344.8	219.48
TRICHLOROETHANE(1,1,1)	7.80E-02	75.0	8.643	2136.6	302.8
TRICHLOROETHANE(1,1,2)	7.80E-02	74	6.951	1314.4	209
TRICHLOROETHYLENE	7.90E-02	87.0	6.518	1018.6	192.7
TRICHLOROFLUOROMETHANE	8.70E-02	23.8	6.884	1043.004	236.88
TRICHLORO(1,1,2)TRIFLUORO	7.80E-02	48.0	6.88	1099.9	227.5
VINYL CHLORIDE	1.06E-01	-13.9	3.425008		
VINYLDIENE CHLORIDE	9.00E-02	31.9	6.972	1099.4	237.2
XYLENE(-o)	8.70E-02	144.4	6.998	1474.679	213.69

EXHIBIT A-10 (continued)

COMPOUND NAME	BIORATE mgv0/g-hr	K1 L/g-hr.
ACETALDEHYDE	82.42	0.196681
ACETONE	1.30	1.15
ACROLEIN	7.80	0.34
ACRYLONITRILE	18.00	0.75
ALLYL ALCOHOL	0.00	0.156655
BENZENE	19.00	1.4
BENZYL CHLORIDE	17.75	1.010395
BUTANOL-1	7.80	0.11
CARBON DISULFIDE	0.00	1
CARBON TETRACHLORIDE	1.50	1.5
CHLOROBENZENE	0.39	10
CHLOROFORM	2.94	0.79
CHLOROPRENE	0.00	1
CRESOL (-o)	23.21	17
CRESOL (-o)	22.78	17
CRESOL (-p)	23.21	17
CRESYLIC ACID	15.00	1
CUMENE (isopropylbenzene)	0.00	1
CYCLOHEXANONE	11.49	0.274276
DICHLOROBENZENE(1,2) (-o)	2.50	0.58
DICHLOROBENZENE(1,4) (-p)	6.40	2.3
DICHLOROETHANE(1,2)	2.10	0.98
DICHLOROPROPANE(1,2)	17.00	1.4
DIMETHYL NITROSAMINE	0.00	0.081977
DIOXIN	0.00	1
EPICHLOROHYDRIN	0.00	0.138991
ETHYLACETATE	17.58	1
ETHYLBENZENE	6.80	2.1
ETHYLENEOXIDE	4.20	0.91
ETHYLETHER	0.77	0.279119
FORMALDEHYDE	5.00	0.25
FREONS	0.00	1
HEXACHLOROBUTADIENE	0.00	3.562786
HEXACHLOROCYCLOPENTADIENE	0.00	4.434115
HHB	29.2	1
HHB	0.30	1
HHB	2.71	1
ISOBUTYL ALCOHOL	7.80	0.11
LHB	23.2	1
LHB	3.55	1
MALEIC ANHYDRIDE	4.08	1
METHANOL	18.00	0.2
METHYL ACETATE	19.87	0.124775
METHYL ETHYL KETONE	2.00	0.2
METHYL ISOBUTYL KETONE	0.74	0.451677
METHYLENE CHLORIDE	22.00	0.403106
METHYLNAPHTHALENE (1)	24.03	1
HHB	40.1	1
HHB	0.39	1
HHB	11.2	1
NAPHTHALENE	42.47	1
NITROBENZENE	11.00	2.3
NITROSONORPHOLINE	0.00	0.091855
PHENOL	97.00	13
PHTHALIC ANHYDRIDE	0.00	0.078467
POLYCHLORINATED BIPHENYLS	0.00	1
PROPYLENE OXIDE	0.00	1
PYRIDINE	35.03	0.238439
TETRACHLOROETHANE(1,1,2,2)	6.20	0.68
TETRACHLOROETHYLENE	6.20	0.68
TOLUENE	73.48	2.4
TRICHLOROETHANE(1,1,1)	3.50	0.74
TRICHLOROETHANE(1,1,2)	3.50	0.74
TRICHLOROETHYLENE	3.90	0.88
TRICHLOROFLUOROMETHANE	0.00	1.235682
TRICHLORO(1,1,2)TRIFLUORO	0.00	1
VINYL CHLORIDE	0.00	0.142278
VINYLDENE CHLORIDE	0.00	1
XYLENE(-o)	40.79	1.784578

EXHIBIT A-10 (continued)

NON-AERATED WASTEWATER TREATMENT

WINDSPEED 4.47 m/s
 depth 1.8 m
 AREA 1500 m2
 FLOW 0.00156 m3/s
 VO inlet conc. 100 mg/l
 TOTAL ORGANICS IN 250 mg/l
 ACTIVE BIOMASS 0 g/l
 BIOMASS SOLIDS IN 0 g/l
 TEMPERATURE 25 deg. C

TOTAL AIR EMISSIONS 3.94 Mg/yr

COMPOUND NAME	EFFLUENT EMISS.	BIOL.	PHOTOL.	ADSORB.	air & HYDRO. emiss. (Mg/yr)
BENZENE	0.199	0.801	0.000	0.000	0.000 3.9432

EXHIBIT A-11. NONAERATED MODEL INPUT PARAMETERS AND EMISSION RESULTS

AERATED WASTEWATER TREATMENT

WINDSPEED	4.47 m/s
DEPTH	4 m
AREA	27 m2
FLOW	0.0075 m3/s
ACTIVE BIOMASS	4 g/l
BIOMASS SOLIDS IN	0 g/l
VO INLET CONC.	100 mg/l
TOTAL ORGANICS IN	250 mg/l
TOTAL BIORATE	19 mg/g bio-hr
FRACT. AGITATED	0.7
SUBMERGED AIR FLOW	0.04 m3/s
Number impellers	1
Oxygen trans. rat.	3 lbO2/h-hp
POWR (total)	7.5 HP
Power efficiency	0.83
Temperature	25 deg C
impeller dia	61 cm
impeller speed	126 rad/s

TOTAL AIR EMISSIONS 9.745 Mg/yr

COMPOUND NAME	RELATIVE AERATED WASTEWATER VOC PATHWAYS					emiss. (Mg/yr)
	EFFLUENT EMISS.	BIOL.	PHOTOL.	ADSORB. & HYDRO.	air	
BENZENE	0.030	0.412	0.557	0.000	0.001	9.7448

EXHIBIT A-12. AERATED MODEL INPUT PARAMETERS AND EMISSION RESULTS FOR DIFFUSED AIR-ACTIVATED SLUDGE UNIT

LAND TREATMENT MODEL DATA
 (open landfill, waste pile)
 L, Loading (g oil/cc soil) 0.464
 Concentration in oil(ppmw) 500000
 l, Depth of tilling (cm) 229
 Total porosity 0.5
 Air Porosity(0 if unknown) 0.25
 MW oil 147
 For aqueous waste, enter 1 0
 Time of calc. (days) 365.25
 For biodegradation, enter 1 0
 Temperature (Deg. C) 25
 Wind Speed (m/s) 4.47
 Area (m2) 14200

COMPOUND NAME	LANDTREATMENT FRACTION LOST		INTERMEDIATE TIME 365.25 days	
	AIR	BIOL.	AIR	BIOL.
CHLOROFORM	1.000	0.000	0.053	0.000

EXHIBIT A-13. LAND TREATMENT MODEL INPUT PARAMETERS AND EMISSION
 RESULTS FOR OPEN LANDFILLS AND WASTEPILES

DISPOSAL IMPOUNDMENT
(no outlet flow)

WINDSPEED	4.47 m/s
DEPTH	1.8 m
AREA	9000 m ²
ACTIVE BIOMASS	0.05 g/l
BIOMASS SOLIDS IN	0 m ³ /s
VO INLET CONC.	100 mg/l
TOTAL ORGANICS IN	250 mg/l
TOTAL BIORATE	19 mg/g bio-hr
TIME FOR EMISSIONS	6 months
SUBMERGED AIR FLOW	0 m ³ /s
TEMPERATURE	25 deg. C

TOTAL AIR EMISSIONS 1.77E+00 Mg/yr

COMPOUND NAME	DISPOSAL IMPOUND. VO EMISSIONS 6 MONTHS				air emiss. (Mg/yr)	
	AQUEOUS	EMISS. BIOL.	PHOTOL. ADSORB. & HYDRO.			
BENZENE	0.000	0.546	0.454	0.000	0.000	1.769547

EXHIBIT A-14. DISPOSAL IMPOUNDMENT INPUT PARAMETERS AND EMISSION RESULTS

OIL FILM SURFACE

WINDSPEED	(m/s)	4.47
depth	(m)	0.072
AREA	(m ²)	25000
FLOW	(m ³ /s)	0
VO conc. in oil	(mg/l)	200
oil (fraction of waste)		1
molecular weight of oil		282
density of oil (g/cc)		1
Enter months for disposal		0.033
Temperature (deg C)		25

COMPOUND NAME	OIL FILM emissions fraction
BUTANOL-1	0.494451

EXHIBIT A-15. OIL FILM SURFACE MODEL INPUT PARAMETERS AND EMISSION RESULTS

Y99: 'GENERAL HELP : IF THESE HELP SCREENS DO NOT ANSWER YOUR QUEST MENU
GENERAL MODEL DATA LANDTREAT LANDFILL BIO-RATE IMPOUND QUIT

View general information

 Y Z AA AB AC AD AE
99 GENERAL HELP : IF THESE HELP SCREENS DO NOT ANSWER YOUR QUESTIONS,
100 CONSULT THE MANUAL.

101 for ...

102

103 all compounds same concent. enter concentration on data form.

104 compounds different concent. enter values using CONC on DATA-FORM

105 select compounds use the SORT menu option

106 which data form to use? press M for model help screen

107 General use of spreadsheet Consult your Lotus 123 manual.

108

109 New compounds can be added by copying other data over the existing

110 compound data base. [b14..w82] See data help screen.

111 Go to a cell for data entry by pressing [F5] and typing cell label

112 New compounds can be added by selecting [IMPORT]. See DATA-FORM menu.

113 New compounds can be added by selecting [SORT][DATA].

114 hold {ALT} down and press {O} to restore borders for screens.

115

116 calculate: press {F9} before you use the predictions!

117 last use:

118 20-Mar-89

20-Mar-89 02:51 PM

VER 7.06 3-20-89 cca

CMD CALC

EXHIBIT A-16. GENERAL HELP SCREEN

Y139:	Model Assistance Help Screen						MENU
GENERAL	<u>MODEL</u>	DATA	LANDTREAT	LANDFILL	BIO-RATE	IMPOUND	QUIT
Help in selecting the data form for your unit							
	Y	Z	AA	AB	AC	AD	AE
139	Model Assistance Help Screen						
140							
141	to model..						
142							
143	Treatment Tanks	Use the non-aerated model.					
144	Lagoon sludges	Use the landtreatment model.					
145	Open waste piles	Use the landtreatment model.					
146	For municipal waste treatment assume that approximately 1% of flow						
147	is inlet solids by volume.						
148							
149	Disposal	for non flow system you specify time for calc.					
150	Non-aerated and aerated impoundments are flow through systems at						
151	steady state conditions.						
152	Plug flow lagoon	Use disposal model: first order kinetics are					
153	assumed for a batch. The equations for plug flow						
154	can be simulated by non-flow, since there is						
155	no backmixing. Emission time is residence time.						
156	Plug flow aeration	Not supported since strong backmixing is assumed.					
157	You can run multiple times for series reactors.						
158	Use conc. out for inlet conc. for next reactor.						
20-Mar-89	02:52 PM			CMD		CALC	

EXHIBIT A-17. MODEL ASSISTANCE HELP SCREEN

AV138: 'DATA ENTRY CONCERNS: CONSULT MANUAL FOR ADDITIONAL DETAILS MENU
 GENERAL MODEL DATA LANDTREAT LANDFILL BIO-RATE IMPOUND QUIT
 View general information
 AV AW AX AY AZ BA BB BC BD
 138 DATA ENTRY CONCERNS: CONSULT MANUAL FOR ADDITIONAL DETAILS
 139
 140 New compounds can be added by copying other data over the existing
 141 compound data base. [b14..w82]
 142 Sometimes data omissions cause errors. The following are needed:
 143
 144 COLUMNS OF DATA REQUIRED
 145 Landfills E F G I K L M O
 146 landtreatment F G I K L M N O
 147 non aerated G H I K L M N O P
 148 aerated G H I K L M N O P
 149 disposal G H I K L M N O P
 150 oil film F I K L M
 151 impoundment G H I O P
 152
 153 Where no data is available and the pathway is not evaluated, non-zero
 154 values can be inserted and the effect of the value is canceled in data
 155 entry. Example: put any positive value for O and P and set biomass=0.
 156
 157 column [C]: A is aromatic, N nitrogen, O oxygenate, C chlorinated, P phenol
 20-Mar-89 02:53 PM CMD CALC

EXHIBIT A-18. DATA ENTRY HELP SCREEN

AI118: 'LANDTREATMENT HELP SCREEN: CONSULT MANUAL FOR ADDITIONAL DETAILS. MENU
GENERAL MODEL DATA LANDTREAT LANDFILL BIO-RATE IMPOUND QUIT

View general information

AI AJ AK AL AM AN AO

118 LANDTREATMENT HELP SCREEN: CONSULT MANUAL FOR ADDITIONAL DETAILS.

119

120 The model used simulates the diffusion of a gas out of a porous
121 solid. The rate of diffusion depends on the concentration in the
122 gas phase. The concentration in the gas phase is reduced by
123 absorption in the oil which is mixed with the porous solid. The
124 wind resistance to diffusion is assumed to be unimportant for the
125 long term emissions.

126

127

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EXHIBIT A-19. LAND TREATMENT HELP SCREEN

AP118: 'LANDFILL HELP SCREEN: CONSULT MANUAL FOR ADDITIONAL DETAILS
GENERAL MODEL DATA LANDTREAT LANDFILL BIO-RATE IMPOUND QUIT

MENU

View general information

AP AQ AR AS AT AU AV

118 LANDFILL HELP SCREEN: CONSULT MANUAL FOR ADDITIONAL DETAILS

119

120 The landfill model is based upon air emissions from these
121 pathways:

122

123 diffusion through cap

124 gas generation by biodegradation

125 barometric pumping

126

127 Open landfills can be modeled using the equations
128 in the landtreatment model.

129

130

131

132

133

134

135

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137

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EXHIBIT A-20. LANDFILL HELP SCREEN

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AW118:      'BIORATE HELP SCREEN FOR AQUEOUS SYSTEMS
GENERAL MODEL DATA LANDTREAT LANDFILL BIO-RATE IMPOUND QUIT
View general information
      AW      AX      AY      AZ      BA      BB      BC      BD      BE
118 BIORATE HELP SCREEN FOR AQUEOUS SYSTEMS
119
120 The rate of biooxidation is estimated by the Monod kinetic model.
121   The value of K1 is specified in column V14...V84.
122   If K1 is not estimated empirically, there is an equation
123   based upon the octanol- water partition coefficient.
124   If the octanol-water partition coefficient is absent, K1 is
125   assumed to equal 1.
126 The value of Kv is determined empirically and is identical to the
127 previously used biorate constants in CHEMDAT6.
128
129      [biomass] [ Kv ] [concentration]
130 RATE = -----
131      [concentration] + [ Kv/K1 ]
132
133 The effect of temperature on the rate is as follows:
134      ( 1.046 )^(deg C-25deg)
135
136
137
20-Mar-89  02:55 PM                      CMD          CALC

```

MENU

EXHIBIT A-21. BIORATE HELP SCREEN

BF118: 'IMPOUNDMENTS HELP SCREEN: FOR ADDITIONAL DETAILS CONSULT THE MENU
 GENERAL MODEL DATA LANDTREAT LANDFILL BIO-RATE IMPOUND QUIT

View general information

BF BG BH BI BJ BK BL BM BN

118 IMPOUNDMENTS HELP SCREEN: FOR ADDITIONAL DETAILS CONSULT THE MANUAL.

119 The flow in the aerated and non-aerated impoundments is assumed to be
 120 well mixed. Plug flow can reduce effluent concentrations, but may not
 121 significantly change fraction emitted.

122 If only the exit concentration is known, the inlet may be estimated
 123 by trial and error.

124 Aerated processes can describe impoundments as well as processes.

125

126 Both surface aeration and submerged aeration are available at the same
 127 time. Submerged aeration is expected to agitate the surface in a
 128 similar manner to surface aeration. If there is no agitated surface
 129 available for air transport, the fract. agitated can be set to 0.

130

131 The component properties of mass transfer are based on the ratio to
 132 a reference compound. The following exponents are used with the ratios.

133

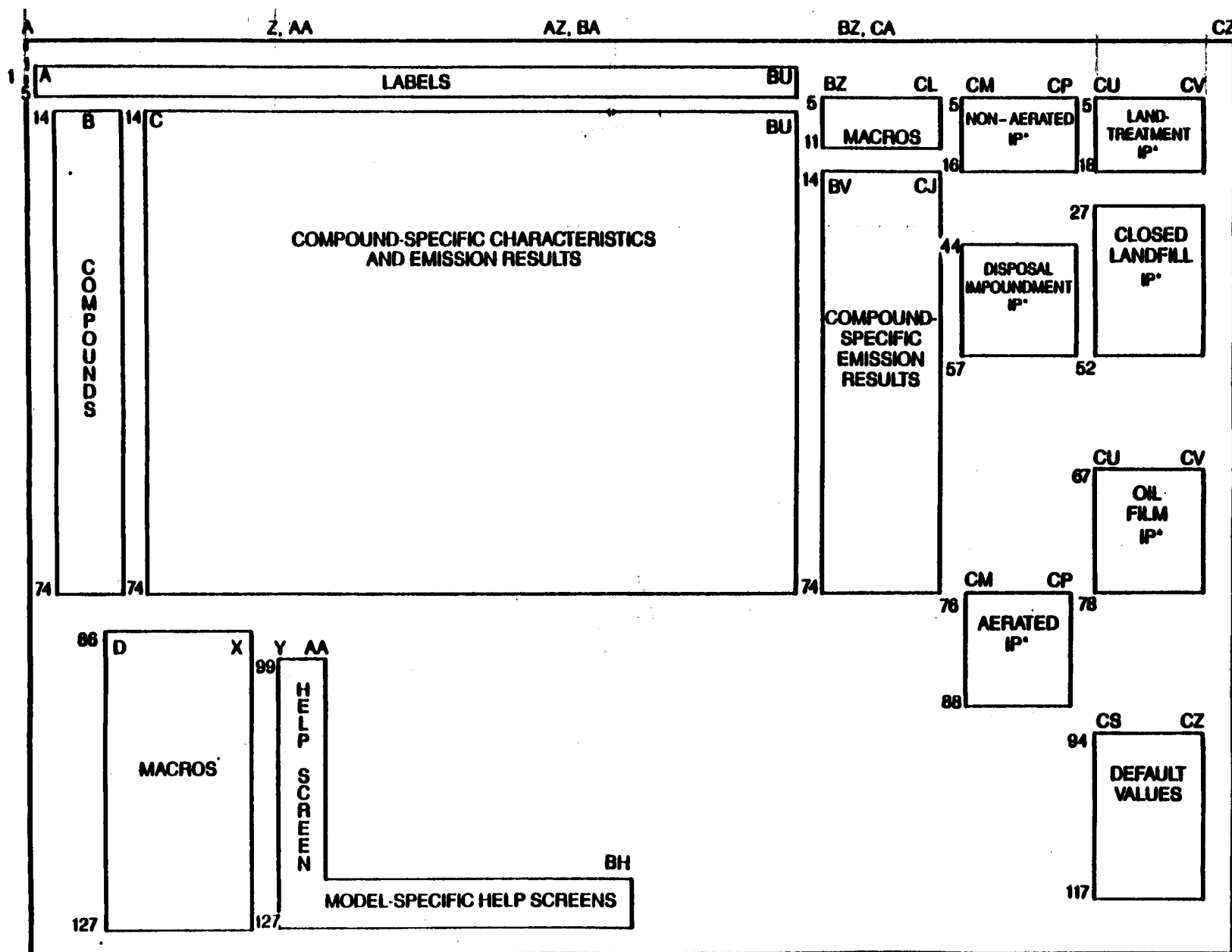
		nonaerated	aerated
	temp.	dif.coef.	dif.coef.
135 D			
136 liquid	1	0.67	0.5
137 gas	1.75	0.67	0.5

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EXHIBIT A-22. IMPOUNDMENTS HELP SCREEN



* = Input parameters.

ATTACHMENT A. GENERALIZED LAYOUT OF CHEMDAT7 SPREADSHEET

APPENDIX B
A GUIDE THROUGH THE LITERATURE

APPENDIX B

A GUIDE THROUGH THE LITERATURE

B.1 INTRODUCTION

There is concern that volatilization of organic compounds (VO) from hazardous waste treatment, storage, and disposal facilities (TSDF) poses a public health problem. These VO emissions may adversely affect ambient air quality in or around a hazardous waste TSDF. However, there are other competing mechanisms or pathways through which VO can leave hazardous waste facilities. These include adsorption, migration, runoff, biological or chemical degradation, hydrolysis, oxidation, and hydroxyl radical reaction. Consequently, the potential hazard of volatilization should be assessed in relation to the potential importance of these other pathways.

Hazardous waste TSDF include, among others, surface impoundments, landfills, and wastewater treatment plants. The important competing pathways for each hazardous waste site have been identified in earlier sections of this report. This evaluation has been based on field and laboratory measurements as well as predictive or mathematical models of these pathways. This appendix supplements the body of the report; it serves to direct the reader through the literature concerning VO pathways from hazardous waste TSDF.

For the convenience of the reader, a comprehensive source list is presented in Appendix C of this report. In addition to the references cited in this appendix and in the individual sections of the report, this bibliography lists other literature of interest for VO pathways and TSDF emission models.

B.2 SURFACE IMPOUNDMENTS

B.2.1 Volatilization

Direct measurement of volatilization rates from surface impoundments is extremely complicated. Hwang and Thibodeaux¹ reviewed the concentration

profile and plume mapping technique and proposed a new method requiring fewer concentration measurements. This latter technique has yet to gain popularity. Thibodeaux et al.² used the concentration profile technique to measure the rate at which selected VO were emitted to the air from basins in the pulp and paper industry. The ranges of the average flux for methanol and acetone were 1.4 to 3.8 ng/cm²•s and 0.028 to 0.10 ng/cm²•s, respectively, which were higher than background. Radian³ obtained emission rates from four different hazardous waste sites containing surface impoundments as well as landfills and landfarms. They used the concentration profile, transect, materials balance, and vent sampling approaches. These methods^{4,5} are also applicable to other nonpoint source hazardous waste facilities such as landfills and landfarms.

Volatilization rates from surface impoundments can be estimated via mathematical models. Mackay and Leinonen⁶ predicted air emissions from nonaerated surface impoundments receiving influent irregularly (unsteady state). The liquid and gas mass transfer coefficients in this model were modified by Mackay and Yeun.⁷ Thibodeaux et al.⁸ developed predictive models for both aerated and nonaerated steady-state surface impoundments. DeWolf and Wetherold⁹ critiqued these models and presented a protocol for their proper use. Shen¹⁰ modified the nonaerated model of Thibodeaux et al.¹¹ In an extensive review of these and other predictive models, GCA¹² judged the theoretical work of Thibodeaux et al.¹³ and Mackay and Leinonen¹⁴ as most suitable for predicting air emissions from surface impoundments.

The use of these mathematical models for predicting volatile emissions is less expensive and faster than actual field measurements. However, to be cost-effective, these mathematical models must provide accurate estimates of volatilization rates. It is disappointing to note that relatively few validation studies are reported in the literature. A description of these follows.

Hwang¹⁵ compared predicted and measured emission rates of toluene and 1,1,1-trichloroethane from an evaporation pond. The transect method was used for field measurements, and the models summarized in Hwang¹⁶ provided the predicted rates. For each organic compound, the predicted result was

within the confidence limits of the average measured result. Balfour et al.¹⁷ used the Thibodeaux et al.¹⁸ model to predict emissions from these surface impoundments. Emission rates were measured via the flux chamber in all three ponds as well as via the concentration profile in the third pond. In this latter pond, the emission rate of most compounds as determined using the flux chamber was statistically greater than that determined using the concentration profile. Furthermore, results of a comparison of measured versus predicted emission rates were inconclusive. Vaught¹⁹ used the Springer et al.²⁰ and Mackay and Yeun²¹ approaches to predict air emissions from quiescent surface impoundments. He concluded that volatilization rates predicted from the Mackay and Yeun model were comparable to rates measured via the flux chamber. In contrast, the predicted rates from Springer et al. exceeded the measured rates.

B.2.2 Other Pathways

The role of other pathways in the removal of VO from surface impoundments has not been addressed extensively in the literature. However, biological removal mechanisms associated with stabilization ponds and lagoons²² will be applicable where conditions of pH, temperature, and nutrient levels are suitable for biological growth.

B.3 LAND TREATMENT

For the past 25 years, the petroleum industry has operated land treatment, sludge farming, and land disposal facilities. The pharmaceutical and organic chemical manufacturers recently have considered this method of hazardous waste disposal because of its comparatively reasonable cost, simplicity, and use of natural processes. How does a land treatment effectively and safely treat and dispose of VO? The purpose of land treatment is to exploit the microbiological actions of the upper soil layer to degrade toxic organic material at a controlled rate. Although photodegradation takes place in a land treatment facility,²³ the short time that the materials are exposed to direct sunlight and the screening effect of the oil in which hazardous materials are suspended make this pathway negligible. Several other pathways may exist: volatilization, runoff, and leaching.^{24,25} However, these latter mechanisms do not occur at a properly sited, operated, and maintained RCRA-permitted land treatment facility.

B.3.1 Degradation

The chemical structure of the hazardous waste, application and mixing techniques, and soil characteristics (texture, temperature, moisture content, oxygen level, nutrient level, pH, and the kind and number of microbes) affect biodegradation.^{26,27} Although biodegradation is purported to be the principal mechanism for removal of organic compounds by land treatment, only a few experiments have attempted to quantify the resulting removal. A laboratory simulation of land treatment of oily sludge revealed that 85 percent of the polynuclear aromatics degraded.²⁸ Results from Snyder et al.²⁹ are comparable: oil removal on fertilized plots approached 80 percent at 1 year postapplication.

Mathematical models for degradation could not be found in the literature.

B.3.2 Volatilization

Techniques for direct measurement of volatilization at landfarms^{30,31} were discussed previously. Exogenous factors affecting volatilization in land treatment include properties of the soil, waste application techniques, mixing schedules, and atmospheric conditions.^{32,33,34}

Farmer and Letey³⁵ proposed five gradient models for pesticide volatilization rates from the soil based on diffusion laws. The models accommodate soil-incorporated pesticides with and without significant mobility in flowing water. With nonincorporated pesticides, vapor density relationships and air flow rate rather than movement in the soil control the volatilization rate. These approaches do not, however, accommodate subsurface injection. Thibodeaux and Hwang³⁶ developed a gradientless model of air emissions from petroleum waste landfarms. Their approach accurately predicted the volatilization of dieldrin reported in Farmer and Letey³⁷ and is considered most suitable for estimating air emissions from land treatment.

B.3.3 Migration and Runoff

Migration and runoff of VO from a landfarm may occur after improper application or treatment of the hazardous waste. A description of factors affecting these two pathways appeared in Reference 38. Results from a laboratory study of refinery and petrochemical sludge³⁹ suggested that the

presence of hazardous waste in runoff decreases with time after application. In addition, leachate water collected 1.5 meters below the subsurface was essentially free of toxic components.

However, as previously mentioned, migration and runoff do not occur at a properly sited, operated, and maintained RCRA-permitted land treatment facility. This paragraph is included for the sake of completeness.

B.4 LANDFILLS

B.4.1 Volatilization

Direct measurements of VO emissions from landfills are possible. During field tests conducted for EPA's Office of Air Quality Planning and Standards (OAQPS), Radian⁴⁰ measured air emissions from landfills at three of the four monitored hazardous waste TSDF. Markle et al.⁴¹ collected air samples from three landfills representative of those used by the polyvinyl chloride industry for health hazard evaluations. To compare the efficiencies of water and soil coverings in reducing volatilization, Farmer et al.⁴² measured emission rates from simulated landfills.

Numerous equations also have been developed to model VO emissions from hazardous waste landfills. The procedure of Farmer et al.⁴³ based on Fick's Law for steady-state diffusion, estimates emission from covered or buried landfills. This was later modified by Shen.⁴⁴ Thibodeaux's⁴⁵ emission models differentiate covered landfills by the presence or absence of internal gas generation. Another approach⁴⁶ incorporates time-varying atmospheric pressure into the emission model. Volatilization rates from landfills with no covering, i.e., open dumps, were modeled by Shen.⁴⁷ DeWolf and Wetherold⁴⁸ recommend Shen's⁴⁹ emission model for covered landfills. GCA, in their excellent comprehensive review of these and other emission models, prefers the work of Farmer et al.⁵⁰ and Thibodeaux.⁵¹

Field validation of these mathematical models has not been reported in the literature. Despite this, Baker and Mackay⁵² employed Shen's⁵³ model in their protocol to evaluate toxic air pollution downwind of hazardous waste landfills.

B.4.2 Migration

Several scientists have investigated the potential problem of migration of toxic contaminants from landfills. Rovers and Farquhar⁵⁴

suggested that the production of leachate within a landfill is not uncommon. However, the migration of harmful compounds through adjacent soils is not significant. Shen and Tofflemire⁵⁵ reported that annual losses of PCB to migration from uncovered landfills in the Hudson River Basin (New York) were substantially less than losses to volatilization.

B.4.3 Other Pathways

The impact of other pathways is not discussed quantitatively in the literature.

B.5 WASTEWATER TREATMENT PLANT EFFLUENT

A description of the pathways operating in a wastewater treatment plant is complicated by the number of different treatment systems. There are closed tanks and open tanks (with and without mixing). Air emissions from closed tanks occur during venting.

E. C. Jordan⁵⁶ and Burns and Roe^{57,58} examined the fate of priority toxic pollutants in publicly owned treatment plants. They observed a decrease in VO concentrations across the activated sludge process and a lack of pollutant accumulation in the waste-activated sludge. This suggests that VO are substantially air-stripped or biodegraded during secondary treatment. Results from the controlled laboratory experiments of Roberts et al.⁵⁹ imply that organic solutes more likely volatilize during wastewater treatment with surface aeration than with bubble aeration. Lurker et al.⁶⁰ examined how aeration rate, suspended particle concentration, and detergent concentration influence aerial organic chemical release from an activated sewage treatment process.

The nonaerated open tank system is similar to the nonaerated surface impoundment discussed previously; see Section B.2.1 for a discussion of the corresponding emission rate models. Similarly, open tank wastewater treatment processes with mixing can be estimated from Thibodeaux et al.⁶¹ Hwang⁶² went a step further in his activated sludge surface aeration model. His approach was to estimate pollutant removal by degradation, adsorption, and air stripping via a mass balance equation. Like Hwang and Thibodeaux et al., Freeman⁶³ considered air stripping losses at the air-water interface. Unlike Hwang, however, he viewed the adsorption pathway as insignificant and, thus, ignored it. In an entirely different approach,

Freeman^{64,65} modeled the mass transfer of a toxic compound into the bubbles of the aerated system (diffused air [subsurface] activated sludge model). The structure of these and other models was critiqued in GCA.⁶⁶

Allen et al.⁶⁷ presented models of VO losses at each process encountered in wastewater treatment systems. The models include a methodology for estimating the relative importance of competing pathways. Additionally, these investigators compared the loss of volatiles obtained from field tests at several treatment facilities⁶⁸ and from these mathematical models. The models predict VO losses due to biodegradation or volatilization in close agreement with the field data. Results from other validation studies are not as consistent. Predicted emission rates from aerated surface impoundments at two wastewater treatment plants were generally larger than measured values.⁶⁹ The difference between measured and predicted emission rates in Cox et al.⁷⁰ appears to be a function of the type of compound and the presence of aerators.

B.6 SUMMARY

This appendix serves to guide the reader through the literature concerning VO pathways from hazardous waste TSDF. The pathways of interest include volatilization, adsorption, migration, runoff, biological or chemical degradation, hydrolysis, oxidation, and hydroxyl radical reaction. The hazardous waste TSDF considered are surface impoundments, landfills, landfarms, and wastewater treatment plants. The body of this report expands on the pathways and models most applicable to current research.

B.7 REFERENCES

1. Hwang, S. T., and L. J. Thibodeaux. Measuring Volatile Chemical Emission Rates from Large Waste Disposal Facilities. Environmental Progress. 2:81-86. 1983.
2. Thibodeaux, L. J., D. G. Parker, and H. H. Heck (University of Arkansas). Measurement of Volatile Chemical Emissions from Wastewater Basins. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. R805534. December 1981.
3. Radian Corporation. Hazardous Waste Treatment, Storage, and Disposal Facility Area Sources: VOC Air Emissions. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-02-3850. January 1985.

4. Reference 1.
5. Reference 3.
6. Mackay, D., and P. J. Leinonen. Rate of Evaporation of Low-Solubility Contaminants from Water Bodies to Atmosphere. Environmental Science and Technology. 13:1178-1180. 1975.
7. Mackay, D., and A. T. K. Yeun. Mass Transfer Coefficient Correlations for Volatilization of Organic Solutes from Water. Environmental Science and Technology. 17:211-217. 1983.
8. Reference 2.
9. DeWolf, G. B., and R. G. Wetherold (Radian Corporation). Protocols for Calculating VOC Emissions from Surface Impoundments Using Emission Models, Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-02-3850. September 1984.
10. Shen, T. T. Estimation of Organic Compound Emissions from Waste Lagoons. Journal of the Air Pollution Control Association. 32. 1982.
11. Reference 2.
12. GCA Corp. Evaluation and Selection of Models for Estimating Air Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-450/8-84-020. December 1984.
13. Reference 2.
14. Reference 6.
15. Hwang, S. T. Model Prediction of Volatile Emissions. Environmental Progress. 4:141-144. 1985.
16. Hwang, S. T. Toxic Emissions From Land Disposal Facilities. Environmental Progress. 1:46-52. 1982.
17. Balfour, W. D., C. E. Schmidt, R. G. Wetherold, D. L. Lewis, J. I. Steinmetz, and R. C. Hanish. Field Verification of Air Emission Models for Hazardous Waste Disposal Facilities. In: Proceedings of the Tenth Annual Research Symposium. Publication No. EPA-600/9-84/007. Fort Mitchell, Kentucky. April 1984.
18. Reference 2.
19. Vaught, C. C. (GCA). Air Emissions for Quiescent Surface Impoundments--Emissions Data and Model Review, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-01-6871. August 1985.

20. Springer, S., P. D. Lunney, K. T. Valsaraj, and L. J. Thibodeaux (University of Arkansas and Louisiana State University.) Emissions of Hazardous Chemicals from Surface and Near Surface Impoundments to Air, Draft Final Report. Prepared for U.S. Environmental Protection Agency. Washington, DC. Project No. 808161-2. December 1984.
21. Reference 7.
22. Metcalf and Eddy, Inc. Wastewater Engineering: Collection, Treatment, Disposal. New York, McGraw-Hill. 1972.
23. Kaufman, D. D. Fate of Organic Compounds in Land-Applied Wastes. In: Land Treatment of Hazardous Wastes, Parr, J. F., P. B. Marsh, and J. M. Kla (eds.). 1983. p. 77-151.
24. U.S. Environmental Protection Agency. Hazardous Waste Land Treatment, Technical Resource Document. EPA Contract Nos. 68-03-2940 and 68-03-2943. April 1983.
25. Brown, K. W. Chapter 36, Land Treatment of Hazardous Wastes. In: Proceedings of the Fourth Life Sciences Symposium, Environment and Solid Wastes. Gatlinburg, TN. October 4-8, 1981. p. 449-482.
26. Reference 24.
27. Reference 23.
28. Bossert, I., W. M. Kachel, and R. Barth. Fate of Hydrocarbons During Oily Sludge Disposal in Soil. Applied and Environmental Microbiology. 47:763-767. 1984.
29. Snyder, H. J., G. B. Rice, and J. J. Skujins. Residual Management by Land Disposal. In: Proceedings of the Hazardous Waste Research Symposium, Fuller, W. H., (ed.) Publication No. EPA-600/9-76-015. July 1976.
30. Reference 1.
31. Reference 3.
32. Wetherold, R. G., J. L. Randall, and K. R. Williams (Radian Corporation). Laboratory Assessment of Potential Hydrocarbon Emissions from Land Treatment of Refinery Oily Sludges. Prepared for U.S. Environmental Protection Agency. Washington, DC. Publication No. EPA-600/2-84-108. June 1984.
33. Reference 24.
34. Farmer, W. J., and J. Letey. Volatilization Losses of Pesticides from Soil. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-660/2-74-054. August 1974.

35. Reference 34.
36. Thibodeaux, L. J., and S. T. Hwang. Land Farming of Petroleum Wastes--Modeling the Air Emissions Problem. Environmental Progress. 1:42-46. 1982.
37. Reference 34.
38. Reference 23.
39. Reference 25.
40. Reference 3.
41. Markle, R. A., R. B. Iden, and F. A. Sliemers (Battelle). A Preliminary Examination of Vinyl Chloride Emissions from Polymerization Sludges during Handling and Land Disposal. Prepared for U.S. Environmental Protection Agency. Washington, DC. Publication No. EPA-660/2-74-054. February 1976.
42. Farmer, W. J., M. Yang, J. Letey, and W. F. Spencer. Problems Associated with the Land Disposal of an Organic Industrial Hazardous Waste Containing HCB. In: Residual Management by Land Disposal, Proceedings of the Hazardous Waste Research Symposium, Fuller, W. H. (ed.). Publication No. EPA-600/9-76-015. July 1976.
43. Farmer, W. J., M. S. Yang, and J. Letey. Land Disposal of Hazardous Wastes: Controlling Vapor Movement in Soils. In: Fourth Annual Research Symposium. Publication No. EPA-600/9-78-016. August 1978.
44. Shen, T. T. Estimating Hazardous Air Emissions from Disposal Sites. Pollution Engineering. 31-34. August 1981.
45. Thibodeaux, L. J. Estimating the Air Emissions of Chemicals from Hazardous Waste Landfills. Journal of Hazardous Materials. 4:235-244. 1981.
46. Thibodeaux, L. J., C. Springer, and L. M. Riley. Models of Mechanisms for the Vapor Phase Emission of Hazardous Chemicals from Landfills. Journal of Hazardous Materials. 7:63-74. 1982.
47. Reference 44.
48. DeWolf, G. B., and R. G. Wetherold (Radian Corporation). Protocols for Calculating VOC Emissions from Land Applications Using Emission Models, Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-02-3850. December 1984.
49. Reference 44.
50. Reference 43.

51. Reference 2.
52. Baker, L. W., and K. P. Mackay. Hazardous Waste Management: Screening Models for Estimating Toxic Air Pollution Near a Hazardous Waste Landfill. Journal of the Air Pollution Control Association. 35:1190-1195. 1985.
53. Reference 44.
54. Rovers, F. A., and G. J. Farquhar. Evaluating Contaminant Attenuation in the Soil to Improve Landfill Selection and Design. In: Proceedings of the International Conference on Land for Waste Management. 1974. p. 161-173.
55. Shen, T. T., and T. J. Tofflemire. Air Pollution Aspects of Land Disposal of Toxic Wastes. Environmental Engineering Division Journal. 106:211-226. 1980.
56. E. C. Jordan Co. Fate of Priority Toxic Pollutants in Publicly Owned Treatment Works, 30 Day Study. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-440/1-82-302. August 1982.
57. Burns and Roe Industrial Services Corp. Fate of Priority Pollutants in Publicly Owned Treatment Works, Final Report, Volume II. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-440/1-82-303. July 1982.
58. Burns and Roe Industrial Services Corp. Fate of Priority Pollutants in Publicly Owned Treatment Works, Final Report, Volume I. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-440/1-82-303. September 1982.
59. Roberts, P. V., C. Munz, P. Dandliker, and C. Matter-Muller. Volatilization of Organic Pollutants in Wastewater Treatment-Model Studies, Project Summary. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-600/52-84-047. April 1984.
60. Lurker, P. A., C. S. Clark, V. J. Elia, P. S. Gartside, and R. N. Kinman. Aerial Organic Chemical Release from Activated Sludge. Water Research. 18:489-494. 1984.
61. Reference 2.
62. Hwang, S. T. Treatability and Pathways of Priority Pollutants in Biological Wastewater Treatment. (Presented at the American Institute of Chemical Engineers Symposium. Chicago, Illinois. November 1980.)
63. Freeman, R. A. Stripping of Hazardous Chemicals from Surface Aerated Waste Treatment Basins. In: APCA/WPCF Speciality Conference on Control of Specific Toxic Pollutants. Gainesville, Florida. February 13-16, 1979.

64. Freeman, R. A. Comparison of Secondary Emissions From Aerated Treatment Systems. (Presented at AIChE Meeting. Paper 5c. Orlando, Florida. February 1982.)
65. Freeman, R. A. Secondary Emissions from Subsurface Aerated Treatment Systems. Environmental Progress. 1:117-119. 1982.
66. Reference 12.
67. Allen, C. C., S. Simpson, and G. Brant (RTI and Associated Technologies, Inc.). Field Evaluations of Hazardous Waste Pretreatment as an Air Pollution Control Technique. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-02-3992. April 1985.
68. Alsop, G. M., R. L. Berglund, T. W. Siegrist, G. M. Whipple, and B. E. Wilkes. Fate of Specific Organics in an Industrial Biological Wastewater Treatment Plant, Draft Report. Prepared for U.S. Environmental Protection Agency, Industrial Environmental Research Laboratory. Research Triangle Park, NC. June 29, 1984.
69. Meisenheimer, D. C. (GCA). Emissions Data and Model Review for Wastewater Treatment Operations, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-01-6871. August 1985.
70. Cox, R. D., Lewis, R. G. Wetherold, and J. I. Steinmetz (Radian Corporation). Evaluation of VOC Emissions from Wastewater Systems (Secondary Emissions). Prepared for U.S. Environmental Protection Agency. Washington, DC. Project No. 68-03-3038. July 1983.

APPENDIX C
COMPREHENSIVE SOURCE LIST

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Addendum to memorandum dated September 6, 1985, from Eichinger, Jeanne, GCA Corporation, to Hustvedt, K. C., EPA/OAQPS. September 12, 1985. TSDF model source parameters and operating practices data base.

Allen, C. C. Prediction of Air Emissions from Surface Impoundments, Paper 31a. (Presented at 1986 Summer Meeting of AIChE. Boston, MA. August 1986.) 26 p.

Allen, C. C., and D. A. Green (Research Triangle Institute). Review of VOC Pathway Models, Draft Report. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-01-6826. 1985.

Allen, C. C., D. A. Green, and J. B. White (Research Triangle Institute). Preliminary Assessment of Aerated Waste Treatment Systems at TSDFs - Phase I, Draft Final Report. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-03-3149. May 1985.

Allen, C. C., D. A. Green, and J. B. White (Research Triangle Institute). Preliminary Assessment of Aerated Waste Treatment Systems at TSDFs--Phase I. Draft. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. EPA Contract No. 68-03-3149, Task 54-01F. 1985.

Allen, C. C., S. Simpson, and G. Brant (Research Triangle Institute and Associated Technologies, Inc.). Field Evaluations of Hazardous Waste Pretreatment as an Air Pollution Control Technique. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-02-3992. April 1985.

Alsop, G. M., R. L. Berglund, T. W. Siegrist, G. M. Whipple, and B. E. Wilkes. Fate of Specific Organics in an Industrial Biological Wastewater Treatment Plant, Draft Report. Prepared for U.S. Environmental Protection Agency, Industrial Environmental Research Laboratory. June 29, 1984.

Armstrong, N. E., E. F. Gloyna, and O. Wyss. Biological Countermeasures for the Control of Hazardous Material Spills, Project Summary. Publication No. EPA-600/S2-84-071. March 1984.

- Arnold, J. H. Studies in Diffusion: III. Unsteady-State Vaporization and Absorption. Transactions of the American Institute of Chemical Engineers. 40:361-379. 1944.
- Bailey, J. E., and D. F. Ollis. Biochemical Engineering Fundamentals. New York, McGraw-Hill. 1977. p. 343-349.
- Baker, L. W., and K. P. Mackay. Hazardous Waste Management: Screening Models for Estimating Toxic Air Pollution Near a Hazardous Waste Landfill. Journal of the Air Pollution Control Association. 35: 1190-1195. 1985.
- Balfour, W. D., C. E. Schmidt, R. G. Wetherold, D. L. Lewis, J. I. Steinmetz, and R. C. Hanish. Field Verification of Air Emission Models for Hazardous Waste Disposal Facilities. In: Proceedings of the Tenth Annual Research Symposium. Publication No. EPA-600/9-84/007. Fort Mitchell, Kentucky. April 1984.
- Bossert, I., W. M. Kachel, and R. Barth. Fate of Hydrocarbons during Oily Sludge Disposal in Soil. Applied and Environmental Microbiology. 47: 763-767. 1984.
- Branscome, M., and A. Gitelman. Sensitivity Analysis: Emission Estimates for Surface Impoundments. Prepared for the U.S. Environmental Protection Agency. (Docket Number ____). March 1986. 67 p.
- Brown, K. W. Chapter 36, Land Treatment of Hazardous Wastes. In: Proceedings of the Fourth Life Sciences Symposium, Environment and Solid Wastes. Gatlinburg, TN. October 4-8, 1981. p. 449-482.
- Burns and Roe Industrial Services Corp. Fate of Priority Pollutants in Publicly Owned Treatment Works, Final Report, Volume II. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-440/1-82-303. July 1982.
- Burns and Roe Industrial Services Corp. Fate of Priority Pollutants in Publicly Owned Treatment Works, Final Report, Volume I. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-440/1-82-303. September 1982.
- Canter, L., and R. C. Knox. Evaluation of Septic Tank System Effects on Ground Water Quality. Publication No. EPA-600/S2-84-107. September 1984.
- Clark, J. W., W. Viessman, Jr., and M. J. Hammer. Water Supply and Pollution Control. Scranton, PA, International Textbook Company. 1971.
- Cox, R. D., D. L. Lewis, R. G. Wetherold, and J. I. Steinmetz (Radian Corporation). Evaluation of VOC Emissions from Wastewater Systems (Secondary Emissions). Prepared for U.S. Environmental Protection Agency. Washington, DC. EPA Project No. 68-03-3038. July 1983.

- Cox, R. D., J. L. Steinmetz, and D. L. Lewis (Radian Corporation). Evaluation of VOC Emissions from Wastewater Systems (Secondary Emission), Volume II, Appendices. Prepared for U.S. Environmental Protection Agency. Washington, DC. EPA Project No. 68-03-3038. July 1983.
- Crank, J. The Mathematics of Diffusion. London, England, Oxford University Press. 1970. p. 45-47.
- DeWolf, G. B., and R. G. Wetherold (Radian Corporation). Protocols for Calculating VOC Emissions from Surface Impoundments Using Emission Models, Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-02-3850. September 1984.
- DeWolf, G. B., and R. G. Wetherold (Radian Corporation). Protocols for Calculating VOC Emissions from Land Applications Using Emission Models, Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-02-3850. December 1984.
- E. C. Jordan Co. Fate of Priority Toxic Pollutants in Publicly Owned Treatment Works, 30 Day Study. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-440/1-82-302. August 1982.
- Ely, R. L., G. L. Kingsbury, M. R. Branscome, L. J. Goldman, C. M. Northeim, J. H. Turner, and F. O. Mixon, Jr. (Research Triangle Institute). Performance of Clay Caps and Liners for Disposal Facilities. Prepared for U.S. Environmental Protection Agency. Cincinnati, OH. EPA Contract No. 68-03-3149. March 1983.
- Farino, W., P. Spawn, M. Jasinski, and B. Murphy (GCA/Technology). Evaluation and Selection of Models for Estimating Air Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Prepared for U.S. Environmental Protection Agency. Publication No. EPA 450/3-84-020. 1984.
- Farmer, W. J., and J. Letey. Volatilization Losses of Pesticides from Soil. Publication No. EPA-660/2-74-054. August 1974.
- Farmer, W. J., M. S. Yang, and J. Letey. Land Disposal of Hazardous Wastes: Controlling Vapor Movement in Soils. In: Fourth Annual Research Symposium. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-600/9-78-016. August 1978.
- Farmer, W. J., M. S. Yang, and W. F. Spencer. Land Disposal of Hexachlorobenzene: Controlling Vapor Movement in Soil. Publication No. EPA-600/2-80-119. August 1980.
- Farmer, W. J., M. S. Yang, J. Letey, W. F. Spencer, and M. H. Roulier. Land Disposal of Hexachlorobenzene Wastes: Controlling Vapor Movement in Soils. Fourth Annual Research Symposium. U.S. Environmental Protection Agency. Publication No. EPA-600/9-78-016. August 1978.

- Farmer, W. J., M. Yang, J. Letey, and W. F. Spencer. Problems Associated with the Land Disposal of an Organic Industrial Hazardous Waste Containing HCB. In: Residual Management by Land Disposal, Proceedings of the Hazardous Waste Research Symposium, Fuller, W. H. (ed.). U.S. Environmental Protection Agency. Publication No. EPA-600/9-76-015. July 1976.
- Freeman, R. A. Comparison of Secondary Emissions From Aerated Treatment Systems. (Presented at AIChE Meeting. Paper 5c. Orlando, Florida. February 1982.)
- Freeman, R. A. Secondary Emissions from Subsurface Aerated Treatment Systems. Environmental Progress. 1:117-119. 1982.
- Freeman, R. A. Stripping of Hazardous Chemicals from Surface Aerated Waste Treatment Basins. In: APCA/WPCF Speciality Conference on Control of Specific Toxic Pollutants. Gainesville, Florida. February 13-16, 1979.
- Galegar, J. (East Central Oklahoma State University). Annotated Literature Reference on Land Treatment of Hazardous Waste. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-600/2-84-098. May 1984.
- GCA Corporation. Evaluation and Selection of Models for Estimating Air Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-450/8-84-020. December 1984.
- GCA/Technology Division. Evaluation and Selection of Models for Estimating Air Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. Publication No. EPA-450/3-84-020. December 1984. pp. 85-89.
- GCA Corporation. Hazardous Waste TSDF Waste Process Sampling. 1985. p. 1-11.
- GCA Corporation. Air Emission Estimation Methods for Transfer, Storage, and Handling Operations, Draft Technical Note. Prepared for U.S. Environmental Protection Agency, Office of Air Quality Planning and Standards. Research Triangle Park, NC. EPA Contract No. 68-01-6871. August 1985.
- GCA Corporation. Air Emissions for Quiescent Surface Impoundments--Emissions Data and Model Review, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Contract No. 68-01-6871, Assignment 49. August 1985.
- GCA Corporation. Air Emissions from Land Treatment--Emissions Data and Model Review, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. August 1985. 120 pp.

GCA Corporation. Air Emissions of VOC from Waste Piles at Hazardous Waste Treatment, Storage, and Disposal Facilities. Prepared for U.S. Environmental Protection Agency, Office of Air Quality Planning and Standards. Research Triangle Park, NC. EPA Contract No. 68-01-6871. August 1985.

GCA Corporation. Air Emissions from Land Treatment--Emissions Data and Model Review. Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Research Triangle Park, NC. September 1985. Chapter 4.

Gerachty, J. J., D. W. Miller, F. Vander Leeden, and F. L. Troise. Water Atlas of the United States. Port Washington, NY, Water Information Center, Inc. 1973. Plate 30.

Ghassemi, M., M. Haro, and L. Fargo. Assessment of Hazardous Waste Surface Impoundment Technology: Case Studies and Perspectives of Experts, Project Summary. Publication No. EPA-600/S2-84-173. January 1985.

Graver Standard Cone-Roof, Flat-Bottom Tanks. Sizes and Capacities.

Hornick, S. B. In: Land Treatment of Hazardous Waste, Parr, J. F. (ed). Park Ridge, NJ, Noyes Data Company. 1982.

Huddleston, R. L. Solid-Waste Disposal: Land Farming. Chemical Engineering. 119-124. February 26, 1979.

Hwang, S. T. Treatability and Pathways of Priority Pollutants in Biological Wastewater Treatment. (Presented at the American Institute of Chemical Engineers Symposium. Chicago, Illinois. November 1980.)

Hwang, S. T. Toxic Emissions from Land Disposal Facilities. Environmental Progress. 1:46-52. February 1982.

Hwang, S. T. Model Prediction of Volatile Emissions. Environmental Progress. 4:141-144. 1985.

Hwang, S. T., and L. J. Thibodeaux. Measuring Volatile Chemical Emission Rates from Large Waste Disposal Facilities. Environmental Progress. 2:81-86. 1983.

ICF, Inc. The RCRA Risk-Cost Analysis Model Phase III Report, Appendix E. Chemical and Physical Processes Affecting Decay Rates of Chemicals in Aquatic Environments, Draft. Prepared for U.S. Environmental Protection Agency, Office of Solid Waste, Economic Analysis Branch. Washington, DC. 1984.

Kaufman, D. D. Fate of Organic Compounds in Land-Applied Wastes. In: Land Treatment of Hazardous Wastes, Parr, J. F., P. B. Marsh, and J. M. Kla (eds.). 1983. p. 77-151.

- Kincannon, D. F. Evaluation of Biological Tower Design Methods.
(Presented at First International Conference on Fixed-Film Biological Processes. Kings Island, OH. April 1982.)
- Land Farming Fills on HPI Need. Hydrocarbon Processing 60:97-103. 1980.
- Letter and attachment from Shen, T., New York State Department of Environmental Conservation, to McDonald, R., EPA/OAQPS. Modification of Arnold's equation. February 16, 1986.
- Loehr, R. C., and J. F. Malina, Jr. (eds.) Land Treatment, A Hazardous Waste Management Alternative. In: Proceedings of the Water Resources Symposium Number Thirteen. Austin, Texas, Center for Research in Water Resources, The University of Texas at Austin. 1986.
- Lunney, P. D. Characterization of Wind and Depth Effects Upon Liquid Phase Mass Transfer Coefficients: Simulation Studies. Master's thesis, University of Arkansas, Fayetteville, AR. January 1983.
- Lurker, P. A., C. S. Clark, V. J. Elia, P. S. Gartside, and R. N. Kinman. Aerial Organic Chemical Release from Activated Sludge. Water Research. 18:489-494. 1984.
- Lyman, W. J., et al. Research and Development of Methods for Estimating Physicochemical Properties of Organic Compounds of Environmental Concern. Phase II, Part I. NTIS AD 11875A. 1981.
- Lyman, Warren J., William F. Reehl, and David H. Rosenblatt. Handbook of Chemical Property Estimation Method--Environmental Behavior of Organic Compounds. New York, NY, McGraw-Hill Book Company. 1982. Chapter 9.
- Mabey, W. R., T. Mill, and D. G. Hendry. Photolysis in Water. In: Laboratory Protocols for Evaluating the Fate of Organic Chemicals in Air and Water, Draft. Prepared for U.S. Environmental Protection Agency. EPA Contract 68-03-2227. 1980.
- Mackay, D., and A. Yeun. Mass Transfer Coefficient Correlations for Volatilization of Organic Solutes from Water. Environmental Science and Technology. 17:211-217. 1983.
- Mackay, D., and P. J. Leinonen. Rate of Evaporation of Low-Solubility Contaminants from Water Bodies to Atmosphere. Environmental Science and Technology. 13:1178-1180. 1975.
- Mackay, D., W. Y. Shiu, A. Bobra, J. Billington, E. Chau, A. Yeun, C. Ng, and F. Szeto (University of Toronto). Volatilization of Organic Pollutants from Water. Prepared for U.S. Environmental Protection Agency. Publication No. EPA-600/3-82-019. April 1982.

- Markle, R. A., R. B. Iden, and F. A. Sliemers (Battelle Laboratories). A Preliminary Examination of Vinyl Chloride Emissions from Polymerization Sludges during Handling and Land Disposal. Prepared for U.S. Environmental Protection Agency. Washington, DC. Publication No. EPA-660/2-74-054. February 1976..
- Matter-Muller, C., W. Gujer, W. Geiger, and W. Stumm. The Prog. Wat. Tech. (Toronto). 12:299-314. Great Britain, IAWPR/Pergamon Press, Ltd. 1980.
- Meisenheimer, D. C. (GCA). Emissions Data and Model Review for Wastewater Treatment Operations, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-01-6871. August 1985.
- Memorandum and attachment from Wright, M., Research Triangle Institute, to Thorneloe, S., EPA/OAQPS. Selection of an emission model for land treatment. May 30, 1986.
- Memorandum from Thorneloe, S., EPA/OAQPS, to Durham, J., EPA/OAQPS. January 31, 1986. Land treatment data base.
- Metcalf and Eddy, Inc. Wastewater Engineering: Collection, Treatment, Disposal. New York, McGraw-Hill. 1972. 782 p.
- Meyers, J. D., and R. L. Huddleston. Treatment of Oily Refinery Wastes by Land Farming. In: Proceedings of the Industrial Waste Conference. Lafayette, Indiana. May 8-10, 1979. p. 686-698.
- Millington, R. J., and J. P. Quirk. Permeability of Porous Solids. Trans. Faraday Society. 57:1200-1207. 1961.
- Ollis, D. F. Contaminant Degradation in Water. ES&T. 19(6):480-484. 1985.
- Parr, J. F., P. B. Marsh, and J. M. Kla (eds.). Land Treatment of Hazardous Wastes. Park Ridge, NJ, Noyes Data Corporation. 1983.
- Perry, R. H., and C. H. Chilton. Chemical Engineers' Handbook, Fifth Edition. New York, McGraw-Hill. 1973.
- Pitter, P. Determination of Biological Degradability of Organic Substances. Water Research. 10:231-235. 1976.
- Radian Corporation. Hazardous Waste Treatment, Storage, and Disposal Facility Area Sources: VOC Air Emissions. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-02-3850. January 1985.
- Radiation Technologies, Inc. Treatability Data in Support of a Treatment Zone Model for Petroleum Refining Land Treatment Facilities. Prepared for American Petroleum Institute, Washington, DC. March 1986. 154 pp.

- Reinhardt, J. R. Gas-Side Mass-Transfer Coefficient and Interfacial Phenomena of Flat-Bladed Surface Agitators. Ph.D. dissertation, University of Arkansas, Fayetteville, AR. 1977.
- Roberts, P. V., C. Munz, P. Dandliker, and C. Matter-Muller. Volatilization of Organic Pollutants in Wastewater Treatment-Model Studies, Project Summary. Publication No. EPA-600/52-84-047. April 1984.
- Rovers, F. A., and G. J. Farquhar. Evaluating Contaminant Attenuation in the Soil to Improve Landfill Selection and Design. In: Proceedings of the International Conference on Land for Waste Management. 1974. p. 161-173.
- Research Triangle Institute. Exponential Biological Decomposition Model, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Contract No. 68-01-6826. August 15, 1985.
- Research Triangle Institute. RTI Review of VOC Pathway Models, Draft Report. Prepared for U.S. Environmental Protection Agency. Contract No. 68-01-6826. October 31, 1985.
- Savage, G. M., L. F. Diaz, and C. G. Golueke. Disposing of Organic Hazardous Wastes by Composting. BioCycle. 26:31-34. 1985.
- Shen, T. T. Emission Estimation of Hazardous Organic Compounds from Waste Disposal Sites. (Presented at the Air Pollution Control Association Annual Meeting. Montreal, Quebec. June 1980.)
- Shen, T. T. Estimating Hazardous Air Emissions from Disposal Sites. Pollution Engineering. 31-34. August 1981.
- Shen, T. T. Estimation of Organic Compound Emissions from Waste Lagoons. Journal of the Air Pollution Control Association. 32. 1982.
- Shen, T. T. and T. J. Tofflemire. Air Pollution Aspects of Land Disposal of Toxic Wastes. Environmental Engineering Division Journal. 106: 211-226. 1980.
- Snyder, H. J., G. B. Rice, and J. J. Skujins. Residual Management by Land Disposal. In: Proceedings of the Hazardous Waste Research Symposium, Fuller, W. H. (ed.). Publication No. EPA-600/9-76-015. July 1976.
- Spivey, J. J., C. C. Allen, D. A. Green, J. P. Wood, and R. L. Stallings (Research Triangle Institute). Preliminary Assessment of Hazardous Waste Pretreatment as an Air Pollution Control Technique. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-03-3149, Task 12-5. 1984.

- Springer, S., P. D. Lunney, K. T. Valsaraj, and L. J. Thibodeaux
(University of Arkansas and Louisiana State University.) Emissions of
Hazardous Chemicals from Surface and Near Surface Impoundments to Air,
Draft Final Report. Prepared for U.S. Environmental Protection
Agency. Washington, DC. EPA Project No. 808161-2. December 1984.
- Springer, C., P. D. Lunney, and K. T. Valsaraj. Emission of Hazardous
Chemicals from Surface and Near Surface Impoundments to Air. U.S.
Environmental Protection Agency, Solid and Hazardous Waste Research
Division. Cincinnati, OH. EPA Project Number 808161-02. December
1984.
- Stallings, R. L., T. N. Rogers, and M. E. Mullins. Air Stripping of
Volatile Organics. In: Proceedings of the Institute of Environmental
Sciences. Las Vegas, NV. April 30-May 2, 1985. p. 170-174.
- Stiver, W., and D. Mackay. Evaporation Rate of Spills of Hydrocarbons and
Petroleum Mixtures. Environmental Science Technology. 18:834-840.
1984.
- Telecon. Goldman, Len, Research Triangle Institute, with Hannak, Peter,
Alberta Environmental Center. April 4, 1986.
- Telecon. Goldman, Len, Research Triangle Institute, with Hughes, John,
National Climatic Center, Asheville, NC. May 15, 1986.
- Telecon. Goldman, Len, Research Triangle Institute, with Wallis, Al,
National Climatic Center, Asheville, NC. May 30, 1986.
- Telecon. Goldman, Len, Research Triangle Institute, with Borden, Roy,
Department of Civil Engineering, North Carolina State University,
Raleigh, NC. August 13, 1986.
- Thibodeaux, L. J. Air Stripping of Organics from Wastewater. A
Compendium. Air/Water. p. 373. (no date yet).
- Thibodeaux, L. J. Air Stripping of Organics From Wastewater: A
Compendium. Proceedings of the Second National Conference on Complete
Water Use. Chicago, IL. May 4-8, 1978.
- Thibodeaux, L. J. Estimating the Air Emissions of Chemicals from Hazardous
Waste Landfills. Journal of Hazardous Materials. 4:235-244. 1981.
- Thibodeaux, L. J., and S. T. Hwang. Land Farming of Petroleum
Wastes--Modeling the Air Emissions Problem. Environmental Progress.
1:42-46. 1982.
- Thibodeaux, L. J., and S. T. Hwang. Toxic Emissions from Land Disposal
Facilities. Environmental Progress. 1(1):46. February 1982.

- Thibodeaux, L. J., C. Springer, and L. M. Riley. Models of Mechanisms for the Vapor Phase Emission of Hazardous Chemicals from Landfills. *Journal of Hazardous Materials*. 7:63-74. 1982.
- Thibodeaux, L. J., D. G. Parker, and H. H. Heck (University of Arkansas). Measurement of Volatile Chemical Emissions from Wastewater Basins. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. R805534. December 1981.
- Trip Report. Goldman, Len, Research Triangle Institute, with Chemical Waste Management, Sulphur, Louisiana. February 25, 1986.
- Truong, K. N., and J. W. Blackburn. The Stripping of Organic Compounds in Biological Treatment Processes. *Environmental Progress*. 3:153-152. 1984.
- Turner, J. H., C. C. Allen, and D. A. Green (Research Triangle Institute). VOC Pathways from Certain Hazardous Waste Sites, Draft Interim Report. Prepared for U.S. Environmental Protection Agency. EPA Contract No. 68-01-6826. September 1985.
- U.S. Environmental Protection Agency. Transportation and Marketing of Petroleum Liquids. In: AP-42, Compilation of Air Pollutant Emission Factors, Third Edition, Supplement 12, Section 4.4. Office of Air Quality Planning and Standards. Research Triangle Park, NC. July 1979. 13 p.
- U.S. Environmental Protection Agency. Tank and Drum Cleaning. In: AP-42, Compilation of Air Pollutant Emission Factors, Third Edition, Supplement 12, Section 4.8. Office of Air Quality Planning and Standards. Research Triangle Park, NC. February 1980. 4 p.
- U.S. Environmental Protection Agency. Storage of Organic Liquids. In: AP-42, Compilation of Air Pollutant Emission Factors, Third Edition, Supplement 12, Section 4.3. Office of Air Quality Planning and Standards. Research Triangle Park, NC. April 1981. 25 p.
- U.S. Environmental Protection Agency. Hazardous Waste Land Treatment, Technical Resource Document. EPA Contract Nos. 68-03-2940 and 68-03-2943. April 1983.
- U.S. Environmental Protection Agency. EPA Design Manual: Municipal Wastewater Stabilization Ponds. Publication No. EPA-625/1-83-015. October 1983. 327 p.
- U.S. Environmental Protection Agency. Assessing the Release and Costs Associated with Truck Transport of Hazardous Wastes. Publication No. PB 84-224-468 (Contract No. 68-01-0021). Washington, DC. January 1984. 151 p.

U.S. Environmental Protection Agency. Control of Volatile Organic Compound Leaks from Synthetic Organic Chemical and Polymer Manufacturing Equipment. Research Triangle Park, NC. Publication No. EPA-450/3-86-006. March 1984.

U.S. Environmental Protection Agency. Evaluation and Selection of Models for Estimating Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Office of Air Quality Planning and Standards. Research Triangle Park, NC. Publication No. EPA-450/3-84-020. December 1984.

U.S. Environmental Protection Agency. Hazardous Waste Ranking - Assessment of Air Emissions from Hazardous Waste Treatment, Storage, and Disposal Facilities. Publication No. EPA-450/3-85-006. February 1985.

Vaught, C. C. (GCA). Air Emissions for Quiescent Surface Impoundments-- Emissions Data and Model Review, Draft Technical Note. Prepared for U.S. Environmental Protection Agency. Washington, DC. Contract No. 68-01-6871. August 1985.

Westat Corporation. National Survey of Hazardous Waste Generators and TSD's Regulated Under RCRA in 1981. Prepared for the U.S. Environmental Protection Agency. Contract No. 68-01-6861. April 1984.

Wetherold, R. G., J. L. Randall, and K. R. Williams (Radian Corporation). Laboratory Assessment of Potential Hydrocarbon Emissions from Land Treatment of Refinery Oily Sludges. Prepared for U.S. Environmental Protection Agency. Washington, DC. Publication No. EPA-600/2-84-108. June 1984.

Zepp, R. G. Quantum Yields for Reaction of Pollutants in Dilute Aqueous Solution. Environ. Sci. Technol. 12(3):327-329. 1979.

Zepp, R. G., and D. M. Cline. Rate of Direct Photolysis in Aquatic Environment. Environ. Sci. Technol. 11(4):359-366. 1977.

APPENDIX D
PROPERTIES FOR COMPOUNDS OF INTEREST

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PROPERTIES FOR COMPOUNDS OF INTEREST

This appendix contains compound-specific properties of about 700 chemicals, most of which are not included in CHEMDAT7. These data, presented as a source of information, can be easily incorporated into CHEMDAT7. Consequently, this appendix greatly increases the utility of CHEMDAT7. The chemical "universe" tested in this appendix represents those chemicals that could be encountered in TSDf and that are useful for calculating emission rates for the facilities modeled in the body of this report. The list was extracted from the GCA Physical/Chemical Database, WET model stream compositions, and the Industrial Studies Database.

The compounds listed in this appendix were not all included in CHEMDAT7 because their inclusion would seriously slow the execution time of the program, the memory requirements would prevent the program from being run on many machines, and because the data are not complete for some of the chemicals. Compounds included in the CHEMDAT7 program were selected on the basis of the estimated frequency with which they are found in hazardous wastes and on their position in prioritized lists of pollutants.

Due to the limited number of compounds for which adequate biodegradation rate data were available to determine appropriate biodegradation rate constants, these constants are provided separately as Table D-1. It is recognized that biodegradation rates can vary widely from site to site. Therefore, the following priority schedule is provided as guidance in determining the appropriate biodegradation rate constants to be employed in the emission models:

- Use site-specific biodegradation rate data where available.
- Use the rate constants suggested in Table D-1 as available.

TABLE D-1. SUMMARY OF RECOMMENDED BIODEGRADATION RATE CONSTANTS

Compound	K_{max} , mg/g-h	K_1 , L/g-h	$K_s = K_{max}/K_1$, mg/L
Acenaphthene		2.7	
Acetic acid	14	0.98	14
Acetone	1.3	1.15	1.1
Acrolein	7.8	0.36	22
Acrylonitrile	18	0.75	24
Aniline	7.1	21	0.34
Anthracene		2.2	
Arochlor (1254)		0.95	
Benzene	19	1.4	13.6
bis(2-Ethylhexyl)phthalate	0.77	0.35	2.2
Butanol (iso)	7.8	0.11	71
Butylbenzylphthalate		2.2	
Carbon tetrachloride	1.5	1.5	1.0
Chloroaniline(s)	0.27	0.86	0.31
Chlorobenzene	0.39	10	0.039
Chloroform	2.94	0.79	3.7
Chlorophenol(o)	15		
Chlorophenol(p)	6.5		
Chrysene		1.4	
Cresol(s)	23	17	1.4
Dibromochloromethane		0.035	
Dibutylphthalate	0.40	1.0	0.40
Dichlorobenzene(1,2)	2.5	0.58	4.4
Dichlorobenzene(1,3)	6.4	2.3	2.8
Dichloroethane(1,1)		2.3	
Dichloroethane(1,2)	2.1	0.98	2.1
Dichloroethene(1,1)		1.8	

(continued)

TABLE D-1 (Continued)

Compound	K_{max} , mg/g-h	K_1 , L/g-h	$K_S = K_{max}/K_1$, mg/L
Dichlorophenol (2,4)	25	3.3	7.5
Dichloropropane (1,2)	17	1.4	12
Diethylphthalate		2.7	
Dimethylphenol (2,4)	10.7	4.7	2.3
Dimethylphenol (3,4)	5.5	1.05	5.2
Dimethylphthalate	2.2	3.1	0.71
Dinitrophenol (2,4)	8.0	0.62	13
Ethanol	8.8	0.9	9.8
Ethylbenzene	6.8	2.1	3.2
Ethylene oxide (oxirane)	4.2	0.91	4.6
Fluoranthene		1.5	
Fuorene		3.5	
Formaldehyde	5.0	0.25	20
Lindane		0.031	
Methanol	18	0.20	90
Methyl ethyl ketone	2.0	0.20	10
Methylene chloride	22	0.40	55
Naphthalene	42.5	1.0	42.5
Nitrobenzene	11	2.3	4.8
Pentachlorophenol	130	3.4	38
Phenanthrene	1.5	2.2	0.67
Phenol	97	13	7.6
Propanol (2)	15	0.075	200
Styrene		0.11	
Tetrachloroethane (& ethene)	6.2	0.68	9.1
Thiobismethane	0.16	0.93	0.17
Toluene	73.5	2.4	30.6

(continued)

TABLE D-1 (Continued)

Compound	K_{max} , mg/g-h	K_1 , L/g-h	$K_s = K_{max}/K_1$, mg/L
Trichloroethane(1,1,1)	3.5	0.74	4.73
Trichloroethene	3.9	0.88	4.43
Trichlorophenol(2,4,6)		0.26	
Trinitrotoluene (alpha)	4.4	0.45	9.8
Xylenes (total)	40.8	1.8	22.7

Estimate the biodegradation rate constants using the following methodology:

- Approximate K_{\max} from available data for K_{\max} for compounds of similar structure and/or functional groups; and
- Approximate K_1 either by using the following correlation:

$$K_1 (\text{L/h/g}) = 0.135 K_{ow}^{0.38}$$

where K_{ow} = octanol-water partitioning coefficient

or by using the default (average) value for K_1 , which is $K_1 = 1 \text{ L/h/g}$, and then calculate K_s as: $K_s = K_{\max}/K_1$.

The following properties are given for each chemical (listed by name and Chemical Abstract Source [CAS] number):

- Molecular weight
- Density
- Vapor pressure at 25 °C
- Solubility
- Henry's law constant
- Diffusion coefficient in water
- Diffusion coefficient in air
- Boiling point
- Coefficients for the Antoine equation for estimating vapor pressure at temperatures other than 25 °C.

To estimate vapor pressures at temperatures other than 25 °C, the Antoine equation coefficients are used with the following equation:

$$\log_{(10)} \text{ Vapor Pressure (mm Hg)} = A - \frac{B}{T + C}$$

where

A, B, and C = the Antoine equation coefficient

T = temperature in °C.

Two approaches may be used to introduce a new compound and its properties into CHEMDAT7. First, the data for one compound in CHEMDAT7 may be replaced with data for the compound of interest in the

columns specified above. With this approach, the list of compounds in CHEMDAT7 remains constant at 62. The second approach involves appending the new compound and its properties to the existing list of chemicals in CHEMDAT7. All the equations/calculations must then be copied from one of the existing rows via Lotus 1,2,3 into the appropriate cells in the new row of the spreadsheet. As mentioned above, the inclusion in CHEMDAT7 of all or a large part of the chemicals listed in this appendix could result in increasing the time required to exercise CHEMDAT7 and could prevent its use on some microcomputers.

The properties of interest listed above, with the exception of the CAS number, mimic those in columns B, D-M, and Q of the CHEMDAT7 spreadsheet.

name	type	mw	dens	vp	hi	dl	dv	boil pt	A	B	C
2,4,6 T	P	285.48	0	5.25E-09	0	0	0	0	0	0	0
ACENAPHTHENE	A	184.21	1.07	.005	.00771	0	0	278	7.728	2534.234	241
ACENAPHTHYLENE	A	182.21	0	0	.000114	0	0	285	7.72819	2534.234	241
ACETALDEHYDE	O	44	.79	700	.000005	.000141	.124	20.8	8.005	1000.017	291
ACETAMIDE	N	0	0	0	1.2E-07	0	0	80.1	7.150005	1300.440	224
ACETATE	O	0	0	0	0	0	0	0	0	0	0
ACETIC ACID	O	60.05	1.05	15.4	.0027	.00012	.113	118	7.307	1533.313	221
ACETIC ANHYDRIDE	O	102.09	1.04	5.29	5.91E-08	9.23E-08	.236	139.6	7.149	1444.718	191
ACETONE	O	58	.79	285	.000025	.000114	.124	56.2	7.117	1210.595	221
ACETONITRILE	N	41.05	.78	90	.0000003	.000108	.128	81.6	7.119	1314.4	231
ACETOPHENONE	O	120.16	1.03	1	.000141	0	0	202.3	9.135	2878.8	271
ACETYL CHLORIDE	C	78.5	1.11	200	0	.000115	.000	52	6.948	1115.954	221
ACETYL DIETHYLMALONATE	O	194	0	0	.000194	0	0	0	0	0	0
ACETYLAMINOPROPYLENE, 2-	A	223.27	0	0	0	0	0	0	0	0	0
ACETYLENE	H	26.04	0	40	.00126	0	0	0	0	0	0
ACETYLFURAN 2		110.08	0	0	.00011	0	0	0	0	0	0
ACETYLMETHYLPHthalate 4	A	238	0	0	0	0	0	0	0	0	0
ACETYLPYRIDINE 3		125	0	0	1.25E-07	0	0	0	0	0	0
ACETYLPYRIDINE 3	O	121.15	0	0	.300	0	0	0	0	0	0
ACETYL-3-THIOUREA, 1-	N	118.15	0	0	0	0	0	0	0	0	0
ACETYL-5-HYDROXYPIRIDINE 3	O	140	0	0	.0000007	0	0	0	0	0	0
ACRIDINE ORANGE	N	285.30	0	0	0	0	0	0	0	0	0
ACRIDINE YELLOW	N	237.03	0	0	0	0	0	0	0	0	0
ACROLEIN	O	58.1	.84	244	.0000000	.000122	.105	53	0	0	0
ACRYLAMIDE	N	71.09	1.12	.012	5.2E-10	.000100	.007	0	11.23015	3030.077	27
ACRYLIC ACID	O	0	0	5.2	.0000001	.000100	.000	141.0	5.052	648.000	15
ACRYLONITRILE	N	53.1	.81	114	.000000	.000134	.122	77.4	7.038	1232.53	22
ADAMANTANE DICARBOXYLIC ACID	O	180.25	0	0	0	0	0	0	0	0	0
ADAMANTANE DICHLORIDE	C	287.1	0	0	.00104	0	0	0	0	0	0
ADONIS	N	135.13	0	0	0	0	0	220	0	0	0
ADIPIC ACID	O	146.14	1.37	.0000225	5.00E-11	0.04E-03	.0000	205	0	0	0
AFLATOXINS		324.3	0	0	0	0	0	0	0	0	0
ALACHLOR	P	280.0	0	.000022	0	0	0	0	0	0	0
ALDICARB	P	190.29	0	.0001	3.17E-08	0	0	207	6.927004	1900.701	11
ALDRIN	P	304.08	0	.000000	.000400	0	0	0	90	0	0
ALKYLAMINE CARBOXYLIC ACID N,SUB	O	200	0	0	.00001	0	0	0	0	0	0
ALLYL ALCOHOL	O	58.1	.85	23.3	.000010	.000114	.114	97	0	0	0
ALLYL CHLORIDE	C	76.53	.94	300	.371	0	0	45	0	0	0
ALPHA METHYL STYRENE	A	110	0	.070	.00001	.000114	.204	-30.0	6.923	1400.00	21
ALPHA METHYL STYRENE DIMERS	A	230	0	0	.0110	0	0	0	0	0	0
alpha-CHLORO-beta-METHYLNAPHTHALENE	A	178.05	0	0	.00003	0	0	0	0	0	0
ALPHA-HYDROXYACETALDEHYDE	O	60	0	1	.000005	0	0	0	0	0	0
ALPHA-HYDROXYADIPALDE	N	162	0	0	.000102	0	0	0	0	0	0
alpha-METHYLSTYRENE (-4)	A	110.19	0	0	.00001	0	0	0	7.0112	1535.1	2
alpha-PICOLINE		98.14	0	0	4.00E-07	0	0	0	0	0	0
AMINOBOZOIC ACID (-9)	O	137.15	0	0	4.00E-03	0	0	0	0	0	0
AMINOETHENYL, 4-	N	100.23	0	0	0	0	0	302	0	0	0
AMINOCYCLOHEXANE	N	98.10	0	0	.000240	0	0	0	0	0	0
AMINOMETHYL-3-ISOXAZOLE 5	N	114.1	0	.1	.0000701	0	0	0	0	0	0
AMINOPIHCL (-9)	N	100.12	0	.011	3.07E-05	0.04E-03	.0774	174	0	0	0
AMINOPIHCL (-9)	N	100.12	0	.000	.0000197	2.30E-03	.0774	164	-3.357	000.107	-3
AMINOPIHCL, 3-	N	100.13	0	0	0	0	0	164	0	0	0
AMINOPROPIONITRILE 3	N	75.11	0	2	9.23E-03	0	0	0	0	0	0
AMINOPYRIDINE, 4-	N	94.12	0	.002	0	0	0	273	0	0	0
AMINO-3-CHLOROTOLUENE 4	N	140	0	0	.007	0	0	0	0	0	0
AMINO-3-CHLORO-5-PHENYLCYCLOHEXANONE 2	N	163.5	0	3	.000004	0	0	0	0	0	0
AMINO-4'-CHLOROBIHENYL 4	N	205.5	0	0	0	0	0	0	0	0	0
AMINO-4-CHLOROPYRIDINE 2	N	120.50	0	0	0	0	0	0	0	0	0
AMINO-4-CHLORO-6-CYANOPYRIDINE 2	N	155	0	0	.00001	0	0	0	0	0	0
AMINO-4-NITROBIHENYL ALCOHOL 2	N	152	0	0	0.30E-03	0	0	0	0	0	0
AMINO-4-NITROTOLUENE 2	N	152.17	0	0	.00701	0	0	0	0	0	0
AMINO-5-CHLOROPYRIDINE 2	N	120.50	0	0	.000057	0	0	0	0	0	0
AMINO-9'-METHLAZOBIHENE P	N	212	0	0	.0100	0	0	0	0	0	0
AMITROLE	P	94.00	0	0	0	0	0	0	0	0	0
AMONIA	N	17.00	0	7470	.000020	.0000000	.250	-33.4	7.55400	1002.711	
AMPHETAMINE	N	135.23	0	0	.000135	0	0	0	0	0	0
AMYL ACETATE (-9)	O	130.10	.80	5.42	.000404	.000012	.004	140.4	0	0	0
AMTROLE		140.21	0	0	0	0	0	0	0	0	0
ANILINE	N	93.1	1.02	1	.000020	.0000003	.07	184	7.32	1731.515	
ANISIDINE, o-		0	0	0	0	0	0	0	0	0	0

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
ANISOLE		106.13	0	16	.711	0	0	0	0	0	0
ANTHRACENE	A	178.23	1.25	.0000013	.0575	0	0	0	8.91	3781	273.16
ANTHRAQUINONE	Q	0	1.43	3E-08	3.2E-09	0	0	388	7.289244	2411.734	157.0593
ARAMITE		334.85	0	0	0	0	0	0	0	0	0
ARSENIC ACID	I	141.98	0	0	0	0	0	168	0	0	0
ATRAZINE	P	215.08	0	.0000008	0	0	0	0	0	0	0
AURACINE	N	257.37	0	0	0	0	0	0	0	0	0
AZASERINE	N	178.13	0	0	0	0	0	0	0	0	0
AZEPINE		333.3	0	0	.00033	0	0	0	0	0	0
AZIRIDINE		43.1	0	168	.000464	0	0	0	0	0	0
BENZAL CHLORIDE	C	127	1.28	.87	.00741	0	0	287	0	0	0
BENZALDEHYDE	Q	106.13	0	1	.0000423	0	0	178.9	0	0	0
BENZALKONIUM CHLORIDE	C	192	0	0	1.92E-05	0	0	0	0	0	0
BENZEN SULFONATE	S	0	0	0	0	0	0	0	0	0	0
BENZENE	A	78.1	.87	95.2	.0055	.0000006	.008	78	6.905	1211.033	228.79
BENZENE ARSONIC ACID	Q	282.84	0	.00001	0	0	0	0	0	0	0
BENZENE SULFONIC ACID	Q	188.17	0	0	7.91E-07	0	0	0	0	0	0
BENZETHONIUM CHLORIDE	C	448.15	0	0	.0000224	0	0	0	0	0	0
BENZIDINE	A	184.23	0	0	0	.000015	0	488	7.5424	2825.8	163.258
BENZIDINE DIHYDROCHLORIDE	C	257.18	0	0	0	0	0	0	0	0	0
BENZODIOXANE-1,3		233.3	0	0	4.57E-05	0	0	0	0	0	0
BENZOFURANTHENE,3,4-	A	252.32	0	0	0	0	0	0	0	0	0
BENZOIC ACID	Q	122.13	1.27	.00784	1.82E-08	7.97E-05	0	0	9.833	3333.3	273
BENZONITRILE	N	188.07	0	1	.0000138	0	0	0	6.74631	1438.72	181
BENZOPHENONE	Q	182.23	0	0	.00011	0	0	0	7.34958	2331.4	185
BENZOPYRENE 3,4	A	252.32	0	0	.000128	0	0	0	0	0	0
BENZOQUINONE, p-	Q	188.1	0	.89	0	0	0	0	0	0	0
BENZOTHAZOLE	S	135.19	0	0	.000136	0	0	231	0	0	0
BENZOTRICHLORIDE	C	195.47	1.38	.2	.00219	0	0	221	0	0	0
BENZOYL CHLORIDE	C	113.57	0	214	0	0	0	0	7.9245	2372.1	273.16
BENZO(A)ANTHRACENE	A	228.3	1.11	1.5E-07	1.38E-09	.000000	.051	435	6.9824	2428.8	155.6
BENZO(A)PYRENE	A	252.3	1.11	.000508	1.38E-09	.000000	.043	312	9.245505	3724.353	273.16
BENZO(B)FLUORANTHENE	A	252.32	0	0	.0000281	0	0	4	0	0	0
BENZO(g,h)PERYLENE	A	278.34	0	1E-10	0	0	0	0	0	0	0
BENZO(j)FLUORANTHENE	A	252.32	0	0	0	0	0	0	0	0	0
BENZO(k)FLUORANTHENE	A	252.32	0	9.55E-11	0	0	0	488	0	0	0
BENZYL ALCOHOL	Q	108.15	0	.15	6.1E-07	0	0	0	7.19817	1632.593	172.79
BENZYL CHLORIDE	C	126.6	1.1	1.21	.000433	.0000078	.075	179.4	0	0	0
BENZYL METHYL ETHER	Q	122.16	.987	0	0	0	0	0	0	0	0
BENZ(c)ACRIDINE	A	229.29	0	0	0	0	0	0	0	0	0
BHC, alpha-	P	298.83	0	.06	0	0	0	288	0	0	0
BHC, beta-	P	298.83	0	.17	0	0	0	0	0	0	0
BHC, delta-	P	298.83	0	.82	0	0	0	0	0	0	0
BHC, gamma-	P	298.83	0	.83	4.93E-07	0	0	323.4	0	0	0
BICYCLO[2.2.1]-2,5-HEPTADIENE DICHLORIDE		158	0	0	.079	0	0	0	0	0	0
BIPHENYL	B	154.2	1.18	1	.181	0	0	254	7.245	1998.725	282.733
BIS (2-CHLOROETHOXY) METHANE	C	173.1	0	0	0	0	0	218.1	0	0	0
BISPHENOL(A)	Q	228.31	0	0	.00228	0	0	0	99	0	0
BIS(1,1,2,2-TETRACHLOROPROPYL) ETHER	C	377.7	0	875	43.5	0	0	21.3	0	0	0
BIS(2-CHLOROETHYL) ETHER	C	143	1.22	1.4	.000013	7.53E-05	.0092	178	0	0	0
BIS(2-CHLOROISOPROPYL) ETHER	C	171.1	1.11	.95	.00011	6.41E-05	.0082	189	0	0	0
BIS(2-ETHYLHEXYL) PHTHALATE		381.87	0	.0000002	.0000003	3.60E-05	.0351	384	0	0	0
BIS(CHLOROMETHYL) ETHER	C	115	1.32	38	.00021	9.38E-05	0	184	0	0	0
BROMACETONE	Q	136.98	0	0	0	0	0	136	0	0	0
BROMOBENZENE	C	157.82	1.4952	0	0	0	0	0	6.86054	1438.817	205.441
BROMOBENZYL ALCOHOL -(m)	Q	187.84	0	0	3.74E-08	0	0	0	0	0	0
BROMOBENZYL ALCOHOL -(o)	Q	187.84	0	0	3.74E-08	0	0	0	0	0	0
BROMOBENZYL ALCOHOL -(p)	Q	187.84	0	0	3.74E-08	0	0	0	0	0	0
BROMOCHLOROBENZENE P	C	191.46	0	0	239	0	0	0	0	0	0
BROMOCHLOROBENZYL ALCOHOL	C	285	0	0	.0000002	0	0	0	0	0	0
BROMOCHLOROMETHANE	C	129.39	0	0	25.9	0	0	0	6.49005	942.257	192.587
BROMOCHLOROMETHYL ACETATE	C	187	0	0	1.87E-07	0	0	0	0	0	0
BROMODICHLOROMETHANE	C	183.8	1.97	59.2	.285	0	0	98	0	0	0
BROMOETHYL ACETATE	C	167.82	0	0	.000418	0	0	0	0	0	0
BROMOETHYLENE	C	186.96	0	0	13.4	0	0	0	0	0	0
BROMOFORM	C	252.77	2.89	5.6	.000584	0	0	149.5	0	0	0
BROMOMETHANE	C	94.95	0	1508	.221	0	0	4.6	0	0	0
BROMOPHENYL PHENYL ETHER, 4-	C	249.11	0	0	0	0	0	318.1	0	0	0
BROMOPROPIONITRILE 3	C	133.98	1.615	0	0	0	0	0	0	0	0
BROMOTOLUENE 4	C	171	1.3559	0	0	0	0	0	7.00782	1612.35	206.36

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
BROMACETYL-5-	C	198.98	0	0	0	0	0	0	0	0	0
BROMO-3-CHLOROBUTADIENE 2	C	180	0	0	.00045	0	0	0	0	0	0
BROMO-4-CHLOROCYCLOHEXONE 1	C	198.5	0	0	.0000	0	0	0	0	0	0
BROMO-4-CHLORO-6-CYANOBENZYL ALCOHOL 2	C	180	0	0	.0000100	0	0	0	0	0	0
BROMO-4-CYANOMETHYL BENZOATE 2	C	246	0	0	.12	0	0	0	0	0	0
BROMO-4-CYANOMETHYL BENZOATE 3	C	241	0	0	.0041	0	0	0	0	0	0
BROMO-(1)-CHLOROETHANE-2	C	148.42	0	0	170	0	0	0	0	0	0
BUTADIENE-(1,3)	H	54.09	0	21.00	.142	.0000100	.240	-4.4	0.840	930.540	238.85
BUTANE	H	58.12	0	0	.201	0	0	0	0	0	0
BUTANEDINITRILE	N	98.09	0	0	0	0	0	205	0	0	0
BUTANENITRILE	N	68.11	0	10	0	0	0	117.9	0	0	0
BUTANOL ISO	O	74	.79	10	.0000022	.0000000	.000	0	7.47431	1314.19	186.55
BUTANOL(S)	O	74.14	0	10	.0000127	0	0	0	7.4700	1302.30	178.77
BUTANOL-1	O	74.1	.81	0.5	.0000000	.0000000	.00	0	7.4700	1302.30	178.77
BUTYL ACETATE(-m)	O	116	.80	15	.000104	0	0	0	7.127	1430.418	210.74
BUTYL ACRYLATE	O	120.2	.9	5.8	.000011	0	0	146	0	0	0
BUTYL BENZENE	A	134.22	0	1	.0003	0	0	0	0.90317	1577.905	201.37
BUTYL BENZYL PHTHALATE	A	232.30	0	.0000000	.0100	0	0	370	0	0	0
BUTYL CARBITOL		102.28	.90	.00400	.0011	0	0	0	7.74114	2050.904	195.05
BUTYL CELLULOSE	H	118.2	.9	1.01	2.00E-07	0	0	170	0.908	1300.908	172.15
BUTYL MERCAPTAN	S	98.10	0	0	.225	0	0	0	0	0	0
BUTYLAMINE	N	73.14	.7327	72	0	0	0	77.9	0.840454	2025.130	273.16
BUTYLENE GLYCOL-(1,3)	O	90.14	0	.00	3.00E-05	0	0	0	0	0	0
BUTYL-3-METHOXY PYRAZINE,3- ISO	N	100.22	0	0	0	0	0	0	0	0	0
BUTYL-0-CRESYL MONO T	O	104.25	0	0	0	0	0	0	0	0	0
BUTYL-p-CRESYL MONO T	O	104.25	0	0	0	0	0	0	0	0	0
BUTYRALDEHYDE	O	72.11	0	0	.000000	0	0	74.8	0.350044	913.50	185.40
BUTYRALDEHYDE ISO	O	72.12	0	170	.000147	0	0	0	0	0	0
BUTYRIC ACID	O	88.1	0	.04	1.70E-05	0	0	103.5	0	0	0
CACODYLIC ACID	O	137.90	0	0	0	0	0	0	0	0	0
CAMPHENE		136.28	.84	0	.00100	0	0	150	0	0	0
CANARYL	P	201.22	0	.00004	0	0	0	0	0	0	0
CARBAZOLE		157.2	0	1	44	0	0	0	0	0	0
CARBON DISULFIDE	S	76.1	1.20	300	.0100	.00001	.104	46.3	0.942	1100.11	241.50
CARBON DIFLUORIDE	I	66.02	0	0	0	0	0	-83.1	0	0	0
CARBON TETRACHLORIDE	C	153.8	1.50	113	.00	.0000000	.070	70.0	0.9330	1242.43	230
CATECHOL	P	110.11	0	10	0	0	0	240.5	0	0	0
CELLULOSE	O	804.27	0	0	.0000004	0	0	0	0	0	0
CHLORAL		147.4	1.51	50	.00007	0	0	97.0	0	0	0
CHLORALBUCEL	C	304.23	0	0	0	0	0	0	0	0	0
CHLORDANE	P	410	1.11	.00001	.0000007	0	0	175	0	0	0
CHLOROACIC ANHYDRIDE	O	0	0	0	0	0	0	0	0	0	0
CHLORINATED TARS	C	300	0	0	.000170	0	0	0	0	0	0
CHLORINAPHAZINE	C	200.2	0	0	0	0	0	0	0	0	0
CHLOROACETALDEHYDE	C	78.5	1.11	00	.000020	.0000115	.000	05	0	0	0
CHLOROACETIC ACID	C	0	0	0	1.11E-00	0	0	0	7.50016	1723.306	179.90
CHLORALLYL ALCOHOL 2	C	92.03	0	1.5	.0000100	0	0	140	0	0	0
CHLORANILINE(2)	C	127.0	0	1	.0100	0	0	0	7.50205	1900.0	220
CHLORANILINE(3)	C	127.07	0	1	.0100	0	0	0	7.50000	2073.75	215
CHLORANILINE,p-	C	127.07	0	.010	0	0	0	232	0	0	0
CHLORAZOBENZENE	C	210.7	0	0	.0100	0	0	0	0	0	0
CHLOROBENZENE	C	112.0	1.11	11.0	.00000	.0000007	.073	0	0.970	1431.05	217.50
CHLOROBENZENESULFONIC ACID (-p)	C	176.02	0	0	0.00E-00	0	0	0	0	0	0
CHLOROBENZYLATE	P	325.2	0	.0000022	0	0	0	0	0	0	0
CHLOROBENZOIC ACID,2	O	156.07	0	0	7.40E-00	0	0	0	0	0	0
CHLOROBENZOIC ACID,3-	O	156.07	0	0	0	0	0	0	0	0	0
CHLOROBENZOIC ACID,4-	O	156.07	0	0	0	0	0	0	0	0	0
CHLOROBENZOPHENONE (PARA)	C	210.07	0	.0000020	3.4E-00	0	0	330	0	0	0
CHLOROBIS(2)CHLORIDE P	C	220.07	0	0	.115	0	0	0	0	0	0
CHLOROBENZYL ALCOHOL -(m)	C	142.00	0	5.00E-00	2.00E-00	0	0	237	0.230450	2300.110	194.20
CHLOROBENZYL ALCOHOL -(o)	C	142.00	0	0	2.00E-00	0	0	0	0	0	0
CHLOROBENZYL ALCOHOL -(p)	C	142.00	0	0	2.00E-00	0	0	0	0	0	0
CHLOROBIPHENYL (-p)	C	180	0	0	.0004	0	0	0	0	0	0
CHLOROCUMARIN 2	C	180.00	0	0	.00000	0	0	0	0	0	0
CHLOROCYCLOBENZENE (1,4)	C	117.0	0	0	17.2	0	0	0	0	0	0
CHLOROCYCLOHEXANE	C	110.01	0	0	14.0	0	0	0	0	0	0
CHLOROCYCLOHEXANOL 2	C	134.01	0	0	.000200	0	0	0	0	0	0
CHLOROCYCLOHEXANOL 4	C	134.0	0	0	.00100	0	0	0	0	0	0
CHLORODIACETYL	C	120.0	0	0	0	0	0	0	0	0	0
CHLORODIMETHYL PHTHALATE 3	C	220	0	0	.115	0	0	0	0	0	0

name	type	wrt	dens	vp	hl	dl	dv	boil pt	A	B	C
CHLORDIPHENYL THIOETHER P	C	284.57	0	0	.0182	0	0	0	0	0	0
CHLORETHANE	C	64.52	.9214	0	0	0	0	0	6.98347	1636.01	238.01
CHLORETHANOL (ETHYLENE CHLORHYDRIN) 2	C	88.52	0	10	.0000100	0	0	0	0	0	0
CHLORETHYLENE	C	0	0	0	0	0	0	0	6.89117	905.01	239.48
CHLOROETHYL (2-) VINYL ETHER	C	100.55	0	24.7	.0040	0	0	100	0	0	0
CHLOROFUOROBENZENE P	C	130	0	0	103	0	0	0	0	0	0
CHLOROFUORMETHANE	C	86.40	0	0	1.71	0	0	0	0	0	0
CHLOROFORM	C	119.4	1.49	200	.00330	.00001	.104	0	6.408	929.44	196.03
CHLOROHYDROXYPHENYL 4 METHYL BENZOIC ACID	C	202.5	0	0	0	0	0	0	0	0	0
CHLOROMETHYL ACETYLENE	C	74.5	0	0	0	0	0	0	0	0	0
CHLOROMETHYL BENZOATE P	C	170.6	0	0	.0063	0	0	0	0	0	0
CHLOROMETHYL ETHYL KETONE	C	100.55	0	0	.00206	0	0	0	0	0	0
CHLOROMETHYL METHYL ETHER	C	80.51	0	224	0	0	0	59.1	6.831075	1134.219	228.0306
CHLOROMETHYL PHENYL KETONE	C	154.6	0	0	3.00E-08	0	0	0	0	0	0
CHLOROMETHYL PHENYLHYDRAZINE P	C	157	0	0	.00014	0	0	0	0	0	0
CHLOROMETHYLAMINOIMINE	C	78.5	0	0	0	0	0	0	0	0	0
CHLOROPHTHALENE, 2-	C	102.02	0	.017	.0182	0	0	250	0	0	0
CHLORONITROALKOXIDINE	C	230	0	0	.000023	0	0	0	0	0	0
CHLORONITROBENZENE (-o)	C	157.00	0	0	.00700	0	0	0	0	0	0
CHLOROPHENOL POLYMERS	C	2000	0	0	.0000001	0	0	0	0	0	0
CHLOROPHENOL-2	C	120.6	1.28	3	.0000170	0	0	175.6	6.877	1471.01	193.17
CHLOROPHENOL-3	C	120.6	1.24	.5	3.25E-08	0	0	214	0	0	0
CHLOROPHENOL-4	C	120.6	1.31	.18	1.12E-08	0	0	217	0	0	0
CHLOROPHENYL PHENYL ETHER, 4-	C	204.00	0	.0027	0	0	0	204	0	0	0
CHLOROPHENYLETHANOL 1,1	C	160.53	0	0	.00700	0	0	0	0	0	0
CHLOROPHTHALIC ANHYDRIDE 4	C	180.6	0	0	3.07E-08	0	0	0	0	0	0
CHLOROPRENE	C	68.5	.950	273	.301	.00001	.104	0	6.161	783.45	179.7
CHLOROPROPANE-1	C	70.54	.80	350	.013	0	0	0	6.92840	1110.19	227.94
CHLOROPROPANE-2	C	70.54	.87	523	.017	0	0	0	7.771	1.582	206
CHLOROPROPENE 3	C	70.6	0	351	.350	0	0	0	5.29716	418.375	128.100
CHLOROPROPIONITRILE, 3-	C	80.53	0	2.5	0	0	0	0	7.32973	1732.55	211.70
CHLOROPROPYLENE-2	C	70.53	0	0	.350	0	0	0	0	0	0
CHLOROPYRIDINE 2	C	113.55	0	1	.00140	0	0	0	0	0	0
CHLOROSTYRENE (-o)	C	130.6	0	0	.00000	0	0	0	0	0	0
CHLOROTETRAHYDROFURAN 3	C	121	0	0	.000000	0	0	0	0	0	0
CHLOROTHIOPHENOL P	C	144.02	0	0	.0723	0	0	0	0	0	0
CHLOROTOLUENE-4	C	120.6	1.07	2.8	.00400	0	0	162	0	0	0
CHLORURACIL, 5-	C	0	0	0	0	0	0	0	0	0	0
CHLORO (-p) CRESOL (-m)	C	142.6	0	.0006	1.04E-07	0	0	236	0	0	0
CHLORO (-p) PHENYLHYDRAZINE	C	142	0	0	.000204	0	0	0	0	0	0
CHLORO-1,2-ETHANE DIOL	C	90.52	0	0	9.05E-08	0	0	0	0	0	0
CHLORO-1,3-CYCLOPENTADIENE 5	C	100	0	0	.05	0	0	0	0	0	0
CHLORO-2,2-DIBROMOETHANE 1	C	222.5	0	0	.000700	0	0	0	0	0	0
CHLORO-2,3-EPOXYPROPANE, 1-	C	92.53	1.18	17	.0000023	.0000000	.000	117	0	0	0
CHLORO-2,5-DIETOPYRROLIDINE 3	C	133.5	0	0	6.00E-08	0	0	0	0	0	0
CHLORO-2-METHOXYBENZOIC ACID 4	C	187	0	0	.00374	0	0	0	0	0	0
CHLORO-2-NITROBENZYL ALCOHOL 4	C	187	0	0	3.74E-08	0	0	0	0	0	0
CHLORO-3-NITROANILINE 4	C	172.00	0	0	.000173	0	0	0	0	0	0
CHLORO-3-NITRO-5-PHENYLCYCLOHEXANE 2	C	190.5	0	3	4.13E-08	0	0	0	0	0	0
CHLORO-4-AMINOCUMARAN-8-CARBOXYLIC ACID 2C	C	213.5	0	0	0	0	0	0	0	0	0
CHLORO-4-CYANOBENZYL ALCOHOL 2	C	153	0	0	6.12E-08	0	0	0	0	0	0
CHLORO-4-HYDROXYBIPHENYL 3	C	204.00	0	4	.530	0	0	0	0	0	0
CHLORO-4-METHOXY-6-AMINOBENZOIC ACID 2	C	200	0	0	.0004	0	0	0	0	0	0
CHLORO-4-METHYL-N-METHYLBENZAMIDE 3	C	183	0	0	9.15E-08	0	0	0	0	0	0
CHLORO-4-NITROANISOLE 2	C	187.5	0	0	0	0	0	0	0	0	0
CHLORO-4-PHENYLPYRIDINE 2	C	190	0	0	.0000006	0	0	0	0	0	0
CHLORO-5-AMINO-3-PYRIDINE CARB. ACID AMIDE 2C	C	183	0	0	.000000	0	0	0	0	0	0
CHLORO-5-CYANOPHTHALIC ACID 4	C	225.5	0	0	0	0	0	0	0	0	0
CHLORO-5-CYANOTOLUENE 3	C	151	0	0	.00151	0	0	0	0	0	0
CHLORO-5-FLUOROTOLUENE 3	C	144.52	0	0	.000200	0	0	0	0	0	0
CHLORO-5-PHENOXIDIMETHYL PHTHALATE 4	C	320	0	0	.10	0	0	0	0	0	0
CHLORO-N-METHYLBENZAMIDE P	C	100.01	0	0	8.40E-08	0	0	0	0	0	0
CHLORO-p'-METHYLBIPHENYL P	C	202.09	0	0	.0101	0	0	0	0	0	0
CHLORO-p-XYLENE	C	140.01	0	00	.00141	0	0	0	0	0	0
CHOLINE CHLORIDE	C	0	0	0	0	0	0	0	0	0	0
CHROMIUM (TOTAL)	M	51.900	0	0	0	0	0	2072	0	0	0
CHRYSENE	A	228.2	1.11	5.70E-10	1.10E-09	0	0	400	0	0	0
CITRIC ACID	O	192.12	1.006	.001	6.32E-10	0	0	0	0	0	0
CITRUS RED #2	A	300.30	0	0	0	0	0	0	0	0	0
COPPER PHTHALOCYANINE	N	576.1	0	0	.00070	0	0	0	0	0	0

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
CLAMPHAN		146.15	0	1	.0002	0	0	0	0	0	0
CLAROSE	A	485	0	0	.05-00	0	0	200	0	0	0
CLEROL	O	180	1.00	.3	.0000213	0	0	195	0	0	0
CLEROL(-)	O	180.1	1.00	.00	4.43E-07	.00001	.074	202	7.500	1858.36	199.1
CLEROL(-)	O	180.1	1.00	.24	.0000220	.0000000	.074	198.8	6.911	1435.5	185.1
CLEROL(-)	O	180.1	1.00	.11	4.43E-07	.00001	.074	200	7.035	1511.00	151.1
CLEROLIC ACID	O	180	1.00	.3	.0000017	.0000000	.074	235	0	0	0
CLONALDENE	O	70.00	.05	30	.0000154	.0000102	.0000	90	0	0	0
CLONALINE		54.00	0	0	0.70	0	0	0	0	0	0
CLONE (Tetrapropylbenzene)	A	120.2	.00	4.0	.0140	.0000071	.005	153	6.963	1400.793	207.1
CLUMPHENOL-4	O	212.20	0	.000012	1.00E-00	0	0	335	0	0	0
CYANIDE	N	27	0	0	2.7E-00	0	0	0	0	0	0
CYANOBENZYL ALCOHOL P	O	119	0	0	2.30E-00	0	0	0	0	0	0
CYANOGEN	N	52.04	.05	3000	.00400	0	0	-21.2	0	0	0
CYANOGEN BROMIDE	C	195.92	0	100	0	0	0	61.4	0	0	0
CYANOGEN CHLORIDE	C	61.47	0	1000	0	0	0	12.7	0	0	0
CYANOLAMINE	N	84.00	0	0	0	0	0	0	0	0	0
CYANOMETHYL BENZOATE 4	N	0	0	0	1.2E-00	0	0	0	0	0	0
CYANOMETHYLPHthalate 4	N	300	0	0	0	0	0	0	0	0	0
CYANOPYRIDINE 3	N	104.11	0	0	.20	0	0	0	0	0	0
CYANOPYRIDINE (-)	N	104.11	0	0	.20	0.70E-00	.0750	0	0	0	0
CYANTOLINE 4	N	117.4	0	0	14.7	0	0	0	0	0	0
CYANURIC ACID	O	120.00	0	0	2.50E-00	0	0	0	0	0	0
CYCASIN	O	202.22	0	0	0	0	0	0	0	0	0
CYCLOHEXANE	H	98.2	.70	100	.0137	.0000001	.0030	81	6.041	1201.53	222.1
CYCLOHEXANOL	O	100.2	.05	1.22	4.47E-00	0.31E-00	.214	101	6.255	912.87	109.1
CYCLOHEXANONE	O	98.2	.05	4.0	4.13E-00	0.02E-00	.0704	157	7.04010	2137.192	273.1
CYCLOHEXONE	H	98.15	0	0	10.3	0	0	0	0	0	0
CYCLOHEXYL ACETATE	O	140.22	0	0	.0000711	0	0	0	0	0	0
CYCLOHEXYLAMINE	N	90.17	0	0	0	0	0	0	0.00004	1229.42	180
CYCLOHEXYLCYCLOHEXANONE 4	H	90.32	0	0	.00400	0	0	0	0	0	0
CYCLOHEXYL-2,2-DIPHENYLETHYLAMINE 2	N	257	0	0	.000257	0	0	0	0	0	0
CYCLOHEXYL-4,6-DINITROPHENOL-2-	P	200.20	0	0	0	0	0	0	0	0	0
CYCLOPENTADIENE	H	60.1	.02	4	19.3	0	0	40	40.17035	14104.70	273
CYCLOPENTADIENE 1,3	H	60.1	0	0	.00031	0	0	0	0	0	0
CYCLOPHOSPHAMIDE	N	261.1	0	0	0	0	0	0	0	0	0
CYCLOHEXYL o,o-DIMETHYL PHOS.DITHIOATE	SP	240	0	0	0	0	0	0	0	0	0
CYTOSINE	N	111.1	0	0	0	0	0	0	0	0	0
DALOMACIN	A	527.51	0	0	0	0	0	0	0	0	0
DDO,p,p'-	P	320.00	0	1.02E-00	0	0	0	0	0	0	0
DDO,p,p'-	P	310.00	0	.0000000	0	0	0	0	0	0	0
DDT	P	354.40	0	1.0E-07	.114	0	0	200	0	0	0
DIACETYL	O	80.1	0	0	.0000001	0	0	0	0	0	0
DIAMINODIPHENYL METHANE P,P'	N	190.20	0	0	0	0	0	0	0	0	0
DIAMINO-S-SULFONYL BENZYL 2,4	N	105	0	0	0	0	0	0	0	0	0
DIAZINON	P	304.30	0	.00041	0	0	0	0	0	0	0
DIBENZOPYRONE 1,2,7,8	A	302.4	0	0	0	0	0	0	0	0	0
DIBENZO(a,h)ANTHRACENE	A	270.30	0	5.2E-11	3.01E-00	0	0	524	0	0	0
DIBROMOCHLOROMETHANE	C	200.20	2.30	15	.000700	0	0	122	0	0	0
DIBROMOMETHANE-1,2	C	187.00	0	0	.0100	0	0	0	6.72140	1200.62	201
DIBROMOMETHANE	C	173.00	0	40	.000000	0	0	0	0	0	0
DIBROMO-3-CHLOROPROPANE,1,2	C	230.30	0	.0	.0000230	0	0	190	0	0	0
DIBUTYL ETHER	O	130.22	.07	7	.004	0	0	0	0	0	0
DIBUTYLPHthalate	O	270.3	1.47	.0001	2.0E-07	7.00E-00	.0430	340	9.300044	3055.114	270
DICHLORO 2-PROPANOL 1,3	C	120	1.30	.27	.00040	0	0	174	0	0	0
DICHLORO PROPANOL 2,3	C	120	1.30	7	.0000234	0	0	182	0	0	0
DICHLOROBENZENE(1,2) (-)	C	147	1.31	1.5	.00194	.0000079	.000	179	6.002553	1537.072	200
DICHLOROBENZENE(1,3) (-)	C	147.01	1.29	2.20	.00001	7.00E-00	.0002	172	0	0	0
DICHLOROBENZENE(1,4) (-)	C	147	1.29	1.2	.0010	.0000079	.000	173.4	0	0	0
DICHLOROBENZIDINE,3,3'-	C	253.13	0	0	0	0	0	0	0	0	0
DICHLOROBENZOTRIAZOLE,2,6-	P	172.01	0	.0000	0	0	0	270	0	0	0
DICHLOROBENZOPHENONE P,P'	C	261.11	0	2.07E-00	4.74E-00	0	0	353	0	0	0
DICHLOROBIPHENYL (PARA)	C	223.1	0	0	.000	5.04E-00	.003	0	0	0	0
DICHLOROBUTANE (1,4)	C	127.01	0	0	3.10	0	0	0	0	0	0
DICHLORODIFLUOROMETHANE	C	120.92	0	0000	.401	0	0	-29.8	0	0	0
DICHLORODIPHENYL METHANE	C	237	0	0	.0110	0	0	0	0	0	0
DICHLOROETHANE(1,1)	C	90.00	1.17	001	.0154	0	0	0	6.977	1174.02	220
DICHLOROETHANE(1,2)	C	90	1.20	00	.0012	.0000000	.104	0	7.025	1272.3	220
DICHLOROETHYL ETHER	C	0	0	0	0	0	0	0	0	0	0
DICHLOROETHYLENE(1,1)	C	97	1.21	020	.015	0	0	0	6.9722	1000.4	230

no	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
ICHLOROETHYLENE(1,2)	C	98.96	1.28	288	.8819	0	0	0	7.8223	1285.4	238.6
ICHLOROETHYLENE,1,2-TRANS-	C	98.94	0	331	.88532	0	0	0	6.9861	1141.9	231.9
ICHLOROMONOFLOROMETHANE	C	182.92	0	1388	921	0	0	9	0	0	0
ICHLOROPHENOL(2,4)	C	183.81	0	.12	.888848	0	0	218	7.497876	1898.859	199.3596
ICHLOROPHENOL(2,6)	C	183	0	.894	.888848	0	0	228	6.898338	1677.797	197.4596
ICHLOROPHENOXACETIC ACID(2,4)	C	221	0	298	.8821	6.46E-08	.8588	47.2	0	0	0
ICHLOROPROPANE(1,2)	C	113	1.16	48	.8823	8.73E-08	.8782	0	6.98	1388.1	222.8
ICHLOROPROPENE(1,3)	C	111	1.2	43	.88233	0	0	112	0	0	0
ICHLOROPROPYLENE,1,2- (cis)	C	118.97	0	376.841	0	0	0	92.5	6.845289	1245.558	221.6846
ICHLOROPROPYLENE,1,2-(trans)	C	118.97	0	0	0	0	0	77	0	0	0
ICHLOROPROPYLENE-2,3	C	118.97	0	0	.8129	0	0	0	0	0	0
ICHLOROSTYRENE 2,6	C	172	0	0	.8888	0	0	0	0	0	0
ICHLORTETRAHYDROFURAN 3,4	C	141	0	0	1.41E-07	0	0	0	0	0	0
ICHLORO-1,3-CYCLOPENTADIENE 5,5	C	134.5	0	0	.8873	0	0	0	0	0	0
ICHLORO-2-BUTENE 1,2	C	124.9	0	0	0	0	0	0	0	0	0
ICHLORO-2-BUTENE(1,4)	C	126	1.19	2.87	.888250	8.12E-08	.8725	158	0	0	0
ICHLORO-TRANS-ETHYLENE(1,2)	C	98.94	0	488	.886	0	0	0	0	0	0
ICHLORO-(2,6)-NITROANILINE(4)	C	285.98	0	.8888812	6.54E-08	0	0	0	0	0	0
IBLDRON	P	388.98	0	1.8E-07	.8888884	0	0	0	0	0	0
IBETHYL AMINE	N	73.14	0	0	.88731	0	0	0	5.8818	583.3	144.1
IBETHYL ETHER	O	74.14	0	0	.888285	8.61E-08	.8782	246	6.92832	1884.87	228.8
IBETHYL ETHER ACID CHLORIDE	C	138.5	0	0	0	0	0	0	0	0	0
IBETHYL PHTHALATE		222	1.12	0	.8111	0	0	298	0	0	0
IBETHYL PROPIONAMIDE,2,N	P	271.38	0	.888884	0	0	0	0	0	0	0
IBETHYL THIOETHER	S	98	0	0	.46	0	0	0	0	0	0
IBETHYL (N,N) ANILINE	N	148.28	.98	.88283	5.74E-08	5.87E-08	.8513	0	7.488	1983.57	218.5
IBETHYLBENZENE P	A	134.22	0	0	.88871	0	0	0	6.9882	1588.31	281.97
IBETHYLDIPHENYL UREA SYM	N	298.38	0	0	.8134	0	0	0	0	0	0
IBETHYLENE GLYCOL	O	188.14	0	1	.8888814	0	0	0	7.6367	1939.4	182.7
IBETHYLHYDRAZINE N,N		98.2	0	0	0	0	0	0	0	0	0
IBETHYLUREA 1,1	N	118.87	0	0	.8888858	0	0	0	0	0	0
IBHYDRO-5-OXAZOLINE (DIHYDROAZLACTONE)	O	181	0	0	0	0	0	0	0	0	0
IBISOPROPYL BENZENE (PARA)	A	182.28	0	1	.187	0	0	0	6.9833	1853.88	194.41
IBMETHOATE		229.27	0	0	9.17E-07	0	0	0	0	0	0
IBMETHOXY METHANE	O	78.1	0	488	.888121	0	0	0	0	0	0
IBMETHOXY-(3,3')-BENZIDINE	O	244.32	0	0	.88244	0	0	0	0	0	0
IBETHYL AMINE	N	46.89	0	1528	5.24E-08	0	0	6.9	7.88212	988.242	221.87
IBETHYL BENZYLAMINE N,N	N	135.23	0	0	.88135	0	0	0	0	0	0
IBETHYL BENZ(A)ANT 7,12		212.28	0	0	0	0	0	0	0	0	0
IBETHYL DISULFIDE	S	94.2	0	0	.8888815	0	0	0	7.1589	1196.58	242.88
IBETHYL FORMAMIDE	N	73.89	.96	4	.8888192	.8888183	.8939	0	6.928	1488.87	196.43
IBETHYL HYDRAZINE(1,1)		88.1	.8	157	.888124	.8888189	.188	0	7.488	1385.91	225.53
IBETHYL METHYLTHIOCARBAMATE N,N	S	119	0	0	0	0	0	0	0	0	0
IBETHYL NITROSOPROPYLAMINE N,N	N	133	0	0	.888286	0	0	0	0	0	0
IBETHYL NITROSAMINE	N	74.88	1.886	0	0	.88881	.184	153	0	0	0
IBETHYL PHTHALATE		194.2	1.19	.888187	2.15E-08	6.29E-08	.8588	283.8	4.522	788.31	51.42
IBETHYL SULFATE	S	128.14	0	.1	5.88E-07	0	0	0	0	0	0
IBETHYL SULFIDE	S	82.12	0	428	.88546	0	0	0	7.1589	1196.58	242
IBETHYLAMINOAZOBENZENE,4-	N	0	0	0	0	0	0	0	0	0	0
IBETHYLBENZIDINE 3,3	A	212.28	0	0	0	0	0	0	0	0	0
IBETHYLBENZYL HYDROPEROXIDE	H	182.21	0	.24	.888481	0	0	0	0	0	0
IBETHYLBENZ(A)ANTHRACENE(7,12)	A	258.33	0	1.82E-09	2.73E-18	4.98E-08	.8461	477	0	0	0
IBETHYLETHYLAMINE	N	73.19	1.11	28	.888385	0	0	0	7.88212	988.242	221.87
IBETHYLHYDANTOIN,5,5-	N	128.15	0	0	0	0	0	0	0	0	0
IBETHYLHYDRAZINE,1,1-	N	88.1	0	157	0	0	0	63.9	0	0	0
IBETHYLPHENOL(2,4)	O	122.18	1.84	.8873	.888821	0	0	211.5	0	0	0
IBETHYLPHENOL(3,4)	O	122.17	0	0	0	0	0	0	0	0	0
IBETHYLPHENYL CARBINOL	O	138.19	0	0	0	0	0	282	0	0	0
IBETHYL-1-NITROBENZENE 2,4	N	151.18	0	0	.88758	0	0	0	0	0	0
IBNITROBENZENE M	N	188.1	1.58	.86	.888822	7.84E-08	.279	0	4.337	229.2	-137
IBNITROPHENOL 2,4	N	184	1.88	1.178479	1.53E-07	0	0	258	7.518823	2848.872	191.7698
IBNITROTOLUENE 2,6	N	182.14	0	0	9.11E-08	0	0	0	4.372	388	-43.6
IBNITROTOLUENE(2,4)	N	182.1	1.31	.8851	4.87E-08	7.88E-08	.283	0	5.798	1118	61.8
IBNITRO-o-CRESOL(4,6)	N	198	0	.818	.888814	0	0	254.7	0	0	0
IBNOCAP	P	384.4	0	.881	188888	0	0	-188.7	0	0	0
IBNOSEB	P	248.24	0	1	.8812	0	0	0	0	0	0
IBOXANE(1,4)	O	88.2	1.83	37	.8888231	.8888182	.229	0	7.431	1554.88	248.34
IBOXON	C	322	1.83	1.52E-09	.8888812	.8888858	.184	421.4	12.88	6485.5	273
IBPHENYL ETHER	B	178.21	1.875	.82	.88224	0	0	0	0	0	0
IBPHENYL THIOETHER	B	188.28	0	0	.88931	0	0	0	0	0	0

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
DIPHENYLAMINE	B	169.2	1.16	.00075	2.70E-05	6.31E-05	.000	302	0	0	0
DIPHENYLUTIOXINE 1,3	B	265.16	0	0	.00000	0	0	0	0	0	0
DIPHENYLCHLOROMETHANE	B	202	0	0	.0001	0	0	0	0	0	0
DIPHENYLDIBUTYLENE	B	230.34	0	0	.0005	0	0	0	0	0	0
DIPHENYLETHYLENE 1,1	B	182	0	0	.00001	0	0	0	0	0	0
DIPHENYLETHANOL 1,1	B	198.14	0	0	1.30E-05	0	0	0	0	0	0
DIPHENYLHIDRAZINE(1,2)	B	184.22	0	.000023	1E-15	0	0	220	0	0	0
DIPHENYLHIDRAZINE,1,1-	B	184.24	0	40	0	0	0	0	0	0	0
DIPHENYLMETHANE	B	168.20	0	0	0	0	0	0	6.291	1261	185
DIPROPYLAMINE	N	101.22	0	0	.000053	0	0	0	0	0	0
DIPROPYLFORMAMIDE	N	129.23	0	0	0	0	0	0	0	0	0
DIVINYL KETONE	O	70	0	0	.000438	0	0	0	0	0	0
DI-n-OCTYL FTHALATE	N	300.50	0	0	.137	0	0	400	7.001507	2021.431	146.15
DI-tert-BUTYL-p-CRESOL	O	220.30	0	0	0	0	0	0	0	0	0
DOTA	0	0	0	0	0	0	0	0	0	0	0
DIOXOLAN	P	400.95	0	.00001	0	0	0	0	0	0	0
DIOXOLAN SULFATE	P	422.9	0	0	0	0	0	0	0	0	0
DIOXIN	P	300.90	0	.000002	0	0	0	0	0	0	0
EPICHLOROHYDRIN	C	92.5	1.18	17	.000023	.000000	.000	117	6.22943	2005.816	273.10
EPICHRONE	0	100.21	0	0	1.00E-07	0	0	0	0	0	0
ETHANE	H	30	0	0	0	0	0	0	6.82915	983.72	258.60
ETHANOL	O	46.1	.79	80	.000000	.000013	.123	0	6.321	1718.21	237.50
ETHANOLAMINE(mono-)	N	61.09	1.02	.4	1.22E-07	.000014	.167	172	7.400	1577.67	173.30
ETHOXYETHANOL-2	O	80	.9	5.4	6.30E-05	0	0	135	7.874	1843.5	234.2
ETHYL ACETATE PEROXIDE	O	120	0	0	.000	0	0	0	0	0	0
ETHYL ACRYLATE	O	100	.92	40	.00005	.000000	.077	100	7.904453	1897.011	273.10
ETHYL CHLORIDE	C	64.52	0	1200	.014	.0000115	.271	12.4	6.900	1030.01	238.6
ETHYL CYANIDE (PROPIONITRILE)	N	55.09	0	40	.000075	0	0	97.3	0	0	0
ETHYL ETHER	O	74.12	.71	520	.00000	.000000	.074	34.5	6.92	1004.07	228.8
ETHYL ISOPROPYL PEROXIDE	O	104	0	0	.00000	0	0	0	0	0	0
ETHYL S,S-DIPHENYL PHOSPHORODITHIOATE	O S	310.30	0	0	.000105	0	0	0	0	0	0
ETHYLACETATE	O	88.1	.9	100	.000120	9.00E-05	.0732	77	7.101	1244.95	217.8
ETHYLAMINE	N	45	0	0	0	0	0	0	7.05413	907.31	220
ETHYLENE	A	28.02	.87	10	.00044	.0000070	.075	130.2	6.975	1424.258	213.2
ETHYLENE DIAMINE	N	60.11	0	10.10002	0.40E-05	0	0	120	7.337200	1821.051	215.3
ETHYLENE DIBROMIDE	C	187.80	2.7	14	.00000	0	0	131.6	6.721	1200.82	201.7
ETHYLENE GLYCOL	O	0	1.11	.120	1.00E-07	.0000122	.100	0	6.80	2000.9	203.5
ETHYLENE GLYCOL DIMETHYL ETHER	O	90.12	0	40	.000000	0	0	0	0	0	0
ETHYLENE GLYCOL MONOMETHYL ETHER	O	70.1	0	.70	7.61E-07	0	0	0	7.9400	1790.9	236.5
ETHYLENE GLYCOL MONOMETHYL ETHER ACETATED	0	110.15	0	2	1.11E-05	0	0	0	0	0	0
ETHYLENEDIAMINE	N	60.1	0	10	0	0	0	110.5	0	0	0
ETHYLENEDIOXIDE	O	44	.87	1200	.000142	.0000145	.104	0	7.120	1054.54	237.7
ETHYLETHER	O	74.1	.71	520	.00000	.000000	.074	34.5	6.92	1004.07	228.8
ETHYLENEDIACRYLATE,3-	N	0	0	0	0	0	0	0	0	0	0
ETHYLENEDIOL,3-	P	122.17	0	1	0	0	0	0	7.400	1000	187
ETHYL(2) HEXANOL	O	130.22	0	.30	.000017	0	0	0	90	0	0
ETHYL-(2)-PROPYL-(3) ACRYLATE	0	92.5	1.10	17	.000023	.000000	.000	117	0	0	0
ETHYLENE,4-	0	102.20	0	0	0	0	0	100.5	0	0	0
FLUORANTHENE	A	202	0	.0177	.007	0	0	200	6.373	1700	118
FLUORINE	A	100	1.2	.017	.000117	0	0	0	7.701	2037.1	243.0
FLUOROACETIC ACID, SODIUM SALT	O	100.02	0	0	0E-00	0	0	0	0	0	0
FLUORACETAL,5-	0	0	0	0	0	0	0	0	0	0	0
FORMALDEHYDE	O	30	0	3000	.0000070	.0000100	.170	-14	7.195	970.8	244.0
FORMAMIDE	O	45	0	0	0	0	0	0	2.002050	19.02512	204.0
FORMIC ACID	O	0	1.22	42	.000007	1.37E-05	.070	100.7	7.501	1000.2	200.0
FRENIC	C	120.92	1.400	0000	.401	.00001	.104	-29.8	0	0	0
FUMARIC ACID	O	110.07	0	0	1.00E-05	0	0	0	0	0	0
FURAN	O	60.00	.94	500	.00034	.0000122	.104	31.4	6.975	1000.07	227.0
FURFURAL	O	90.00	1.10	2	.0000011	.0000104	.0072	101.7	6.575	1190.7	162.0
FURIC ACID	O	112.00	0	0	0	0	0	0	0	0	0
GEDSIN	0	102.31	0	0	0	0	0	270	0	0	0
GLUTARIC ACID	O	0	0	0	2.00E-05	0	0	340	6.947032	2003.118	174.0
GLYCERIN (GLYCEROL)	O	92.00	1.20	.00016	1.3E-05	0	0	0	6.105	1000	20
GLYCINAMIDE	N	74.1	0	0	1.40E-07	0	0	0	0	0	0
GLUCIDINE, NITROSO	N	90.00	0	0	0.01E-07	0	0	0	0	0	0
GLUCOSE	N	180.13	0	0	0	0	0	0	0	0	0
GLUCIDOL	O	74.00	1.11	.005	9.02E-07	0	0	102	0	0	0
HEPTACHLOR	C	370.35	1.57	.0000	.0023	0	0	19.2	90	0	0
HEPTACHLOR EPOXIDE	P	300.2	0	0	.000016	0	0	0	0	0	0

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
HEPTANE ISO	H	100.21	0	00	4.35	7.11E-03	.187	0	6.89044	1331.53	212.41
HEPTANE(-n)	H	100.22	.68	40	2.82	0	0	0	6.89677	1284.9	218.54
HEXACHLORO CYCLOPENTADIENE	C	272.77	0	.00	0	0	0	239	0	0	0
HEXACHLOROBENZENE	C	284.8	2.84	1	.00000	5.91E-03	.0542	322	0	0	0
HEXACHLOROBUTADIENE	C	280.8	1.67	.15	.0255	6.10E-03	.0501	215	0	0	0
HEXACHLOROCYCLOHEXANE (GAMMA ISOMER)	C	280.83	1.87	1.002946	.0000078	0	0	300	6.900292	2200.180	185.1500
HEXACHLOROCYCLOPENTADIENE	C	272.8	1.71	.001	.016	6.10E-03	.0501	234	0	0	0
HEXACHLOROETHANE	C	237	2.60	.05	.00005	.0000008	.00240	186	0	0	0
HEXACHLOROPENTADIENE	C	275	0	0	.0130	0	0	0	0	0	0
HEXADECANE N	H	226.30	0	0	25.2	0	0	0	0	0	0
HEXAMETHYLENEDIAMINE	N	116.21	0	0	0	0	0	205	0	0	0
HEXANE(-n)	H	86.2	.66	150	.122	7.77E-03	.2	60	6.876	1171.17	224.41
HEXANOIC ACID	O	116.09	0	0	1.00E-03	0	0	0	0	0	0
HEXANOL-1	O	102.18	.82	.812	.0000182	7.53E-03	.050	158	7.86	1761.20	196.68
HEXEN-2-ONE 5	O	100.100	.847	0	0	0	0	0	10.1158	3306.57	-50.43
HHS	surro	98.4	0	1	.00530	.0000004	.0073	0	0	0	0
HHS	surro	98.4	0	1	.00530	.0000004	.0073	0	0	0	0
HLS	surro	144	0	1	.00	9.30E-03	.0070	0	0	0	0
HLS	surro	144	0	1	.00	9.30E-03	.0070	0	0	0	0
HMS	surro	117	0	1	.00118	8.24E-03	.074	0	0	0	0
HMS	surro	117	0	1	.00118	8.24E-03	.074	0	0	0	0
HYDRAZINE	N	32.05	1.0001	14.4	6.00E-07	0	0	0	90	0	0
HYDROCYANIC ACID	O	27	.80	720	4.00E-07	.0000182	.197	25.7	7.528	1329.5	200.6
HYDROFLUORIC ACID	O	0	0	900	.000237	.000033	.300	19.7	7.217	1208.37	273.87
HYDROGEN SULFIDE	C	34.1	0	15200	.023	.0000161	.176	-60.2	7.614	885.319	250.26
HYDROQUINONE	O	0	0	0	1.57E-07	0	0	200	8.137	2461	183
HYDROXY DIMETHYL ETHER	O	78	0	0	.0195	0	0	0	0	0	0
HYDROXYMETHYLPYRIDINE-3-CARBOXYLIC ACID	20	153.14	0	0	.000305	0	0	0	0	0	0
HYDROXYACETIC ACID	O	78.05	0	0	.00019	0	0	0	0	0	0
HYDROXYCYCLOHEXANONE 4	O	114	0	3	4.13E-03	0	0	0	0	0	0
HYDROXYDIMETHYL PHTHALATE 4	O	210	0	0	.105	0	0	0	0	0	0
HYDROXYMETHYL ACETYLENE	O	58	0	0	0	0	0	0	0	0	0
HYDROXYMETHYL ISOPROPYL KETONE	O	90	0	0	.00225	0	0	0	0	0	0
HYDROXYMETHYLPHENYL CARBAMATE N	O	150	0	0	.0000150	0	0	0	0	0	0
HYDROXYMETHYLTHIOBENZENE	O	140	0	0	.007	0	0	0	0	0	0
HYDROXYMETHYL VINYL ETHER	O	130	0	0	.0025	0	0	0	0	0	0
HYDROXYMETHYL, N-METHYLETHYL AMINE N	O	89	0	0	.445	0	0	0	0	0	0
HYDROXYMETHYL-N-CHLOROMETHYLETHYLAMINE NO	NO	100.5	0	0	0	0	0	0	0	0	0
HYDROXPENTANE 3	O	88.7	0	0	.000403	0	0	0	0	0	0
HYDROXY-1,3-CYCLOPENTADIENE 5	O	81	0	0	.00405	0	0	0	0	0	0
HYDROXY-4-METHYLTETRAHYDROFURAN 3	O	100	0	0	.000250	0	0	0	0	0	0
HYDROXY-5-METHYLDIMETHYL PHTHALATE 4	O	226	0	0	.113	0	0	0	0	0	0
HYDROXY-(2)-PROPIONITRILE	O	71.09	0	.00	7.48E-03	0	0	0	0	0	0
INDANOL, 5-	N	0	0	0	0	0	0	0	0	0	0
INDENO(1,2,3-cd)-PYRENE	N	276.34	0	1E-00	5.07E-13	0	0	536	0	0	0
INDOLE	N	117.15	0	0	0	0	0	254	0	0	0
IODOCUMARAN 2		240	0	0	0	0	0	0	0	0	0
ISOPHORONE		130.21	.92	.430	5.70E-03	6.70E-03	.0023	215	0	0	0
ISOKAZOLE, 5-(AMINOMETHYL)-3-	N	114.1	0	0	0	0	0	0	0	0	0
LEAD ACETATE	M	325.28	0	0	7.34E-10	0	0	0	0	0	0
LEAD SUBACETATE	M	607.00	0	0	1.47E-09	0	0	0	0	0	0
LEUCINE		131.2	0	0	5.4E-07	0	0	0	0	0	0
LHS	surro	97.3	0	1	1.58E-07	9.04E-03	.0027	0	0	0	0
LHS	surro	97.3	0	1	1.58E-07	9.04E-03	.0027	0	0	0	0
LHS	surro	78.4	0	1	1.58E-07	.0000113	.1	0	0	0	0
LHS	surro	78.4	0	1	1.58E-07	.0000113	.1	0	0	0	0
LINDANE	P	250.05	0	0	0	0	0	0	0	0	0
MALATHION	P	330.38	0	.00004	0	0	0	0	0	0	0
MALEIC ACID	O	116.07	0	0	1.47E-09	0	0	0	0	0	0
MALEIC ANHYDRIDE	O	98.1	.93	0	4E-08	.0000111	.006	202	0	0	0
MELAMINE	N	126.2	0	50	.003	0	0	0	0	0	0
MERCURY	M	200.59	0	.002	.0114	0	0	350.58	0	0	0
MESTYL OXIDE	O	90.15	0	0	3.51E-07	0	0	0	0	0	0
METHACRYLIC ACID	O	86.1	0	.1	.0000113	0	0	0	0	0	0
METHANE	H	16.04	0	0	1.34	0	0	0	0	0	0
METHANE SULFONIC ACID	O	90.11	0	0	4.81E-07	0	0	0	0	0	0
METHANETHIOL	S	48.11	0	1520	.00418	0	0	0	0	0	0
METHANOL	O	32	.79	114	.0000027	.0000104	.15	0	7.897	1474.00	229.13
METHAPYRILENE	N	251.30	0	3	0	0	0	0	0	0	0
METHOMYL		102.2	0	.00005	8.11E-07	0	0	0	0	0	0

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
METHANOACETIC ACID	O	90	0	0	.000018	0	0	0	0	0	0
METHANOACETONITRILE	N	71.04	0	0	.000170	0	0	0	0	0	0
METHOCHLOR	P	345.06	0	0	0	0	0	0	0	0	0
METHOXYBENZOL P	O	124.15	0	0	3.1E-07	0	0	0	0	0	0
METHYL 1-PHENYLE 2	H	94	0	0	10.5	0	0	0	6.83029	1121.3	229.687
METHYL ACETATE	O	74.1	.92	235	.000102	.00001	.104	54	7.805	1157.63	219.73
METHYL ACRYLATE	O	86.1	0	80	1.44E-07	0	0	77	6.830048	1193.779	224.625
METHYL ACRYLONITRILE	N	67.1	0	85	.302	0	0	0	0	0	0
METHYL AMINE	N	31.05	0	1820	.00030	0	0	0	7.3300	1011.5	233.3
METHYL AMINOACETYLENE	N	54	0	0	.135	0	0	0	0	0	0
METHYL AZIDINE 2	N	57.1	0	0	0	0	0	0	0	0	0
METHYL BENZYL ALCOHOL 4	O	130	0	0	.0000130	0	0	0	0	0	0
METHYL BIPHENYL (-p)	B	160.45	0	0	.00042	0	0	0	0	0	0
METHYL CHLORIDE	C	50.5	.95	3030	.00014	.0000005	.125	-24	7.508	948.53	249.34
METHYL CHLORACETAMIDE N	C	107.5	0	0	.0000105	0	0	0	0	0	0
METHYL CHLOROCARBONATE	C	94.5	0	0	.235	0	0	0	0	0	0
METHYL CHOLANTRINE 3	A	290.38	0	0	.000134	0	0	0	0	0	0
METHYL COLMANN 2	0	100.10	0	0	.00001	0	0	0	0	0	0
METHYL CYCLOHEXANE	H	98.2	0	43	.079	0	0	0	6.823	1270.76	221.42
METHYL ETHER	O	40.00	0	3000	.00010	0	0	0	0	0	0
METHYL ETHYL KETONE	O	72.1	.92	100	.0000435	.0000005	.0000	70.6	6.97421	1200.6	216
METHYL FORMATE	O	60.05	0	800	.13	0	0	0	3.827	3.82	-11.9
METHYL HONAZINE	N	40.00	0	40.0	3.44E-05	0	0	0	6.5702	1207.5	181.4
METHYL IODIDE	N	141.04	0	400	.00003	0	0	42.4	0	0	0
METHYL ISOMAL KETONE	O	114	.8	4.53	.000130	0	0	144	0	0	0
METHYL ISOBUTYL KETONE	O	100.2	.8	15.7	.0000405	.0000070	.075	115.8	6.572	1168.4	191.9
METHYL ISOCYANATE	N	67.05	0	0	0	0	0	30.1	0	0	0
METHYL ISOPROPYL KETONE	0	98.15	0	0	.000400	0	0	0	0	0	0
METHYL METHACRYLATE	O	100.1	.95	30	.000005	.0000000	.077	101	8.400	2050.5	274.4
METHYL METHANESULFONATE	S	130.13	0	0	0	0	0	203	0	0	0
METHYL NAPHTHALENE(1-)	A	142.10	0	.1	.00071	0	0	0	7.83502	1820.945	195
METHYL NAPHTHALENE(2-)	A	142.10	0	.0003	.000000	0	0	0	7.0005	1840.254	198.4
METHYL PARATHION	P	263.23	0	.0000000	0	0	0	0	0	0	0
METHYL PROPENE 2	H	58	0	0	7	0	0	0	6.58405	805.25	234.64
METHYL SULFURIC ACID	S	112.1	0	3	5.61E-07	0	0	0	0	0	0
METHYL TETRAHYDROFURN 2	O	87	0	0	.0210	0	0	0	0	0	0
METHYL THIOPHENOL 4	S	134	0	3	4.4E-07	0	0	0	0	0	0
METHYL THIOURACIL	N	142.10	0	0	0	0	0	0	0	0	0
METHYLACETONITRILE(ACETONE CYANOHYDRIN)	N	85.12	0	.8	.300	0	0	95	0	0	0
METHYLENE CHLORIDE	C	95	1.34	430	.00019	.0000117	.101	30.8	7.400	1325.9	252.0
METHYLENE-BIS (2-CHLOROMETHYLENE), 4,4'-	N	257.15	0	0	0	0	0	0	0	0	0
METHYLFURN 2	O	82.11	0	0	2.74E-05	0	0	0	0	0	0
METHYLFURFURN 5	O	110.11	0	0	2.2E-07	0	0	0	0	0	0
METHYLINDOACETIC ACID	O	100	0	0	.000001	0	0	0	0	0	0
METHYLISOBORNOL, 2-	O	100	0	0	0	0	0	0	0	0	0
METHYLPHENYL CARBONATE N	O	140	0	0	.000014	0	0	0	0	0	0
METHYLTRIN TRICHLORIDE	C	240.00	0	0	.0000024	0	0	0	0	0	0
METHYL-1,3-CYCLOPENTADIENE 5	H	66.14	0	0	.0401	0	0	0	0	0	0
METHYL-2,3,4-TRIMOROQUINOLINE N	O	140	0	0	.0000140	0	0	0	0	0	0
METHYL-3-AMINOETHYLAMINE	A	74.05	0	0	.0105	0	0	0	0	0	0
METHYL-3-MORPHOETHYLAMINE	N	75.11	0	.7	3.40E-05	0	0	0	0	0	0
METHYL-3-METHOXYAZIRIDINE 1	N	67	0	0	4.30E-07	0	0	0	0	0	0
METHYL-3-ACETILCYCLOPENTADIENE 1	H	100	0	0	.0003	0	0	0	0	0	0
METHYL-3-NITROBENZYL ALCOHOL 4	N	160	0	0	6.72E-05	0	0	0	0	0	0
METHYL-4-NITROBENZYL ALCOHOL 2	N	157.10	0	0	3.34E-05	0	0	0	0	0	0
METHYL-5-THIOACETYLDOHNO1,3THIAZOLE 4	N	180	0	0	0	0	0	0	0	0	0
METHYL-PHENYLETHYLAMINE N	N	135.23	0	0	.00135	0	0	0	0	0	0
METHYL-p'-METHYLTRIPHENYL PHOSPHINE P	B	292.3	0	0	.0140	0	0	0	0	0	0
METHYL-TERTIARY-BUTYL ETHER	O	0	0	0	0	0	0	0	0	0	0
METHYL-TRIMORO-1,3-THIAZOLE 4	N	105	0	0	.000105	0	0	0	0	0	0
MMS	surro	80.9	0	1	.000008	.0000110	.005	0	0	0	0
MMS	surro	80.9	0	1	.000008	.0000110	.005	0	0	0	0
MMS	surro	112	0	1	.0000222	.0000000	.070	0	0	0	0
MMS	surro	112	0	1	.0000222	.0000000	.070	0	0	0	0
MMS	surro	57	0	1	.0000400	.0000110	.115	0	0	0	0
MMS	surro	57	0	1	.0000400	.0000110	.115	0	0	0	0
METONACIN C	N	334.37	0	0	0	0	0	0	0	0	0
MMS	N	147.00	0	0	0	0	0	0	0	0	0
MONOCHLOROFLUOROMETHANE	C	80.5	0	7400	.420	0	0	-45.3	0	0	0
MORPHOLINE	N	67.12	1	10	.0000573	.0000000	.001	0	7.71813	1745.8	235

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
MUSTARD GAS		159.97	0	.89	0	0	0	217	0	0	0
NAPHTHALENE	A	128.2	1.14	.23	.88848	.888875	.859	218	7.81	1733.71	281.88
NAPHTHOL, alpha-	O	0	0	0	0	0	0	0	7.28421	2877.58	184
NAPHTHOL, beta-	O	144.17	0	16	0	0	0	0	7.347	2138	183
NAPHTHOQUINONE-1,4	O	158	1.42	.111	.8888231	0	0	188	0	0	0
NAPHTHYLAMINE, alpha-	N	143.19	0	0	0	0	0	388.8	0	0	0
NAPHTHYLAMINE, beta-	N	143.19	0	0	0	0	0	388.1	0	0	0
NEOPENYL GLYCOL	O	184	0	3.418916	1.8E-08	0	0	298	8.219421	2178.788	199.7398
NIACIN		128.12	0	0	8.18E-07	0	0	0	0	0	0
NIACINAMIDE		122.14	0	0	1.22E-08	0	0	0	0	0	0
NICKEL CYANIDE	M	118.75	0	0	.8888564	0	0	0	0	0	0
NITROANILINE P	N	138.14	0	.8815	.888227	0	0	0	9.5596	4839.73	273.18
NITROANILINE(-o)	N	138.14	1.44	.883	.8888885	.888888	.873	0	8.888	3338.5	273.18
NITROBENZENE	N	123.1	1.2	.3	.8888131	.8888888	.878	0	7.115	1748.8	281.8
NITROBENZENESULFONYL CHLORIDE P	N	221.88	0	0	.888443	0	0	0	0	0	0
NITROBENZYL ALCOHOL P	N	153.88	0	16	.888281	0	0	0	0	0	0
NITROBIPHENYL, 4-	B	198.21	0	0	0	0	0	348	0	0	0
NITROCELLULOSE	N	188.88	0	0	.881	0	0	0	0	0	0
NITROGEN MUSTARD N-OXIDE	O	172.87	0	0	0	0	0	0	0	0	0
NITROGLYCERIN	N	227.89	1.6	.8888	8E-19	0	0	288	0	0	0
NITROMETHANE	N	81.85	0	27.8	.8836	0	0	0	7.28188	1448.94	227.8
NITROPHENOL, 2-	N	139.11	0	0	0	0	0	218	0	0	0
NITROPHENOL, 4-	N	139.11	1.4	2.2	.88834	0	0	278	0	0	0
NITROPROPANE 2	N	89.89	0	0	.888223	0	0	0	0	0	0
NITROSOBENZYL ALCOHOL 4	N	136	0	0	.8888136	0	0	0	0	0	0
NITROSDIMETHYLAMINE N	N	74.88	0	0	0	0	0	154	0	0	0
NITROSDIPHENYLAMINE N	N	198.23	0	0	0	0	0	0	0	0	0
NITROSDI-n-PROPYLAMINE N	N	138.19	0	0	0	0	0	288	0	0	0
NITROSMORPHOLINE	N	118.14	0	0	0	.88881	.859	225	0	0	0
NITROSPYRROLIDINE N	N	188.12	0	0	12.5	0	0	0	0	0	0
NITROSO-N-METHYLUREA N	N	188.1	0	0	.8888518	0	0	0	0	0	0
NITROTOLUENE (-p)	N	137.13	0	1	.888488	0	0	0	8.9948	1728.39	184.9
NITRO-4-METHYLBENZOATE 3	N	196	0	0	0	0	0	0	0	0	0
NONYLPHENOL	O	228.39	0	0	0	0	0	315	0	0	0
OCTAMETHYLPYROPHOSPHORAMIDE	P	288.25	0	.8888	0	0	0	0	0	0	0
OCTANE	H	114.3	.7	17	3.87	0	0	125.7	6.918	1351.99	289.16
OCTANOL 1	O	138.3	.8269	.124	.8888434	0	0	0	12.8781	4888.8	319.9
OCTANOL 2	O	138.3	.8287	0	0	0	0	0	6.3888	1888.4	122.5
OCTANOL 3	O	138.3	.8216	0	0	0	0	0	5.2215	588.3	84.7
OCTANOL 4	O	138.3	.8192	0	0	0	0	0	5.7388	788.5	89.5
OIL	H	178	0	0	0	0	0	0	0	0	0
ODDIETHYLOETH. THIOETH. PHOSPHOROTHIOATE N		288.38	0	1.3E-13	2.21E-14	0	0	549.5	0	0	0
OXALIC ACID	O	98.84	0	0	4.5E-09	0	0	0	0	0	0
OXAMIC ACID	O	89.88	0	0	.888889	0	0	0	0	0	0
PARABROMOPHENOL	P	173.81	0	0	0	0	0	238	0	0	0
PARAFORMALDEHYDE	O	188	0	0	.881	0	0	0	0	0	0
PARALDEHYDE	O	132.3	.99	25.3	.8888367	0	0	125	0	0	0
PARATHION	P	291.3	1.28	.888	.888848	0	0	375	0	0	0
PCB 1878	B	257.9	0	0	.8136	0	0	0	0	0	0
PCB 1221	B	288.7	0	0	.888228	0	0	0	0	0	0
PCB 1232	B	232.2	0	0	0	0	0	0	0	0	0
PCB 1242	B	288.5	0	0	0	0	0	0	0	0	0
PCB 1248	B	298.5	0	0	0	0	0	0	0	0	0
PCB 1254	B	225.1	1.11	.88888	.888298	0	0	0	0	0	0
PCB 1288	B	375.7	0	0	0	0	0	0	0	0	0
PCB'S	B	0	0	.88488	.888884	.888888	0	0	0	0	0
PENTACHLOROBENZENE	C	288.34	1.61	.8848	.8873	.8888883	.857	277	0	0	0
PENTACHLOROETHANE	C	282.3	1.67	4.4	.821	.8888873	.888	162	6.74	1378	197
PENTACHLORONITROBENZENE	C	296.38	0	0	0	0	0	0	0	0	0
PENTACHLOROPHENOL	C	288.4	1.98	.885	.8888882	.8888881	.858	318	0	0	0
PENTADISE 1,2	H	87.1	0	0	0	0	0	0	6.9182	1184.991	228.85
PENTAERYTHRITOL		138.15	0	0	0	0	0	0	0	0	0
PENTAERYTHRITOL TETRANITRATE	N	318.17	0	0	.8888316	0	0	0	0	0	0
PERCHLOROMETHYL MERCAPTAN	C	188.88	0	0	0	0	0	147	0	0	0
PERYLENE	A	252.3	0	0	0	0	0	0	0	0	0
PHENACETIN	A	178.24	0	.88872	2.23E-07	0	.857	288	0	0	0
PHENANTHRENE	A	178.22	1.18	.88821	.88885	0	0	348	0	0	0
PHENOL	O	94.1	1.87	.341	4.54E-07	.8888881	.882	182	7.133	1516.79	174.95
PHENOL, 3-(1,1-DIMETHYLETHYL)-	O	0	0	0	0	0	0	0	0	0	0
PHENOTHAZINE		198.28	0	0	.8199	0	0	0	0	0	0

name	type	mw	dens	vp	nl	dl	dv	boil	ph	A	B	G
PHENYL ISOCYANATE	N	119.13	0	0	.0019	0	0	0	0	0	0	0
PHENYL MERCURIC ACETATE	M	336.7	0	.00000	1.2E-08	0	0	0	0	0	0	0
PHENYLACETIC ACID	O	136.14	0	0	8.2E-07	0	0	0	0	0	0	0
PHENYLACETIC PEROXIDE	O	182	0	0	.0000182	0	0	0	0	0	0	0
PHENYLCYCLOHEXANONE 4	O	176	0	0	.00076	0	0	0	-7.700	188.4	-146.6	0
PHENYLENE DIAMINE(-o)	N	108.14	1.14	.020	1.12E-08	0	0	294	0	0	0	0
PHENYLENE DIAMINE(-p)	N	108.14	0	.00797	1.12E-08	0	0	258	0	0	0	0
PHENYLENE DIAMINE(-m)	N	108.14	0	.0046	1.12E-08	0	0	257	0	0	0	0
PHENYLHYDRAZINE	N	168.18	0	0	.000168	0	0	0	0	0	0	0
PHENYLPHENOL P	O	170.2	0	10	.0002	0	0	0	0	0	0	0
PHENYLTHIOUREA	S	182.22	0	0	0	0	0	320	0	0	0	0
PHOSITE		250.4	0	24.7	4.37E-07	0	0	118.5	0	0	0	0
PHOSGENE		98.92	0	1300	.171	1.12E-08	.100	8.1	0	0	0	0
PHOSPHINE		34	0	2000	.227	0	0	-87.70	0	0	0	0
PHthalic ACID	O	148.14	1.50	121	.0132	.0000000	.004	72.8	0	0	0	0
PHthalic ANHYDRIDE	O	148.1	1.38	.0015	.0000000	.0000000	.071	204	0	0	0	0
PHthalimIDE	N	147.1	0	118	.0114	0	0	60.5	0	0	0	0
PICOLINE(2-)		98.12	.98	10.4	.000127	.0000000	.076	128.8	7.802	1415.73	211.63	0
PINE(alpha-)		130.2	.86	5	.448	0	0	0	0	0	0	0
PIPERAZINE		88.14	0	3.72	.0000422	0	.001	140	0	0	0	0
POLYCHLORINATED BIPHENYLS	S	200	1.46	0	.000204	.00001	.104	0	0	0	0	0
POLYCYCLIC KETONE 0	O	1000	0	0	.05	0	0	0	0	0	0	0
PROPANE	H	44.09	0	700	.002	0	0	0	0	0	0	0
PROPANE SULFONE,1,3-		122.14	0	0	0	0	0	180	0	0	0	0
PROPANE,2,2'-OXYBIS(2-CHLORO-	C	0	0	0	0	0	0	0	0	0	0	0
PROPANOIC ACID	O	74.08	0	0	0	0	0	141.1	0	0	0	0
PROPANOL	O	60.1	.0007	0	0	0	0	97.2	7.84767	1400.21	204.6-	0
PROPANOL ISO	O	60.09	.70	42.8	.00015	.0000104	.000	82.4	0.117	1500.92	219.6-	0
PROPIOLACTONE b	O	72.1	0	3.4	9.22E-07	0	0	0	0	0	0	0
PROPIONALDEHYDE	O	58.08	.51	300	.00115	.0000114	.102	40.5	16.2315	2050.02	-44.15	0
PROPIONIC ACID	O	74.08	0	10	.0000407	0	0	0	0	0	0	0
PROPIONITRILE	N	55.08	0	40	.000275	0	0	0	5.270-	605.52	159.1	0
PROPYL ACETATE ISO	O	102.13	.87	73.95302	.000017	0	0	86.2	0.843416	1231.175	222.5	0
PROPYL AMINE ISO	N	59.08	0	400	.000050	0	0	0	0	0	0	0
PROPYL ETHER ISO	O	102.10	0	100	.00224	0	0	68.4	0.834001	1105.131	226.2	0
PROPYL THIOURACIL	S	170.23	0	0	0	0	0	0	0	0	0	0
PROPYLENE	H	42.12	0	7000	2.11	0	0	0	0	0	0	0
PROPYLENE GLYCOL	O	76.11	1.04	.3	.0000015	.0000102	.003	180	0.200243	2005.9	203.5	0
PROPYLENE OXIDE	O	58.1	.83	825	.00134	.00001	.104	33.9	0.276775	1658.884	273.1	0
PROPYLBENZENE,1,2-	N	57.09	0	0	0	0	0	65	0	0	0	0
PROPYL(-n) ACETATE	O	102.13	.80	36	.000204	0	0	101.6	7.816	1282.29	208.0	0
PROPYL(-n) BENZENE	A	120.19	.80	2.5	.00000	0	0	0	0	0	0	0
PROPYL-3-METHOXY PYRAZINE,2- ISO	N	152.2	0	0	0	0	0	0	0	0	0	0
PROPYN-1-OL 2(PROPARGOL)	O	58.08	0	11.6	0.0000000	0	0	113.6	0	0	0	0
PTERINE	A	202.3	1.27	4.2E-00	7E-00	0	0	404	0	0	0	0
PYRIDINE	O	79.1	.98	20	.0000230	.0000070	.001	115.5	7.841	1373.8	214.5	0
PYRIDINIUM BROMIDE	S	330	0	0	1.00E-00	0	0	0	0	0	0	0
QUINALDINE	A	0	0	0	0	0	0	0	0	0	0	0
QUINOLINE	A	120.10	0	0	0	0	0	237.7	0	0	0	0
RESERPINE	A	600.7	0	.0002	.00200	0	.002	205	0	0	0	0
RESORCINOL	O	0	0	.00020	1.00E-00	.0000007	.070	200	0	0	0	0
SACCH. CYCLOHEX. DIMETH. PHOS. DITHIATE	S	346	0	0	.000173	0	0	0	0	0	0	0
SACCHARIN	O	183.10	0	0	0	0	0	0	0	0	0	0
SAPPOL		162.10	0	0	0	0	0	234.5	0	0	0	0
SILVEX	P	200.51	0	0	0	0	0	0	0	0	0	0
SDIAZINE		201.7	0	0.1E-00	0.00E-10	0	0	0	0	0	0	0
SODIUM ACETATE	M	82.04	0	0	4.1E-00	0	0	0	0	0	0	0
SODIUM DODECYL SULFATE-	M	288.38	0	0	0	0	0	0	0	0	0	0
SODIUM DODECYLBENZENE SULFONATE	M	294.1	0	0	.0000142	0	0	0	0	0	0	0
SODIUM FORMATE	M	60.01	0	0	1.7E-00	0	0	0	0	0	0	0
STREPTOZOTOCIN	N	205.22	0	0	0	0	0	0	0	0	0	0
STRYCHNIDIN-10-ONE,2,3-DIMETHOXY-		304.45	0	0	1.40E-00	0	0	0	0	0	0	0
STRYCHNINE	P	334.42	0	1E-00	0	0	0	0	0	0	0	0
STYRENE	A	104.2	.9	7.3	.00201	.000000	.071	145	7.14	1574.51	224.1	0
SUCCINIC ACID	O	118.09	0	0	1.74E-00	0	0	0	0	0	0	0
SUCCINIMIDE	O	90.1	0	0	3.3E-00	0	0	0	0	0	0	0
SULFANILIC ACID	S	173.2	0	0	.0000010	0	0	0	0	0	0	0
SULFIDE	S	34.08	0	0	0	0	0	-0	0	0	0	0
TAB	A	400	0	0	.002	0	0	0	0	0	0	0
TETRAFLOROQUINONE	C	240	0	0	0	0	0	0	0	0	0	0

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
TETRACHLOROBENZENE(1,2,3,4)	C	218.9	0	.019	.0027	0	0	254	0	0	0
TETRACHLOROBENZENE(1,2,3,5)	C	218.9	0	.00	.00428	0	0	246	0	0	0
TETRACHLOROBENZENE(1,2,4,5)	C	218.9	1.00	.00	.00428	0	0	246	0	0	0
TETRACHLORODIBENZOFURAN 2,3,7,8	C	306	0	0	.000153	0	0	0	0	0	0
TETRACHLORODIBENZOFURAN 2,3,7,8-	C	306.96	0	0	0	0	0	0	0	0	0
TETRACHLORODIBENZO-p-DIOXIN,2,3,7,8-	C	321.96	0	7.4E-16	0	0	0	0	0	0	0
TETRAETHYLENE(1,1,1,2)	C	100	1.0	0.5	.002	.0000079	.071	140.3	0.898	1305.88	200.74
TETRAETHYLENE(1,1,2,2)	C	100	1.50	0.5	.00030	.0000079	.071	140.2	0.631	1228.1	179.9
TETRAETHYLENE	C	100.83	1.024	19	.029	.0000002	.072	121.4	0.978	1306.92	217.53
TETRAETHYLENE	C	100.83	1.024	19	.029	.0000002	.072	121.4	0.978	1306.92	217.53
TETRAETHYLENE(2,3,4,6)	C	231.9	0	.00	4.53E-00	0	0	184	0	0	0
TETRAETHYLENE(2,3,5,6)	C	231.9	0	.01	111	0	0	0	0	0	0
TETRAETHYLENE(1,1,2,3)	C	179.86	0	0	.00000	0	0	0	0	0	0
TETRAETHYL LEAD	M	323.46	1.053	.36	.0000	0	0	0	0	0	0
TETRAETHYLDITHIOPYROPHOSPHATE	P	322.31	0	.00017	0	0	0	0	0	0	0
TETRAETHYLDITHIOPYROPHOSPHATE	S	322.34	0	3.0E-07	7.03E-00	0	0	0	0	0	0
TETRAETHYLENE GLYCOL	O	194.20	0	1	.0000120	0	0	0	0	0	0
TETRAETHYLPYROPHOSPHATE	P	290.2	0	.000155	0	0	0	0	0	0	0
TETRAHYDROFURAN	O	72.12	.00	72.1	.000040	.0000105	.000	67	0.906	1202.29	220.26
TETRAHYDRONAPHTHALENE,1,2,3,4-	O	132.2	0	0	0	0	0	207.6	0	0	0
TETRALIN	0	0	0	0	0	0	0	0	7.07055	1741.3	200.20
TETRAHYDROMETHANE	N	100.00	0	0	0	0	0	120	0	0	0
THIOACETAMIDE	S	75.13	0	0	0	0	0	0	0	0	0
THIOBENZYL ALCOHOL P	S	124.21	0	0	.0000124	0	0	0	0	0	0
THIOCYANATE (TOTAL AS SCN-)	S	0	0	0	0	0	0	0	0	0	0
THIOFANOL	P	210.36	0	.00017	0	0	0	315	0	0	0
THIOMETHANOL	S	40.11	0	0	0	0	0	0	0	0	0
THIOPHENOL	O	110.17	0	0	0	0	0	100.7	0	0	0
THIOPROPIONAMIDE 2	S	100	0	0	5.15E-00	0	0	0	0	0	0
THIOSEMICARBAZIDE	S	91.14	0	.01	0E-00	0	0	320	0	0	0
THIOUREA	N	76.12	1.41	145	.00016	0	.107	102	0	0	0
THIOUREA,1-(o-CHLOROPHENYL)-	N	100.00	0	.000002	0	0	0	323	0	0	0
THURAM	P	240.41	0	20	0	0	0	0	0	0	0
THYMINE	N	126.12	0	0	0	0	0	0	0	0	0
TOLUENE	A	92.4	.07	30	.00000	.0000000	.007	110.7	0.964	1344.8	219.48
TOLUENE DIAMINE(2,4)	N	122	1.11	.001	0.00E-00	0	0	0	0	0	0
TOLUENE DIISOCYANATE(2,4)	N	174.16	1.2	.00	.0000000	.0000002	.001	251	0	0	0
TOLUENE2-DIAZOBIIS-METATOLUENEDIAMINE HCIN	N	420	0	0	.0000420	0	0	0	0	0	0
TOLUENEDIAMINE,2,4-	N	122.17	0	3.254333	0	0	0	140	7.427020	1632.941	211.1300
TOLUENEDIAMINE,2,6-	N	122.17	0	0	0	0	0	0	0	0	0
TOLUENEDIAMINE,3,4-	N	122.17	0	0	0	0	0	206	0	0	0
TOLUIC ACID (para-)	O	130.10	0	.00031	5.0E-00	0	0	276	0	0	0
TOLUIC ALDEHYDE	O	120.14	1.00	.10	.000253	0	0	204	0	0	0
TOLLUDINE HYDROCHLORIDE,o-	C	143.02	0	0	0	0	0	242	0	0	0
TOLLUDINE P	A	107.17	1.040	1	.0000191	0	0	200.0	7.77229	2317.300	273.10
TOLLUDINE (-0)	A	107.17	.900	.242	0	0	0	200.4	0.825311	2015.070	273.10
TOXAPHENE	0	414	1.11	.3	.00400	0	0	0	90	0	0
TRIBROMOMETHYLPHOSPHATE	C	340.07	0	0	.0000347	0	0	0	0	0	0
TRIBUTYL PHOSPHOTRITHIOATE SSS	O	314.04	0	0	.000157	0	0	0	0	0	0
TRIBUTYLPHOSPHATE	O	200.32	0	127	0	0	0	200	0	0	0
TRICHLORANISOLE,2,3,6-	C	211.40	0	0	0	0	0	227	0	0	0
TRICHLOROBENZENE 1,2,3	C	181.40	0	0	0	0	0	221	0	0	0
TRICHLOROBENZENE 1,2,4	C	181.5	0	.10	.00142	0	0	213	0	0	0
TRICHLOROBENZENE 1,3,5	C	181.5	0	.23	.0200	0	0	200.5	0	0	0
TRICHLOROBUTANE 1,2,3	C	161.40	0	4.30	4.00	.0000072	.000	0	0	0	0
TRICHLOROETHANE 1,1,1	C	133.4	1.33	123	.0172	.0000000	.070	75	0.043	2130.0	302.0
TRICHLOROETHANE 1,1,2	C	133.4	1.3	25	.00074	.0000000	.070	74	0.051	1314.4	209
TRICHLOROETHYLENE	C	131.4	1.4	75	.0001	.0000001	.079	87	0.510	1010.0	192.7
TRICHLOROFLUOROMETHANE	C	137.4	1.49	700	.0000	.0000007	.007	23.0	0.004	1043.004	230.00
TRICHLOROPHENOL (2,4,6-	P	197.46	0	400	0	0	0	253	0	0	0
TRICHLOROPHENOL (2,4,6)	0	197.46	0	.0073	.0000177	0	0	244.5	0	0	0
TRICHLOROPROPANE(1,1,1)	C	147.43	0	3.1	.029	.0000079	.071	107	0	0	0
TRICHLOROPROPANE(1,1,2)	C	147.43	0	0.04	.029	0	0	140	0	0	0
TRICHLOROPROPANE(1,2,2)	C	147.30	0	1.37	.029	0	0	124	0	0	0
TRICHLOROPROPANE(1,2,3)	C	147.4	0	3	.020	.0000079	.071	100	0.003	700.2	243.23
TRICHLOROPROPENE (1,1,2)	C	145.4	0	0	.00727	0	0	0	0	0	0
TRICHLORO(1,1,2)TRIFLUORO	C	0	0	24200	1.40E-10	0	0	0	0	0	0
TRICHLORO(1,1,2)TRIFLUOROETHANE(1,2,2)	C	107.30	1.50	300	.435	.0000000	.070	40	0.00	1000.0	227.5
TRICHLORO-1,2,2-TRIFLUOROETHANE,1,1,2-	C	107.30	0	270	0	0	0	47.7	0	0	0
TRICHLORO-1,3,5-TRIAZINE 2,4,6	C	104.41	0	0	.000022	0	0	0	0	0	0

name	type	mw	dens	vp	hl	dl	dv	boil pt	A	B	C
TRICOSANE N		184.31	0	0	.0000022	0	0	0	0	0	0
TRIDECYLAMINE	N	140.19	0	.01	0	0	0	328.4	0	0	0
TRIDECYLAMINE	N	162.22	0	400	.00000	0	0	0	0	0	0
TRIDECYLAMINE GLYCOL	O	180.2	0	1	9.00E-08	0	0	0	0	0	0
TRIDECYLPHOSPHONATE, o, o, o-	O	198.22	0	10	0	0	0	0	0	0	0
TRIFLUOROETHANE (1,1,1)	C	84	0	9040	04	0	0	-47.3	0	0	0
TRIGALLIC ANHYDRIDE	O	192.13	0	.0001	.0000041	0	0	242.5	0	0	0
TRIDECYLAMINE	N	50.11	0	1000	0	0	0	0	0	0	0
TRIDECYLSENE (1,3,5)	A	120.2	0	1.00	.147	0	0	0	0	0	0
TRIDECYLSENE 2,2,4	H	114.22	0	40.0	10.0	0	0	0	0	0	0
TRIDECYL-4-NITROLINE 2,3,5	N	180	0	0	.000	0	0	0	0	0	0
TRINITROBENZENE, sym-	A	213.11	0	0	0	0	0	315	0	0	0
TRINITROTOLUENE (2,4,6)	N	227.1	0	.040	.0000137	0	0	0	0	0	0
TRIPHENYL PHOSPHINE	A	202.3	0	0	.000131	0	0	0	0	0	0
TRIPHENYLMETHANE	H	244.32	0	0	0	0	0	300	0	0	0
TRIPHENYLPHOSPHINE NICKEL CARBONYL	O	377	0	0	.00077	0	0	0	0	0	0
TRIS (2-ACRIDONYL) PHOSPHINE SULFIDE	N	100.23	0	0	0	0	0	0	0	0	0
TRIS (2,3-DIBROMOPROPYL) PHOSPHATE	C	607.07	0	0	0	0	0	0	0	0	0
TRIOLEIN NITROLOTRACETATE	N	0	0	0	0	0	0	0	0	0	0
TRIPAN BLUE	A	900.00	0	0	0	0	0	0	0	0	0
URACIL	N	112.09	0	0	0	0	0	0	0	0	0
URACIL MUSTARD	A	202.1	0	0	0	0	0	0	0	0	0
UREA	N	60.00	1.34	0.00	.0000004	.0000137	.122	130	0	0	0
URETHANE	N	60.00	0	10	.0000000	.0000100	0	-40.3	7.421	1750.23	205
VALERIC ACID	O	102.13	0	1	.0000072	0	0	0	0	0	0
VINYL ACETATE	O	80.00	.90	115	.00002	.0000002	.000	73	7.21	1290.10	220.0
VINYL CHLORIDE	C	62.5	.91	2000	.000	.0000123	.100	-13.9	0	0	0
VINYL CYCLOHEXANE 4	H	100.2	0	20.0	1.04	0	0	0	0	0	0
VINYLIDENE CHLORIDE	C	97	1.21	001	.010	.0000104	.00	31.9	0.972	1000.4	237.2
WAFRIN	P	300.33	0	.00001	0	0	0	300	0	0	0
WATER	I	18.015	1	0	0	0	0	0	10.3000	3010.44	-40.13
XYLENE	A	100.2	0	0.0	.00000	0	0	0	0	0	0
XYLENE(-o)	A	100.10	.00	0	.00002	.0000070	.07	130	7.000	1420.200	210.1
XYLENE(-m)	A	100.2	.00	7	.000027	.00001	.007	144.4	0.900	1474.070	213.0
XYLENE(-p)	A	100.10	.00	0.0	.000027	0	0	130.4	0.90	1453.43	210.3
XYLENOL (3,4)	O	122.17	0	0	0	0	0	220	7.07010	1021.40	150.2
XYLYL CHLORIDE M	C	140.0	0	0	.00141	0	0	0	0	0	0
XYLYL CHLORIDE O	C	140.0	0	0	.00141	0	0	0	0	0	0
ZINC ACETATE	M	183.40	0	0	0.17E-00	0	0	0	0	0	0

name	Ln(OH)	SDOV	k1	Rt H/D	CAS	SQL	unicode
2,4,6 T	0	0	0	0	98-78-8	278	82:1;1J1[2f888
ACENAPHTHENE	3.92	0	0	0	82-22-0	3.42	3228:4;8888888
ACENAPHTHYLENE	4.87	0	2.7	0	288-88-8	3.98	3188:4;8888888
ACETALDEHYDE	.48	82.42	0	0	75-07-0	0	2111E8888888888
ACETAMIDE	0	0	0	0	65-35-5	888888	21C1N8888888888
ACETATE	0	0	0	0	NA	0	
ACETIC ACID	-.31	14	.98	0	64-19-7	0	2111[888888888
ACETIC ANHYDRIDE	0	0	0	0	108-24-7	0	
ACETONE	-.34	1.3	1.15	0	67-64-1	0	2211C8888888888
ACETONITRILE	-.34	0	0	0	75-05-0	0	11Y8888888888888
ACETOPHENONE	1.58	0	0	0	98-09-2	5588	
ACETYL CHLORIDE	0	0	0	0	79-36-5	0	21C1_88888888888
ACETYL DIETHYLMALONATE	0	0	0	0		285 1888	
ACETYLAMINOFLUORENE,2-	0	0	0	0	52-05-3	0	
ACETYLENE	0	0	0	0	74-86-2	1888	
ACETYLURAN 2	0	0	0	0	1192-82-7	188	
ACETYLMETHYLPHthalate 4	0	0	0	0		2	
ACETYLPYRIDINE 3	0	0	0	0	618-42-8	188888	
ACETYLPYRIDINE 3	0	0	0	0	1122-54-9	188	
ACETYL-2-THIOUREA,1-	0	0	0	0	551-88-2	0	
ACETYL-5-HYDROXYPIPERIDINE 3	0	0	0	0		475 28888	
ACRIDINE ORANGE	0	0	0	0	484-38-2	0	
ACRIDINE YELLOW	0	0	0	0	92-25-2	0	
ACROLEIN	-.88	7.8	.38	0	187-82-8	288888	2151E888888888888
ACRYLAMIDE	0	0	0	0	79-05-1	2888888	21510888888888888
ACRYLIC ACID	.31	0	0	0	79-10-7	0	21\158888888888888
ACRYLONITRILE	-.92	18	.75	0	187-12-1	773888	21212888888888888
ADAMANTANE DICARBOXYLIC ACID	0	0	0	0	828-51-3	2	
ADAMANTANE DICHLORIDE	0	0	0	0		171 .2	
ADENINE	-.18	0	0	0	73-24-5	588	
ADIPIC ACID	.88	0	0	0	124-04-9	18888	
AFLATOXINS	0	0	0	0	1482-88-2	0	
ALACHLOR	0	0	0	0	15872-88-2	242	
ALDICARB	0	0	0	0	116-88-3	8888	521141G1P1888
ALDRIN	0	0	0	0	588-88-2	.817	61243162*2_188
ALKYLIMINE CARBOXYLIC ACID N,SUB	0	0	0	0		482 2888	
ALLYL ALCOHOL	.17	0	0	0	187-18-6	0	31512178888888888
ALLYL CHLORIDE	0	0	0	0	187-85-1	3888	2151]8888888888888
ALPHA METHYL STYRENE	0	0	0	0	98-83-0	2	411185:1;88888
ALPHA METHYL STYRENE DIMERS	0	0	0	0		88 2	
alpha-CHLORO-beta-METHYLNAPHTHALENE	0	0	0	0	88-52-2	2	
ALPHA-HYDROXYACETALDEHYDE	0	0	0	0		0	
ALPHA-HYDROXYADIPIC ACID	0	0	0	0		429 1888	
alpha-METHYLSTYRENE (-4)	0	0	0	0	98-83-0	2	411185:1;88888
alpha-PICOLINE	0	0	0	0	1333-41-1	28888	
AMINO BENZOIC ACID (-p)	0	0	0	0	158-13-8	3488	
AMINOBIOPHENYL,4-	0	0	0	0	92-67-1	0	
AMINOCYCLOHEXANE	0	0	0	0	108-91-8	188888	
AMINOMETHYL-3-ISOKAZOLE 8	0	0	0	0	2763-88-4	288	
AMINO PHENOL(-o)	0	0	0	0	95-55-8	28888	44:1;171LE888888
AMINO PHENOL(-p)	0	0	0	0	181-88-4	8888	44:1;171LE88888
AMINO PHENOL,3-	.15	0	0	0	551-27-5	28888	
AMINO PROPNITRILE 3	0	0	0	0	151-18-8	28888	
AMINO PYRIDINE,4-	.28	0	0	0	584-24-5	0	33:171LE888888888
AMINO-2-CHLOROTOLUENE 4	0	0	0	0		784 2	
AMINO-3-CHLORO-5-PHENYLCYCLOHEXANONE 2	0	0	0	0		0	
AMINO-4'-CHLOROBIPHENYL 4	0	0	0	0		2	
AMINO-4-CHLOROPYRIDINE 2	0	0	0	0	1872-88-5	18888	
AMINO-4-CHLORO-5-CYANOPYRIDINE 2	0	0	0	0	598	58	
AMINO-4-NITROBENZYL ALCOHOL 2	0	0	0	0	118	2588	
AMINO-4-NITROTOLUENE 2	0	0	0	0	98-55-8	2	
AMINO-5-CHLOROPYRIDINE 2	0	0	0	0	1872-88-5	58	
AMINO-p'-METHYLAMAZOBENZENE P	0	0	0	0		125 2	
AMITROLE	0	0	0	0	61-82-5	288888	
AMONIA	0	0	0	0	7884-41-7	0	
AMPHETAMINE	0	0	0	0	98-15-1	188	
AMYL ACETATE(-n)	0	0	0	0	628-63-7	0	41132101I888888
AMTHOLE	0	0	0	0	184-48-1	0	
ANILINE	.9	7.1	21	0	62-53-3	35888	25:1LE88888888888
ANISIDINE,o-	0	0	0	0	98-84-8	0	

name	Ln(OW)	BIOV	k1	Rt HYD	CAS	SOL	unicode
ANISOLE	0	0	0	0	105-09-3	2	
ANTHRACENE	4.46	0	2.2	0	120-12-7	1.20	
ANTHRAQUINONE	0	0	0	0	84-05-1	0	38:4;2E000000
ARSENITE	0	0	0	0	145-67-8	0	
ARSENIC ACID	0	0	0	0	7770-30-4	167000	
ATRAZINE	0	0	0	0	1912-24-0	20	
ALUMAZINE	0	0	0	0	402-88-0	0	
AZASERINE	0	0	0	0	115-02-0	100000	
AZEPINE	0	0	0	0	111-40-0	10000	
AZIRIDINE	0	0	0	0	151-08-4	20000	21Q12000000000
BENZAL CHLORIDE	0	0	0	0	90-07-3	2	35:1;1]000000
BENZALDEHYDE	1.40	0	0	0	105-02-7	3300	
BENZALKONIUM CHLORIDE	0	0	0	0		130	10000
BENZEN SULFONATE	0	0	0	0	NA	0	
BENZENE	2.15	19	1.4	0	71-43-2	1700	18:0000000000
BENZENE ARSONIC ACID	0	0	0	0	90-05-5	25000	
BENZENE SULFONIC ACID	0	0	0	0		707	20000
BENZETHONIUM CHLORIDE	0	0	0	0	121-64-0	2000	
BENZIDINE	1.01	0	0	0	92-07-5	400	32:0;2,000000
BENZIDINE DIHYDROCHLORIDE	0	0	0	0	531-05-1	0	
BENZODIAXAS-1,3	0	0	0	0		151	5000
BENZOFULLORANTHENE,3,4-	0.57	0	0	0	205-00-2	0	
BENZOIC ACID	1.00	0	0	0	65-05-0	2000	
BENZONITRILE	0	0	0	0	100-47-0	10000	35:1;1T000000
BENZOPHENONE	0	0	0	0	119-01-0	2	3::2;J0000000
BENZOPYRENE 3,4	0	0	0	0	50-32-0	2	
BENZQUINONE,p-	.2	0	0	0	100-51-4	0	
BENZOTHAZOLE	2.01	0	0	0	95-10-0	100	
BENZOTRICHLORIDE	2.92	0	0	0	90-07-7	0	31d1;5:000000
BENZYL CHLORIDE	0	0	0	0	90-09-4	0	25:1"00000000
BENZO(A)ANTHRACENE	5.61	0	0	0	50-65-3	.01	2::;;00000000
BENZO(A)PYRENE	5.90	0	0	0	50-32-0	.003	2<:0;00000000
BENZO(B)FLUORANTHENE	0	0	0	0	205-00-2	0	2<:0;00000000
BENZO(ghi)PERYLENE	7.23	0	0	0	191-24-2	.00020	
BENZO(j)FLUORANTHENE	0	0	0	0	205-02-3	0	
BENZO(k)FLUORANTHENE	0.84	0	0	0	207-00-0	0	18:0000000000
BENZYL ALCOHOL	0	0	0	0	100-51-0	35000	4125:1;1?0000
BENZYL CHLORIDE	2.3	17.75	0	0	100-44-7	0	20:1]00000000
BENZYL METHYL ETHER	0	0	0	0	530-00-3	10	
BENZ(c)ACRIDINE	0	0	0	0	225-51-4	0	
BHC,alpha-	3.01	0	0	0	319-04-0	1.03	
BHC,beta-	3.0	0	0	0	319-05-7	.7	
BHC,delta-	4.14	0	0	0	319-06-0	21.3	
BHC,gamma-	3.72	0	0	0	50-00-0	17	
BICYCLO[2.2.1]-2,5-HEPTADIENE DICHLORIDE	0	0	0	0		172	.2
BIPHENYL	0	0	0	0	92-52-4	2	2::2;00000000
BIS (2-CHLOROETHOXY) METHANE	1.20	0	0	0	111-01-1	01000	
BISPHENOL(A)	0	0	0	0	00-05-7	10	
BIS(1,1,2,2-TETRACHLOROPROPYL) ETHER	0	0	0	0		100	10
BIS(2-CHLOROETHYL)ETHER	1.50	0	0	0	111-44-4	10000	32[2a2b000000
BIS(2-CHLOROISOPROPYL)ETHER	2.50	0	0	0	100-00-1	1700	32]1J12000000
BIS(2-ETHYLHEXYL)PHYTHALATE	5.3	.77	.35	0	117-01-7	.4	
BIS(CHLOROMETHYL)ETHER	-1.30	0	0	0	542-00-1	22000	
BROMACETONE	0	0	0	0	500-31-2	0	
BROMOBENZENE	0	0	0	0	100-00-1	400	35:1;1f000000
BROMOBENZYL ALCOHOL -(m)	0	0	0	0	15052-73-	5000	
BROMOBENZYL ALCOHOL -(o)	0	0	0	0	10002-34-	5000	
BROMOBENZYL ALCOHOL -(p)	0	0	0	0	073-75-0	5000	
BROMOCHLOROBENZENE P	0	0	0	0	100-30-0	.2	
BROMOCHLOROBENZYL ALCOHOL	0	0	0	0		114	2500
BROMOCHLOROMETHANE	0	0	0	0	74-07-5	10	11'0000000000
BROMOCHLOROMETHYL ACETATE	0	0	0	0		300	100000
BROMODICHLOROMETHANE	1.00	0	0	0	75-27-4	200	
BROMOETHYL ACETATE	0	0	0	0	927-00-4	100000	
BROMOETHYLENE	0	0	0	0	543-00-2	2	
BROMOFORM	2.3	0	0	0	75-25-2	3100	
BROMOMETHANE	1.1	0	0	0	74-83-0	000	
BROMOPHENYL PHENYL ETHER,4-	4.20	0	0	0	101-55-3	0	
BROMOPROPIONITRILE 3	0	0	0	0	2417-00-5	10000	
BROMOTOLLENE 4	0	0	0	0	100-30-7	10	34:1<1f000000

name	Ln(00)	SDOV	k1	Rb HYD	CAS	SL	united
BROMACIL, 5-	0	0	0	0	51-25-7	0	
BROMO-3-CHLOROBUTADENE 2	0	0	0	0		21 2	
BROMO-4-CHLOROCYCLOHEXANE 1	0	0	0	0	725	.2	
BROMO-4-CHLORO-6-CYANOBENZYL ALCOHOL 2	0	0	0	0		112 1000	
BROMO-4-CYANOMETHYL BENZOATE 2	0	0	0	0		80 .2	
BROMO-4-CYANOMETHYL BENZOATE 3	0	0	0	0		535 1	
BROMO-(1)-CHLOROTHANE-2	0	0	0	0	107-04-0	.2	
BUTADENE-(1,3)	0	0	0	0	100-00-0	0	12000000000000
BUTANE	0	0	0	0	100-07-0	400	
BUTANEDINITRILE	.35	0	0	0	110-01-2	120000	12000000000000
BUTANENITRILE	0	0	0	0	100-74-0	30000	
BUTANOL ISO	.75	7.0	.11	0	70-03-1	0	42112131700000
BUTANOL(S)	.75	7.0	.11	0	70-03-2	77000	31132170000000
BUTANOL-1	.83	7.0	.11	0	71-30-3	77000	31132170000000
BUTYL ACETATE(-n)	0	0	0	0	123-00-4	14000	31132170000000
BUTYL ACRYLATE	0	0	0	0	141-32-2	1000	
BUTYL BENZENE	0	0	0	0	104-61-0	2	411225:1-000000
BUTYL BENZYL PHTHALATE	4.70	0	2.2	0	05-00-7	2.0	
BUTYL CARBITOL	0	0	0	0	112-34-0	.2	41152172.000000
BUTYL CELLOSOLVE	0	0	0	0	111-70-2	40000	41142171.000000
BUTYL MERCAPTAN	0	0	0	0		001 100	
BUTYLAMINE	.30	0	0	0	100-73-0	0	311221N0000000
BUTYLENE GLYCOL-(1,3)	0	0	0	0	107-00-0	2000	
BUTYL-3-METHOXY PYRAZINE,2- ISO	0	0	0	0	24003-00-0	0	
BUTYL-p-CRESOL MONO T	0	0	0	0	1333-13-7	0	
BUTYL-p-CRESOL MONO T	0	0	0	0	25007-40-0	0	
BUTYRALDEHYDE	0	0	0	0	123-72-0	70000	311221E0000000
BUTYRALDEHYDE ISO	0	0	0	0	70-04-2	110000	
BUTYRIC ACID	0	0	0	0	107-02-0	00200	311221[00000000
CACODYLIC ACID	0	0	0	0	70-00-0	070000	
CAMPHENE	0	0	0	0	70-02-0	10	
CARBARYL	2.30	0	0	0	03-25-2	120	
CARBAZOLE	0	0	0	0	00-74-0	2	
CARBON DISULFIDE	2.10	0	0	0	75-15-0	2000	
CARBON OXYFLORIDE	0	0	0	0	353-00-4	0	
CARBON TETRACHLORIDE	2.72	1.5	1.5	0	00-23-0	000	11.000000000000
CATECHOL	.34	0	0	0	120-00-0	430000	
CELLULOSE	0	0	0	0	0000-11-7	10	
CHLORAL	0	0	0	0	302-17-0	10000	21E1.0000000000
CHLORALBUCL	0	0	0	0	300-00-3	0	02;4:2]421[170
CHLORDANE	2.70	0	0	0	07-74-0	.000	
CHLORIDIC ANHYDRIDE	0	0	0	0	115-27-0	0	
CHLORINATED TARS	0	0	0	0		000	
CHLORINAPHAZINE	0	0	0	0		0	03;7:12172]000
CHLOROACETALDEHYDE	0	0	0	0	107-20-0	0	21E1]0000000000
CHLOROACETIC ACID	0	0	0	0	70-11-0	000000	21[1]0000000000
CHLOROALLYL ALCOHOL 2	0	0	0	0	0070-47-0	10000	41217171]00000
CHLORANILINE(2)	0	.27	.00	0	00-01-2	10	34:111700000000
CHLORANILINE(3)	0	.27	.00	0	100-02-0	10	
CHLORANILINE,p-	1.00	.27	.00	0	100-47-0	0	
CHLORAZOBENZENE	0	0	0	0		120 2	
CHLOROBENZENE	2.5	.30	10	0	100-00-7	400	25:1*0000000000
CHLOROBENZENESULFONIC ACID (-p)	0	0	0	0	100-00-0	2000	
CHLOROBENZYLATE	0	0	0	0	010-15-0	10	
CHLOROBENZOIC ACID,2	1.00	0	0	0	110-01-2	2100	
CHLOROBENZOIC ACID,3-	2.00	0	0	0	030-00-0	400	
CHLOROBENZOIC ACID,4-	2.00	0	0	0	74-11-3	77	
CHLOROBENZOPHENONE (PARA)	0	0	0	0	134-00-0	0	40:2;101]000000
CHLOROBENZOTRICHLORIDE P	0	0	0	0	0210-25-1	.2	
CHLOROBENZYL ALCOHOL -(n)	0	0	0	0	073-03-2	0000	0124:1;17170000
CHLOROBENZYL ALCOHOL -(o)	0	0	0	0	17040-30-	0000	
CHLOROBENZYL ALCOHOL -(p)	0	0	0	0	073-70-7	0000	
CHLOROBIPHENYL (-p)	0	0	0	0	2001-02-0	0.0	
CHLOROCUMARIN 2	0	0	0	0	2001-00-4	2	
CHLOROCYANOBENZENE (1,4)	0	0	0	0	073-32-0	2	
CHLOROCYCLOHEXANE	0	0	0	0	042-10-7	2	
CHLOROCYCLOHEXANOL 2	0	0	0	0	1001-00-0	00	
CHLOROCYCLOHEXANOL 4	0	0	0	0		721 10	
CHLORODIACETYL	0	0	0	0		0	
CHLORODIMETHYL PHTHALATE 3	0	0	0	0		270 .2	

NAME	Ln(DW)	SDV	kl	Rt HYD	CAS	SCL	unocode
CHLORDIPHENYL THIOETHER P	0	0	0	0	7005-72-3	2	
CHLORETHANE	0	0	0	0		0	2111]00000000
CHLORETHANOL (ETHYLENE CHLORHYDRIN) 2	0	0	0	0	107-07-3	100000	
CHLORETHYLENE	0	0	0	0		0	2111-04000000
CHLOROMETHYL(2-) VINYL ETHER	1.20	0	0	0	110-70-0	100	
CHLOROFUOROBENZENE P	0	0	0	0	352-33-0	.2	
CHLOROFUORMETHANE	0	0	0	0	500-70-4	10	
CHLOROFORM	1.90	2.94	.70	0	67-68-3	9000	110000000000
CHLOROHYDROXYPHENYL 4 METHYL BENZOIC ACID	0	0	0	0		2000	
CHLOROMETHYL ACETYLENE	0	0	0	0		10	
CHLOROMETHYL BENZOATE P	0	0	0	0	1120-40-1	.2	
CHLOROMETHYL ETHYL KETONE	0	0	0	0		200	10000
CHLOROMETHYL METHYL ETHER	-.21	0	0	0	107-30-2	0	21]1100000000
CHLOROMETHYL PHENYL KETONE	0	0	0	0	532-27-4	0000	
CHLOROMETHYL PHENYLHYDRAZINE P	0	0	0	0		02	00
CHLOROMETHYLAMINOIMINE	0	0	0	0		2000	
CHLORONAPHTHALENE, 2-	4.12	0	0	0	91-50-7	0.74	37:1.72;000000
CHLORONITROALKOXIMINE	0	0	0	0		400	1000
CHLORONITROBENZENE(-o)	0	0	0	0	00-73-3	2	
CHLOROPHENOL POLYMERS	0	0	0	0		000	20000
CHLOROPHENOL-2	2.15	10	0	0	90-07-0	20000	34:101f000000
CHLOROPHENOL-3	0	0	0	0	100-43-0	20000	34:101]000000
CHLOROPHENOL-4	0	0.5	0	0	100-40-0	27100	34:101f000000
CHLOROPHENYL PHENYL ETHER, 4-	4.00	0	0	0	7005-72-3	3.3	
CHLOROPHENYLETHANOL 1,1	0	0	0	0	3301-10-4	2	
CHLOROPHTHALIC ANHYDRIDE 4	0	0	0	0		500	5000
CHLOROPRENE	.57	0	0	0	120-60-0	0	31v1710000000
CHLOROPROPANE-1	0	0	0	0	540-54-0	0	311121]000000
CHLOROPROPANE-2	0	0	0	0	70-29-0	0	2211]00000000
CHLOROPROPENE 3	0	0	0	0	557-00-2	100	2151]00000000
CHLOROPROPIONITRILE, 3-	0	0	0	0	542-70-7	4000	2121]00000000
CHLOROPROPYLENE-2	0	0	0	0	507-00-2	0	
CHLOROPYRIDINE 2	0	0	0	0	100-00-1	100	
CHLOROSTYRENE (-4)	0	0	0	0	1331-20-0	2	
CHLORTETRAHYDROFURAN 3	0	0	0	0		504	10000
CHLORTHIOPHENOL P	0	0	0	0	100-54-7	.2	
CHLORTOLLENE-4	0	0	0	0	100-40-4	100	34:101f000000
CHLORURACIL, 5-	0	0	0	0	1020-01-1	0	
CHLORO(-p) CRESOL (-m)	3.1	0	0	0	50-00-7	4000	
CHLORO(-p) PHENYLHYDRAZINE	0	0	0	0		541	00
CHLORO-1,2-ETHANE DIOX	0	0	0	0		402	10000
CHLORO-1,3-CYCLOPENTADIENE 0	0	0	0	0		240	.2
CHLORO-2,2-DIBROMOETHANE 1	0	0	0	0		2	
CHLORO-2,3-EPOXYPROPANE, 1-	.00	0	0	0	100-00-0	00000	
CHLORO-2,5-DIKETOPYRROLIDINE 3	0	0	0	0		420	200000
CHLORO-2-METHOXYBENZOIC ACID 4	0	0	0	0	57479-70-0	0	
CHLORO-2-NITROBENZYL ALCOHOL 4	0	0	0	0	22900-10-	0000	
CHLORO-3-NITROANILINE 4	0	0	0	0	030-22-3	100	
CHLORO-3-NITRO-5-PHENYLCYCLOHEXANE 2	0	0	0	0		0	
CHLORO-AMINOOCUMARAN-0-CARBOXYLIC ACID 2	0	0	0	0		2000	
CHLORO-4-CYANOBENZYL ALCOHOL 2	0	0	0	0		110	2000
CHLORO-4-HYDROXYBIPHENYL 3	0	0	0	0	92-04-0	2	
CHLORO-4-METHOXY-0-AMINOBENZOIC ACID 2	0	0	0	0		01	00
CHLORO-4-METHYL-N-METHYLBENZAMIDE 3	0	0	0	0		400	2000
CHLORO-4-NITROANISOLE 2	0	0	0	0		2	
CHLORO-4-PHENYLPYRIDINE 2	0	0	0	0		400	2000
CHLORO-0-AMINO-3-PYRIDINE CARB. ACID AMIDE 2	0	0	0	0		500	00
CHLORO-0-CYANOPHTHALIC ACID 4	0	0	0	0		1000	
CHLORO-0-CYANTOLLENE 3	0	0	0	0		00	10
CHLORO-0-FLUOROTOLLENE 3	0	0	0	0	443-03-4	00	
CHLORO-0-PHENOXIDIMETHYL PHTHALATE 4	0	0	0	0		271	.2
CHLORO-N-METHYLBENZAMIDE P	0	0	0	0		470	2000
CHLORO-p'-METHYLBIPHENYL P	0	0	0	0	1007-11-4	2	
CHLORO-p-XYLENE	0	0	0	0	104-02-0	.2	
CHOLINE CHLORIDE	0	0	0	0	07-40-1	0	
CHROMIUM (TOTAL)	0	0	0	0	7440-47-3	0	
CHRYSENE	5.01	0	1.4	0	210-01-0	.000	20:0;00000000
CITRIC ACID	0	0	0	0	77-02-0	400000	
CITRUS RED #2	0	0	0	0	0300-03-0	0	
COPPER PHTHALOCYANINE	0	0	0	0	147-14-0	10	

name	Ln(OH)	BDDV	k1	Rb H/D	CAS	SQL	unicode
CLAMARIN	0	0	0	0	91-04-5	2	
CRESOTE	0	0	0	0	8851-00-0	2000	
CRESOL	0	23	17	0	1319-77-3	2000	
CRESOL(-o)	.97	23.21	17	0	100-30-4	25000	34:1(180000000)
CRESOL(-o)	1.90	22.70	17	0	90-40-7	31000	
CRESOL(-p)	1.94	23.21	17	0	100-44-5	24000	
CRESYLIC ACID	0	15	1	0		0	
CROTONALDEHYDE	0	0	0	0	470-30-5	0	31116150000000
CROTONALINE	0	0	0	0	800-17-5	2	
CLAMINE (Isopropylbenzene)	0	0	0	0	90-02-0	0	
CUMYLPHENOL-4	0	0	0	0	27570-00-2		
CYANIDE	0	0	0	0	57-12-5	100000	
CYANOBENZYL ALCOHOL P	0	0	0	0		116	0000
CYANOGEN	0	0	0	0	400-10-5	0000	12200000000000
CYANOGEN BROMIDE	0	0	0	0	500-00-3	0	
CYANOGEN CHLORIDE	.04	0	0	0	500-77-4	30000	
CYANOLIMEDINE	0	0	0	0	401-00-5	10000	
CYANOMETHYL BENZOATE 4	0	0	0	0		770	2
CYANOMETHYLPHTHALATE 4	0	0	0	0			2
CYANOPYRIDINE 3	0	0	0	0	100-54-0	100	
CYANOPYRIDINE (-o)	0	0	0	0	100-40-1	100	
CYANTOLLINE 4	0	0	0	0		700	2
CYANURIC ACID	0	0	0	0	100-00-5	0000	
CYOSIN	0	0	0	0	14001-00-0	0	
CYCLOHEXANE	0	0	0	0	110-02-7	0	10200000000000
CYCLOHEXANOL	0	0	0	0	100-00-0	0	30213170000000
CYCLOHEXANONE	.01	11.40	0	0	100-04-1	23000	21D42000000000
CYCLOHEXENE	0	0	0	0	110-00-0	0	
CYCLOHEXYL ACETATE	0	0	0	0	022-45-7	200	
CYCLOHEXYLAMINE	0	0	0	0	100-01-0	0	20210000000000
CYCLOHEXYLCYCLOHEXANONE 4	0	0	0	0	50025-00-2		
CYCLOHEXYL-2,2-DIPHENYLETHYLAMINE 2	0	0	0	0		300	100
CYCLOHEXYL-4,6-DINITROPHENYL-2-	0	0	0	0	131-00-5	0	
CYCLOPENTADIENE	0	0	0	0		200	
CYCLOPENTADIENE 1,3	0	0	0	0	722	2	21220000000000
CYCLOPHOSPHAMIDE	.03	0	0	0	50-10-0	40000	
CYCLOHEXYL o,p-DIMETHYL PHOS.DITHIOATE 3	0	0	0	0		2	
CYTOSINE	-1.40	0	0	0	71-30-7	7700	
DALNACIN	0	0	0	0	20030-01-0	0	
DDO,p,p'-	5.00	0	0	0	72-54-0	.02	
DDO,p,p'-	5.00	0	0	0	72-55-0	.04	
DDT	0.10	0	0	0	50-29-3	.0001	00:1(10271;000)
DIACETYL	0	0	0	0	431-00-0	200000	
DIAMINODIPHENYL METHANE P,P'	0	0	0	0	100-77-0	100	
DIAMINO-6-SULFONYL BENZYL 2,4	0	0	0	0		25000	
DIAZON	0	0	0	0	330-01-5	40	
DIBENZOPIRENE 1,2,7,8	0	0	0	0		0	2::;00000000
DIBENZO(a,h)ANTHRACENE	5.97	0	0	0	53-70-3	.0000	
DIBROMOCHLOROMETHANE	2.00	0	.000	0	124-40-1	.2	
DIBROMOETHANE-1,2	0	0	0	0	100-00-4	4310	2102_00000000
DIBROMOMETHANE	0	0	0	0	74-00-3	11000	
DIBROMO-3-CHLOROPROPANE,1,2	0	0	0	0	90-12-0	1000	22]1^00000000
DIBUTYL ETHER	0	0	0	0	140-00-1	300	
DIBUTYLPHTHALATE	0.2	.4	1	0	04-74-2	400	041222(0;2)000
DICHLORO 2-PROPANOL 1,3	0	0	0	0	90-23-1	100000	31317(0)000000
DICHLORO PROPANOL 2,3	0	0	0	0	010-23-0	0	41217(1)1^0000
DICHLOROBENZENE(1,2) (-o)	3.30	2.5	.00	0	90-00-1	140	24:270000000000
DICHLOROBENZENE(1,3) (-o)	3.30	0.4	2.3	0	041-73-1	120	
DICHLOROBENZENE(1,4) (-p)	3.30	0.4	2.3	0	100-40-7	70	
DICHLOROBENZIDINE,3,3'-	3.51	0	0	0	01-04-1	0	42;0:272.00000
DICHLOROBENZOTRIAZOLE,2,0-	0	0	0	0	1194-00-0	10	33:122700000000
DICHLOROBENZOPHENONE P,P'	0	0	0	0	90-00-2	.2	40:2;0^270000
DICHLOROBIPHENYL (PARA)	0	0	0	0	213029-00	.02	
DICHLOROBUTANE (1,4)	0	0	0	0	110-00-0	10	
DICHLORODIFLUOROMETHANE	2.10	0	0	0	70-71-0	200	
DICHLORODIPHENYL METHANE	0	0	0	0	2001-00-3	2	
DICHLOROETHANE(1,1)	1.70	0	2.3	0	70-34-3	0000	21110000000000
DICHLOROETHANE(1,2)	1.40	2.1	.00	0	107-00-2	0000	12]0000000000
DICHLOROETHYL ETHER	0	0	0	0		0	32]1.1120000000
DICHLOROETHYLENE(2,1)	1.40	0	1.0	0	70-30-4	210	2102_00000000

name	Ln(DN)	SDV	k1	Rs HYD	CAS	SOL	united
DICHLOROETHYLENE(1,2)	0	0	0	0	155-54-2	000	
DICHLOROETHYLENE,1,2-TRANS-	1.48	0	0	0	155-55-5	000	
DICHLORODIFLUOROMETHANE	0	0	0	0	75-43-4	.2	11<0000000000
DICHLOROPHENOL(2,4)	2.75	25	3.3	0	128-85-2	4000	33:1827000000
DICHLOROPHENOL(2,6)	0	0	0	0	87-85-6	0	33:1827000000
DICHLOROPHENYLACETIC ACID(2,4)	0	0	0	0	94-75-7	000	63:1;1D1(27000
DICHLOROPROPANE(1,2)	2.28	17	1.4	0	78-57-5	0	3111~1]0000000
DICHLOROPROPANE(1,3)	1.98	0	0	0	542-75-6	2700	3111~1]0000000
DICHLOROPROPYLENE,1,2- (cis)	0	0	0	0		0	311162~0000000
DICHLOROPROPYLENE,1,2-(trans)	0	0	0	0	853-54-2	0	
DICHLOROPROPYLENE-2,3	0	0	0	0	78-88-6	2100	
DICHLOROSTYRENE 2,6	0	0	0	0	000	2	
DICHLORTETRAHYDROFURAN 3,4	0	0	0	0	3511-19-1	100000	
DICHLORO-1,3-CYCLOPENTADIENE 5,5	0	0	0	0	247	.2	
DICHLORO-2-BUTENE 1,2	0	0	0	0	835	.02	22]1000000000
DICHLORO-2-BUTENE(1,4)	0	0	0	0	784-41-6	0	22]1000000000
DICHLORO-TRANS-ETHYLENE(1,2)	0	0	0	0	548-59-6	000	
DICHLORO-(2,6)-NITROANILINE(4)	0	0	0	0	99-35-0	50	
DIELDRON	0	0	0	0	65-57-1	.100	
DIETHYL AMINE	0	0	0	0	109-89-7	20000	3211210000000
DIETHYL ETHER	0	0	0	0	682-87-6	60000	321121~0000000
DIETHYL ETHER ACID CHLORIDE	0	0	0	0		100	
DIETHYL PHTHALATE	2.98	0	2.7	0	84-69-2	2	
DIETHYL PROPIONATE,2,N	0	0	0	0	18298-69-	73	
DIETHYL THIOETHER	0	0	0	0	352-88-2	50	
DIETHYL (N,N) ANILINE	0	0	0	0	91-68-7	14400	621225:1;1000
DIETHYLBENZENE P	0	0	0	0	25346-17-	2	3214:2~0000000
DIETHYLDIPHENYL UREA SYN	0	0	0	0	85-88-3	2	
DIETHYLENE GLYCOL	-1.98	0	0	0	111-45-6	100000	
DIETHYLHYDRAZINE N,N	0	0	0	0		0	2212000000000
DIETHYLUREA 1,1	0	0	0	0	634-05-7	2000	
DIDYDRO-5-OXAZALONE (DIDYDROAZLACTONE)	0	0	0	0		5000	
DIDISOPROPYL BENZENE (PARA)	0	0	0	0	185-18-5	2	3474:2~0000000
DIMETHOATE	0	0	0	0	65-51-5	25000	
DIMETHOXY METHANE	0	0	0	0	188-87-5	330000	
DIMETHOXY-(3,3')-BENZIDINE	0	0	0	0	119-68-4	10	
DIMETHYL AMINE	0	0	0	0	124-48-3	0	2111~000000000
DIMETHYL BENZYLAMINE N,N	0	0	0	0	188-83-3	10	
DIMETHYL BENZ(A)ANT 7,12	0	0	0	0		0	32<6;::0000000
DIMETHYL DISULFIDE	0	0	0	0	624-92-6	6300	
DIMETHYL FORMAMIDE	0	0	0	0	68-12-2	20000	
DIMETHYL HYDRAZINE(1,1)	0	0	0	0	57-14-7	100000	
DIMETHYL METHYLTHIOCARBAMATE N,N	0	0	0	0		2000	
DIMETHYL NITROISOPROPYLAMINE N,N	0	0	0	0	612	50	
DIMETHYL NITROSAMINE	-.57	0	0	0		0	
DIMETHYL PHTHALATE	1.87	2.2	3.1	0	131-11-3	6000	
DIMETHYL SULFATE	0	0	0	0	77-78-1	0	
DIMETHYL SULFIDE	0	0	0	0	75-18-3	6300	
DIMETHYLAZOBENZENE,4-	0	0	0	0	68-11-7	0	
DIMETHYLBENZIDINE 3,3	0	0	0	0		0	4212(2;5:00000
DIMETHYLBENZYL HYDROPEROXIDE	0	0	0	0	85-15-0	100	
DIMETHYLBENZ(A)ANTHRAcene(7,12)	0	0	0	0	57-87-6	2	3:;6;2<0000000
DIMETHYLETHYLAMINE	0	0	0	0	75-84-0	5000	321121S00000000
DIMETHYLHYDANTOIN,5,5-	0	0	0	0	77-71-4	0	
DIMETHYLHYDRAZINE,1,1-	-1.94	0	0	0	57-14-7	0	12~00000000000
DIMETHYLPHENOL(2,4)	2.42	19.7	4.7	0	105-67-0	10	32<183:0000000
DIMETHYLPHENOL(3,4)	0	5.5	1.05	0		0	32<183:0000000
DIMETHYLPHENYLACETOL	0	0	0	0	617-84-7	0	
DIMETHYL-1-NITROBENZENE 2,4	0	0	0	0	25168-84-	2	
DINITROBENZENE M	0	0	0	0	98-05-6	0	22]4:000000000
DINITROPHENOL 2,4	1.54	0	.02	0	51-28-5	5000	32]183:0000000
DINITROTOLUENE 2,6	2.85	0	0	0	605-25-2	20	31a1]000000000
DINITROTOLUENE(2,4)	2.61	0	0	0	121-14-2	270	33:1C2]0000000
DINITRO-4-CRESOL(4,6)	2.85	0	0	0	534-82-1	250	
DINOCAP	0	0	0	0	30888-45-	2	
DINOSES	0	0	0	0	88-85-7	50	
DIOXANE(1,4)	0	0	0	0	123-61-1	0	22.22000000000
DIOXON	0	0	0	0	628-88-2	35-11	
DIPHENYL ETHER	0	0	0	0	101-84-8	2	
DIPHENYL THIOETHER	0	0	0	0	138-68-2	2	

NAME	Ln(OW)	SDV	W1	Rn HO	CAS	SL	unlcoda
DIPHENYLAMINE	3.22	0	0	0	122-29-4	389	
DIPHENYLSTADIENE 1,2	0	0	0	0	989-65-7	16	
DIPHENYLOLOMETHANE	0	0	0	0	95-69-3	2	
DIPHENYLOXETONE	0	0	0	0	134-61-6	2	
DIPHENYLETHANE 1,1	0	0	0	0	739	29	
DIPHENYLETHANOL 1,1	0	0	0	0	999-67-7	19999	
DIPHENYLHIDRAZINE(1,2)	3.88	0	0	0	122-69-7	0	
DIPHENYLHIDRAZINE,1,1-	0	0	0	0	529-69-7	0	
DIPHENYLMETHANE	0	0	0	0	101-61-6	3	3:1;1;0000000
DIPROPYLAMINE	0	0	0	0	149-64-7	199999	3102221.0000000
DIPROPYLFORMAMIDE	0	0	0	0	6292-65-4	0	
DIXDAL. METONE	0	0	0	0		24	49999
DI-n-OCTYL PHYMALATE	9.2	0	0	0	117-64-6	.285	521>24;2;2)000
DI-tert-BUTYL-p-CRESOL	0	0	0	0	129-27-0	0	
DTA	0	0	0	0	95-69-4	0	
DUCOLFIN	3.85	0	0	0	115-29-7	.29	
DUCOLFIN SULFATE	3.88	0	0	0	1991-67-6	.117	
DUCON	5.6	0	0	0	72-29-6	.29	
EPICHLOROHYDIN	.99	0	0	0	199-69-6	99999	31.1131)0000000
EPINEPHRINE	0	0	0	0	51-43-4	899	
ETHNE	0	0	0	0		0	121999999999999
ETHNOL	-.22	8.8	.3	0	64-17-5	199999	3111217.0000000
ETHYLAMINE(mono-)	0	0	0	0	141-43-5	199999	31217.0000000
ETHYLETHANOL-2	0	0	0	0	116-69-6	199999	41122171.00000
ETHYL ACETATE PEROXIDE	0	0	0	0		282	19999
ETHYL ACRYLATE	0	0	0	0	149-69-5	0	4111215U.00000
ETHYL CHLORIDE	1.43	0	0	0	75-69-3	5749	
ETHYL CYANIDE (PROPIDETROL)	.94	0	0	0	197-12-6	99999	
ETHYL ETHER	0	0	0	0	69-29-7	0	321121.0000000
ETHYL ISOPROPYL PEROXIDE	0	0	0	0		283	199999
ETHYL S,S-DIPHENYL PHOSPHORODITHIATE 0	0	0	0	0	1799-49-6	2	
ETHYLACETATE	0	17.99	0	0	141-79-6	0	
ETHYLAMINE	0	0	0	0		0	2111N0000000000
ETHYLENE	3.15	6.8	2.1	0	199-41-4	152	31=415:0000000
ETHYLENE DIAMINE	0	0	0	0	74-85-1	131	
ETHYLENE DIIMINE	0	0	0	0	197-15-3	199999	120999999999999
ETHYLENE DIISOCYANIDE	1.6	0	0	0	199-69-4	4399	
ETHYLENE GLYCOL	0	0	0	0	197-21-1	199999	22227.000000000
ETHYLENE GLYCOL DIMETHYL ETHER	0	0	0	0	116-71-4	199999	
ETHYLENE GLYCOL MONOMETHYL ETHER	0	0	0	0	199-69-4	199999	41112171.00000
ETHYLENE GLYCOL MONOMETHYL ETHER ACETATE	0	0	0	0	116-49-6	199999	
ETHYLENE DIAMINE	0	0	0	0	197-15-3	0	120999999999999
ETHYLENE DIISOCYANIDE	-.391	4.2	.91	0	75-21-6	0	21.1209999999999
ETHYLETHYR	.99	.77	0	0		0	
ETHYLENYLACRYLATE,2-	0	0	0	0	199-11-7	0	
ETHYLPHENOL,3-	2.4	0	0	0	629-17-7	0	511124:1;19999
ETHYL(2) HEDROL	0	0	0	0	194-79-7	1999	421521317.00000
ETHYL-(2)-PROPYL-(3) ACRYLEIN	0	0	0	0	645-62-5	0	
ETHYLENE,4-	0	0	0	0	4995-62-0	0	
FLUORANTHENE	5.33	0	1.5	0	299-44-6	.3	2:16;000000000
FLUORINE	4.19	0	3.5	0	99-79-7	2	
FLUORACETIC ACID, SODIUM SALT	0	0	0	0	62-74-6	29999	
FLUORURACIL,5-	0	0	0	0	51-21-6	0	
FORMALDEHYDE	1.94	5	.25	0	95-69-6	599999	
FORMAMIDE	0	0	0	0		0	21E1N0000000000
FORMIC ACID	0	0	0	0	64-19-6	0	11\0000000000000
FREDS	0	0	0	0		0	
FUMARIC ACID	0	0	0	0	119-17-6	7999	2162(00000000000
FURAN	0	0	0	0	119-69-9	0	2291K0000000000
FURFURAL	0	0	0	0	99-61-1	0	416171E1.00000
FURIC ACID	0	0	0	0	99-14-2	39999	
GEOSIN	0	0	0	0	19799-21-	0	
GLUTARIC ACID	0	0	0	0	119-64-1	649999	2322(00000000000
GLYCERIN (GLYCEROL)	0	0	0	0	99-61-6	19999	3221337.0000000
GLYCINAMIDE	0	0	0	0	529-41-4	99999	
GLYCIDINE, NITROSD	0	0	0	0	674-61-7	19999	
GLYIDE	-1	0	0	0	73-45-5	0	
GLUCIDOL	0	0	0	0	999-62-5	199999	41213171D00000
HEPTACHLOR	5.95	0	0	0	79-44-9	.999	
HEPTACHLOR EPOXIDE	4.6	0	0	0	1924-67-3	.35	

NAME	Ln(OH)	SDV	k1	Rb HYD	CAS	SL	unlcoda
HEPTANE ISO	0	0	0	0	31304-64-	2	33132130000000
HEPTANE(-n)	0	0	0	0	142-82-6	3	22152000000000
HEXACHLORO CYCLOPENTADIENE	3.30	0	0	0	77-47-4	.005	
HEXACHLOROBENZENE	5.47	0	0	0	112-74-1	.11	10700000000000
HEXACHLOROBUTADIENE	3.74	0	0	0	67-69-3	2	2250-0000000000
HEXACHLOROCYCLOHEXANE (GAMA ISOMER)	0	0	0	0	50-69-0	17	16700000000000
HEXACHLOROCYCLOPENTADIENE	3.30	0	0	0	77-47-4	50	24710000000000
HEXACHLOROETHANE	0	0	0	0	67-72-1	50	12000000000000
HEXACHLOROPENTADIENE	0	0	0	0		415 2	
HEXACEANE N	0	0	0	0	544-70-3	.0000	
HEXAMETHYLENEDIAMINE	0	0	0	0	124-69-4	1000000	
HEXONE(-n)	0	0	0	0	118-54-3	0	22142000000000
HEXANOIC ACID	0	0	0	0	142-82-1	11000	
HEXANOL-1	0	0	0	0	111-27-3	0	31152170000000
HEXEN-3-ONE S	0	0	0	0		0	31225100000000
HHS	0	0	0	0	surrogate	0	
HHS	0	29.2	1	0		0	
HLS	0	.3	1	0		0	
HLS	0	0	0	0	surrogate	0	
HMS	0	2.71	1	0		0	
HMS	0	0	0	0	surrogate	0	
HYDRAZINE	-1.37	0	0	0	302-61-2	0	
HYDROCYANIC ACID	0	0	0	0	74-85-0	0	
HYDROFLUORIC ACID	0	0	0	0	7664-39-3	100000	
HYDROGEN SULFIDE	0	0	0	0	7782-62-4	0	
HYDROQUINONE	.50	0	0	0	123-31-0	70000	24:200000000000
HYDROXY DIMETHYL ETHER	0	0	0	0		270 1000	
HYDROXYMETHYLPYRIDINECARBOXYLIC ACID 2	0	0	0	0	30116-61-	00	
HYDROXYACETIC ACID	0	0	0	0	79-14-1	100000	
HYDROXYCYCLOHEXANONE 4	0	0	0	0		0	
HYDROXYDIMETHYL PHTHALATE 4	0	0	0	0		272 .2	
HYDROXYMETHYL ACETYLENE	0	0	0	0		10	
HYDROXYMETHYL ISOPROPYL KETONE	0	0	0	0		281 10000	
HYDROXYMETHYLPHENYL CARBAMATE N	0	0	0	0		471 1000	
HYDROXYMETHYLTHIOBENZENE	0	0	0	0		63 2	
HYDROXYMETHYL VINYL ETHER	0	0	0	0		277 1000	
HYDROXYMETHYL, N-METHYLETHYL AMINE N	0	0	0	0		611 50	
HYDROXYMETHYL-N-CHLOROMETHYLETHYLAMINE N	0	0	0	0		50	
HYDROXYPENTANE 3	0	0	0	0		737 5000	
HYDROXY-1,3-CYCLOPENTADIENE S	0	0	0	0		723 2	
HYDROXY-4-METHYL TETRAHYDROFURAN 3	0	0	0	0		500 100000	
HYDROXY-5-METHYLODIMETHYL PHTHALATE 4	0	0	0	0		273 .2	
HYDROXY-(2)-PROPIONITRILE	0	0	0	0		100-78-4 100000	
INDANOL, 5-	0	0	0	0		1470-04-0 0	
INDENO(1,2,3-cd)-PYRENE	7.00	0	0	0		190-30-6 0	20:9;0000000000
INDOLE	2.14	0	0	0		120-72-0 0	
IODOCUMARAN 2	0	0	0	0		2	
ISOPHORONE	1.7	0	0	0		70-60-1 12000	
ISOUAZOLE, 5-(AMINOETHYL)-3-	0	0	0	0		2703-00-4 0	
LEAD ACETATE	0	0	0	0		301-64-2 0	
LEAD SUBACETATE	0	0	0	0		1336-22-0 0	
LEUCINE	0	0	0	0		61-00-5 24300	
LHS	0	0	0	0	surrogate	0	
LHS	0	23.2	1	0		0	
LHS	0	3.50	1	0		0	
LHS	0	0	0	0	surrogate	0	
LINDANE	0	0	0	0		0	15700000000000
MALATHION	0	0	0	0		121-75-6 145	
MALEIC ACID	0	0	0	0		110-16-7 700000	
MALEIC ANHYDRIDE	0	4.00	1	0		100-31-0 700	
MELANINE	0	0	0	0		100-70-1 100	
MERCURY	0	0	0	0		7439-97-0 0	
MESITYL OXIDE	0	0	0	0		141-79-7 20000	
METHACRYLIC ACID	0	0	0	0		79-41-4 1000	
METHANE	0	0	0	0		74-82-6 24	
METHANE SULFONIC ACID	0	0	0	0		70-78-2 20000	
METHANETHIOL	0	0	0	0		74-88-1 23000	
METHANOL	-.7	10	.2	0		67-58-1 0	11000000000000
METHAPYRILENE	0	0	0	0		91-00-5 0	
METHOYL	0	0	0	0		15762-77- 20000	

NAME	Ln(DN)	SDV	L2	RA MD	CAS	SQL	unloads
METHOXYACETIC ACID	0	0	0	0	278	0000	
METHOXYACETONITRILE	0	0	0	0	1720-20-0	100000	
METHOXYCHLOR	0	0	0	0	73-43-6	.04	
METHOXYPHENYL P	0	0	0	0	188-78-6	40000	
METHYL 1-PENTENE 2	0	0	0	0	780-20-1	2	32122170000000
METHYL ACETATE	-0.00	19.87	0	0	78-26-0	2000	
METHYL ACRYLATE	0	0	0	0	96-33-3	00000	2161P000000000
METHYL ACRYLONITRILE	0	0	0	0	120-69-7	25	
METHYL AMINE	0	0	0	0	74-89-6	11630	11A000000000000
METHYL AMINOETHYLENE	0	0	0	0	04	100	
METHYL AZIRIDINE 2	0	0	0	0	0	0	31013110000000
METHYL BENZYL ALCOHOL 4	0	0	0	0	780	1000	
METHYL BIPHENYL (-p)	0	0	0	0	844-00-0	2	
METHYL CHLORIDE	1.02	0	0	0	74-87-3	0000	11J00000000000
METHYL CHLOROACETAMIDE N	0	0	0	0	481	1000	
METHYL CHLOROCARBONATE	0	0	0	0	78-23-1	100	
METHYL CHOLANTHRENE 3	0	0	0	0	88-40-6	2	4119:9;2200000
METHYL COLUMBIN 2	0	0	0	0	687-71-0	2	
METHYL CYCLOHEXANE	0	0	0	0	100-67-2	14	31152130000000
METHYL ETHER	0	0	0	0	115-10-6	70000	
METHYL ETHYL KETONE	.38	2	.2	0	78-66-3	270000	31112100000000
METHYL FORMATE	0	0	0	0	107-51-3	304	2112H000000000
METHYL MORAZINE	0	0	0	0	68-34-4	0	11P00000000000
METHYL IODIDE	1.00	0	0	0	74-88-4	14000	2111p000000000
METHYL ISOMAL KETONE	0	0	0	0	116-12-3	5400	
METHYL ISOBUTYL KETONE	1.30	.74	0	0	100-10-1	10000	41C12132100000
METHYL ISOCYANATE	0	0	0	0	624-68-0	0	
METHYL ISOPROPYL KETONE	0	0	0	0	800-80-4	47000	
METHYL METHACRYLATE	0	0	0	0	96-63-6	10000	
METHYL METHANESULFONATE	0	0	0	0	60-27-3	200000	
METHYL NAPHTHALENE(1-)	0	24.00	0	0	90-12-0	0	
METHYL NAPHTHALENE(2-)	0	0	0	0	91-67-0	0	
METHYL PARATHION	2.04	0	0	0	200-00-0	00	
METHYL PROPENE 2	0	0	0	0	115-11-7	2	
METHYL SULFURIC ACID	0	0	0	0	400	100000	
METHYL TETRAHYDROFURN 2	0	0	0	0	705	1000	
METHYL THIOFENOL 4	0	0	0	0	100-45-0	0	
METHYL THIOURACIL	0	0	0	0	50-04-2	6700	
METHYLACETONITRILE2(ACETONE CYANOHYDRON)	0	0	0	0	78-08-6	0	
METHYLENE CHLORIDE	1.25	22	.4	0	78-09-2	16700	11'000000000000
METHYLENE-BIS (2-CHLOROMETHYLENE),4,4'-	0	0	0	0	101-14-4	0	
METHYLFURN 2	0	0	0	0	524-22-6	3000	
METHYLFURNAL 5	0	0	0	0	620-02-0	50000	
METHYLDIOXACETIC ACID	0	0	0	0	64	10000	
METHYLISOBORNOL,2-	0	0	0	0	NA	0	
METHYLPHENYL CARBONATE N	0	0	0	0	472	1000	
METHYL TIN TRICHLORIDE	0	0	0	0	900-10-0	100	
METHYL-1,3-CYCLOPENTADIENE 5	0	0	0	0	20519-01-	.2	
METHYL-2,3,4-TRIMETHOXYMETHYLENE N	0	0	0	0	478	1000	
METHYL-2-AMINOETHANOLAMINE	0	0	0	0	100-01-0	1000	
METHYL-2-HYDROXYETHANOLAMINE	0	0	0	0	100-00-1	20000	
METHYL-2-METHOXYAZIRIDINE 1	0	0	0	0	474	20000	
METHYL-2-ACETILCYCLOPENTADIENE 1	0	0	0	0	253	2	
METHYL-2-NITROBENZYL ALCOHOL 4	0	0	0	0	48070-00-	2500	
METHYL-4-NITROBENZYL ALCOHOL 2	0	0	0	0	23670-13-	0000	
METHYL-5-THIOACETYLDIHYDRO-1,3THIAZOLE 4	0	0	0	0	50	0	
METHYL-PHENYLETHANOLAMINE N	0	0	0	0	500-00-2	10	
METHYL-p'-METHYLTHIOFENYL PHOSPHINE P	0	0	0	0	200	2	
METHYL-TERTIARY-BUTYL ETHER	0	0	0	0	1634-04-4	0	
METHYL-TRIMORO-1,3-THIAZOLE 4	0	0	0	0	404	100	
MMS	0	40.1	1	0	0	0	
MMS	0	0	0	0	surrogate	0	
MMS	0	.30	1	0	0	0	
MMS	0	0	0	0	surrogate	0	
MMS	0	0	0	0	surrogate	0	
MMS	0	11.2	1	0	0	0	
METAMCIN C	0	0	0	0	68-07-7	0	
MMS	0	0	0	0	78-25-7	0	
METHOCHLOROFLUOROMETHANE	0	0	0	0	75-45-6	2	11c00000000000
MORPHOLINE	-1.00	0	0	0	110-01-0	20000	

NAME	Ln(OH)	RDIV	kl	Rs H/D	CAS	SL	unicode
PHENYL ISOCYANATE	0	0	0	0	100-71-0	1	
PHENYL MERCURIC ACETATE	0	0	0	0	00-30-4	1700	
PHENYLACETIC ACID	0	0	0	0	100-02-2	10000	
PHENYLACETIC PEROXIDE	0	0	0	0	70	1000	
PHENYLACETOXYBENZENE 4	0	0	0	0	4004-75-1	2	30:2;100000000
PHENYLENE DIAMINE(-o)	0	0	0	0	100-45-2	0	24:2;1000000000
PHENYLENE DIAMINE(-o)	0	0	0	0	95-54-5	0	24:2;1000000000
PHENYLENE DIAMINE(-p)	0	0	0	0	100-00-3	0	24:2;1000000000
PHENYLHYDRAZINE	0	0	0	0	100-00-0	100	
PHENYLPHENOL P	0	0	0	0	90-00-3	700	30:2;100000000
PHENYLTHIOUREA	.70	0	0	0	100-00-5	2000	
PHOSATE	0	0	0	0	200-00-2	00	
PHOSGENE	1.170	0	0	0	75-44-5	0	21224000000000
PHOSPHINE	0	0	0	0	7000-01-2	300	
PHTHALIC ACID	0	0	0	0	90-00-3	2000	34:2;2;10000000
PHTHALIC ANHYDRIDE	-.02	0	0	0	90-44-0	0200	
PHTHALIMIDE	0	0	0	0	90-41-0	2000	24:2;1000000000
PICOLINE(2-)	1.00	0	0	0	100-00-0	10000	
PINENE(alpha-)	0	0	0	0	90-00-0	2	53122231410000
PITERAZINE	0	0	0	0	110-00-0	10000	2222200000000000
POLYCHLORINATED BIPHENYLS	0	0	0	0		0	
POLYCYCLIC KETONE 0	0	0	0	0	010	2	
PROPANE	0	0	0	0	74-00-0	2000	2211200000000000
PROPANE SULTONE,1,3-	-2.00	0	0	0	1120-71-4	0	
PROPANE,2,2'-OXYDIS(2-CHLORO-	0	0	0	0	30000-32-0	0	
PROPANOIC ACID	0	0	0	0	70-00-4	0	
PROPANOL	0	0	0	0		0	3111220000000000
PROPANOL ISO	-.10	10	.070	0	07-00-0	100000	31714210000000
PROPIOLACTONE b	0	0	0	0	07-07-0	300000	
PROPIONALDEHYDE	0	0	0	0	123-20-0	20000	3111210000000001
PROPIONIC ACID	0	0	0	0	70-00-4	20000	311121;10000000
PROPIONITRILE	.04	0	0	0	107-12-0	00000	
PROPYL ACETATE ISO	0	0	0	0	100-21-4	30000	321131F000000000
PROPYL AMINE ISO	0	0	0	0	70-21-0	100000	
PROPYL ETHER ISO	0	0	0	0	100-20-3	0000	3411310000000000
PROPYL THIOURACIL	0	0	0	0	01-02-0	1100	
PROPYLENE	0	0	0	0	110-07-1	200	2111000000000000
PROPYLENE GLYCOL	0	0	0	0	07-00-0	20000	41112132700000
PROPYLENE OXIDE	0	0	0	0	70-00-0	30000	2111000000000000
PROPYLENEDINE,1,2-	0	0	0	0	70-00-0	0	
PROPYL(-n) ACETATE	0	0	0	0	100-00-4	10000	311221F000000000
PROPYL(-n) BENZENE	0	0	0	0	100-00-1	00	411120;1-000000
PROPYL-3-METHOXY PYRAZINE,2- ISO	0	0	0	0	20773-40-0	0	
PROPYL-1-OL 2(PROPARGYL)	0	0	0	0	107-19-7	100000	312171-00000000
PYRENE	0.10	0	0	0	120-00-0	2	
PYRIDINE	.00	30.00	0	0	110-00-1	20000	1100000000000000
PYRIDINIUM BROMIDE	0	0	0	0	000	20000	
QUINALDINE	0	0	0	0	01-03-4	0	
QUINOLINE	2.00	0	0	0	01-23-0	0	
RESERFINE	0	0	0	0	00-00-0	2	
RESORCINOL	.0	0	0	0	100-00-3	2000	24:200000000000
SACCH. CYCLO-HEX. DIMETH. PHOS. DITHIOATE	0	0	0	0	040	2	
SACCHARIN	.01	0	0	0	01-07-2	3400	
SAPROLE	0	0	0	0	04-00-7	0	
SILVEX	0	0	0	0	00-72-1	140	22:30700111(1;0
SIMAZINE	0	0	0	0	122-34-0	2	
SODIUM ACETATE	0	0	0	0	704	20000	
SODIUM DODECYL SULFATE	1.0	0	0	0	101-21-3	100000	
SODIUM DODECYLBENZENE SULFONATE	0	0	0	0	20100-30-20		
SODIUM FORMATE	0	0	0	0	141-03-7	40000	
STREPTOTOCIN	-1.40	0	0	0	10003-00-0	0	
STRYCHNIDIN-10-ONE,2,3-DIMETHOXY-	1.00	0	0	0	307-07-3	700	
STRYCHNINE	1.00	0	0	0	07-24-0	100	
STYRENE	2.10	0	.11	0	100-42-0	300	3100;1;00000000
SUCCINIC ACID	0	0	0	0	110-10-0	00000	2222;1000000000
SUCCINIMIDE	0	0	0	0	120-00-0	300000	
SULFANILIC ACID	0	0	0	0	121-47-1	10000	
SULFIDE	0	0	0	0	7700-00-4	4132.2	
TARS	0	0	0	0	000	.2	
TETRACHLOROQUINONE	0	0	0	0		2	

NAME	Ln (00)	SDV	K1	Rb HND	CAS	SQL	unccode
MUSTARD GAS	0	0	0	0	505-60-2	0	
NAPHTHALENE	2.37	42.47	1	0		0	28:2;000000000
NAPHTHOL, alpha-	0	0	0	0	90-15-3	0	37:2;100000000
NAPHTHOL, beta-	2.84	0	0	0	120-19-3	1000	
NAPHTHOLONE-1,4	0	0	0	0	120-15-4	1000	
NAPHTHOLAMINE, alpha-	2.22	0	0	0	134-30-7	1700	31:7;2;0000000
NAPHTHOLAMINE, beta-	2.25	0	0	0	91-00-0	0	31:7;2;0000000
NEPHTHYL GLYCOL	0	0	0	0	120-30-7	000000	42:22142;00000
NEACIN	0	0	0	0	50-67-0	20000	
NEACINAMIDE	0	0	0	0	90-00-0	10000	
NEOCL CYANIDE	0	0	0	0	507-19-7	2	
NETRONLINE P	0	0	0	0	100-01-0	000	34:111;0000000
NETRONLINE(-o)	1.00	0	0	0	80-74-4	1200	
NITROBENZENE	1.04	11	2.3	0	90-00-0	1000	25:1;000000000
NITROBENZENESULFONYL CHLORIDE P	0	0	0	0	90-74-0	00	
NITROBENZYL ALCOHOL P	0	0	0	0	610-73-0	10000	
NITROBIPHENYL, 4-	0	0	0	0	90-00-0	0	
NITROCELLULOSE	0	0	0	0	9004-70-0	10	
NITROGEN MUSTARD N-OXIDE	0	0	0	0	120-05-2	0	
NITROGLYCERIN	0	0	0	0	50-00-0	0	22:1;000000000
NITROMETHANE	0	0	0	0	75-50-5	90	11;00000000000
NITROPHENOL, 3-	1.70	0	0	0	90-70-5	2100	34:101;0000000
NITROPHENOL, 4-	1.01	0	0	0	100-00-7	10000	34:101;0000000
NITROPROPANE 2	0	0	0	0	70-40-0	100000	22:1;000000000
NITROSODIMETHYL ALCOHOL 4	0	0	0	0	701	1000	
NITROSODIMETHYLAMINE N	-0.47	0	0	0	60-70-0	1000000	
NITROSODIPHENYLAMINE N	2.13	0	0	0	90-30-0	0	
NITROSODI-n-PROPYLAMINE N	1.31	0	0	0	021-04-7	9000	
NITROSOMORPHOLINE	-0.44	0	0	0		0	
NITROSPYRROLIDINE N	0	0	0	0	900-00-2	2	
NITROSO-N-METHYLUREA N	0	0	0	0	004-00-0	2	
NITROTOLUENE (-p)	0	0	0	0	90-00-0	442	34:1<1;0000000
NITRO-4-METHYLBENZATE 3	0	0	0	0		20	
NONYLPHENOL	0	0	0	0	25104-02-0	0	
OCTAMETHYLPHOSPHONAMIDE	0	0	0	0	102-10-0	0	
OCTANE	0	0	0	0	111-00-0	.00	22:02000000000
OCTANOL 1	0	0	0	0	111-07-0	300	31:721;0000000
OCTANOL 2	0	0	0	0		0	42:152131;00000
OCTANOL 3	0	0	0	0		0	42:152131;00000
OCTANOL 4	0	0	0	0		0	42:152131;00000
OIL	0	0	0	0		0	22:1;20000000000
ODIMETHYLETH. THIOETH. PHOSPHOROTHIOATE	0	0	0	0	120-70-0	2	
OXALIC ACID	0	0	0	0	144-02-7	20000	12;00000000000
OXAMIC ACID	0	0	0	0	471-07-0	100	
PARABENOPHENOL	2.00	0	0	0	100-01-2	140000	
PARAFORMALDEHYDE	0	0	0	0	30020-00-0	100	
PARALDEHYDE	0	0	0	0	120-00-7	120000	23:13;000000000
PARATHION	0	0	0	0	50-20-2	0	42:14;2<1;00000
PCB 1010	4.30	0	0	0	12074-11-	.040	
PCB 1221	2.0	0	0	0	11104-20-	.00	
PCB 1222	3.2	0	0	0	11141-10-	1.40	
PCB 1240	4.11	0	0	0	03400-21-	.1	
PCB 1240	0.70	0	0	0	12072-20-	.004	
PCB 1254	0.00	0	.00	0	11007-00-	.007	
PCB 1260	7.14	0	0	0	11000-02-	.00	
PCB'S	0	0	0	0		1.40	
PENTACHLOROBENZENE	0	0	0	0	000-00-0	0	21:5;000000000
PENTACHLORETHANE	0	0	0	0	70-01-7	0	21:0;000000000
PENTACHLORONITROBENZENE	0	0	0	0		0	25:1;000000000
PENTACHLOROPHENOL	0.01	130	3.4	0	07-00-0	14	21:05;000000000
PENTAZINE 1,2	0	0	0	0		0	21:27;000000000
PENTAZOTHRITOL	0	0	0	0	110-77-0	00000	
PENTAZOTHRITOL TETRANITRATE	0	0	0	0	70-11-0	10	
PERCHLOROMETHYL MERCAPTAN	0	0	0	0	004-02-0	0	
PERYLENE	0	0	0	0	100-00-0	0	
PHENACETIN	1.00	0	0	0	02-44-2	700	
PHENANTHRENE	4.40	1.0	2.2	0	85-01-0	0	
PHENOL	1.40	97	13	0	100-00-2	00000	25:10;000000000
PHENOL, 2-(1,1-DIMETHYLETHYL)-	0	0	0	0	000-34-2	0	
PHENOTHAZINE	0	0	0	0	00-04-2	1	

NAME	Ln(OM)	SDV	K1	Rb HYD	CAS	SL	unicode
TETRACHLOROBENZENE(1,2,3,4)	0	0	0	0	634-88-2	2	22:4f00000000
TETRACHLOROBENZENE(1,2,3,5)	0	0	0	0	634-88-2	2	22:4f00000000
TETRACHLOROBENZENE(1,2,4,5)	4.51	0	0	.42	95-04-3	.3	22:4f00000000
TETRACHLORODIBENZOFURAN 2,3,7,8	0	0	0	0	688	2	
TETRACHLORODIBENZOFURAN,2,3,7,8-	0	0	0	0	51287-21-	0	
TETRACHLORODIBENZOP-DIOXIN,2,3,7,8-	6.64	0	0	0	1740-61-6	.00002	
TETRACHLOROETHANE(1,1,1,2)	0	0	0	0	630-20-6	200	12:0000000000
TETRACHLOROETHANE(1,1,2,2)	2.88	6.2	.00	0	78-34-6	2300	2194:00000000
TETRACHLOROETHYLENE	0	0	0	0	127-18-4	0	2194:00000000
TETRACHLOROETHYLENE	2.6	6.2	.00	0	127-18-4	150	2194:00000000
TETRACHLOROPHENOL(2,3,4,6)	0	0	0	.35	88-08-2	0	31:1B4f000000
TETRACHLOROPHENOL(2,3,5,6)	0	0	0	0	98-05-5	2	31:1B4f000000
TETRACHLOROPROPENE(1,1,2,3)	0	0	0	.30	10430-30-	2	
TETRAETHYL LEAD	0	0	0	0	78-08-2	.8	
TETRAETHYLDITHIOPYRPHOSPHATE	0	0	0	0	3000-24-5	25	
TETRAETHYLDITHIOPYRPHOSPHATE	0	0	0	0	3000-24-5	25	
TETRAETHYLENE GLYCOL	0	0	0	0	112-09-7	20000	
TETRAETHYLPYRPHOSPHATE	0	0	0	0	107-40-3	0	
TETRAHYDROFURAN	0	0	0	.004	100-00-0	0	2321:00000000
TETRAHYDRONAPHTHALENE,1,2,3,4-	0	0	0	0	119-04-2	0	
TETRALIN	0	0	0	0	0	0	3424:2:000000
TETRAMETROMETHANE	0	0	0	0	880-14-0	0	14:0000000000
THIOACETAMIDE	0	0	0	0	62-55-6	140000	
THIOBENZYL ALCOHOL P	0	0	0	0	100-63-0	1000	
THIOCYANATE (TOTAL AS SCN-)	0	0	0	0	NA	0	
THIOFANOL	0	0	0	0	30190-19-	5200	
THIOETHANOL	0	0	0	0	74-08-1	23300	
THIOPHENOL	2.82	0	0	0	100-00-5	470	
THIOPROPIONAMIDE 2	0	0	0	0	91	2000	
THIOSEMICARBAZIDE	-1.14	0	0	.000	70-19-0	20000	
THIOUREA	0	0	0	.45	62-56-6	91000	2201:00000000
THIOUREA,1-(o-CHLOROPHENYL)-	0	0	0	0	5344-82-1	0	44:10111f0000
THURAM	0	0	0	0	137-26-0	30	
THYMINE	-.44	0	0	0	66-71-4	4000	
TOLUENE	2.00	73.48	2.4	0	100-00-3	515	25:1:00000000
TOLUENE DIAMINE(2,4)	0	0	0	.07	95-00-7	2000	33:1:00000000
TOLUENE DIISOCYANATE(2,4)	0	0	0	0	504-04-0	0	
TOLUENE-2-OIAZOIS-METATOLUENE-DIAMINE HCl	0	0	0	0	123	1000	
TOLUENEDIAMINE,2,4-	0	0	0	0	95-00-7	0	33:1:00000000
TOLUENEDIAMINE,2,6-	0	0	0	0	823-46-5	0	33:1:00000000
TOLUENEDIAMINE,3,4-	0	0	0	0	400-72-0	0	33:1:00000000
TOLUIC ACID (para-)	0	0	0	.51	90-04-6	0	44:1:1:000000
TOLUIC ALDEHYDE	0	0	0	0	122-70-1	100	44:1:1:000000
TOLLIDINE HYDROCHLORIDE,o-	0	0	0	0	630-21-6	0	
TOLLIDINE P	1.30	0	0	0	100-40-0	7400	34:1:00000000
TOLLIDINE (-0)	0	0	0	0	95-53-4	0	34:1:00000000
TOLUENE	3.3	0	0	0	8001-35-2	.5	
TRIFLUOROMETHYLPHOSPHATE	0	0	0	.30	544	10	
TRIBUTYL PHOSPHOTRITHIODATE SSS	0	0	0	0	70-40-0	2	
TRIBUTYLPHOSPHATE	0	0	0	0	120-73-0	5000	23192000000000
TRICHLORANISOLE,2,3,6-	0	0	0	0	50075-10-	0	
TRICHLOROBENZENE 1,2,3	0	0	0	0	87-61-6	0	23:3f00000000
TRICHLOROBENZENE 1,2,4	3.98	0	0	.05	120-82-1	19	23:3f00000000
TRICHLOROBENZENE 1,3,6	0	0	0	0	100-70-3	0	23:3f00000000
TRICHLOROBUTANE 1,2,3	0	0	0	0	10330-40-	.2	
TRICHLOROETHANE 1,1,1	2.40	3.5	.74	.30	71-55-6	4400	2111:00000000
TRICHLOROETHANE 1,1,2	2.17	3.5	.74	0	70-00-6	4500	11a1]00000000
TRICHLOROETHYLENE	2.29	3.9	.00	.12	70-01-6	0	21:1b00000000
TRICHLOROFLOROMETHANE	2.53	0	0	0	75-00-4	0	
TRICHLOROPHENOL(2,4,6)	0	0	0	0	95-05-4	1100	32:1B3f000000
TRICHLOROPHENOL(2,4,6)	3.00	0	.20	0	80-00-2	000	31B3f2:000000
TRICHLOROPROPANE(1,1,1)	0	0	0	.07	7700-00-1	0	22]1"00000000
TRICHLOROPROPANE(1,1,2)	0	0	0	.000	500-77-6	0	3111:1a000000
TRICHLOROPROPANE(1,2,2)	0	0	0	0	3175-23-3	0	3111]1b000000
TRICHLOROPROPANE(1,2,3)	0	0	0	0	90-10-4	0	22]1"00000000
TRICHLOROPROPENE (1,1,2)	0	0	0	0	003	2	
TRICHLORO(1,1,2) TRIFLUORO	0	.070	0	0	0	0	
TRICHLORO(1,1,2) TRIFLUOROETHANE(1,2,2)	0	0	0	0	70-13-1	170	
TRICHLORO-1,2,2-TRIFLUOROETHANE,1,1,2-	0	0	0	0	70-13-1	170	12:0000000000
TRICHLORO-1,3,5-TRIAZINE 2,4,6	0	0	0	0	100-77-0	.2	

NAME	Ln(OH)	RDV	k1	Rb H2O	CAS	SL	united
TRICORNE N	0	0	0	0	629-85-5	2	
TRIMELAMINE	-1.75	0	0	0	100-71-0	100000	
TRIMELAMINE	0	0	0	0	121-44-0	20000	
TRIMELINE GLYCOL	0	0	0	.002	112-27-0	20000	
TRIMELPHOSPHORIC ACID, o, o, o-	0	0	0	0	120-69-1	0	
TRIFLUOROMETHANE(1,1,1)	0	0	0	0	000	2	2111:000000000
TRIGLYLIC ANHYDRIDE	0	0	0	.700	552-25-7	0	
TRIMELAMINE	0	0	0	0	75-05-5	100000	
TRIMELAMINE(1,3,5)	0	0	0	0	100-67-3	2	
TRIMELPHOSPHINE 2,2,4	0	0	0	0	540-04-1	.00	
TRIMEL-4-NITROGLYCOLINE 2,1,5	0	0	0	0	00	2	
TRIMETHYLENE, o, o, o-	0	0	0	0	90-35-4	300	
TRIMETHYLENE(2,4,6)	0	4.4	.45	0	110-05-7	1000	3111:02:0000000
TRIPHENYL PHOSPHINE	0	0	0	0	000-35-0	2	
TRIPHENYLAMINE	0	0	0	0	510-73-3	0	
TRIPHENYLPHOSPHINE NICKEL CARBONYL	0	0	0	0	207	10	
TRIS (1-ACETOXYMETHYL) PHOSPHINE SULFIDE	0	0	0	0	00-04-4	100000	
TRIS (2,3-DIBROMOPROPYL) PHOSPHATE	0	0	0	0	120-72-7	0	
TRISOLM NITROTRICACETATE	0	0	0	0	0004-21-3	0	
TRYPAN BLUE	0	0	0	0	72-67-1	0	
URACIL	-1.07	0	0	0	00-22-0	2000	
URACIL MUSTARD	0	0	0	0	00-75-1	0	
UREA	0	0	0	.15	57-13-0	2000	2120:000000000
URETHANE	0	0	0	.01	01-70-0	20000	411121:0000000
VALERIC ACID	0	0	0	0	100-52-4	2000	
VINYL ACETATE	0	0	0	0	100-05-4	0	2151:000000000
VINYL CHLORIDE	.00	0	0	.00	75-01-4	1.1	
VINYLCYCLOHEXANE 4	0	0	0	0	100-05-3	2	
VINYLIDENE CHLORIDE	0	0	0	0	0	0	2172:000000000
WAFARIN	0	0	0	0	01-01-2	0	011220:3;30000
WATER	0	0	0	0	0	0	1140:000000000
XYLENE	1.15	40.0	1.0	.001	130-05-7	0	24:2:000000000
XYLENE(-o)	2.2	0	0	.07	100-30-3	200	24:2:000000000
XYLENE(-p)	2.05	40.70	1.0	.12	05-47-6	170	24:2:000000000
XYLENE(-m)	1.15	0	0	0	100-07-0	100	24:2:000000000
XYLENOL(3,4)	.1	0	0	0	05-05-0	0	4213:2;100000
XYLYL CHLORIDE M	0	0	0	0	020-10-0	10	
XYLYL CHLORIDE O	0	0	0	0	020-45-4	10	
ZINC ACETATE	0	0	0	0	000	20000	

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	16. ABSTRACT <p>Analytical models are presented for estimating air emissions from hazardous waste treatment, storage, and disposal facilities (TSDF). Air emission models have been developed for aerated and nonaerated surface impoundments, land treatment facilities, landfills, and wastepiles. Emission model predictions are compared to available field data. This report also includes emission factors for transfer, storage, and handling operations at TSDF.</p> <p>The models have been assembled into a spreadsheet that is included in this report as floppy diskette for use on a microcomputer.</p> <p>Appendices include a list of physical-chemical properties for approximately 700 compounds and a comprehensive source list of pertinent literature in addition to that cited in the report.</p>	
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