



SI 17/18 EN 3003 ENVIRONMENTAL TRANSPORT PROCESS.

Date

YEO PEI QI

No.

Q1 a) (i)  $x = 0.1 \text{ mm/s} \times 10 \text{ hr} \times 3600 \frac{\text{s}}{\text{hr}}$   
 $= 3600 \text{ mm}$   
 $= 3.6 \text{ m}.$

(ii)  $4\sigma = 4\sqrt{2D_{eff}t}$   
 $= 4\sqrt{2(5.21 \times 10^{-8})(10 \times 3600)}$   
 $= 24.499 \text{ cm}.$

(iii)  $C(3.5 \text{ m}, 10 \text{ hr}) = \left( \frac{M}{\sqrt{4\pi D_{eff}t}} \right) e^{-\frac{(x-ut)^2}{4D_{eff}t}}$   
 $= \left[ \frac{10^{-5}}{\sqrt{4\pi(5.21 \times 10^{-8} \text{ m}^2/\text{s})(36000 \text{ s})}} \right] e^{-\frac{[3.5 - (1 \times 10^{-4} \text{ m/s})(36000)]^2}{4(5.21 \times 10^{-8})(36000)}}$   
 $= \frac{3.1831}{0.1535} e^{-1.3329}$   
 $= 5.4677 \text{ g/m}^3.$

b) (i) time = 10 hr - 0.5 hr = 9.5 hr = 34200 s.

and injection:

$$C = \left( \frac{3.1831}{\sqrt{4\pi(5.21 \times 10^{-8})(34200)}} \right) e^{-\frac{[3.5 - (1 \times 10^{-4})(34200)]^2}{4(5.21 \times 10^{-8})(34200)}}$$

$$= \frac{3.1831}{0.1496} e^{-0.898}$$

$$= 8.6684 \text{ g/m}^3.$$

Salt concentration = 1st injection + 2nd injection.

$$= 5.4677 + 8.6684$$

$$= 14.1361 \text{ g/m}^3.$$

(ii) Advection flux

$$J = uc$$

$$= (1 \times 10^{-4} \text{ m/s})(14.1361 \text{ g/m}^3)$$

$$= 1.4136 \times 10^{-3} \text{ g/m}^2 \cdot \text{s}$$

(iii) Diffusive flux

$$J_m = -D_m \frac{dc}{dx}$$

Differentiate equation  $C = \frac{M}{\sqrt{4\pi D_{eff}t}} e^{-\frac{(x-ut)^2}{4D_{eff}t}}$  with respect to  $x$ ,  
to find  $\frac{dc}{dx}$ .

Diffusive flux due to the two injections will be in same direction. Substance diffuse from high conc. to lower conc. Both injection started from  $x=0$ , which are transported in same direction.



Q2

a) (i)

Equation

zeroth order

$$[A] = [A]_0 - kt$$

Second order

$$\frac{1}{[A]} = \frac{1}{[A]_0} + kt$$

unit of k

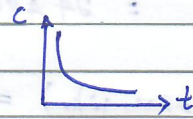
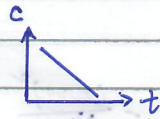
$$\frac{\text{mol}}{\text{L}\cdot\text{s}}$$

$$\frac{\text{L}}{\text{mol}\cdot\text{s}}$$

half time

$$\frac{[A]_0}{2k}$$

$$\frac{1}{k[A]_0}$$



zero order reaction has constant rate that is independent of reactant's concentration.  
rate = k

Second order reaction has a rate proportional to the product of 2 reactants or square of the concentration of a single reactant.  
rate = k[A]<sup>2</sup> or rate = k[A][B]

(ii) zero order

$$[A] = [A]_0 - kt$$

$$0.1C_0 = C_0 - kt$$

$$t = \frac{0.9C_0}{k}$$

Second order

$$\frac{1}{[A]} = \frac{1}{[A]_0} + kt$$

$$\frac{1}{0.1C_0} = \frac{1}{C_0} + kt$$

$$\frac{1}{C_0} (10-1) = kt$$

$$t = \frac{9}{kC_0}$$

b) (i)

$$DO = DO_{\text{sat}} - (DO_{\text{sat}} - DO_0) e^{-\frac{k_1}{u}x} - \frac{k_1 L_0}{k_2 - k_1} \left( e^{-\frac{k_1}{u}x} - e^{-\frac{k_2}{u}x} \right)$$

$$= 8.4 \frac{\text{g}}{\text{m}^3} - [8.4 - (6-1)] e^{-\frac{1.736 \times 10^{-6}}{0.1} (10 \times 1000)} - \frac{1.736 \times 10^{-6} (20909)}{2.014 \times 10^{-6} - 1.736 \times 10^{-6}} \left( e^{-0.1736} - e^{-0.2014} \right)$$

$$= 8.4 - 2.8582 - 130.568 (0.023)$$

$$= 2.5387 \text{ g/m}^3$$

$$k_1 = 0.15/\text{d}$$

$$= 1.736 \times 10^{-6} / \text{s}$$

$$k_2 = 0.174/\text{d}$$

$$= 2.014 \times 10^{-6} / \text{s}$$

$$L_0 = \frac{Q_1 L_1 + Q_2 L_2}{Q_1 + Q_2}$$

$$= \frac{0.5(20) + 0.05(30)}{0.55}$$

$$= 20909 \text{ g/m}^3$$

(ii) 
$$x_c = \frac{u}{k_2 - k_1} \ln \left\{ \frac{k_2}{k_1} \left[ 1 - \left( \frac{k_2}{k_1} - 1 \right) \left( \frac{D_0}{L_0} \right) \right] \right\}$$

$$= \frac{0.1}{2.014 \times 10^{-6} - 1.736 \times 10^{-6}} \ln \left\{ \frac{2.014 \times 10^{-6}}{1.736 \times 10^{-6}} \left[ 1 - \left( \frac{2.014 \times 10^{-6}}{1.736 \times 10^{-6}} - 1 \right) \left( \frac{5}{20909} \right) \right] \right\}$$

$$= 359712.23 \ln \{ 1.1601 (0.9617) \}$$

$$= 39377 \text{ m}$$

$$= 39.377 \text{ km}$$



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(iii)

$$DO = DO_{sat} - (DO_{sat} - DO_0) e^{-\frac{k_1}{u} x} - \frac{k_1 L_0}{k_2 - k_1} \left( e^{-\frac{k_1}{u} x} - e^{-\frac{k_2}{u} x} \right) \quad (39377)$$

$$3 = 8.4 - [8.4 - 5] e^{-\frac{1.736 \times 10^{-6} (39377)}{0.1}} - \frac{1.736 \times 10^{-6} (L_0)}{(2.014 - 1.736) \times 10^{-6}} \left[ e^{-\frac{1.736 \times 10^{-6}}{0.1} (39377)} - e^{-\frac{2.014 \times 10^{-6}}{0.1}} \right]$$

$$-5.4 = -3.4 (0.5048) - 6.2446 L_0 (0.5048 - 0.4525)$$

$$-3.68368 = -0.3266 L_0$$

$$L_0 = 11.279 \text{ g/m}^3$$

If able to bring wastewater stream BOD to zero.

$$L_0 = \frac{Q_1 L_1 + Q_2 L_2}{Q_1 + Q_2}$$

$$= \frac{0.5(20) + 0}{0.5 + 0.05}$$

$$= 18.18 \text{ g/m}^3 > 11.279 \text{ g/m}^3$$

∴ Cannot achieve. BOD conc. in open channel flow alone is higher than requirement.

(iv) Instead of calculating  $L_0$  using weighted average of both streams, can use channel BOD for  $L_0$ .



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Q3.

a) 
$$t = \frac{x}{u}$$

$$= \frac{540 \text{ m}}{0.2 \text{ m/s}}$$

$$= 2700 \text{ s}$$

$$= 45 \text{ mins.}$$

b) 
$$c = \frac{3 \text{ m}^3 \times 1000 \text{ kg/m}^3}{1200 \text{ m}^3}$$

$$= 2.5 \text{ kg/m}^3$$

c) (i) case 1

$$c = c_0 e^{-kt}$$

$$0.05 (2.5) = 2.5 e^{-\frac{0.25 \text{ m}^2/\text{min}}{1200 \text{ m}^2} t}$$

$$\ln 0.05 = -2.083 \times 10^{-4} t$$

$$t = 14381.816 \text{ min.}$$

$$= 9.987 \text{ days.}$$

(ii) case 2.

$$c = c_0 e^{-\frac{t}{\tau} - kt}$$

$$0.05 (2.5) = 2.5 e^{-\frac{t}{80} - 0.1 t}$$

$$\ln 0.05 = -\frac{t}{80} - 0.1 t$$

$$\ln 0.05 = -0.1125 t$$

$$t = 26.63 \text{ hrs.}$$

$$= 1.1 \text{ days.}$$

$$\bar{t} = \frac{V}{Q} = \frac{1200}{0.25} = 4800 \text{ mins.} = 80 \text{ hr}$$



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$$\begin{aligned} Q4a)(i) \quad K_{oc} &= 0.63 K_{ow} = 0.63 (10^{1.2}) \text{ L/kg} \\ &= 9.985 \text{ L/kg} \\ &= 9.985 \times 1 \times 10^{-6} \frac{\text{m}^3}{\text{g}} \\ &= 9.985 \times 10^{-6} \frac{\text{m}^3}{\text{g}} \end{aligned}$$

$$\begin{aligned} K_d &= f_{oc} K_{oc} \\ &= 0.05 (9.985 \times 10^{-6}) \\ &= 4.9925 \times 10^{-7} \frac{\text{m}^3}{\text{g}} \end{aligned}$$

$$\begin{aligned} (ii) \quad K_p &= K_d \cdot \rho_s \\ &= 4.9925 \times 10^{-7} (1500 \times 10^3 \text{ g/m}^3) \\ &= 0.7489 \end{aligned}$$

$$\begin{aligned} (iii) \quad \frac{1}{F_s} &= 1 + \frac{V_w}{V_s} \frac{1}{K_p} + \frac{V_g}{V_s} \frac{H_{cc}}{K_p} \\ &= 1 + \frac{20}{1} \frac{1}{0.7489} + \frac{5}{1} \frac{0.02}{0.7489} \\ &= 27.839 \\ F_s &= 0.0359 \end{aligned}$$

$$\begin{aligned} \frac{1}{F_w} &= \frac{V_s K_p}{V_w} + 1 + \frac{V_g H_{cc}}{V_w} \\ &= \frac{1(0.7489)}{20} + 1 + \frac{5(0.02)}{20} \\ &= 1.0424 \\ F_w &= 0.9593 \end{aligned}$$

$$\begin{aligned} \frac{1}{F_g} &= \frac{V_s}{V_g} \frac{K_p}{H_{cc}} + \frac{V_w}{V_g} \frac{1}{H_{cc}} + 1 \\ &= \frac{1}{5} \frac{0.7489}{0.02} + \frac{20}{5} \frac{1}{0.02} + 1 \\ &= 7.489 + 200 + 1 \\ F_g &= 4.796 \times 10^{-3} \end{aligned}$$

(iv)  $F_s$  will be twice as calculated above.  $F_w$  and  $F_g$  will reduce but it is insignificant. Because the differences between the ratios are still very big.

- b)
- ① Remove contaminant from sediment. from river bed.
  - ② Treating sludge containing organic contaminants.
  - ③ Treating solid wastes such as synthetic rubber, to remove harmful compound.



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Q5 a) Accumulation = In - out ± reaction

$$A \Delta x \frac{dc}{dt} = -Aq(C_x - C_{x+\Delta x}) - kAS(C - C_s)$$

$$\frac{dc}{dt} = -q \frac{dc}{dx} - k_a(C - C_s)$$

steady state,  $0 = -q \frac{dc}{dx} - k_a(C - C_s)$

$$q \frac{dc}{dx} = -k_a(C - C_s)$$

$$v \frac{dc}{dx} = -k \left( \frac{2}{R} \right) (C - C_s)$$

$$v \frac{dc}{dx} = \frac{2k}{R} (C_s - C)$$

b)  $v \frac{dc}{dx} = \frac{2k}{R} (C_s - C)$

$$\frac{1}{C_s - C} \frac{dc}{dx} = \frac{2k}{RV}$$

$$\int \frac{1}{C_s - C} dC = \int \frac{2k}{RV} dx$$

$$-\ln \frac{C_s - C}{C_s - C_0} = \frac{2k}{RV} x$$

$$\frac{C_s - C}{C_s - C_0} = e^{-\frac{2k}{RV} x}$$

$$C_s - C = (C_s - C_0) e^{-\frac{2k}{RV} x}$$

$$C = C_s - (C_s - C_0) e^{-\frac{2k}{RV} x}$$

c)  $\frac{k_d}{D} = 0.023 Re^{0.83} Sc^{0.33}$

$$\frac{k(2.5)}{1 \times 10^{-5}} = 0.023 \left[ \frac{(20)(2.5)(1)}{0.01} \right]^{0.83} \left[ \frac{0.01}{1(1 \times 10^{-5})} \right]^{0.33}$$

$$2.5 \times 10^5 k = 0.023 (1175.29)(9.7724)$$

$$k = 1.0567 \times 10^{-3}$$

d)  $C = C_s - (C_s - C_0) e^{-\frac{2k}{RV} x}$

$$= 10 \text{ mg/L} - (10 - 0) e^{-\frac{2(1.0567 \times 10^{-3})}{0.25 \text{ cm} \cdot 200 \text{ cm}^2/\text{s}} (5 \text{ cm})}$$

$$= 10 - 10 e^{-4.2268 \times 10^{-4}}$$

$$= 4.226 \times 10^{-3} \text{ mg/L}$$

Assume  $C_0$ , the concentration of pb in the bulk water at upstream is zero.

Assume Pb at the soldered surface is uniformly distributed and dissolves at same  $k$ .