



WINTER-16 EXAMINATION
Model Answer

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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-A	Any 4	
1A-a	Sensible Heat: - Sensible heat is the heat that must be transferred to raise or lower the temperature of a substance or mixture of substance. Heat capacity: It is the amount of heat required to increase the temperature of one gram of substance by one degree.	1 1
1A-b	Law of conservation of mass: It states that For any process input= output+accumulation	2
1A-c	Value of R: 1.987 Kcal/kmol.K 8.315 J/mol.K	1 1
1A-d	$\text{CO} + 2\text{H}_2 \rightarrow \text{CH}_3\text{OH}$ Stoichiometric coefficient of $\text{CO}:\text{H}_2 = 1:2$	2
1 A-e	Law for real gas is Vander Waal's equation of state: $(P+a/V^2)(V-b)= nRT$ Where a & b are constants	1 1
1A-f	Law of conservation of energy: Energy input= energy output + accumulation	1 1
1-B	Any 2	12
1B-a	Basis: 100 kmoles of feed A fed= 60 kmol B fed=30 kmol Inert fed = 10 kmol Reaction is $2\text{A} + \text{B} \rightarrow \text{C}$	2

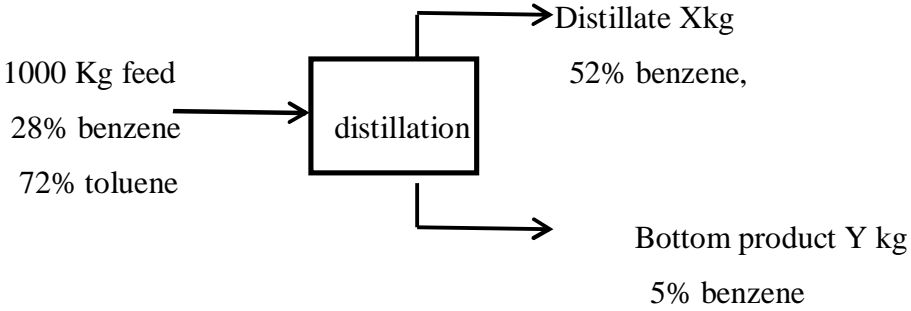


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	<p>Conversion of A =80%</p> <p>A converted= $(80/100) *60 = 48 \text{ kmol}$</p> <p>A unreacted= $60-48 = 12 \text{ kmol}$</p> <p>B reacted = 24 kmol</p> <p>B unreacted = $30-24 = 6 \text{ kmol}$</p> <p>C formed= 24 kmol</p> <p>Inert= 10 kmol</p> <p>Composition of product stream:</p> <table border="1" data-bbox="282 858 1304 1199"> <thead> <tr> <th>component</th> <th>kmol</th> <th>Mol%</th> </tr> </thead> <tbody> <tr> <td>A</td> <td>12</td> <td>23.07</td> </tr> <tr> <td>B</td> <td>6</td> <td>11.54</td> </tr> <tr> <td>C</td> <td>24</td> <td>46.15</td> </tr> <tr> <td>Inert</td> <td>10</td> <td>19.23</td> </tr> <tr> <td>total</td> <td>52</td> <td></td> </tr> </tbody> </table>	component	kmol	Mol%	A	12	23.07	B	6	11.54	C	24	46.15	Inert	10	19.23	total	52		<p>2</p> <p>2</p>
component	kmol	Mol%																		
A	12	23.07																		
B	6	11.54																		
C	24	46.15																		
Inert	10	19.23																		
total	52																			
<p>1B-b</p>	<p>Basis: 1000 kg benzene-toluene</p> <div style="text-align: center;">  </div> <p>Overall balance is</p> <p>$1000= X+ Y$</p> <p>Benzene balance is</p>	<p>1</p> <p>1</p> <p>1</p>																		



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	(28/100)* 1000= 0.52X + 0.05Y Solving X= 489.36kg and Y = 510.64 Kg Distillate= 489.36kg Bottom product = 510.64 Kg	1 2
1B-c	Basis: 1 Kg HCl acid $4\text{HCl} + \text{O}_2 \rightarrow 2\text{Cl}_2 + 2\text{H}_2\text{O}$ HCl fed = $1/36.5 = 0.027$ kmoles From reaction Oxygen theoretically required = $0.027/4 = 6.85 \times 10^{-3}$ % excess = 30 Oxygen fed = 8.9×10^{-3} kmoles Air fed = 0.042 kmoles = 1.22 kg	1 1 1 1 1
2	Any 4	16
2-a	Basis: 1000 Kg/hr HCHO formed $\text{CH}_3\text{OH} \rightarrow \text{HCHO} + \text{H}_2$ HCHO formed = $1000/30 = 33.33$ kmole/h From reaction, CH_3OH reacted = 33.33 kmole/h Conversion = 65% Conversion= (kmoles of CH_3OH reacted/kmoles of CH_3OH fed)* 100 Therefore CH_3OH fed = 51.28 Feed rate of methanol= 51.28×32 kmol/h = 1640.96 Kg/h	1 1 1 1
2-b	Hess's law:	



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	<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> $A \xrightarrow{\Delta T1} B$ $B \xrightarrow{\Delta T2} C$ $C \xrightarrow{\Delta T3} D$ $A \xrightarrow{\Delta T} D$ </p> <p>Then</p> <p>$\Delta T = \Delta T1 + \Delta T2 + \Delta T3$</p>	<p>2</p> <p>2</p>
2-c	<p>Basis: 4.73 kg coal</p> <p>$C + O_2 \rightarrow CO_2$</p> <p>At NTP, P= 101.325 KPa</p> <p>T= 273 K</p> <p>$PV = nRT$</p> <p>Or $n = PV/RT$</p> <p>Moles of $CO_2 = 0.237$ kmoles</p> <p>Weight of $CO_2 = 0.237 * 44 = 10.428$ kg</p> <p>Carbon in the sample = $(12/44) 10.428 = 2.844$ kg</p> <p>Carbon content in the sample = $(2.844 / 4.73) 100$ $= 60.13\%$</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p>
2-d	<p>Basis: 2000 kg wet solid</p> <p> <div style="text-align: center;"> </div> </p>	<p>1</p>



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	1% moisture 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	1 1 1
2-e	Basis: 15 kg propane Moles $n = 15/44 = 0.341 \text{ kmoles}$ At NTP, $P = 101.325 \text{ KPa}$ $T = 273 \text{ K}$ $PV = nRT$ Or $V = nRT/P$ $= 0.341 * 8.314 * 273/101.325$ $= 7.64 \text{ m}^3$	1 1 1 1
2-f	Basis: 1 mol liquid C_5H_{12} $\Delta H^0_R = \Sigma \Delta H^0_{f(\text{pr})} - \Sigma \Delta H^0_{f(\text{react})}$ $= [(-393.51 * 5) + (-285.83 * 6)] - (-173.49)$ $= -3509.04 \text{ KJ}$	2 2
3	Any 4	16
3-a	Basis: Fixed mass of gas at constant temperature Initial volume $V_1 = 1 \text{ m}^3$ Initial pressure $P_1 = P_1 \text{ atm}$ Final volume $V_1 = V_2 \text{ m}^3$	1



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	<p>Final pressure $P_2 = 1.85P_1 \text{ atm}$</p> <p>$P_1V_1/T_1 = P_2V_2/T_2$</p> <p>$P_1 \cdot 1/T = 1.85P_1 \cdot V_2/T$</p> <p>$V_2 = \mathbf{0.5405 \text{ m}^3}$</p>	<p>1</p> <p>1</p> <p>1</p>
3-b	<p>Basis: 1 kmol methane gas</p> <p>$Q = n[19.2494(T_2 - T_1) + 52.1135 \cdot 10^{-3}/2(T_2^2 - T_1^2) + 11.973 \cdot 10^{-6}/3 (T_2^3 - T_1^3) - 11.3173 \cdot 10^{-9}/4 (T_2^4 - T_1^4)]$</p> <p>$Q = 1[19.2494(523 - 303) + 52.1135 \cdot 10^{-3}/2(523^2 - 303^2) + 11.973 \cdot 10^{-6}/3 (523^3 - 303^3) - 11.3173 \cdot 10^{-9}/4 (523^4 - 303^4)]$</p> <p>$= 4234.86 + 4735.03 + 459.9 - 187.84$</p> <p>$= \mathbf{9241.96 \text{ KJ}}$</p>	<p>2</p> <p>1</p> <p>1</p>
3-c	<p>Basis: 100 kg coal</p> <div style="text-align: center;"> </div> <p>Component balance for ash</p> <p>$24 = 0.93 Y$</p> <p>Or $Y = 25.80 \text{ Kg}$</p> <p>Balance for carbon</p> <p>$63 = X + 0.07 \cdot 25.80$</p> <p>$X = 61.194 \text{ Kg}$</p>	<p>1</p> <p>1</p> <p>1</p>



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	Unburnt carbon = $0.07 \times 25.80 = 1.806 \text{ Kg}$ % of original C unburnt = $(1.806 / 63) \times 100$ = 2.867%	1
3-d	$\text{N}_2 + 3\text{H}_2 \rightarrow 2\text{NH}_3$ 1 kmol $\text{N}_2 = 3 \text{ kmol H}_2$ 25 “ = ? Molal flow rate of $\text{H}_2 = 75 \text{ kmol/hr}$ % conversion of $\text{N}_2 = (\text{N}_2 \text{ reacted} / \text{N}_2 \text{ fed}) \times 100$ $25 = (\text{N}_2 \text{ reacted} / 25) \times 100$ $\text{N}_2 \text{ reacted} = 6.25 \text{ kmol/h}$ From reaction NH_3 formed = 12.50 kmol/h Kg NH_3 formed = $12.50 \times 17 = 212.5 \text{ Kg}$	1 1 1 1
3-e	Basis: 1 kmol NH_3 $Q = n[C_{pm2}(T_2 - T_0) - C_{pm1}(T_1 - T_0)]$ = $1[37.7063(422 - 298) - 35.8641(311 - 298)]$ = 4209.35 KJ	2 2
3-f	Basis: 100 kmol air Avg. mol.wt of air = $M_1X_1 + M_2X_2$ = $28 \times 0.79 + 32 \times 0.21$ = 28.84 Density = $P \times M_{av} / RT$ = $101.325 \times 28.84 / 8.314 \times 273$ = 1.287 Kg/m³	1 1 1 1
4	Any 2	16



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4-a	<p>Basis: Mixture of N₂, CO₂, O₂</p> <p>Let X₁, X₂, X₃ be the mol fraction of N₂, CO₂, O₂ respectively.</p> <p>1st person 28 X₁ + 44 X₂ + 32 X₃ = 30.08</p> <p>2nd person 14 X₁ + 44 X₂ + 32 X₃ = 30.08</p> $X_1 + X_2 + X_3 = 1$ <p>Solving the equations, we get</p> <p>X₁ = 0.81</p> <p>X₂ = 0.11</p> <p>X₃ = 0.08</p> <p>Volume % of N₂ = 81%</p> <p>Volume % of CO₂ = 11%</p> <p>Volume % of O₂ = 8%</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>2</p> <p>2</p>
4-b	<p>Basis: Mixture of N₂, CO₂</p> $X_{CO_2} = 1 - X_{N_2}$ <p>Avg. mol.wt = M₁X₁ + M₂X₂</p> $31 = 28 X_{N_2} + 44(1 - X_{N_2})$ <p>Solving, X_{N₂} = 0.8125</p> $X_{CO_2} = 0.1875$ <p>Partial pressure of N₂ = Total pressure * mol.fr</p> $= 101.325 * 0.8125$ $= \mathbf{82.33 \text{ KPa}}$	<p>1</p> <p>1</p> <p>1</p> <p>2</p> <p>1</p> <p>2</p>
4-c	<p>Basis: 100 kg flaked seeds</p> <pre> graph LR Hexane --> Extractor Seeds[seeds] --> Extractor Extractor --> HexaneOil[hexane+oil] Extractor --> Cake[cake Y kg 0.8% oil, 87.7% solid] </pre> <p>18.6% oil, 69% solid</p>	<p>1</p> <p>1</p>



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	Component balance for solids $69 = 0.877 Y$ Or $Y = 78.68 \text{ Kg}$ Oil in cake = $(0.8/100)78.68$ $= 0.63 \text{ Kg}$ Oil balance is $18.6 = 0.63 + \text{oil recovered}$ Oil recovered = 17.97 kg % oil recovery = $(17.97/18.6) * 100 = 96.61\%$	1 1 1 1 1 1
5	Any 2	16
5-a	Basis : 100 Kg of coke Amount of carbon in coke = $0.9 \times 100 = 90 \text{ Kg}$ Amount of C = $90/12 = 7.5 \text{ katom}$ Reaction : $C + O_2 \rightarrow CO_2$ From reaction , 1 katom C = 1 kmol O_2 12 Kg C = 32 Kg O_2 90 Kg C = $(32/12) \times 90 \text{ Kg } O_2$ O_2 theoretically required = $(32/12) \times 90 = 240 \text{ Kg}$ O_2 theoretically required = $(240 \times 1000) / 32 = 7500 \text{ mol}$ Moles of O_2 theoretically required for 100 kg coke for complete combustion = 750 mol ----- ans (a) O_2 theoretically required = $240 / 32 = 7.5 \text{ kmol}$ Air theoretically required = $7.5 * (100/21) = 35.71 \text{ kmol}$ % excess	1 1 1



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	<p>Air actually supplied = Air theoretically required ($1 + \frac{50}{100}$)</p> <p>Air actually supplied = $35.71 \times (1 + \frac{50}{100}) = 53.57$ kmol</p> <p>Air actually supplied = 53.57 kmol</p> <p>O₂ in air supplied = $53.57 \times 0.21 = 11.25$ kmol</p> <p>N₂ in air supplied = $53.57 \times 0.79 = 42.32$ kmol</p> <p>O₂ reacted for complete combustion of coke = 7.5 kmol</p> <p>Material Balance of O₂</p> <p>O₂ in air supplied = O₂ reacted + O₂ Unreacted</p> <p>O₂ Unreacted = $11.25 - 7.5 = 3.75$ kmol</p> <p>CO₂ produced = $(1/1) \times 7.5 = 7.5$ kmol</p> <p>Analysis of Gases at the end of Combustion:</p> <table border="1" data-bbox="282 1098 1141 1381"> <thead> <tr> <th>Component</th> <th>Quantity , Kmol</th> <th>Mole%</th> </tr> </thead> <tbody> <tr> <td>CO₂</td> <td>7.5</td> <td>14</td> </tr> <tr> <td>N₂</td> <td>42.32</td> <td>79</td> </tr> <tr> <td>O₂</td> <td>3.75</td> <td>07</td> </tr> <tr> <td>Total</td> <td>53.57</td> <td>100</td> </tr> </tbody> </table>	Component	Quantity , Kmol	Mole%	CO ₂	7.5	14	N ₂	42.32	79	O ₂	3.75	07	Total	53.57	100	<p>1</p> <p>1</p> <p>1</p> <p>2</p>
Component	Quantity , Kmol	Mole%															
CO ₂	7.5	14															
N ₂	42.32	79															
O ₂	3.75	07															
Total	53.57	100															
<p>5-b</p>	<p>Basis : 15000 mol/h of N₂- H₂ mixture</p> <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>$C_P^{\circ} \text{ mix} = \sum C_P^{\circ} \text{ mix} \cdot X_i = X_{N_2} \cdot C_P^{\circ} N_2 + X_{H_2} \cdot C_P^{\circ} H_2$</p> <p>$= 0.25 (29.5909 - 5.141 \times 10^{-3} T + 13.1829 \times 10^{-6} T^2 - 4.968 \times 10^{-9} T^3)$</p> <p>$+ 0.75 (28.6105 + 1.0194 \times 10^{-3} T - 0.1476 \times 10^{-6} T^2 + 0.769 \times 10^{-9} T^3)$</p>	<p>1</p> <p>2</p>															



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	$= 28.8556 - 0.5207 \times 10^{-3} T + 3.185 \times 10^{-6} T^2 - 0.6652 \times 10^{-9} T^3$ <p>Q = Heat added</p> $Q = n \int_{T_1}^{T_2} C_p^{mix} dT$ $Q = n \int_{T_1}^{T_2} [28.8556 - 0.5207 \times 10^{-3} T + 3.185 \times 10^{-6} T^2 - 0.6652 \times 10^{-9} T^3] dT$ $= n \left[28.8556 (T_2 - T_1) - \frac{0.5207 \times 10^{-3}}{2} (T_2^2 - T_1^2) + \frac{3.185 \times 10^{-6}}{3} (T_2^3 - T_1^3) - \frac{0.6652 \times 10^{-9}}{4} (T_2^4 - T_1^4) \right]$ <p>Where , n= 15 Kmol/h , T₂ = 473 K , T₁ = 298 K</p> $= 15 \left[28.8556 (473 - 298) - \frac{0.5207 \times 10^{-3}}{2} (473^2 - 298^2) + \frac{3.185 \times 10^{-6}}{3} (473^3 - 298^3) - \frac{0.6652 \times 10^{-9}}{4} (473^4 - 298^4) \right]$ <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>	<p>1</p> <p>2</p> <p>2</p>
5-c	<p>Basis : 1 Kg of petrol</p> <p>Amount of H₂ in petrol = 0.15 Kg, Amount of C in petrol = 0.85 kg</p>	



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$H_2 + \frac{1}{2} O_2 \rightarrow H_2O$	1
$C + O_2 \rightarrow CO_2$	
From reactions : 1 kmol of $H_2 = 0.5$ kmol O_2	
2 kg of $H_2 = 16$ kg O_2	
1 katom of C = 1 kmol O_2	
12 kg of C = 32 kg O_2	1
Theretical requirment of O_2 for $H_2 = 0.15 \times (16/2) = 1.20$ Kg	
Theretical requirment of O_2 for C = $0.85 \times (32/12) = 2.27$ Kg	
Total theretical requirment of $O_2 = 1.20 + 2.27 = 3.47$ Kg	
Amount of air required for combustion = $(3.47/0.23) = 15.09$ kg --- ans	1
(Air contais 23% O_2 and 77 % N_2 on weight basis)	
Amount of air supplied = $15.09 \times 1.15 = 17.35$ kg	
N_2 in supplied air = $(0.77 \times 17.35) / 28 = 0.477$ kmol	
O_2 in supplied air = $(0.23 \times 17.35) / 32 = 0.125$ kmol	1
O_2 in dry product = $[0.23 \times (17.35 - 15.09)] / 32 = 0.016$ kmol	
We have , 1 katom of C = 1 kmol CO_2	
$(0.85/12)$ katom C = ?	1
CO_2 produced = $(1/1) \times (0.85/12) = 0.071$ kmol	
The product flue gases contain CO_2 , O_2 and N_2	1
For ideal gases, Volume % = Mole %	
Composition of Dry Product Gases :	



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	<table border="1"> <thead> <tr> <th>Component</th> <th>Quantity , Kmol</th> <th>Mole%</th> </tr> </thead> <tbody> <tr> <td>CO₂</td> <td>0.071</td> <td>12.6</td> </tr> <tr> <td>N₂</td> <td>0.016</td> <td>2.8</td> </tr> <tr> <td>O₂</td> <td>0.477</td> <td>84.6</td> </tr> <tr> <td>Total</td> <td>0.564</td> <td>100</td> </tr> </tbody> </table>	Component	Quantity , Kmol	Mole%	CO ₂	0.071	12.6	N ₂	0.016	2.8	O ₂	0.477	84.6	Total	0.564	100	2
Component	Quantity , Kmol	Mole%															
CO ₂	0.071	12.6															
N ₂	0.016	2.8															
O ₂	0.477	84.6															
Total	0.564	100															
6	Any 2	16															
6-a	<p>Solve any TWO of the following</p> <p>Basis: 1000 kg of Desired mixed acid Waste acid, 55 % H₂SO₄, 20% HNO₃</p> <p>Con.nitric acid 90% HNO₃</p> <p>Con.sulphuric acid 95% H₂SO₄</p> <p>Desired mixed acid 1000 kg 60% H₂SO₄, 26% HNO₃</p> <p>Let x,y and Z be the kg of waste acid ,concentrated sulphuric acid and concentrated nitric acid to make 1000 kg desired acid.</p> <p>Overall material Balance : X+ Y+ Z = 1000 ----- (1)</p> <p>Material Balance Of H₂SO₄ 0.55X+ 0.95 Y = 0.60 x 1000 0.55X+ 0.95 Y = 600 600 – 0.55 X Y= -----</p>	1 1 1 1															



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	<p>0.95</p> $Y = 631.58 - 0.58 X \text{ ----- (2)}$ <p>Material Balance Of HNO₃</p> $0.2X + 0.90 Z = 0.26 \times 1000$ $0.2X + 0.95Y = 260$ $260 - 0.2 X$ $Z = \text{-----}$ <p>0.90</p> $Z = 288.9 - 0.222 X \text{ ----- (3)}$ <p>Putting values of Y and Z in equation (1) from (2) and (3) and solve for X</p> $X + (631.58 - 0.58 X) + (288.9 - 0.222 X) = 1000$ $0.198 X = 79.52$ $X = 401.6 \text{ kg}$ <p>We have $Y = 631.58 - 0.58 X$</p> $Y = 631.58 - 0.58 \times 401.6 = 398.65 \text{ Kg}$ <p>We have $Z = 288.9 - 0.222 X$</p> $Z = 288.9 - 0.222 \times 401.6 = 199.75 \text{ Kg}$ <p>Waste acid = 401.6 kg</p> <p>Concentrated sulphuric acid = 398.65 kg</p> <p>Concentrated nitric acid = 199.75 kg</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
6-b	<p>Basis: 1 Km³ /h benzene fed to reactor</p> <p>Benzene, HNO₃, H₂SO₄ → Reactor</p> <p>----→ Product stream NB, DNB, HNO₃ etc</p>	



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<p style="text-align: center;">H_2SO_4</p> <p>Reaction 1: $\text{C}_6\text{H}_6 + \text{HNO}_3 \xrightarrow{\text{H}_2\text{SO}_4} \text{C}_6\text{H}_5\text{NO}_2 + \text{H}_2\text{O}$</p>	1
<p>Reaction 2: $\text{C}_6\text{H}_5\text{NO}_2 + \text{HNO}_3 \xrightarrow{\text{H}_2\text{SO}_4} \text{C}_6\text{H}_4(\text{NO}_2)_2 + \text{H}_2\text{O}$</p> <p>Benzene reacted = $0.9 \times 1 = 0.9 \text{ kmol/h}$</p> <p>Let X be the dinitrobenzene formed per hour</p> <p>$(\text{kmol nitrobenzene /h}) / (\text{kmol dinitrobenzene /h}) = 17/1$</p> <p>Nitrobenzene produced = $17 \times \text{kmol}$</p> <p>From reaction 1 and 2</p> <p>1 kmol of $\text{C}_6\text{H}_6 = 1 \text{ kmol of } \text{C}_6\text{H}_5\text{NO}_2$</p> <p style="padding-left: 40px;">$= 1 \text{ kmol of } \text{C}_6\text{H}_4(\text{NO}_2)_2$</p> <p>Benzene reacted to produce nitrobenzene = $(1/1) \times 17 \times \text{kmol/hr}$</p> <p>Benzene reacted to dinitrobenzene = $x \text{ kmol}$</p> <p>Benzene reacted = $0.9 = 17x + x$</p> <p style="padding-left: 40px;">$x = 0.05 \text{ kmol/hr}$</p> <p>Nitrobenzene produce = $17(0.05) = 0.85 \text{ kmol/h}$</p> <p>1 kmol of $\text{C}_6\text{H}_6 = 1 \text{ kmol HNO}_3$</p> <p>Stoichiometric requirement of nitric acid for</p> <p>1 kmol benzene /h = $1/1 \times 1 = 1 \text{ kmol/h}$</p>	1
<p>Nitric acid fed to reactor = $1 [1 + (65/100)] = 1.65 \text{ kmol/h}$</p> <p>Nitric acid fed to reactor = $1.65 \times 63 = 104.95 \text{ kg/h}$</p> <p>Nitrobenzene produce = $17(0.05) = 0.85 \times 123 = 104.55 \text{ kg/h}$</p> <p>Benzene fed to reactor = $1 \times 78 = 78 \text{ kg/hr}$</p>	1
<p>Nitric acid fed to reactor fed to produce of 2000 kg/hr of nitrobenzene =</p> <p>$(103.95/ 104.55) \times 2000 = \mathbf{1988.5 \text{ kg/hr}}$</p>	1



WINTER-16 EXAMINATION
Model Answer

Subject code :

17315

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	<p>Water in wet solids fed to first dryer = $0.5 * 1000 = 500 \text{ Kg}$</p> <p>% of original water removed in first dryer</p> <p style="text-align: center;">$= (375/500) * 100 = 75 \%$</p> <p>% of original water removed in Second dryer</p> <p style="text-align: center;">$= (114.8/500) * 100 = 22.96 \%$</p> <p>% of original water removed in first dryer = 75 %</p> <p>% of original water removed in Second dryer = 22.96 %</p> <p style="text-align: right;">----- ans. (a)</p>	<p>1</p> <p>2</p>
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