

- 11.9.** A fermentation broth with a protein concentration of $C_0 = 100$ mg/l and flow rate of $Q = 4$ m³/h is passed through two downflow adsorption columns connected in series. The adsorption isotherm is: $q = 4C^{1/4}$. Assuming the system is in equilibrium, calculate the following:
- Minimum amount of activated carbon required for two days of operation if removal efficiency for the column is $E = 50\%$.
 - Protein concentration in the effluent of the second column is desired to be $C = 0.5$ mg/L. Determine the minimum amount of activated carbon required for two days of operation.
- 11.10.** In a cross-flow ultrafiltration system used for filtration of proteins from a fermentation broth, gel resistance increases with protein concentration according to the following equation:

$$R_g = 0.5 + 0.01(C), \text{ where } C \text{ is in mg/l.}$$

Pressure at the entrance of the system is $P_i = 6$ atm and at the exit is $P_0 = 2$ atm. The shell side of the filter is open to the atmosphere, resulting in $P_f = 1$ atm. The membrane resistance is $R_M = 0.5$ atm/(mg/m² · h), and protein concentration in the broth is $C = 100$ mg/l. Determine:

- The pressure drop across the membrane.
 - Filtration flux.
 - Rejection coefficient of the membrane for effluent protein concentration of $C_f = 5$ mg/l.
- 11.11.** A solute protein is to be separated from a liquid phase in a chromatographic column. The adsorption isotherm is given by the following equation:

$$C_s = kC_L^2$$

where C_s is the solute concentration in solid phase (mg solute/mg adsorbent) and C_L is the liquid phase concentration of solute (mg solute/ml liquid). Use the following information:

$$k = 0.4, \quad \epsilon = 0.3, \quad A = 25 \text{ cm}^2,$$

$$\mu = 10 \text{ g ads/100 ml column} = 100 \text{ mg/ml}$$

- For $V = 400$ ml and $X = 25$ cm determine the equilibrium solute concentrations in liquid and solid phases.
 - Determine the ratio of travel distances of solute to solvent, R_f .
- 11.12.** Consider the scale-up of a chromatography column for purification of a protein. A column of 40 cm length is used with a superficial velocity of 40 cm/h. The peak concentration of the target protein exits at a time of 100 min. The standard deviation of the peak is 14 min.
- How long must you wait to collect 90% of the protein?
 - If the same column is used, but velocity is increased to 60 cm/h and external or Taylor dispersion controls, what will be the value of σ ?
 - If the column is lengthened to 60 cm while the velocity is at 60 cm/h, how will σ change? Will the peak be sharper or broader than at 40 cm/h with a 40-cm column?