



SUMMER-17 EXAMINATION
Model Answer

Subject Title: Stoichiometry

Subject code : 17315

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Important Instructions to examiners:

- 1) The answers should be examined by key words and not as word-to-word as given in the model answer scheme.
- 2) The model answer and the answer written by candidate may vary but the examiner may try to assess the understanding level of the candidate.
- 3) The language errors such as grammatical, spelling errors should not be given more Importance (Not applicable for subject English and Communication Skills).
- 4) While assessing figures, examiner may give credit for principal components indicated in the figure. The figures drawn by candidate and model answer may vary. The examiner may give credit for any equivalent figure drawn.
- 5) Credits may be given step wise for numerical problems. In some cases, the assumed constant values may vary and there may be some difference in the candidate's answers and model answer.
- 6) In case of some questions credit may be given by judgement on part of examiner of relevant answer based on candidate's understanding.
- 7) For programming language papers, credit may be given to any other program based on equivalent concept.

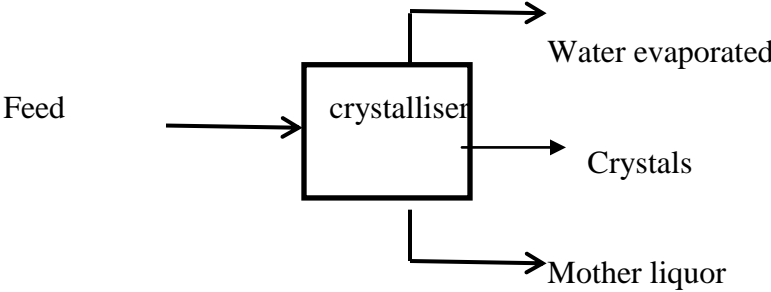


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Q No.	Answer	marks
1	Any 10	20
1-a	Ideal Gas law: $PV=nRT$ where P - pressure, V - volume, n- moles, K-absolute temperature and R – universal gas constant.	1 1
1-b	Vander Waal's equation of state: $(P+a/V^2)(V-b)= nRT$ Where a & b are constants	2
1-c	% Conversion: It is the ratio of amount of limiting reactant reacted to the amount of limiting reactant totally charged. Express in percentage.	2
1-d	Excess component: It is the reactant which is in excess of the theoretical or stoichiometric requirement. OR It is the component which decides the extend of a reaction.	2
1-e	Sensible Heat : Sensible heat is the heat that must be transferred to raise or lower the temperature of a substance or mixture of substance. Latent Heat : When matter undergoes a phase change, the enthalpy change associated with unit amount of matter at constant temperature and pressure is known as Latent Heat of phase change.	1 1
1-f	Material balance diagram for crystallisation 	2



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1-g	Standard heat of combustion: It is the amount of heat liberated when one mol of a compound is combusted or burned in oxygen at standard conditions.(25 ⁰ C and 1atm pressure)	2
1-h	Heat capacity at constant volume: It is the amount of heat required to increase the temperature of one gram of substance by one degree.(under constant volume)	2
1-i	Raoult's law: It states that at a given temperature, the equilibrium partial pressure of a component of a solution in the vapour phase is equal to the product of the mole fraction of the component in the liquid phase and the vapour pressure of the pure component. $p_A = p_A^0 X_A$ p_A - partial pressure p_A^0 - vapour pressure vapour X_A - mole fraction of the component in the liquid phase	1 1
1-j	Law of conservation of energy: Energy input= energy output + accumulation	2
1-k	1)Stoichiometric Equation : The stoichiometric equation of a chemical reaction is the statement indicating relative moles of reactant and products that take part in the reaction. 2) Stoichiometric ratio : it is the ratio of stoichiometric coefficient of two molecular species or Components in the balanced reaction	1 1
1-l	Dalton's law: It states that the total pressure exerted by a gas mixture is equal to the sum of partial pressures	1

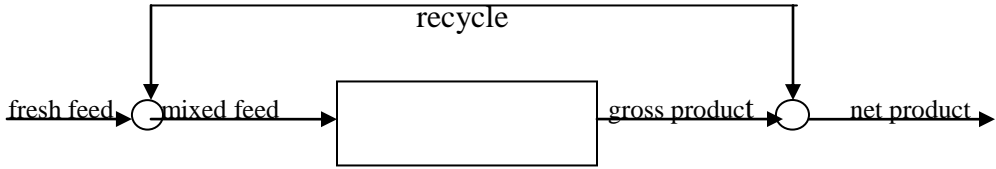
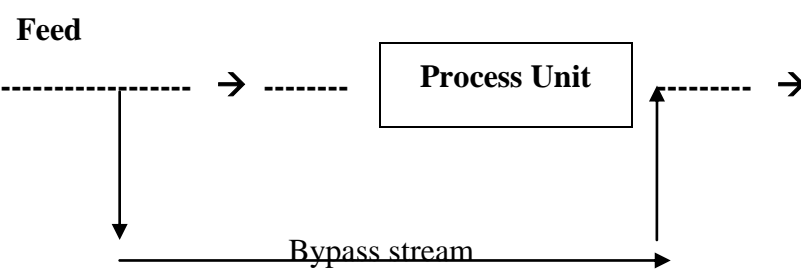


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	<p>Mathematical Statement: $P = P_1 + P_2 + P_3$ where P is the total pressure of gas mixture, P_1, P_2, P_3 are partial pressures</p>	1
2	Any 2	16
2-a	<p>Recycling: It is returning back a portion of stream leaving a process unit to the entrance of the process unit for further processing</p>  <p>Recycling is used for maximum utilisation of the valuable reactants, improvement of the performance of the operation, utilization of the heat being lost in the exit stream, improvement in the selectivity of the products, maintaining process rate at a high value and enrichment of a product.</p> <p>Bypass Operation :</p> <p>In these operations, a fraction of the feed stream to a process unit is diverted around and combined with the output stream.</p>  <p>- Bypassing is practiced industrially whenever accurate control of the</p>	<p>2</p> <p>2</p> <p>2</p>



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	<p>composition or concentration of the process exit stream is expected.</p> <p>- The composition and properties of the product may be varied by varying the fraction of the feed that is bypassed.</p> <p>Example: A juice concentration process in which the dehydration process runs most efficiently by removing more water than is desired. A portion of the feed may be directed around the dehydrator in a bypass loop, to be mixed with unprocessed feed. Or any other example</p>	
2-b	<p>Basis: 100 kmol air</p> <p>Avg. mol.wt of air = $M_1X_1 + M_2X_2$</p> $= 28 * 0.79 + 32 * 0.21$ $= \mathbf{28.84}$ <p>Density = $P * M_{av} / RT$</p> $= 101.325 * 28.84 / 8.314 * 273$ $= \mathbf{1.287 \text{ Kg/m}^3}$	<p>2</p> <p>2</p> <p>2</p> <p>2</p>
2-c	<p>Case –I: Basis: 100 Kg/hr of solid handling capacity of the evaporator.</p> <div style="text-align: center;"> </div> <p>Let X be the be the Kg/hr of weak liquor then we have.</p> $0.05 X = 100$ $X = 2000 \text{ Kg/hr}$ <p>Y be the be the Kg/hr of thick liquor then we have.</p> $0.5 Y = 100$	<p>1</p> <p>1</p>



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	<p>$Y = 200 \text{ kg/hr}$</p> <p>Overall Material Balance:</p> <p>Kg/hr of weak liquor = Kg/hr of thick liquor + Kg/hr of Water evaporated</p> <p>$2000 = 200 + \text{Kg/hr of Water evaporated}$</p> <p>Kg/hr of Water evaporated = 1800 Kg/hr</p> <p>Case –II: Basis: 1800 Kg/hr of Water evaporated</p> <p>Let A be the be the Kg/hr of weak liquor</p> <p>B be the be the Kg/hr of thick liquor</p> <p>Overall Material Balance:</p> <p>$A = B + 1800$</p> <p>Material Balance of Solids :</p> <p>$0.06 A = 0.36 B$</p> <p>$A = 6 B$</p> <p>Putting in above equation</p> <p>$6 B = B + 1800$</p> <p>$B = 360 \text{ Kg/hr}$</p> <p>$A = 360 + 1800 = 2160 \text{ Kg/hr}$</p> <p>Solid in weak liquor = $0.06 \times 2160 = 129.6 \text{ Kg/hr}$</p> <p>Solid handling Capacity = 129.6 Kg/hr ----- Ans.</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
3	Any 2	16
3-a	<p>SOLUTION :</p> <p>BASIS : 1000 kg of desired mixed acid.</p> <p>Waste acid, 28 % H_2SO_4, 35% HNO_3</p> <p style="text-align: center;"> </p>	<p>1</p> <p>1</p>



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<div style="text-align: center;"> </div> <p style="text-align: center;">Block diagram for fortifying waste acid with concentrated acids</p> <p>Let x, y and z be the kg of waste acid , concentrated sulphuric acid and concentrated nitric acid required to make 1000 kg desired acid.</p> <p>Overall material Balance:</p> $x + y + z = 1000 \dots (i)$ <p>Material Balance of H₂SO₄ :</p> $0.28 x + 0.98 y = 0.4 \times 1000 \dots (ii)$ $0.28 x + 0.98 y = 400$ <p>Material Balance of HNO₃ :</p> $0.35 x + 0.72 z = 0.41 \times 1000$ $0.35 x + 0.72 z = 410 \dots (iii)$ <p>Solving the above three equations, we get X= 98.14 Y= 380.12 Z= 521.74</p> <p>Amount of waste acid required = 98.14kg</p> <p>Amount of concentrated sulphuric acid required = 380.12 kg</p> <p>Amount of concentrated nitric acid require = 521.74 kgAns</p>	<p>1</p> <p>1</p> <p>1</p> <p>3</p>
<p>3-b Basis: 50 kmoles /hr butane</p>	



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	$\text{C}_4\text{H}_{10} + 6.5 \text{O}_2 \rightarrow 4\text{CO}_2 + 5 \text{H}_2\text{O}$ <p>100 kmol air fed = 21 kmol O₂ fed 2100 kmol air fed = ? O₂ fed = 2100*21/100= 441 kmoles 1 kmol C₄H₁₀ fed = 6.5 kmol O₂ theoretically required 50 kmol C₄H₁₀ fed = ? O₂ theoretically required = 325 kmol % excess= (O₂ fed-O₂ theoretical)*100/ O₂ theoretical = (441-325)*100/325 = 35.69%</p>	<p>1</p> <p>2</p> <p>2</p> <p>1</p> <p>2</p>												
3-c	<p>Steps involved in solving material balance calculations:</p> <ol style="list-style-type: none"> 1. Assume suitable basis of calculation as given in problem. 2. Adopt weight units in case of problem of process without chemical reaction. 3. Draw block diagram of process 4. Show input and output streams 5. Write overall material balance 6. Write individual material balance 7. Solve above two algebraic equations 8. Get values of two unknown quantities. 9. Write balances as follows: <table border="1" data-bbox="282 1575 1305 1848"> <thead> <tr> <th></th> <th>feed</th> <th>product</th> <th>Component removed</th> </tr> </thead> <tbody> <tr> <td>Unchanging component</td> <td></td> <td></td> <td></td> </tr> <tr> <td>Outgoing</td> <td></td> <td></td> <td></td> </tr> </tbody> </table>		feed	product	Component removed	Unchanging component				Outgoing				<p>1 mark each</p>
	feed	product	Component removed											
Unchanging component														
Outgoing														



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component																					
4	Any 2		16																		
4-a	$C_2H_4 + \frac{1}{2} O_2 \longrightarrow C_2H_4O$ $C_2H_4 + 3O_2 \longrightarrow 2CO_2 + 2H_2O$ Basis : 100 kmol ethylene and 100 kmol O_2 fed Conversion of ethylene = 85% Conversion of ethylene = (kmol ethylene reacted / kmol ethylene fed) * 100 $85 = (\text{kmol ethylene reacted} / 100) * 100$ kmol ethylene reacted = 85 % yield of $C_2H_4O = (\text{kmol } C_2H_4 \text{ reacted to form } C_2H_4O / \text{total } C_2H_4 \text{ reacted}) * 100$ $94.12 = (\text{kmol } C_2H_4 \text{ reacted to form } C_2H_4O / 85) * 100$ kmol C_2H_4 reacted to form $C_2H_4O = 80 \text{ kmol}$ kmol C_2H_4 reacted for 2 nd reaction = $85 - 80 = 5 \text{ kmol}$ CO_2 formed = $2 * 5 = 10 \text{ kmol}$ H_2O formed = $2 * 5 = 10 \text{ kmol}$ O_2 reacted = $(80 * 0.5 + 3 * 5) = 55 \text{ kmol}$ O_2 unreacted = $(100 - 55) = 45 \text{ kmol}$ C_2H_4 unreacted = $(100 - 85) = 15 \text{ kmol}$		1 1 1 1 1 1																		
	<table border="1"> <thead> <tr> <th>Component</th> <th>Kmol</th> <th>Mole %</th> </tr> </thead> <tbody> <tr> <td>C_2H_4</td> <td>15</td> <td>9.38</td> </tr> <tr> <td>C_2H_4O</td> <td>80</td> <td>50</td> </tr> <tr> <td>O_2</td> <td>45</td> <td>28.12</td> </tr> <tr> <td>CO_2</td> <td>10</td> <td>6.25</td> </tr> <tr> <td>H_2O</td> <td>10</td> <td>6.25</td> </tr> </tbody> </table>	Component	Kmol	Mole %	C_2H_4	15	9.38	C_2H_4O	80	50	O_2	45	28.12	CO_2	10	6.25	H_2O	10	6.25		2
Component	Kmol	Mole %																			
C_2H_4	15	9.38																			
C_2H_4O	80	50																			
O_2	45	28.12																			
CO_2	10	6.25																			
H_2O	10	6.25																			
4-b	Basis : 15000 mol/h of N_2 - H_2 mixture																				



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<p>Molal flow rate of gas mixture = 5 kmol/h</p> <p>$X_{N_2} = 25/100 = 0.25$</p> <p>$X_{H_2} = 75/100 = 0.75$</p> <p>$C_p^{\circ} \text{ mix} = \sum C_p^{\circ} \text{ mix} \cdot X_i = X_{N_2} \cdot C_p^{\circ} N_2 + X_{H_2} \cdot C_p^{\circ} H_2$ $= 0.25 (29.5909 - 5.141 \times 10^{-3} T + 13.1829 \times 10^{-6} T^2 - 4.968 \times 10^{-9} T^3)$ $+ 0.75 (28.6105 + 1.0194 \times 10^{-3} T - 0.1476 \times 10^{-6} T^2 + 0.769 \times 10^{-9} T^3)$ $= 28.8556 - 0.5207 \times 10^{-3} T + 3.185 \times 10^{-6} T^2 - 0.6652 \times 10^{-9} T^3$</p> <p>Q = Heat added</p> <p>$Q = n \int_{T_1}^{T_2} C_p^{\circ} \text{ mix} dT$</p> <p>$Q = n \int_{T_1}^{T_2} [28.8556 - 0.5207 \times 10^{-3} T + 3.185 \times 10^{-6} T^2 - 0.6652 \times 10^{-9} T^3] dT$</p> <p>$= n [28.8556 (T_2 - T_1) - \frac{0.5207 \times 10^{-3}}{2} (T_2^2 - T_1^2) + \frac{3.185 \times 10^{-6}}{3} (T_2^3 - T_1^3) - \frac{0.6652 \times 10^{-9}}{4} (T_2^4 - T_1^4)]$</p> <p>Where , n= 15 Kmol/h , T₂ = 473 K , T₁ = 298 K</p> <p>$= 15 [28.8556 (473 - 298) - \frac{0.5207 \times 10^{-3}}{2} (473^2 - 298^2) + \frac{3.185 \times 10^{-6}}{3} (473^3 - 298^3) - \frac{0.6652 \times 10^{-9}}{4} (473^4 - 298^4)]$</p>	<p>1</p> <p>2</p> <p>1</p> <p>2</p>
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	$+ \frac{\dots\dots\dots (473^3 - 298^3)}{3} - \frac{\dots\dots\dots (473^4 - 298^4)}{4}]$ <p>Q = 15(5049.73 - 35.13 - 7.01)</p> <p>Q = 76377.6 KJ/h = 21.216 kJ/s = 21.216 kW ----- ans</p>	2
4-c	<p>Basis: 1 mol liquid C₅H₁₂</p> $\Delta H^0_R = \Sigma \Delta H^0_{f(\text{pr})} - \Sigma \Delta H^0_{f(\text{react})}$ $= [(-393.51*5)+(-285.83*6)] - (-173.49)$ <p>= -3509.04KJ</p>	1 2 2 3
5	Any 2	16
5-a	<p>Basis: 10000 kg/hr of feed</p> <div style="display: flex; align-items: center; justify-content: center;"> <div style="margin-right: 20px;"> <p>10000 kg/hr solution 30 % methanol</p> </div> <div style="border: 1px solid black; padding: 5px; text-align: center;"> <p>distillation</p> </div> <div style="margin-left: 20px;"> <p>→ Distillate X kg/hr 97% methanol</p> <p>→ Waste solution Y kg/hr 1% methanol</p> </div> </div> <p>Overall balance is</p> $10000 = X + Y \dots\dots\dots(1)$ <p>Individual balance for CH₃OH is</p> $0.3*10000 = 0.97X + 0.01*Y \dots\dots\dots(2)$ <p>Solving the equations</p> <p>X = 3020.83 Kg/hr</p> <p>Y = 6979.17 kg/hr</p> <p>CH₃OH in bottom product = 0.01*6979.17 = 69.79Kg/hr</p> <p>% loss of CH₃OH = (CH₃OH in bottom product / CH₃OH in feed)*100</p> $= (69.79/3000)*100$ <p>= 2.32%</p>	1 1 1 2 1 1 1
5-b	Henry's law is p _A = H X _A	1



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	$= 94.12\%$	1
6	Any 4	16
6-a	<p>Basis: 2000 kg wet solid</p> <p>Overall balance is $2000 = X + Y$ Balance for solid $0.70 * 2000 = 0.99 * Y$ $Y = 1414.14 \text{ kg}$ $X = 585.86$ Water removed = 585.86 kg Product obtained = 1414.14 kg</p>	1 1 1 1
6-b	<p>$\text{CO} + 2\text{H}_2 \text{ -----} \rightarrow \text{CH}_3\text{OH}$</p> <p>a) Stoichiometric ratio of H_2 to $\text{CO} = 2:1$ b) Kmoles of CH_3OH produced per kmol CO reacted = 1 c) Weight ratio of CO to $\text{H}_2 = 28:4 = 7$ d) Quantity of CO required to produce 1000 kg CH_3OH $= 1000/32 * 28 = \mathbf{875 \text{ kg}}$</p>	1 1 1 1
6-c	General Material Balance Procedure	4



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	<p>1) Assume Suitable Basis of calculation.</p> <p>2) Adopt weight basis for without chemical reactions and molar basis for with chemical reactions.</p> <p>3) In case of material balance with chemical, write balance chemical reaction and search out limiting component.</p> <p>4) In with chemical reaction, the quantity of a reacting component appearing in the product stream is the quantity of that material remains unreacted.</p> <p>5) Supplied quantity of an excess reactant calculated from the theoretical requirement is based on the quantity of a limiting reactant fed.</p>	
6-d	<p>Hess's law:</p> <p>It states that the heat involved in a chemical reaction is same whether the reaction takes place in a single or in several steps.</p> <p>A \longrightarrow B ΔT_1</p> <p>B \longrightarrow C ΔT_2</p> <p>C \longrightarrow D ΔT_3</p> <p>A \longrightarrow D ΔT</p> <p>Then</p> <p>$\Delta T = \Delta T_1 + \Delta T_2 + \Delta T_3$</p> <p>Application of Hess's law: Using this law we can calculate the heat of formation of a compound from a series of reactions that do not involve the direct formation of the compound from its elements.</p>	<p>2</p> <p>2</p>
6-e	<p>Basis: 1m^3 fixed mass of gas at constant temperature</p> <p>$P_1 = 1\text{atm}$ $V_1 = 1\text{m}^3$ $T_1 = T = T_2$</p> <p>$P_2 = ?$ $V_2 = 0.5\text{m}^3$</p> <p>$P_1 V_1 / T_1 = P_2 V_2 / T_2$</p> <p>$1 * 1 / T = P_2 * 0.5 / T$</p> <p>Or $P_2 = 2\text{ atm}$</p>	<p>1</p> <p>1</p> <p>1</p>

