# CHEMICAL ENGINEERING

# ONE MARKS QUESTIONS (1-20)

- The ordinary differential equation dy/dt = f(Y) is solved using the approximation Y(t + Δt) = Y(t) + f[Y(t)] Δt. The numerical en-or introduced by the approximation at each step is
  - a. proportional to Δt
  - b. proportional to  $(\Delta t)^2$
  - c. independent of  $\Delta t$
  - d. proportional to (1/Δt)
- The trapezoidal rule of integration when applied to  $\int f(x)dx$  will give the exact

value of the integral

- a. if f(x) is a linear function of x
- b. if f(x) is a quadratic function of x
- c. for any f(x)
- d. for no f(x)
- 3. The value of a for which the following three vectors are coplanar is

$$a = i + 2j + k$$
  

$$b = 3j + k$$
  

$$c = 2i + \alpha j$$

- a. 4
- b. zero
- c. -2
- d. -10
- 4. The derivative of |x| with respect to x when x≠0 is
  - a |x|/x
  - b. -I
  - c. I
  - d Undefined
- At a given temperature and pressure, a liquid mixture of benzene and toluene is in equilibrium with its vapor. The available degree(s) of freedom is (are)
  - a. Zero
  - **b**.
  - c. 2
  - d. 3
- A heat engine operates at 75% of the maximum possible efficiency. The ratio of the heat source temperature (in K) to the heat sink temperature (in K) is 5/3. The fraction of the heat supplied that is converted to work is

- a. 0.2
- b. 0.3
- c. 0,4
- d 0.6
- For the isentropic expansion of an ideal gas from the initial conditions  $P_1$ ,  $T_1$  to the final conditions  $P_2$ ,  $T_2$ , which ONE of the following relations is valid? ( $y = C_0/C_0$ )
  - $\mathbf{a.} \quad \left(\frac{P_1}{P_2}\right) = \left(\frac{T_2}{T_1}\right)^2$
  - $\mathbf{b} = \left(\frac{P_1}{P_2}\right) = \left(\frac{T_1}{T_2}\right)^{\frac{T}{T-1}}$
  - $e_{1} \left(\frac{P_{1}}{P_{2}}\right) = \left(\frac{T_{1}}{T_{2}}\right)$
  - $\mathbf{d}, \quad \left(\frac{P_1}{P_2}\right) = \left(\frac{T_1}{T_2}\right)^{\frac{p-1}{p}}$
- Match the following for a centrifugal pump with impeller speed n

List !

- A. Capacity
- B, Head

List II

- 1. proportional to n
- 2. proportional to n<sup>2</sup>
- 3. proportional to n<sup>3</sup>
  - A B
- a 2 1
- b. 1
- c. 2 3
- d 1 2
- 9 The magnitude of the force (in N) required to hold a body of volume 0.05 m<sup>3</sup> and mass 40kg in water (density 1000 kg/m<sup>3</sup>) at a depth of 0.1 m is (g = 9.81 m/s<sup>2</sup>)
  - a. Zero
  - 6. 98.1
  - c. 490,5
  - d. 882.9
- 10. A stagnant liquid film of 0.4 mm thickness is held between two parallel plates. The top plate is maintained at 40°C and the bottom plate is maintained at 30°C. If the thermal conductivity of the liquid is 0.14 W/(m K), then the steady state heat flux (in W/m²) assuming one dimensional heat transfer is

- a. 3.5
- b. 350
- c. 3500
- d: 7000
- 11. Let d<sub>n</sub> be the hydrodynamic entrance length for mercury in laminar flow in a pipe under isothermal conditions. Let d<sub>t</sub>, be its thermal entrance length under fully developed hydrodynamic conditions. Which ONE of the following is TRUE?
  - a.  $d_h > d_t$
  - b. dh < di
  - $\mathbf{e}_{t} \cdot \mathbf{d}_{h} = \mathbf{d}_{t}$
  - d. dh < di only if the pipe is vertical
- 12. The Boussinesq approximation for the fluid density in the gravitational force term is given by ONE of the following (pref is the fluid density at the reference temperature T<sub>ref</sub>, and β is the thermal coefficient of volume expansion at T<sub>ref</sub>)
  - a.  $\rho = \rho_{rel} + T_{rel} \beta (\rho \rho_{rel})$
  - b.  $\rho = \rho_{ref} T_{ref} \beta (\rho \rho_{ref})$
  - c.  $\rho = \rho_{ref} T_{ref} \beta \left( T T_{ref} \right)$
  - d.  $\rho = \rho_{iij} T_{iij} \left( \rho \rho_{rej} \right) = \rho_{rej} \left( T T_{rej} \right) / T_{rej}$
- 13. The reaction 2A + B → 2C occurs on a catalyst surface. The reactants A and B diffuse to the catalyst surface and get converted completely to the product C, which diffuses back. The steady state molar fluxes of A, B and C are related by
  - $\mathbf{a}, \ \mathbf{N}_A = 2\mathbf{N}_B = \mathbf{N}_C$
  - b.  $N_A = -(1/2) N_B = -N_C$
  - c.  $NA = 2N_B N_C$
  - d.  $N_A = (1/2) N_B = N_C$
- 14. An ideal single stage extraction process is used to treat 100 molls of an organic feed solution. The solution concentration in this solution is to be reduced from 0.5 mol% to 0.1 mol%. A pure solvent S is used. To reduce the solvent requirement by half for the same separation.
  - a. add one more ideal co-current stage
  - b. use another pure solvent S\* whose partition coefficient is twice that of S
  - e. use solvent S containing 0.02 mole fraction of the solute
  - d. double the residence time of the solvent S in the contactor
- 15. An irreversible gas phase reaction A → 5B is conducted in an isothermal batch reactor at constant pressure in the presence of an inert. The feed contains no B. If the

volume of the gas at complete conversion must not exceed three times the initial volume, the minimum mole percent of the inert in the feed must be

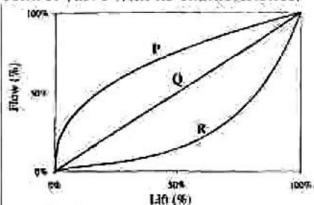
- a, 0
- b. 20
- c. 33
- d. 50
- 16. A first order reversible reaction

occurs in a batch reactor. The exponential decay of the concentration of A has the time constant

- $\mathbf{s}_{i} = \frac{1}{k_{i}}$
- b.  $\frac{1}{k_x}$
- $c = \frac{1}{k_1 + k_2}$
- $d. \frac{1}{k_1 + k_2}$
- 17. If the absolute error in the measurement of A is ΔA and the absolute error in the measurement of B is ΔB, then the absolute error in the estimate of A – B is
  - a. AA+AB
  - b.  $\Delta A \Delta B$
  - e.  $\frac{\Delta A}{A} + \frac{\Delta B}{B}$
  - d.  $\frac{\Delta A}{A} = \frac{\Delta B}{B}$
- 18. The oxo reaction is used for converting
  - a. alcohol to aldehyde
  - b. paraffin to olefin
  - c. olefin to aldehyde
  - d. aldehyde to alcohol
- 19. In a fluid catalytic cracking unit, the nature of the reactions occurring in the reactor and the regenerator is
  - a. Reactor-Exothermic, Exothermic
  - b. Reactor-Exothermic. Regenerator-Endothermic
  - e. Reactor-Endothermie, Regenerator-Exothermie
  - d. Reactor-Endothermic, Regenerator-Endothermic
- 20. The control valve characteristics for three types of control valves (P. Q and R) are

Regenerator-

given in the figure below: Match the control valve with its characteristics.



- a P Quick opening, Q Linear, R -Equal percentage
- b. P Linear, Q Square root, R Equal percentage
- c. P Equal percentage, Q Linear, R -Quick opening
- d. P Square root, Q Quick opening, R Linear

## TWO MARKS QUESTIONS (21-75)

21, If the following represents the equation of a line

$$\begin{vmatrix} x & 2 & 4 \\ y & 8 & 0 \\ 1 & 1 & 1 \end{vmatrix} = 0$$

then the line passes through the point

- a. (0.0)
- b. (3, 4)
- c (4, 3)
- d (4, 4)
- If  $A = \begin{bmatrix} 2 & 1 \\ 2 & 3 \end{bmatrix}$ , then the eigenvalues of  $A^{T}$ 22,

are

- a. 27 and 8
- b. 64 and 1
- 12 and 3
- d 4 and 1
- With  $y = e^{ax}$ , if the sum 23,

$$S = \frac{dy}{dx} + \frac{d^2y}{dx^2} + \dots + \frac{d^ny}{dx^n}$$

approaches 2y as  $n \rightarrow \infty$ , then the value of

- 11 15
- 1/3
- b. 1/2
- 2/3
- d. 2
- Determine the following integral 24.

$$I = \int r dS$$

where r is the position vector field (r = ix +jy + kz) and S is the surface of a sphere of radius R

- a,  $4\pi R^2$
- c, πR<sup>2</sup>
- d. 4πR<sup>3</sup>
- The liquid surface in a cylindrical bucket 25, of radius R rotating about its axis acquires a parabolic profile given by the equation y = a + br2, where y is the height of the liquid surface from the bottom of the bucket at a radial distance r from the bucket axis. If the liquid has density p then the mass of the liquid in the bucket is

a. 
$$\pi \rho R^2 \left( \frac{a + bR^2}{2} \right)$$

- c.  $\pi \rho R^2 a$
- d.  $\pi \rho R^2 \left(a + bR^2\right)$
- The solution to the following equation is 26.

$$x^{2} \frac{d^{3}y}{dx^{3}} + 2x \frac{d^{2}y}{dx^{2}} - 2 \frac{dy}{dx} = 0$$

is given by

- a.  $y = C_1x + C_2x^{-2} + C_3$ b.  $y = C_1x^2 + C_2x^{-2} + C_3$ c.  $y = C_1x^2 + C_2x^{-1} + C_3$ d.  $y = C_1x + C_2x^{-1} + C_3$

- The value of the contour integral 27. where C is the circle |z| = 2 is

  - $b. \quad \frac{1}{2} \left( \frac{1}{e} \frac{1}{e^3} \right)$
  - c. zero
- The Newton-Raphson method is used to 28. solve the equation,  $(x-1)^3 + x - 3 = 0$ . The method will fail in the very first iteration if the initial guess is
  - Zero
  - b. 0.5
  - C. d. 3
- 29. A pair of fair dice is rolled three times. What is the probability that 10(sum of the

numbers on the two faces) will show up exactly once?

- a. 121 1728
- b.  $\frac{363}{1728}$
- e. 121 576
- d.  $\frac{363}{576}$
- 30. A company purchased components from three firms P. Q, and R as shown in the table below:

Firm	Total number of components purchased	Number of components likely to be defective
P	1000	5
Q	2500	5
R	500	2

The components are stored together. One of the components is selected at random and found to be defective. What is the probability that it was supplied by Firm R?

- a.  $\frac{1}{250}$
- b.  $\frac{1}{12}$
- 6.  $\frac{1}{8}$
- d.  $\frac{1}{6}$
- 31. Match the following:

List I

- A. Heat
- B. Internal energy
- C. Work
- D. Entropy

List II

- 1. State Function
- 2. Path Function

	A	B 1 1 2	C	D
a.	A 2 2 2 2 2	1	: 1	D 1 2 1
b.	2	1.	2	2
b. c. d.	2	2	2 1 2	1
d.	2	1	2	1

- 32. For a reversible exothermic gas phase reaction, A + B \( \Bar{\pi} \) C, the equilibrium conversion will increase with
  - increase in pressure and increase in temperature

 b. decrease in pressure and increase in temperature

 increase in pressure and decrease in temperature

d, decrease in pressure and decrease in temperature

33. For a binary mixture of A and B at 400 K and 1 atm, which ONE of the following equilibrium states deviates significantly from ideality?

Given:

$$\ln\left(P_A^{\text{ser}}\right) = 6.2 - \frac{2758}{T}$$

where

 $P_A^{int}$  = vapor pressure of A, atm; T = temperature, K

PA = partial pressure of A, atm

 $x_A$  = mole fraction of A in liquid;  $y_A$  = mole fraction of A in vapor

- a.  $x_A = 0.5$ ;  $y_A = 0.25$
- b.  $x_A = 0.5$ ,  $p_A = 0.25$
- c.  $x_A = 0.5$ ;  $p_A = 0.5$
- d.  $x_A = 0.6$ ;  $y_A = 0.3$
- 34. Pure A at 200°C is fed to a steady slate adiabatic continuous reactor at the rate of 100 kg/hr where it undergoes an exothermic reaction to give its isomer B. The product stream is at temperature 500°C. The heat of reaction is 21 kJ/mol of A and the specific heat of the reaction mixture is constant at 35 J/(mol °C). The conversion in the reactor is
  - a. 25%
  - b. 50%
  - c. 75%
  - d. 100%
- 35. The molar density of water vapor at the normal boiling point of water is 33 mol/m<sup>3</sup>. The compressibility factor under these conditions is close to which ONE of the following? R = 8.314 J/(mol K)
  - a, 0.75
  - 6. 1
  - e. 1.25
  - d. 1,5
- 36. A liquid is pumped at the flow rate Q through a pipe of length L. The pressure drop of the fluid across the pipe is Δ P. Now a leak develops at the mid-point of the length of the pipe and the fluid leaks at the rate of Q/2. Assuming that the friction factor in the pipe remains unchanged, the new pressure drop across the pipe for the same inlet flow rate (Q) will be

- n. (1/2) ΔP
- b. (5/8) AP
- (3/4) AP
- $\Delta P$
- 37. In a laminar flow through a pipe of radius R, the fraction of the total fluid flowing through a circular cross-section of radius R/2 centered at the pipe axis is
  - n. 3/8
  - 7/16
  - 1/2
  - d. 3/4
- 38. A fluid obeying the constitutive equation

$$\tau = \tau_0 + K \left( \frac{dv_y}{dy} \right)^{\frac{1}{2}}, \tau > \tau_0$$

is held between two parallel plates a distance d apart. If the stress applied to the top plate is 370, then the velocity with which the top plate moves relative to the bottom plate would be

- $4 \cdot 2 \left(\frac{\tau_0}{k}\right)^2 d$

- d.  $9\left(\frac{\tau_0}{K}\right)^2 d$
- 39. A bed fluidized by water is used for cleaning sand contaminated with salt. The particles of sand and salt have the same shape and size but different densities (Psaud = 2500 kg/m<sup>3</sup> and  $\rho_{soll}$  = 2000 kg/m<sup>3</sup>). If the initial volume fraction of the salt in the mixture is 0.3 and if the initial value of the minimum fluidization velocity (Umf) is 0.9 m/s, find the final value of the Umf (in m/s) when the sand is washed free of the salt Assume that the bed characteristics (bed porosity and solid surface area per unit volume) do not change during the operation and that the pressure drop per unit length is directly proportional to the fluid velocity
  - a. 0.70
  - b. 0.90
  - e. 1.00
  - d. 1.46
- Two spherical particles have the same 40. outer diameter but are made of different

materials The first one (with material density p<sub>1</sub>) is solid, whereas the second (with material density  $p_2$ ) is a hollow sphere with the inner shell diameter equal to half the outer diameter. If both the spheres have the same terminal velocity in any fluid, then the ratio of their material densities,  $\rho_2/\rho_1$ , is

- a. 1
- b. 8/7
- c. 2
- 4. 8
- 41. filtration is conducted at constant pressure to recover solids from dilute slurry. To reduce the time of filtration, the solids concentration in the feed slurry is increased by evaporating half the solvent. If the resistance of the filter medium is negligible, the filtration time will be reduced by a factor of

  - 2 b.
  - 4 Ċ,
  - d. 8
- 42. One dimensional steady state heat transfer occurs from a flat vertical wall of length 0.1m into the adjacent fluid. The heat flux into this fluid is 21 W/m2. The wall thermal conductivity is 1.73 W/(m K). If the heat transfer coefficient is 30 W/(m<sup>2</sup> K) and the Nusselt number based on the wall length is 20, then the magnitude of the temperature gradient at the wall on the fluid side (in Kim) is
  - a. 0.7
  - b. 12.14
  - 120
  - d. 140
- 43. Experiments conducted with a sparingly dissolving cylinder wall in a flowing liquid yielded the following correlation for the Sherwood number

Assuming the applicability of the Chilton-Colburn analog, the corresponding correlation for heat transfer is

- a.  $Sh = 0.023 (Gr)^{0.83} (Pr)^{1/3}$ b.  $Nu = 0.023 (Re)^{0.83} (Pr)^{1/3}$
- e.  $j_H = 0.023 (Re)^{6.83} (Pr)^{2/3}$ d.  $Nu = 0.069 (We)^{0.5} (Pr)^{4/3}$
- A fluid flows through a cylindrical pipe 44. under fully developed, steady state laminar flow conditions. The tube wall is at constant temperature. maintained

Assuming constant physical properties and negligible viscous heat dissipation, the governing equation for the temperature profile is (z. axial direction; r. radial direction)

**a.** 
$$u_{\text{tot}} \left[ 1 - \left( \frac{r}{R} \right)^2 \right] \left( \frac{\partial T}{\partial z} \right) = \frac{k}{\rho C_s} \left[ \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial T}{\partial r} \right) + \frac{\partial^2 T}{\partial z^2} \right]$$
**b.**  $u_{\text{tot}} \left[ 1 - \left( \frac{r}{R} \right)^2 \right] \left( \frac{\partial T}{\partial r} \right) = \frac{k}{\rho C_s} \left[ \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial T}{\partial z} \right) + \frac{\partial^2 T}{\partial r^2} \right]$ 

$$0. \quad 2u_{\text{max}} \left[ 1 - \left( \frac{r}{R} \right)^{1} \right] \left( \frac{\partial^{2} T}{\partial x^{1}} \right) = \frac{k}{\rho C_{p}} \left[ \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial T}{\partial r} \right) \tau \frac{\partial^{2} T}{\partial r^{2}} \right]$$

$$\mathbf{d} = \mathbf{u}_{\text{exp}} \left[ 1 - \left( \frac{r}{R} \right)^{\text{V}} \right] \left( \frac{\partial T}{\partial z} \right) = \frac{\mathbf{A}}{\rho C_f} \left[ \frac{1}{z} \frac{\partial}{\partial z} \left( z \frac{\partial T}{\partial z} \right) + \frac{\partial^2 T}{\partial z^2} \right]$$

45. An insulated cylindrical pipe of 0.2 m diameter has a surface temperature of 45°C. It is exposed to black body surroundings at 25°C. The emissivity and absorptivity of the insulation surface are 0.96 and 0.93, respectively. The convective heat transfer coefficient outside the insulation surface is 3.25 W/(m² K). The Stefan-Boltzmann constant is 5.67×10° W/(m² K²). The surrounding fluid may be assumed to be transparent. Find the percentage contribution from radiation to the total heat transfer rate to the surroundings

a. 30.9

b. 50.0

c. 57.6

d. 68.4

46. In a multistage countercurrent isothermal stripping column, feed containing 0.05 mol of solute/mol of solute free oil is treated with steam. The absorption factor A = 0.65. The equilibrium relation is given by Y\* = 2X, where Y\* and X refer to the equilibrium mole ratio in the steam and oil phases respectively. The Kremser equation is given as follows ('0' refers to liquid inlet at the top, 'N<sub>p</sub>' refers to the last stage at the bottom).

$$N_{p} = \frac{\log \left[ \left( \frac{X_{0} - \frac{Y_{H_{2p}}}{m}}{X_{H_{2}} - \frac{Y_{H_{2p}}}{m}} \right) (1 - A) + A \right]}{\log \left[ \frac{1}{A} \right]}$$

If the steam is initially free of solute and its exit mole ratio (mol solute/mol steam) is 0.0624, then the number of equilibrium stages required is

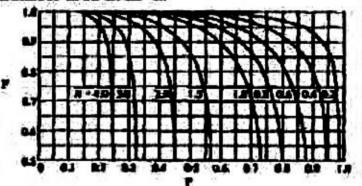
a, 4.2

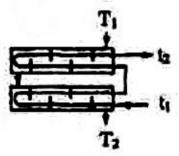
b. 5,2

o. 7.2

d. 8.2

47. A process fluid has to be cooled from 22°C to 2°C using brine in a 2-4 shell-and tube heat exchanger shown below. The brine enters at -3°C and leaves at 7°C. The overall heat transfer coefficient is 500 W/(m² K). The design heat load is 30 kW. The brine flows on the tube side and the process fluid on the shell side. The heat transfer area in m² is





$$P = \frac{t_2 - t_1}{T_1 - t_1}$$

$$R = \frac{T_1 - T_2}{t_2 - t_1}$$

a 1.1

b. 5.77

c. 6.59

d. 7.53

48. 100 moles of a binary mixture F containing 60 mol<sup>4</sup>6 A (more volatile) and 40 mol<sup>5</sup>6 B is treated in a batch distillation still. After 1 hour, 70 moles of the distillate D is collected leaving behind the residue W. Relative volatility α is 2. The governing equation is

$$\log \frac{Fx_{F}}{Wx_{w}} = \alpha \log \frac{F(1-x_{F})}{W(1-x_{w})}$$

The average mole fraction of A in the distillate is

a. 0.43

b. 0.61

c. 0.69

d. 0.73

A drug tablet of mass M<sub>0</sub> administered orally at time t = 0 reaches the intestine at

time  $t = \tau$  without losing any mass. From the intestine, the drug is absorbed into blood. The rate of absorption is found to be proportional to the mass of the drug in the intestine with the proportionality constant k. Assuming no drug is lost from the blood, the total mass of the drug in the blood,  $M_b$ , at time  $t \ge \tau$  is given by

a. 
$$M_0 = M_0 \left[ 1 - \exp \left\{ -k(t-\tau) \right\} \right]$$

b. 
$$M_b = M_0 [1 - exp\{-kt\}]$$

c. 
$$M_b = M_b \exp\{-k(t-\tau)\}$$

d. 
$$M_6 = M_0 \left[ 1 - \exp\{-k(t+\tau)\} \right]$$

50. The rate rat which an antiviral drug acts increases with its concentration in the blood, C, according to the equation

$$r = \frac{kC}{C_{50} + C}$$

where  $C_{50}$  is the concentration at which the rate is 50% of the maximum rate k. Often, the concentration  $C_{50}$ , when the rate is 90% of the maximum, is measured instead of  $C_{50}$ . The rate equation then becomes

$$\mathbf{a}, \quad r = \frac{1.8kC}{\left(C_{10} + C\right)}$$

b. 
$$r = \frac{kC}{\left(\frac{C_{00}}{9} + C\right)}$$

$$\mathbf{e}_{i} \quad r = \frac{kC}{C_{\alpha i}}$$

$$\mathbf{d}_{i} \quad r = \frac{0.9kC}{C_{00}}$$

 Consider the following reactions between gas A and two solid spherical particles, B and C of the same size

$$A + B \xrightarrow{t_i} gaseous product$$

The ash does not leave the particle C. Let t<sub>1</sub> and t<sub>2</sub> be the times required for A to completely consume particles B and C, respectively. If k<sub>1</sub> and k<sub>2</sub> are equal at all temperatures and the gas phase mass transfer resistance is negligible, then

 $a_1 = t_2$  at all temperatures

b.  $t_1 = t_2$  at high temperatures

c. t<sub>1</sub>>t<sub>2</sub> at high temperatures

d, t1 < t2 at high temperatures

52. A reaction A→B is to be conducted in two CSTRs in series. The steady state conversion desired is X<sub>L</sub>. The reaction rate

as a function of conversion is given by r = -1/(1 + X). If the feed contains no B, then the conversion in the first reactor that minimizes the total volume of the two reactors is

a.  $1-X_{\Gamma}$ 

b. 0.2Xr

c. 0.5Xr

d. 0.5(1-X<sub>f</sub>)

53. Consider the following elementary reaction network

$$\begin{array}{ccc} A & \stackrel{1}{\longrightarrow} & B \\ \stackrel{2}{\longrightarrow} & \downarrow^{3} \end{array}$$

The activation energies for the individual reactions are  $E_1 = 100 \text{ kJ/mol}$ ,  $E_2 = 150 \text{ kJ/mol}$ ,  $E_3 = 100 \text{ kJ/mol}$ , and  $E_4 = 200 \text{ kJ/mol}$ . If the feed is pure A and the desired product is C, then the desired temperature profile in a plug flow reactor in the direction of flow should be

a. constant at low temperature

b. constant at high temperature

c, increasing

d. decreasing

The exit age distribution in a stirred reactor is given by

$$E(t) = \frac{1}{\tau}e^{-\frac{1t}{\tau}}$$

Fluid elements  $e_1$  and  $e_2$  enter the reactor at times t = 0 and t = 0 > 0, respectively. The probability that  $e_2$  exists the reactor before  $e_1$  is

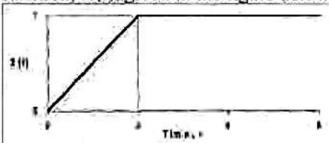
a.  $\frac{1}{2}$ 

 $b. \frac{1}{2}e^{-u/\epsilon}$ 

C. 2 00

d, zero

55. The Laplace transform of the input function, X(t), given in the figure below



is given by

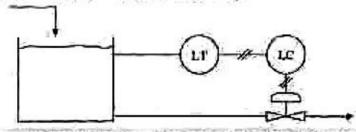
$$a, \frac{1}{2s^2} \left(1 - e^{-cx}\right)$$

b. 
$$\frac{1}{2s^2} (1 + e^{s^2s})$$

$$e_{x} = \frac{1}{s^2} \left( 1 + e^{-2s} \right)$$

d. 
$$\frac{1}{s^2}(1-e^{-2s})$$

56. A liquid level control system is configured as shown in the figure below. If (he level transmitter ILT) is direct acting and the pneumatic control valve is air-to-open, what kind of control action should the controller (LC) have and why?



- Direct acting since the control valve is direct acting
- Reverse acting since the control valve is reverse acting
- Direct acting since the control valve is reverse acting
- d. Reverse acting since the control valve is direct acting
- 57 A 2-input, 2-output process can be described in the Laplace transform domain as given below

$$(\tau_1 s + 1)Y_1(s) = K_1 U_1(s) + K_2 U_2(s)$$
  
$$(\tau_2 s + 1)Y_2(s) = K_3 U_2(s) + K_1 U_1(s)$$

where  $U_1$  and  $U_2$  are the inputs and  $Y_1$  and  $Y_2$  are the outputs. The gams of the transfer functions  $Y_1(s)/U_2(s)$  and  $Y_2(s)/U_2(s)$ , respectively, are

- a. K<sub>2</sub> and K<sub>3</sub>
- b. Kr and K1 + K2 K4
- c. K2 and K3 + K1 K4
- d. K<sub>2</sub> and K<sub>3</sub> + K<sub>2</sub> K<sub>4</sub>
- 58. A process is perturbed by a sinusoidal input,  $u(t) = A\sin\omega t$ . The resulting process output is  $Y(s) = \frac{kA\omega}{(\tau s + 1)(s^2 + \omega^2)}$ . If y(0)

= 0, the differential equation representing the process is

a. 
$$\frac{dy(t)}{dt} + ry(t) = Ku(t)$$

b. 
$$\tau \frac{dy(t)}{dt} + y(t) = KAu(t)$$

e. 
$$\tau \frac{dy(t)}{dt} + y(t) = Ku(t)$$

$$d. \quad r\left(\frac{dy(t)}{dt} + y(t)\right) = KAu(t)$$

59. A weighing machine is calibrated at 25°C and the output reading R (in mm) is related to the weight W (in kg) by the equation

$$R = sW$$

where the sensitivity s = 20 mm/kg. At a temperature of 30°C, the weighing machine undergoes a zero drift (change in instrument output reading at zero value of weight) of +2 mm and its sensitivity changes to 20.5 mm/kg. The weighing machine when used at 30°C shows a reading of 50 mm. The true weight (in kg) of the object is

- a. 2.34
- 6. 2.40
- 2.44
- d. 2,50
- 60. In a desalination plant, an evaporator of area 200 m<sup>2</sup> was purchased in 1996 at a cost of \$3,00,000. In 2002, another evaporator of area 50m<sup>2</sup> was added. What was the cost of the second evaporator (in)? Assume that the cost of evaporator scales as (capacity)<sup>0.54</sup> The Marshall and Swift index was 1048.5 in 1996 and 1116.9 in 2002.
  - a. 1,30,500
  - b. 1,39,100
  - c. 1.41,900
  - d. 1,51,200
- 61 The mixing of rubber latex solution was studied in an unbaffled mixer in the laboratory. The mixer was equipped with a six blade turbine impeller. A tyre company scales this process up using a baffled tank, The baffled tank has 3 times the diameter of the lab scale mixer. It uses the same type of impeller operated at the same speed. The relevant shape factors are also Assuming that ammar the same conditions prevail in both cases, the power requirement in the industrial scale mixer
  - a. is 3 times that of the lab scale mixer
  - is 9 times that of the lab scale mixer
  - c. is 27 times that of the lab scale mixer
  - d. cannot be estimated reliably due to the presence of baffles
- Due to a 20% drop in the product selling price, the pay-back period of a new plant increased to 1.5 times that estimated initially, the production cost and the production rate remaining unchanged. If the production cost is C<sub>p</sub>, and the new selling price is then C<sub>s</sub>, then C<sub>p</sub>/C<sub>s</sub> is

- a. 0.2
- b. 0.4
- c. 0.5
- d: 0.6
- 63. Obtain the optimal diameter of a cylindrical storage vessel of volume V. The curved shell costs C, (in Rs/m²), and the flat top and bottom plates cost C<sub>p</sub> (in Rs/m²)
  - a.  $D = \frac{C_1}{C_p} \left[ \frac{4V}{\pi} \right]^{\frac{1}{2}}$
  - $\mathbf{b.} \quad D = \left[ \frac{8VC_*}{\pi C_p} \right]^{\frac{1}{4}}$
  - $\mathbf{e} \cdot D = \left[ \frac{VC_s}{C_h} \right]^{\frac{1}{2}}$
  - $\mathbf{d.} \quad D = \left[ \frac{4VC_{\pi}}{\pi C_{\mu}} \right]^{\frac{1}{4}}$
- 64. A sale contract signed by a chemical manufacturer is expected to generate a net cash flow of Rs. 2,50,000/- per year at the end of each year for a period of three years. The applicable discount rate (interest rate) is 10%. The net present worth of the total cash flow is Rs.
  - n= 7,50,000
  - b. 6,83,750
  - c. 6.2 1,500
  - d. 3,32,750
- 65. A saturated vapor is fed to a distillation column at 180 kmol/hr. Both the rectifying and stripping sections of the column operate at 60% of their respective flooding velocities. The flooding velocity of the rectifying section is twice that of the stripping section. The assumptions of constant molar overflow and constant molar vapor density throughout the column are valid. If the boil-up rate is 60 kmol/hr, then the relationship between the diameters of the rectifying section (d<sub>t</sub>) and the stripping section (d<sub>t</sub>) is
  - $\mathbf{n}. \ \mathbf{d}_{r} = \sqrt{2} \, \mathbf{d}_{u}$
  - b.  $d_r = \sqrt{3} d_a$
  - $e_r = 2d_r$
  - $d_r = 3d_r$
- 66. Pair the following industrial processes with the catalysts used
  List I

- A. Oxidation of o-xylene to phthalic
- B. Oxidation of ethanol to acetaldehyde
- C. Oxidation of ammonia to oxides of nitrogen
- List II
- 1. V2O5
- 2. Pd
- 3. Ag
- 4. Pi
- A B C a, 1 2 4 b, 2 1 4 c, 4 3 1
- 67. Pair the following reactors with their products
  - List L
  - A. Arc Furnace
  - B. Fermenter
  - C. Hydrogenator
  - List II
  - 1. Citric acid
  - 2. Calcium carbide
  - 3. Saturated fats
  - 4. Alum
  - A B C
    a. 4 1 3
    b. 2 1 3
    c. 4 3 1
- 68. Pair the following polymers with their chain characteristics
  - List I
  - A. HDPE
  - B. LDPE
  - C. LLDPE

#### List II

- Very few branches
- 2. Short and regular branches
- 3. High branching with both short and long chain branches

	A	B	C
a.	1	2	3
b.	2	1	3
и. b. c.	ì	3	2
d	2	3	1

- 69. Choose the most appropriate pairs from the following
  - List I
  - A. Nitration
  - B. Sulphonation
  - C. Carbonation
  - List I

- 1. Detergent
- 2. Pulp and paper
- 3. Explosives
- 4. Sugar

4.4	-24-24-		
	A	$\mathbf{B}$	C
a,	1	2	4
а. Ь.	3	1	2
C.	3	1	4
44			

70. Match the following

List I

- A. Carbon disulphide
- B. Caprolactum
- C. Gypsum

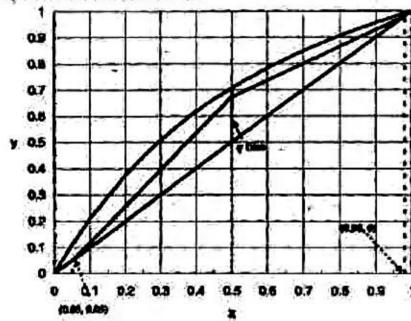
List II

- Nylon-6
- 2. Nylon-66
- 3. Phosphoric acid
- 4. Viscose rayon

	A	В	C
a.	4	2	3
a. b.	2	1	3
c.	3	1	4
ď	4	1	3

### Common Data for Questions (71,72 &73)

A binary distillation column separates 100 mol/hr of a feed mixture into distillate D and residue W. The McCabe-Thiele diagram for this process is given below. The relative volatility for the binary system is constant at 2.4.

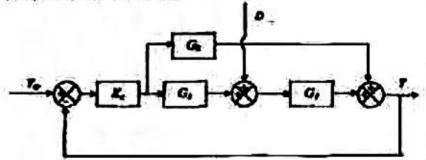


- 71. The distillate and residue flow rates (in mol/hr) are
  - a. D = 48.4, W = 51.6
  - b. D = 51.6, W = 48.4
  - e. D = 54.7, W = 45.3
  - d. D = 45.3, W = 54.7
- 72. The ratio of liquid to vapor molar flow rates in the rectifying section is

- a, 0.64
- b. 1.00
- e. 1.55
- d. 1.80
- 73. The minimum number of theoretical stages (inclusive of reboiler) for this process is
  - a. 5.2
  - b. 6.1
  - c. 7.8
  - d. infinite

## Common Data for Questions (74 & 75)

The block diagram of a closed loop control system is shown in the figure below. Y is the controlled variable. D is disturbance, Y is the set point,  $G_1$ ,  $G_2$ , and  $G_3$ , are transfer functions, and K is the proportional controller.



74. The closed loop transfer function Y(s)/D(s) is given by

a. 
$$\frac{G_{j}G_{j}}{1+(G_{j}G_{j}+G_{j})K_{s}}$$

b. 
$$\frac{G_i}{1+(G_iG_j+G_2)K_e}$$

e. 
$$\frac{G_3}{1+(G_1+G_2)G_1K_1}$$

d. 
$$\frac{G_1}{1+(G_1G_1+G_2)K_2}$$

75. Let G<sub>1</sub>(s) - 1 and G<sub>2</sub>(s) - G<sub>3</sub>(s) - 1/(s+1). A step change of magnitude M is made in the set point. The steady state offset of the closed loop response Y is

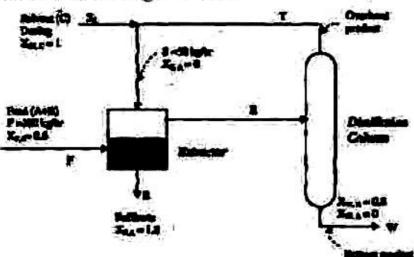
$$a, \frac{M}{1+2K}$$

b. 
$$\frac{M}{1+K_{\epsilon}}$$

$$c. \frac{M(K_c-1)}{1+2K}$$

d. zero

Statement for Linked Answer Question (76 and 77) Solvent C is used to extract solute B selectively from 100 kg/hr feed mixture A+B in a steady state continuous process shown below. The solubility of C in the raffinate and the solubility of A in the extract are negligible. The extract is distilled to recover B in the bottom product. The overhead product is recycled to the extractor. The loss of solvent in the bottoms is compensated by make up solvent S<sub>d</sub>. The total flow rate of the solvent stream S going to the extractor is 50 kg/hr. The mass fractions (X, 's) of some selected streams are indicated in the figure below.



76. Distillation bottoms flow rate W and solvent dosing rate 5d in kg/hr are

a. 
$$W = 50$$
,  $S_d = 50$ 

b. 
$$W = 100, S_3 = 20$$

e. 
$$W = 10$$
,  $S_d = 50$ 

d. 
$$W=50$$
,  $Sd=10$ 

77. Feed rate E to the distillation column and overhead product rate T in kg/hr are

a. 
$$E = 90, T = 40$$

b. 
$$E = 80, T = 40$$

d. 
$$E = 45, T = 20$$

# Statement for Linked Answer Question (78 and 79)

A continuous grinder obeying the Bond crushing law grinds a solid at the rate of 1000 kg/hr from the initial diameter of 10 mm to the final diameter of 1 mm.

- 78. If the market now demands particles of size 0.5 mm, the output rate of the grinder (in kg/hr) for the same power input would be reduced to
  - n. 227
  - b. 474
  - c. 623
  - d. 856
- In order to restore the output back to 1000 kg/hr, an additional grinder was installed.

The two grinders can be operated either in series (configuration-1) or parallel (configuration-2). Compare the two

(configuration-2). Compare the two configurations in terms of the additional power consumption over the case above.

- a. configuration-1 consumes less power than configuration-2
- b. configuration-2 consumes less power than configuration-1 both configurations consume the same power
- configuration-2 consumes less or more power than configuration-1 depending on how the feed is distributed between the two grinders in configuration-2 (the parallel configuration)

# Statement for Linked Answer Question (80 and 81)

Consider the diffusion of a reactant A through a cylindrical catalyst pore of radius R and length L>R. Reactant A undergoes a zeroth order reaction on the cylindrical surface of the pore. The following equation describes changes in the concentration of A within the pore due to the axial diffusion of A and the disappearance of A due to reaction.

$$\frac{d^3c_A}{dx^3} = K$$

where c<sub>A</sub> is the concentration of A at a distance x from the pore entrance, and K is a constant

80. If the concentration of A at the pore entrance (x = 0) is C<sub>AO</sub>, and x = L is a dead end where no reaction occurs, the concentration profile of A in the pore is given by

a. 
$$c_A(x) = \frac{Kx^2}{2} - KLx + c_{All}$$

b. 
$$c_A(x) = \frac{Kx(x-L)}{2} - c_{A0} \frac{x}{L} + c_{A0}$$

$$c: \quad c_A(x) = c_{Ab}$$

d. 
$$c_A(x) = c_{A0} \left( \frac{L-x}{L} \right)$$

81. The minimum pore length for A to be completely converted within the pore is

$$\mathbf{a}. \quad \left(\frac{c_{A0}}{K}\right)^{1/2}$$

b. 
$$\left(\frac{c_{Ali}}{2K}\right)^{U^2}$$

e. 
$$\left(\frac{2c_{40}}{K}\right)$$

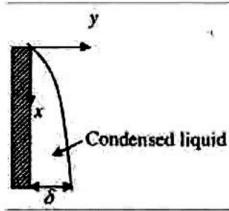
$$d. \left(\frac{2c_{AU}}{K}\right)^{1/2}$$

# Statement for Linked Answer Question (82 and 83)

In film condensation on a vertical plane surface. the x directional velocity distribution is given by

$$u(y) = \frac{g(\rho_1 - \rho_x)}{\mu_1} \left( \delta y - \frac{1}{2} y^2 \right)$$

where & is the film thickness at any x.



82. The mass flow rate of the condensate m(x) through any axial position x per unit width of the plate is given by

a. 
$$m(x) = \frac{g\rho_i(\rho_i - \rho_v)\delta^3}{3\mu_i}$$

b. 
$$m(x) = \frac{g(\rho_i - \rho_s)\delta^z}{3\mu_i}$$

e. 
$$m(x) = \frac{g\rho_r^4 - \delta^3}{\mu_i}$$

c. 
$$m(x) = \frac{g\rho_x^3 - \delta^3}{\mu_1}$$
  
d. 
$$m(x) = \frac{g\rho_1(\rho_x)\delta^3}{3\mu_1}$$

83, Differentiate m(x) with respect to  $\delta$  to get the differential increase in condensate mass dm with film thickness i.e., dm/do. Then obtain dm/dx assuming heat flux through the film to be due to conduction based on a linear temperature profile between the vapor and wall. Hence determine dδ/dx.

> Here  $\mu_1$  is liquid viscosity,  $k_i$  is thermal conductivity, and  $\lambda$  is latent heat of

condensation. T<sub>v</sub> is the vapor temperature and Tw is the wall temperature

a. 
$$\frac{d\delta}{dx} = \frac{\mu_i k_i (T_i - T_{ii})}{g \rho_i (\rho_i - \rho_r) \lambda \delta^2}$$

b. 
$$\frac{d\delta}{dx} = \frac{\mu_i k_i (T_e - T_e)}{g \rho_e (\rho_i - \rho_e) \lambda \delta^3}$$

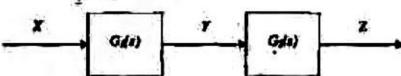
c. 
$$\frac{d\delta}{dx} = \frac{\mu_i (T_i - T_w)}{g \rho_v k_i (\rho_i - \rho_v) \lambda} \frac{1}{\delta^2}$$

d. 
$$\frac{d\delta}{dx} = \frac{\mu_i k_i (T_v - T_w)}{g \rho_v (\rho_i - \rho_v) \lambda_i \delta^3}$$

# Statement for Linked Answer Question (64 and 85)

For the system shown below,  $G_1(s) = \frac{1}{r_1 s + 1}$ 

$$G_2(s) = \frac{1}{\tau_1 s + 1}$$
 and  $\tau_2 = 1\tau_1$ 



When the system is excited by the sinusoidal input  $X(t) = sin\omega t$ , the intermediate response Y is given by

$$Y = A\sin\left(\omega t + \phi\right)$$

84. If the response of Y lags behind the input X by  $45^{\circ}$  and  $\tau_1 = 1$ , then the input frequency w is

For the same input, the amplitude of the 85. output Z will be