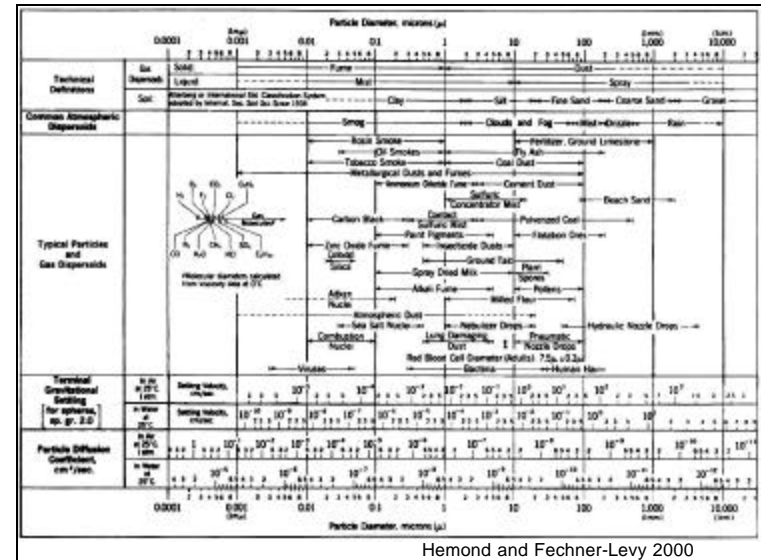


“Movement of Particulates (and Air-Water Exchange)”

Module 2: Surface Waters, Lecture 4

Chemical Fate and Transport in the Environment, 2nd edition. H.F. Hemond and E.J. Fechner-Levy. Academic Press. London. 2000.



Stoke's Law to predict settling of particles (type 1, discrete, non-interacting):

$$w_f = \frac{(2/9) \cdot g \cdot (r_s / r_f - 1) \cdot r^2}{h_f}$$

With the sediment and fluid densities, particle size, and kinematic viscosity.

Downward flux due to settling [M/L²T] is therefore:

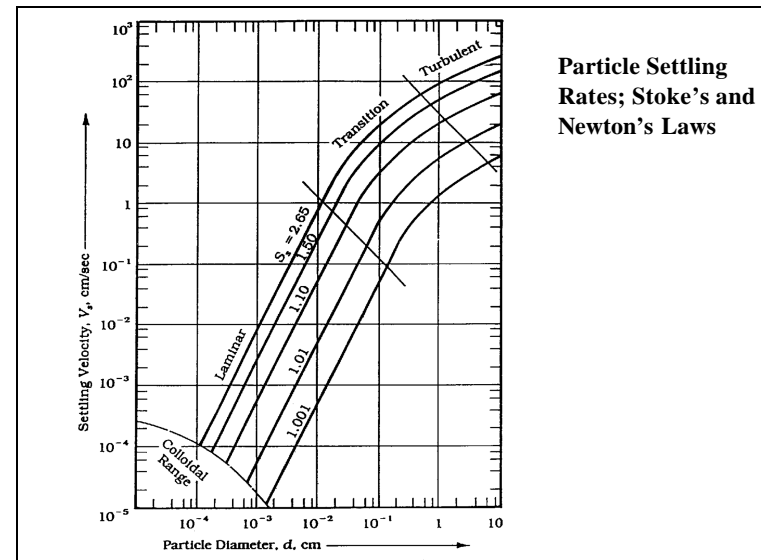
$$J_{Stokes} = C \cdot w_f$$

Corresponding upward flux density:

$$J_{Fickian} = D \cdot dC / dx$$

Vertical concentration profile of particles at steady state:

$$C = C_o \cdot e^{-(w_f / D) \cdot x}$$



Example 2-3

Describe the steady-state distribution of 1 μm clay particles in still water. Assume kinematic viscosity of water at 50°F is 0.013 cm^2/sec and a solid density of 2.6 g/cm^3 . Also calculate the depth when the concentration is halved.

The settling rate for the 1 μm particle is therefore:

$$w_f = \frac{(2/9) \cdot g \cdot (r_p / r_f - 1) \cdot r^2}{h_f} = \frac{(2/9) \cdot 981 \text{ cm/sec}^2 \cdot ((2.6 \text{ g/cm}^3) / (1 \text{ g/cm}^3) - 1) \cdot (5 \times 10^{-5} \text{ cm})^2}{0.013 \text{ cm}^2 / \text{sec}}$$

$$= 6.7 \times 10^{-5} \text{ cm} / \text{sec}$$

The diffusion coefficient for this particle is about $5 \times 10^{-9} \text{ cm}^2/\text{sec}$ (from fig 2-11). The first-order decay coefficient for the large particle is therefore:

$$w_f / D = \frac{6.7 \times 10^{-5} \text{ cm/sec}}{5 \times 10^{-9} \text{ cm}^2 / \text{sec}} = 1.3 \times 10^4 / \text{cm}$$

The distance at which the concentration is halved is:

$$\frac{1}{2} = e^{-(1.3 \times 10^4 / \text{cm}) \cdot x}$$

$$x = 5 \times 10^{-5} \text{ cm} \text{ or } 0.5 \mu\text{m}$$

Example 2-4

What is the minimum distance the particles will travel before settling to the river bottom? Assume a 2-m river depth and 200 μm particles, 2.6 g/cm^3 particle density and $1.3 \times 10^{-2} \text{ cm}^2/\text{sec}$ kinematic viscosity.

Settling rate:

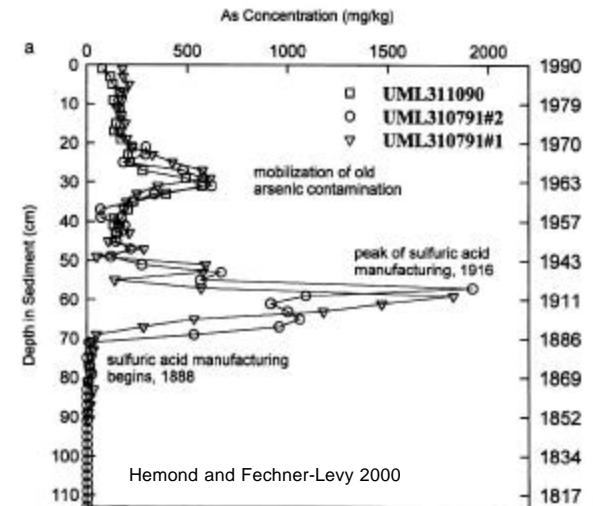
$$w_f = \frac{(2/9) \cdot 981 \text{ cm/sec}^2 \cdot ((2.6 \text{ g/cm}^3) / (1 \text{ g/cm}^3) - 1) \cdot (10^{-2} \text{ cm})^2}{0.013 \text{ cm}^2 / \text{sec}}$$

$$w_f = 2.7 \text{ cm} / \text{sec}$$

The time required to settle 2 m is therefore 75 sec, and the particle will travel:

$$75 \text{ sec} \cdot 0.2 \text{ m} / \text{sec} = 15 \text{ m}$$

Sediment accumulations:



Example 2-5

Sediment from a 10 cm depth of a lake has a ^{210}Pb activity of 2.5 disintegrations per minute (DPM), while surface sediment has a DPM of 4. How rapidly does sediment accumulate in this lake?

Using the basic equation for radioactive decay:

$$t = \frac{-1}{I} \cdot \ln\left(\frac{A_d}{A_o}\right) = \frac{-1}{0.03/\text{year}} \ln\left(\frac{2.5\text{DPM}}{4\text{DPM}}\right) = 16/\text{year}$$

$$\frac{10\text{cm}}{16\text{year}} = 0.6\text{cm}/\text{year}$$

Air-Water Exchange

The concentration of a dissolved gas in water at equilibrium with the atmosphere is:

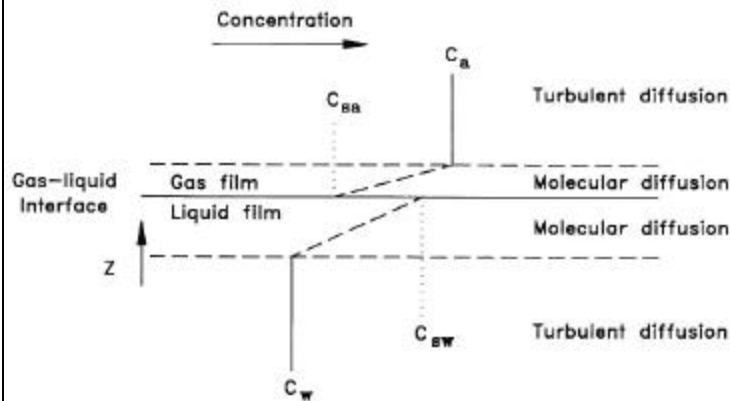
$$C_{\text{equil}} = C_a / H$$

Where C_a is the concentration in air, and H is the Henry's law constant.

The flux density is proportional to the product of the difference between the actual and the equilibrium concentrations :

$$J = -k_w(C_w - C_a / H)$$

Thin Film Model:



Hemond and Fechner-Levy 2000

For water-side control, the flux of the chemical out/in water is:

$$J = -\frac{D_w}{d_w} C_w = -k_w C_w$$

Which considers the molecular diffusion coefficient, the film thickness, and the concentration in the water. The thickness cannot be directly measured, but the k_w factor can, and with an estimate of the diffusion coefficient, the film thickness can also be estimated

The general case (needed when H is on the order of 0.01) and both air and water affect the gas transfer:

$$J = -\left[\frac{1}{d_w / D_w + d_a / (D_a \cdot H)} \right] \left[C_w - \frac{C_a}{H} \right]$$

Estimation of gas exchange coefficients can be made based on the relative molecular weights of a gas tracer and the gas of interest:

$$\frac{k_A}{k_B} = \frac{D_A}{D_B} \approx \frac{\sqrt{MW_B}}{\sqrt{MW_A}}$$

The gas exchange coefficients can also be estimated based on wind speed (measured 10 m above the water). For water side control:

$$k_w (cm/sec) \approx 4 \times 10^{-4} + 4 \times 10^{-5} \cdot u_{10}^2$$

For air-side control (and cm/hr units for k_a):

$$k_a (cm/hr) \approx 1100 \cdot u (m/sec)$$

Example 2-6

The dissolved concentration of trichloroethylene (TCE) (C_2Cl_3H) is 1 μ/L . H is 0.4 and the measured gas exchange rate coefficient for propane (C_3H_8) in water is 3×10^{-3} cm/sec. What is the flux density for TCE from the lake?

MW TCE = 131 g/mol and MW propane = 44 g/mol. Assuming C_a is almost 0. Diffusion through air is not a bottleneck due to fairly high H . D_w for TCE is estimated by the D_w for propane multiplied by the roots of the ratios of the MW:

$$\frac{D_{TCE}}{d_w} = 3 \times 10^{-3} \text{ cm/sec} \cdot \frac{\sqrt{44}}{\sqrt{131}} = 1.7 \times 10^{-3} \text{ cm/sec}$$

$$J = -\frac{D_w}{d_w} C_w = -k_w C_w$$

$$J_{TCE} = (1.7 \times 10^{-3} \text{ cm/sec}) \left(\frac{1 \text{ mg}}{\text{liter}} \right) \left(\frac{1 \text{ liter}}{1,000 \text{ cm}^3} \right) = 1.7 \times 10^{-6} \frac{\text{mg}}{\text{cm}^2 \cdot \text{sec}}$$

Volatilization from Pure Phase Liquids

The concentration at the base of a stagnant air layer, just above the surface of a NAPL, is determined by:

$$C_a = \frac{P}{RT} (MW)$$

Where P is the vapor pressure of the chemical, R is the universal gas constant, T is the absolute temperature, and MW is the molecular weight of the chemical (g/mol).

The volatilization rate from the NAPL surface is:

$$J = \frac{-D_a}{d_a} \cdot C_a$$

Example 2-6

Benzene is spilled in a lake from an overturned tanker truck. The 10m high wind speed is 3 m/sec. What is the flux density from the slick?

The concentration of benzene at the air-NAPL interface is:

$$C_a = \frac{P}{RT} (MW) = \frac{(0.12 \text{ atm})}{(293 \text{ K})} \left(\frac{0.082 \text{ atm} \cdot \text{liter}}{\text{mol} \cdot \text{K}} \right) \cdot \frac{78 \text{ g}}{\text{mol}} = 0.4 \text{ g/liter}$$

The flux density is therefore:

$$J = \left(\frac{3300 \text{ cm}}{\text{hr}} \right) \left(\frac{0.4 \text{ g}}{\text{liter}} \right) \left(\frac{1 \text{ liter}}{1,000 \text{ cm}^3} \right) = \frac{1.3 \text{ g}}{\text{cm}^2 \cdot \text{hr}} \Rightarrow \frac{360 \text{ mg}}{\text{cm}^2 \cdot \text{sec}}$$