1.061 / 1.61 Transport Processes in the Environment Fall 2008

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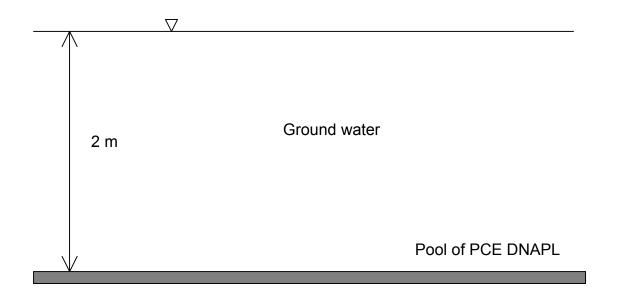
A smokestack of height H = 20 m releases two gases, dichlorodifluoromethane (Freon 12) and trichloroethene (TCE), each at a rate of 5 kg/min. Freon 12 is conservative. TCE undergoes first-order degradation in the atmosphere at a rate of $k_{TCE} = 0.1 \text{ day}^{-1}$, producing the highly toxic chemical phosgene (C(=O)Cl₂). Assume that the wind blows steadily and uniformly at 5 m/s in the positive x direction. The atmospheric turbulence is homogeneous but anisotropic, with the vertical diffusivity, $D_Z = 0.1 \text{ m}^2\text{s}^{-1}$, smaller than the horizontal diffusivities, $D_X = D_Y = 1 \text{ m}^2\text{s}^{-1}$. For both gases the ground acts as a no-flux boundary. Find the maximum concentration of Freon and TCE 10-km downwind of the stack.

Problem 9.2

A small channel is h = 5 cm deep and b = 10 cm wide. It carries flow at U = 10 cms⁻¹. The stream-wise coordinate is x. The vertical coordinate is z, with z = 0 at the bed and positive upward. A continuous source of dye is injected at a rate of $\dot{m} = 1$ gs⁻¹ at middepth and mid-width, and at x = 0. Assume that the channel has no dye upstream of the injection point. The bed of the channel is a perfect absorber for the dye, such that the concentration of dye in equilibrium with the bed is zero, and thus C(z=0) = 0. The molecular diffusivity for the dye is $D = 10^{-5}$ cm²s⁻¹. What is the maximum concentration in the channel 20 m downstream of the source?

Problem 9.3

Dense non-aqueous phase liquids (DNAPLs) are liquids that are heavier than water and have very low solubility in water. Typical DNAPLs include chlorinated solvents like the dry-cleaning fluid perchloroethylene (PCE), also called tetrachloroethylene or tetrachloroethene. When a DNAPL enters an aquifer it will sink under gravity until it encounters a layer of low permeability, such as clay. It then spreads into a thin layer. Consider the pool of PCE depicted below which was created at time t = 0. For t < 0 the PCE concentration in the aquifer is zero. For t > 0, PCE slowly diffuses into the water above. The coefficient of diffusion is D = 4.4 x 10⁻⁹ m²s⁻¹. The ground water is stagnant and the aquifer is 2 meters thick above the DNAPL. PCE has a solubility in water of 150 mg/L. The Maximum Contaminant Level (MCL) for PCE in drinking water is 5 ppb. When will the concentration of PCE throughout the aquifer be above the MCL?



You have identified a point source of TCE that is contaminating a small stream. The stream is h=20 cm deep, b=80 cm wide and flows at U = 10 cm/s. At the source (x = 0) the TCE mixes quickly across the channel depth and width with the resulting initial concentration, $C(x=0)=C_0=10$ ppb. You wish to determine if there are additional sources of TCE to the river. Because TCE is volatile and the concentration in the atmosphere is negligible, you know there is a flux of TCE from the river to the atmosphere. The Henry's Law constant for TCE is $H_{TCE} = 0.42$, indicating that the flux is water-side controlled. Because of the flux to the atmosphere, you expect the TCE concentration to decline downstream. Indeed, 2 km downstream of the known source $C_{2km} = 5$ ppb. To determine the rate of water-air exchange for the TCE you inject and measure the concentration of Propane [$H_{propane} = 0.42$] along the stream. From this study you find that $K_{propane} = 1.5 \times 10^{-4} \text{ s}^{-1}$. The molecular diffusion coefficients for TCE and Propane in water are $D_{w, TCE} = 0.75 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$ and $D_{w, Propane} = 1.3 \times 10^{-5} \text{ cm}^2 \text{ s}^{-1}$.

Determine if additional sources of TCE exist along the reach x = 0 to 2 km.

A small pond is h = 1 m deep with a surface area A. Three chemicals are spilled into the pond and rapidly mixed over the volume. The chemicals are the pesticide Lindane (w/ Henry's Law constant $H_L = 2.2 \times 10^{-5}$), the solvent Toluene ($H_T = 0.28$), and Napthalene ($H_N = 0.04$). Assume that mixing is sufficient to maintain a uniform concentration of each chemical within the bulk of the lake volume, i.e. below the laminar sub-layer at the surface.

Molecular diffusivity in air, $D_a = 10^{-5} \text{ m}^2 \text{s}^{-1}$ for all chemicals

Molecular diffusivity in water, $D_w = 10^{-9} \text{ m}^2 \text{s}^{-1}$ for all chemicals

Turbulent diffusivity in water, $D_{tw} = 10^{-3} \text{ m}^2 \text{s}^{-1}$

Waterside laminar sub-layer, $\delta_w = 100 \ \mu m$

Airside laminar sub-layer, $\delta_a = 10 \text{ mm}$

- a) Sketch the profile of C (z) for each chemical. Indicate the concentration at z = 0, the air-water interface; at $z = -\delta_w$; at $z < -\delta_w$; and at $z = +\delta_a$. Assume that the atmosphere is a perfect sink for each chemical, such that $C_a = 0$ for $z > +\delta_a$.
- b) Write an equation for the mass flux at the air-water interface for each chemical.
- c) For each chemical determine the time at which only 5% of the original mass remains.
- d) For which chemicals is the assumption of a uniform concentration within the bulk fluid appropriate?

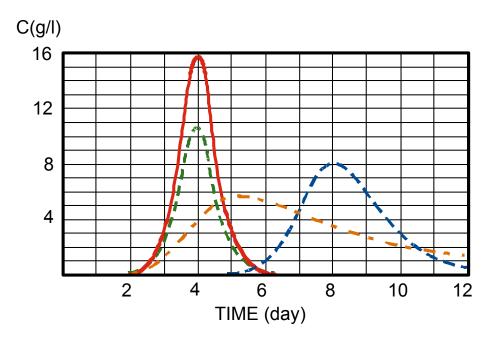
A mixture containing an equal mass of four chemicals is released as a pulse into a groundwater aquifer. The concentration of each chemical is measured in the pore water at a monitoring well located 8-m from the injection. Match the following descriptions to the correct curves.

Chemical 1 is a conservative tracer that does not react or degrade.

Chemical 2 does not adsorb to the grains, but is degraded by microbes living in the aquifer. What is the rate of degradation, $K_d [day^{-1}]$?

Chemical 3 readily adsorbs to organic material. The rate of adsorption/desorption is so rapid that the water/organic matter partitioning is always at equilibrium. What fraction of this chemical is associated with the pore water, *i.e.* is in the mobile phase?

Chemical 4 adsorbs to organic material, but at a rate that is much slower than that of chemical 3, such that the solid/water partitioning is never at equilibrium. What is the order of magnitude of the rate constant describing the adsorption/desorption reaction?



Between midnight and 2 am, illegal dumpers empty two five-gallon drums containing 2 kg of Toluene into a shallow abandoned well. Within 24 hours the Toluene is distributed vertically over the shallow (5 m thick), sandy aquifer. Evidence of the dumping is discovered two days later, and you are asked to assess the risk to a drinking well located 1 km directly downstream, if no remediation is done. Previous tests on this aquifer indicate the following:

Mean pore velocity, u = 1 m /dayIsotropic, Homogeneous Dispersivity, $K = 0.1 \text{ m}^2/\text{day}$. Porosity, n = 0.3Solids density, $\rho_S = 2.6 \text{ g/mL}$.

Toluene partitions rapidly to aquifer solids and has a solid-water partitioning coefficient of

 $K_{d} = 0.5 \frac{g_{toluene} / kg_{solid}}{g_{toluene} / L_{water}}.$

Assume that the partitioning of Toluene is everywhere in equilibrium.

- a) Write an appropriate transport equation.
- b) Estimate the total concentration, C(t), at the drinking well.
- c) Estimate the peak concentration in the pore water at the well and the duration of exposure.

