# Microbial kinetics in reactors II

# Today's lecture

- Nutrient & e⁻ acceptor consumption
- Hydrolysis
- Alternate rate expressions
- Example question

### **Nutrient consumption**

 For the consumption of nutrients for biomass production:

$$r_n = \gamma_n \cdot Y_{obs} \cdot r_{ut} = \gamma_n \cdot Y \cdot r_{ut} \frac{1 + (1 - f_d)b\theta_x}{1 + b\theta_x}$$

 $r_n$  = rate of nutrient consumption [M<sub>n</sub>L<sup>-3</sup>T<sup>-1</sup>]  $\gamma_n$  = the stoichiometric ratio of nutrient mass to VSS for biomass [M<sub>n</sub>M<sub>x</sub><sup>-1</sup>]

Using  $C_5H_7O_2N$  as cell formula:  $\gamma_N = 14 \text{ g N/}113 \text{ g VSS} = 0.124 \text{ g N/g VSS}$  $\gamma_P = 0.2 \times 0.124 = 0.025 \text{ g P/g VSS (generally P = 0.2N)}$ 

### **Nutrient consumption in a CSTR**

Steady-state mass balance:

$$0 = QC_n^0 - QC_n + r_n V$$

$$C_n = C_n^0 + r_n \theta$$

if  $C_n < 0$ , nutrient-limiting

#### e-acceptor consumption

(e⁻ acceptor used in a reactor)

= [(total O.D. in the influent) - (total O.D. in the effluent)] x (conversion factor)

In terms of the use rate for a reactor  $(\Delta S_a/\Delta t)$ :

$$\frac{\Delta S_a}{\Delta t} = \gamma_a [Q(S^0 + 1.42X_v^0) - Q(S + SMP + 1.42X_v)]$$
$$= \gamma_a Q[S^0 - S - SMP + 1.42(X_v^0 - X_v)]$$

 $\gamma_a$  = the stoichiometric ratio of acceptor mass to oxygen demand

for oxygen: 1 g O<sub>2</sub>/g COD

for nitrate: 0.35 g NO<sub>3</sub>-N/g COD

#### e-acceptor consumption

To estimate the required mass rate of acceptor supply (ex: aeration  $[O_2]$  requirement), the calculated  $e^-$  acceptor use rate,  $\Delta S_a/\Delta t$  can be written as:

$$\frac{\Delta S_a}{\Delta t} = \gamma_a \big[ Q \big( S^0 + 1.42 X_v^{\ 0} \big) - Q \big( S + SMP + 1.42 X_v \big) \big]$$

$$= Q \big( S_a^{\ 0} - S_a \big) + R_a$$
(mass flow rate in) – Requirement of e-acceptor addition

# Hydrolysis of particulates & polymers

- Particulates and polymeric substances account for a significant portion of BOD in wastewater
- >50% of BOD in typical sewage is particulates (SS)
- Particulates and large-MW compounds cannot penetrate the cell membrane
  - > needs to be hydrolyzed to smaller molecules
- Catalyzed by extracellular enzymes
- The mechanism and kinetics of hydrolysis it not fully understood

## Hydrolysis of particulates & polymers

One simple way of describing hydrolysis is to assume first-order kinetics for particulates (or polymers):

$$r_{hyd} = -k_{hyd}S_p$$

 $r_{hvd}$  = rate of accumulation of particulates (= $dS_p/dt$ ) [M<sub>s</sub>L<sup>-3</sup>T<sup>-1</sup>]

 $k_{hvd}$  = first-order hydrolysis rate coefficient [T<sup>-1</sup>]

 $S_p$  = concentration of particulates [M<sub>s</sub>L<sup>-3</sup>]

In a steady-state CSTR,

$$0 = Q(S_p^0 - S_p) - k_{hyd}S_pV$$

$$S_p = \frac{S_p^0}{1 + k_{hyd}\theta}$$

# Hydrolysis of particulates & polymers

 Effect of hydrolysis on dissolved substrates mass balance in a steady-state CSTR

$$0 = Q(S^0 - S) - \frac{\widehat{q}S}{K + S}X_aV + k_{hyd}S_pV$$

$$0 = (S^0 - S) - \frac{\widehat{q}S}{K + S} X_a \theta + k_{hyd} S_p \theta$$

- $\rightarrow$  Increase in  $S^0$  by  $k_{hyd}S_p\theta$
- → Increased biomass, but no change in dissolved substrates in the reactor

### Alternate rate expressions

#### Contois equation

$$r_{ut} = -\frac{\hat{q}S}{BX_a + S}X_a$$

$$B = \text{constant } [M_s/M_x]$$

When 
$$X_a \to \infty$$
,  $r_{ut} = -\frac{\hat{q}}{B}S$ 

(at high biomass concentrations substrate utilization depends on S, not  $X_a$ )

### Alternate rate expressions

Moser equation

$$r_{ut} = -\frac{\hat{q}S}{K + S^{-\gamma}}X_a$$
  $\gamma = \text{constant [unitless]}$ 

Tessier equation

$$r_{ut} = -\hat{q}(1 - e^{S/K})X_a$$

Just **REMEMBER** that Monod Eq. is **NOT** the only option!!!

### **Dual Monod equation**

$$r_{ut} = -\hat{q} \frac{S}{K+S} \frac{A}{K_A + A} X_a$$

 $A = e^{-}$  acceptor concentration  $[M_A/L^3]$  $K_A = \text{half-saturation coeff. for } e^{-}$  acceptor  $[M_A/L^3]$ 

- e⁻ acceptor can also be limiting!
- Can be reduced to single Monod Eq. if  $A >> K_A$
- Terms for other limiting substances can be added as well

#### **Analyzing CSTR (Chemostat) Performance**

**Q:** A chemostat having  $V=2,000 \text{ m}^3$  receives a flow rate of  $Q=1,000 \text{ m}^3/\text{d}$  of wastewater containing  $S^0=500 \text{ mg BOD}_L/\text{L}$ . Also included in the wastewater is the inert biomass  $X_i^0=50 \text{ mg VSS/L}$ . The following parameters are found for aerobic biodegradation:

$$\hat{q} = 20 \ g \ BOD_L / g \ VSS_a - d$$
 $Y = 0.42 \ g \ VSS_a / g \ BOD_L$ 
 $K = 20 \ mg \ BOD_L / L$ 
 $b = 0.15 / d$ 
 $f_d = 0.8$ 
 $k_1 = 0.12 \ g \ COD_p / g \ BOD_L$ 

$$k_{2} = 0.09 \ g \ COD_{p} / g \ VSS_{a} - d$$

$$\hat{q}_{UAP} = 1.8 \ g \ COD_{p} / g \ VSS_{a} - d$$

$$K_{UAP} = 100 \ mg \ COD_{p} / L$$

$$\hat{q}_{BAP} = 0.1 \ g \ COD_{p} / g \ VSS_{a} - d$$

$$K_{BAP} = 85 \ mg \ COD_{p} / L$$

#### **Analyzing CSTR (Chemostat) Performance**

#### **Questions:**

- 1. Calculate  $S_{min}$ ,  $\Theta_x^{min}$ , and  $\Theta_x$  of the chemostat
- 2. Calculate effluent VSS, COD, and BOD,
- 3. Calculate the effluent N and P concentrations when influent concentrations are 50 mg  $NH_4^+$ -N/L and 10 mg  $PO_4^{3-}$ -P/L, respectively.
- 4. Calculate the amount of  $O_2$  that should be supplied to the reactor when influent and effluent DO are 6 and 2 mg/L, respectively.
- 5. Assuming that the influent also contains biodegradable particulate organic matter with a concentration of 100 mg COD/L and the hydrolysis rate coefficient is  $k_{hyd} = 0.2/d$ , recalculate the effluent VSS, COD, and BOD<sub>1</sub>.