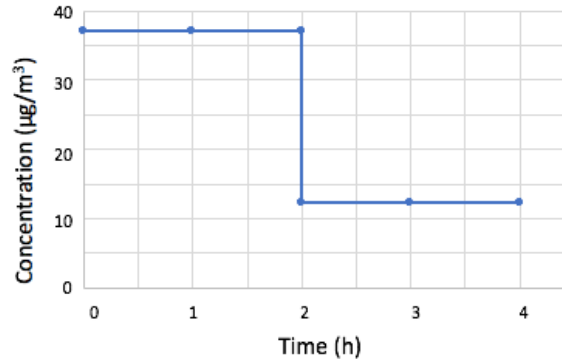
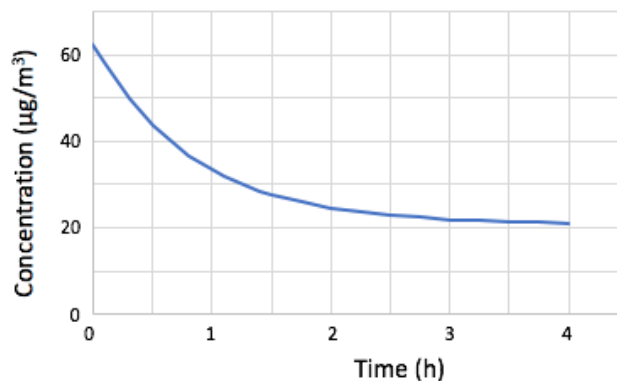


ANSWERS**1. Transient response of reactor models**

- (a) For a PFR with first-order decay, the outlet concentration, C , in steady state is $C_{in} \exp(-k\theta)$, where $\theta = V/Q$. In this problem, $\theta = 2$ h and $k\theta = 1.4$. So, the outlet concentration is $0.247 \times C_{in}$. When $C_{in} = 150 \mu\text{g}/\text{m}^3$, then $C = 37.0 \mu\text{g}/\text{m}^3$. Similarly, when $C_{in} = 50 \mu\text{g}/\text{m}^3$, then $C = 12.3 \mu\text{g}/\text{m}^3$. The step change from high to low happens at the inlet at time $t = 0$. At the outlet, there is a corresponding step change that occurs at time $\theta = 2$ h later. Sketch is presented below.



- (b) For a CMFR with first-order decay, the steady state concentration at the outlet, C , is $C_{in}/(1+k\theta)$. For the given conditions, $1/(1+k\theta) = 0.417$. The initial condition that applies for ($t \leq 0$) is the steady-state solution when $C_{in} = 150 \mu\text{g}/\text{m}^3$; that is $62.5 \mu\text{g}/\text{m}^3$. For $t > 0$, the ultimate steady-state concentration applies when $C_{in} = 50 \mu\text{g}/\text{m}^3$; that is $20.8 \mu\text{g}/\text{m}^3$. The time pattern of response follows an exponential decay from the initial to the final steady state value with a characteristic response time $\tau \sim \theta/(1+k\theta) = 0.83$ h. The governing equation is $d(CV)/dt = C_{in}Q - (Q + kV)C$ or $dC/dt = (Q/V)C_{in} - (Q/V + k)C$. This is of the form $dC/dt = S - LC$ and the characteristic response time is $1/L$. The concentration sketch versus time appears below.

**2. Sedimentation for particle control in drinking water treatment**

- (a) The overflow rate is Q/A_s . Here, $Q = 6 \text{ m} \times 3 \text{ m} \times 600 \text{ m}/\text{d}$ and $A_s = 6 \text{ m} \times 25 \text{ m}$. So, $Q/A_s = 1800/25 = \boxed{72 \text{ m}/\text{d}}$.

- (b) This is just a unit conversion, since the critical settling velocity equals the overflow rate for a sedimentation basin. Since $72 \text{ m} = 7200 \text{ cm}$ and $1 \text{ d} = 86400 \text{ s}$, the critical settling velocity is $7200/86400 = \boxed{0.083 \text{ cm/s}}$.
- (c) A particle will be captured with 75% efficiency if its settling velocity is equal 75% of the critical settling velocity, i.e. 0.0625 cm/s . Let's guess that Stokes law holds. From equation in handout, and ignoring the slip correction factor, we have $d_p^2 = (18 \mu v_s) / [g (\rho - \rho_f)]$. Look up parameter values and use cgs system consistently: $\mu = 0.01$, $v_s = 0.0625$, $g = 980$, $\rho - \rho_f = 1.5$. Substitute and solve for $d_p = 0.0028 \text{ cm} = \boxed{28 \mu\text{m}}$. (Check $Re_p = 0.0028 \times 0.0625 \times 1/0.01 = 0.02 < 0.3$, so Stokes law is okay!)

3. Disinfection performance in drinking water treatment

- (a) At 0.2 mg/L , we can read from the plot that 99% inactivation occurs with a contact time of three minutes. We can recognize that each log removal requires the same amount of time. The 99% inactivation corresponds to 2-log removal; the target is 3-log removal. Each 1-log removal requires $3 \text{ min}/2 = 1.5 \text{ minutes}$, so, to realize the design goal, we would need a hydraulic detention time of $3 \times 1.5 = \boxed{4.5 \text{ min}}$. (Can also solve the problem by computing the rate constant, k , for the given concentration: $N/N_0 = 0.01 = \exp(-kt)$, so at 0.2 mg/L , $k = \ln(100)/3 \text{ min} = 1.53 \text{ per minute}$. To achieve 99.9% inactivation, we need a contact time that satisfies $t = \ln(1000)/1.53 = 4.5 \text{ min}$.)
- (b) For the CMFR, must solve for k . The figure shows that with 3 mg/L concentration, the time needed in a batch reactor to achieve 99% inactivation would be 0.1 min . The corresponding rate constant is $\ln(100)/0.1 = 46 \text{ per minute}$. The steady-state performance of a CMFR yields $N/N_0 = 1/(1+k \theta)$. With given k and $\theta = 5 \text{ min}$, we have $k \theta = 230$ and so $N/N_0 = 1/231 = 0.004$. Consequently, these conditions yield $\boxed{99.6\%}$ inactivation. The corresponding n value is obtained from $-\log(0.004) = \boxed{2.4 \text{ log inactivation}}$.

3. Characteristic response time for a lake following contaminant spill

Perhaps the strongest way to approach this problem is to write a time-dependent material balance equation for the contaminant in the lake: $d(CV)/dt = QC - k_{gl} A C - k V C$. Let's treat V as constant so it can be taken outside the derivative and we can divide both sides by V . The result is of the form $dC/dt = S - LC$, where $S = 0$. The characteristic response time is $1/L$, where $L = Q/V + k_{gl} A/V + k$. That is:

$$\tau \sim \frac{1}{Q/V + k_{gl} A/V + k}$$