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### WINTER-15 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	Total marks
1-A	Any 4		8
1A-a	Ideal Gas law:	1	2
	PV=nRT where P - pressure, V - volume, n- moles, K-absolute temperature	1	
	and R – universal gas constant		
1A-b	Principle involved in solving material balance problems without chemical	2	2
	reaction:		
	Law of conservation of mass: It states that		
	For any process input= output+accumulation		
1A-c	Amagat's law: It states that the total volume exerted by a gas mixture is equal	1	2
	to the sum of pure component volumes		
	Mathematical Statement: $V = V_1 + V_2 + V_3$	1	
	where V is the total volume of gas mixture , $V_1, V_2, V_3$ are pure component		
	volumes		
1A-d	Raoult's law: Raoult's law states that equilibrium partial pressure of a	2	2
	constituent at a given temperature is equal to the product of its vapour pressure		
	in pure state and its mol fraction in the liquid phase.		
	$P_{A} = P_{A}^{0} X_{A}$		
1 A-e	Adiabatic reaction:	1	2
	It is the reaction which proceeds without loss or gain of heat.		
	Adiabatic reaction temperature: Temperature of product under adiabatic	1	
	condition is called adiabatic reaction temperature.		
1A-f	% Conversion: It is the ratio of amount of limiting reactant reacted to the	1	2
	amount of limiting reactant totally charged. Express in percentage.		
	%Yield of desired product = (moles of limiting component reacted to form	1	



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	desired product/ total moles of limiting component reacted)* 100		
1-B	Any 2		12
1B-a	Basis: 100 kmoles of air	1	4
	Average mol.wt of air= 0.79*28+0.21*32= 28.84	1	
	Density of air = $P*Mav/RT$	1	
	At NTP, P= 101.325 KPa, R= 8.314, T=273 K		
	Density = $1.29 \text{ kg/m}^3$	1	
1B-b	Partial pressure of CO <sub>2</sub> in gas phase P <sub>CO2</sub> = H * x <sub>CO2</sub>	2	4
	$=7*10^6*4*10^{-6}$		
	= <b>28 KPa</b>	2	
1B-c	Basis: 0.8m <sup>3</sup> fixed mass of gas at constant temperature	1	4
	$P_1 = 1$ $V_1 = 0.8 \text{ m}^3$ $T_1 = T = T_2$		
	$P_2 = 1.5  V_2 = ? \text{ m}^3$	1	
	$P_1V_1/T_1 = P_2V_2/T_2$	1	
	$1*0.8/T = 1.5*V_2/T$		
	Or $V_2 = 0.533 \text{ m}^3$	1	
2	Any 4		16
2-a	Steps involved in solving material balance calculations:	½ mark	4
	1. Assume suitable basis of calculation as given in problem.	each	
	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 algebric equations		



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	8. Get values of two unknown quantities.		
2-b	Basis: 1 kg air	1	4
	NH3 gas,0.004 Kg NH3  1 Kg air 0.2 kg NH3  NH3 absorbed X Kg NH3	1	
	NH3 balance is 0.2= 0.004+ X X= 0.196 Kg	1	
	% recovery of ammonia = (NH3 absorbed/ NH3 original)*100		
	= (0.196/0.2)*100	1	
	= 98%		
2-c	<b>Excess component:</b> It is the reactant which is in excess of the theoretical or	2	4
	stoichiometric requirement.		
	Limiting component: It is the reactant which would disappear first if a rection	2	
	goes to completion.Or it is the reactant which decides the extent of a reaction.		
2-d	Basis: 100 of ethanol charged.	1	4
	$C_2H_5OH \rightarrow CH_3CHO + H_2$		
	I kmol C <sub>2</sub> H <sub>5</sub> OH reacted = 1kmol CH <sub>3</sub> CHO formed	1	
	C <sub>2</sub> H <sub>5</sub> OH reacted to produce 45 kmol CH <sub>3</sub> CHO = 45 kmol		
	Conversion of $C_2H_5OH = (kmole of C_2H_5OH reacted/kmoles)$	1	
	Of C <sub>2</sub> H <sub>5</sub> OH fed)*100		



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	=( 45/100)*100	1	
	% conversion of ethanol = 45 %		
2-e	Basis – 100Kmol SO <sub>2</sub> charged	1	4
	O <sub>2</sub> fed=75 kmoles		
	Reaction		
	$SO_2 + \frac{1}{2}O_2 = SO_3$		
	Theoretical requirement of O <sub>2</sub>		
	1 Kmol $SO_2 \equiv 0.5$ Kmol $O_2$ theoretical		
	$100 \text{ kmol } SO_2 = \frac{0.5}{1} \times 100$	1	
	Theoretical $O_2 = 50 \text{ Kmol}$		
	∴% excess of O <sub>2</sub> used		
	$= \frac{O_2 \text{ in supplied} - O_2 \text{ theo read}}{O_2 \text{theo read}}$	1	
	$=\frac{75-50}{50}\times 100$		
	=50		
	∴ % excess air used = 50%	1	
2-f	Basis: 100 Kg/hr C <sub>2</sub> H <sub>5</sub> OH		4
	m=100 Kg/hr	1	
	$\lambda$ = 202 kcal/kg		
	Latent heat = $m. \lambda$	2	
	= 100 *202		
	= 20200 Kcal / hr	1	
3	Any 2		16
3-a	SOLUTION:		8



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			_
	<b>BASIS</b> : 100 mol of ethylene		
	Reaction I		
	$C_2H_4 + \frac{1}{2}O_2 \longrightarrow C_2H_4O$		
	Reaction II	2	
	C2H4 + 3O2 → 2CO2 + 2H2O		
	From reaction I,		
	1Kmol of C2H4O formed = 1Kmol C2H4 reacted		
	∴ C2H4 reacted to form 80 kmol C2H4O		
	$=\frac{1}{1} \times 80 = 80 \text{Kmol}$	1	
	From reaction II,		
	2kmol of CO2 formed = 1Kmol C2H4 reacted		
	∴ C2H4 reacted to form 10 kmol CO2		
	$=\frac{1}{2} \times 10 = 5 \text{Kmol}$	1	
	$\therefore$ C2H4 totally reacted = 80 $\div$ 5= 85		
	$\therefore \% \text{ conversion of C2H4} = \frac{85}{100} \times 100$	2	
	= 85%		
	% yield of C2H4O = $\frac{80}{85}$ x100		
	= <b>94.12%</b>	2	
3-b	SOLUTION:		8
	BASIS: 100 kg of mixed acid		
	$\therefore H_2SO_4 = 45 \text{ Kg}$		
	$HNO_3 = 42 \text{ Kg}$		



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H <sub>2</sub> SO <sub>4</sub> 98 %  HNO <sub>3</sub> =42 , H <sub>2</sub> SO <sub>4</sub> =45%,H <sub>2</sub> O =39%	2	
$HNO_3$		
Let, X be the kg of H <sub>2</sub> SO <sub>4</sub> and		
Y be the kg of HNO <sub>3</sub>		
x + y = 100		
Material Balance on H <sub>2</sub> SO <sub>4</sub> :-	1	
0.98x = 45		
$\therefore x = 45/0.98$		
x = 45.91		
$\therefore$ weight of H <sub>2</sub> SO <sub>4</sub> = 45.91 kg		
: weight of $HNO_3 = 100-45.91 = 54.09 \text{ kg}$		
Let , y is strength of HNO <sub>3</sub>	2	
∴ 54.09 y = 42		
y = 42/54.09	1	
y = 0.7764		
$\therefore$ strength of HNO <sub>3</sub> = 0.7764 x 100		
= 77.64 %Ans.	1	
Weight ratio of $H_2SO_4$ 45.91		
Weight ratio of $HNO_3$ 54.09		



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	= 0.8487Ans.	1	
3-с	SOLUTION:		8
	<b>BASIS</b> : 20,000 kg /h of 25 % methanol feed solution to coloumn.		
	Distillate (d)		
	98 % methanol	2	
	20000 kg/h Feed  Distillation Coloumn		
	25 % methanol —		
	Waste solution (w)		
	1 % methanol		
	(bottom product)		
	Block diagram for distillation of 25 % methanol feed		
	Let x and y be the mass flow rates of distillate / product and waste solution		
	(bottom product) respectively.		
		1	
	Overall Material Balance :		
	x + y = 20000(i)		
	Material Balance of Methanol :	1	
	0.98  x + 0.01  y = 0.25  x 20000		
	0.98  x + 0.01  y = 5000(ii)		
	From equation(i),		



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	y = 20000 - x(iii)		T
	y = 20000 - x(iii)		
	Put the value of y from equation (iii) in eqn (ii) and solve for x.		
	0.98 x + 0.01 (20000 - x) = 5000		
	0.98  x - 0.01  x = 5000 - 200	2	
	0.97  x = 4800		
	x = 4948.45  kg/h		
	y = 20000 - x = 20000 - 4948.45 = 15051.55  kg/hr		
	Mass flow rate of distillate = <b>4948.45 kg/h</b>	1	
	Mass flow rate of waste solution = 15051.55 kg/hrAns.		
	Methanol in waste solution = $0.01 \times 15051.55 = 150.51 \text{ kg/h}$		
	Methanol in Feed = $0.25 \times 20000 = 5000 \text{ kg/h}$		
	% loss of methanol = $\frac{\text{Methanol in waste solution}}{\text{Methanol feed solution}} \times 100$		
	$=\frac{150.51}{5000} \times 100$	1	
	= 3.01 %Ans		
4	Any 2		16
4-a	SOLUTION:		8
	<b>Basis:</b> 1 mol of C4H6 (liq)		
	1. $C(s) + O2(g)> CO2(g)$ $\Delta H1 = -393.51 \text{ KJ/mol}$	1	
	2. H2 (g) +1/2 O2(g)> H2O(l) $\Delta$ H1 = - 285.83 KJ/mol	1	



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	3. C4H6 (liq.) +5.5 O2(g)> 4CO2(g)+3 H2O(l)	1	
	$\Delta H0c = -2520.11 \text{ kJ/mol}$		
	4. $4C(s) + 3 H2 (g)> C4H6 (liq.) \Delta H^0 f = ?$	1	
	$\Delta H^0 f$ = Standard heat of formation of gaseous ethyl alcohol at 298.15 K		
	Reaction(4) = $4*$ Reaction (1) + $3*$ Reaction (2) – Reaction (3)	1	
	$\Delta H^0 f = 4*\Delta H1 + 3*\Delta H2 - \Delta H^0 c$		
	= 4(-393.51) + 3*(-285.83) - (-2520.11)	2	
	= (-1547.04) + (-857.49) + 1410.09		
	= 88.58 KJ/mol	1	
	$\Delta H0f = 88.58 \text{ KJ/mol} Ans.$		
4-b	SOLUTION:		
	<b>BASIS</b> : 1000 kg of wet solid	1	
	1000 kg 15 % H <sub>2</sub> O Dried Product 99.5 % solid	1	
	Weight of solid in feed = 1000*0.85		
	= 850  kg		
	Weight of water in feed = 150 kg	1	
	Let, weight of dried producy = $x kg$		



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J			5
	It contain 99.5 % solid and solid is unchanging component.		
	∴ solid balance,	1	
	0.995  x = 850		
	x = 850/0.995		
	x = 854.27  kg		
		1	
	∴ weight of water in dried product = $854.27 - 850 = 4.27 \text{ kg}$		
	Water removed in dryer = $150 - 4.27 = 145.73 \text{ kg}$	1	
	$\therefore$ original water removed = $\frac{145.73}{150}$ x 100	1	
	= 97.15 %Ans.	1	
4-c	SOLUTION:	1	8
	<b>BASIS</b> : 100 kmol of SO <sub>2</sub>		
	$SO_2 + \frac{1}{2}O_2 \longrightarrow SO_3$	1	
	∴ 1 kmol of $SO_2 \equiv \frac{1}{2}$ kmol $O_2$		
	$\therefore$ kmol O <sub>2</sub> required = 100 * 0.5		
	= 50 kmol	1	
	$\therefore$ O <sub>2</sub> supplied = 50*1.8		
	= 90 kmol	1	
	$N_2 \text{ supplied} = \frac{79}{21} \times 90$		
	= 338.57 kmol		
	$SO_2$ reacted = $100 \times 0.70$	1	
	= 70 kmol		
	$O_2$ required = 70 x 0.5	1	
	= 35 kmol		
	SO <sub>3</sub> formed = 70 kmol		
	Composition of gases leaving reactor :-	1	



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	Product	Quantity,Kmo	Mole %		
		1		1	
	$SO_2$	100 - 70 = 30	6.07		
	$O_2$	90 - 35 = 55	11.14		
	$N_2$	338.57	68.6		
	SO <sub>3</sub>	70	14.18		
	Total	493.57			
5	Any 2	•			16
5-a	Basis: 100 kg gas				8
	Avg mol wt of gas =				
	Total moles= 100/39.	8 = 2.512 kmoles			
	Kmoles of ethane= 1.	6328		1	
	Kmoles of butane= 0				
	Reactions are:				
	$C_2H_6+7/2 O_2 \longrightarrow$	$2\text{CO}_2 + 3\text{H}_2\text{O}$		1	
	$C_4H_{10}+ 13/2 O_2$	$4\text{CO}_2 + 5_2\text{O}$			
	Oxygen theoretically	required = 7/2*1.16	328 + 13/2 * 0.879	1	
		= 11.42			
	$O_2$ fed = 11.42 + 0.2*11.42 = 13.704 kmoles				
	$O_2$ unreacted = 13.70	4 – 11.42= 2.284km	oles	1	
	$N_2$ fed= 51.54 kmoles	S		1	
	$CO_2$ formed = 3.27 +	3.516 = 6.786  kmol	es	1	
	$H_2O$ formed = 4.898	+4.395 = 9.293  km	oles	1	
	Weight of gas $= 2.28$	4* 32 + 51.54*28+ 6	6.786*44+9.293*18	3	
	= 198	1.87 Kg		1	



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5-b Ca	ase –I: Basis: 100 Kg/hr o	f solid handling capaci	ty of the evaporator.		8
		→ Water evap	porated		
		<u> </u>	1		
	Weak liquor →	Evaporator		1	
	5% solids				
		↓ → Thick liqu	or 50% solids		
		7 Thick hqu	01,5070 301143		
Le	et X be the be the Kg/hr of	of weak liquor then we	have.		
	0.05  X = 100				
	X = 2000	Kg/hr		1	
	Y be the be the Kg/hr	of thick liquor then we	e have.	1	
	0.5  Y = 100				
	Y = 200  kg/hr				
0	Overall Material Balance:				
Kg	g/hr of weak liquor = Kg	/hr of thick liquor + K	g/hr of		
		Wat	ter evaporated		
	2000 = 20	00 + Kg/hr of Water e	vaporated	1	
Kg	g/hr of Water evaporated =	= 1800 Kg/hr			
Ca	ase –II: Basis: 1800 Kg/hr	of Water evaporated			
	Let A be the be the Kg/	hr of weak liquor			
	B be the be the Kg/h	nr of thick liquor			
Ov	Overall Material Balance:				
	A = B + 1	800		1	
Ma	Material Balance of Solids:				
	0.04 A = 0.35				
Pu	A = 8.75 atting in above equation	В		1	
				-	



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,000 000	ic .(17313)		age 14 01 17
	8.75 B = B + 1800		
	B = 232.26  Kg/hr		
	A = 232.26 + 1800 = 2032.26  Kg/hr		
	Solid in weak liquor = 0.04 X 2032.26 = 81.3 Kg/hr	2	
	Solid handling Capacity = 81.3 Kg/hr <b>Ans.</b>		
5-c	Basis: 1 mol of ethylene gas		8
	$C_2H_4(g) + \frac{1}{2}O_2 \longrightarrow C_2H_4O(g)$		
	$\Delta H_{R}^{o} = Standard heat of reaction$		
	= $[\Sigma \Delta H^{o}_{c}]$ product - $[\Sigma \Delta H^{o}_{c}]$ reactant	2	
	$= [1 \times \Delta H^{o}_{c}) C_{2}H_{4}O(g) ] - [1 \times \Delta H^{o}_{c} C_{2}H_{4}(g) +$		
	$ \begin{array}{c} 1 \times \Delta H^{o}_{c} O_{2} \\ = [1 \times (-12.58)] - [1 \times (12.50) + 1/2 \times (0.0)] \end{array} $		
	= -12.58 - 12.50	3	
	= $-25.08$ KJ per mol $C_2H_4O(g)$ produced.		
	Change in the enthalpy for 5 mol C <sub>2</sub> H <sub>4</sub> O(g) produced		
	= -25.08 X 5 = <b>-125.40 KJAns</b>	3	
6	Any 4		16
6-a	1)Stoichiometric Equation :		4
	The stoichiometric equation of a chemical reaction is the statement indicating	2	
	relative moles of reactant and products that take part in the reaction .		
	For example, the stoichiometric equation		
	$CO + H_2 \rightarrow CH_3OH$		
	Indicates that one molecule of CO react with two molecules of hydrogen to		
	indicates that one molecule of CO feact with two molecules of hydrogen to		



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	42.19		
	% recovery of Oil = * 100		
	45		
		1	
	% recovery of Oil = 93.75% ans.		
6-c	Basis:1000 Kg of desired acid		4
	Basis: 1000 kg of desired acid.		
	Dilute acid  Mixing  Desired acid, 1000 kg 65% H <sub>2</sub> SO <sub>4</sub>		
	Commercial Grade acid	1	
	98% H <sub>2</sub> SO <sub>4</sub>		
	and a seid		
	Let X and y be the kg of dilute acid and Commercial Grade required to		
	prepare desired acid.		
	propure desired detail.		
	Overall Material balance		
	Kg dilute acid + kg Commercial Grade = Kg of desired acid		
	X + Y = 1000(1)		
	Material Balance of H <sub>2</sub> SO <sub>4</sub>		
	Francisco Dalunce of 11/2004		
	$H_2SO_4$ in dilute acid + $H_2SO_4$ in Commercial Grade = $H_2SO_4$ in desired acid		
	0.25  X + 0.98  Y = 0.65 * 1000		



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3			. ago 17 0. 17
	0.25X + 0.98Y = 650		
	650 - 0.25 X		
	Y =	1	
	0.98		
	Y=663.23 - 0.255 X put in equation (1)		
	X + (663.26 - 0.255 X) = 1000		
	X=452  kg	1	
	We have,		
	X + Y = 1000		
	452 + Y = 1000		
	Y = 548  Kg		
	Dilute acid required = 452 Kg	1	
	Commercial Grade Acid required =548 kg		
6-d	Basis: 50 kmol of butane fed to combustion reactor		4
	Reaction,		
	$C_4H_{10} + 13/2 O_2 \rightarrow 4CO_2 + 5H_2O$	1	
	From reaction,		
	1 Kmol of C <sub>4</sub> H <sub>10</sub> =6.5 kmol O <sub>2</sub>		



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Feed contains 60 kmol A, 30 kmol B and 10 kmol inerts

Let X be the kmol of A reacted by reaction:

From reaction 2 kmol A = 1 kmol B = 1 kmol C

B reacted = 
$$(1/2)^* X = 0.5 X$$
 kmol

C formed = 
$$(1/2)^* X = 0.5 X$$
 kmol

Material Balance of A give

A unreacted = 
$$(60 - X)$$
 kmol

**Material Balance of Inerts:** 

Inerts in feed = Inert in product = 10 kmol

C formed = 
$$(1/2)* X = 0.5 X$$
 kmol

B unreacted = 
$$(30 - 0.5 \text{ X}) \text{ kmol}$$

Total moles of product stream = (60-X) + (30-0.5X) + 10=0.5X

$$= 100 - X \text{ Kmol}$$

Mole % of A in product stream = 2%

Kmol A in product stream

Mole % of A = ----- \* 100

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Total kmol of product stream		
60 - X		
2 = * 100		
100 – X		
	1	
X = 59.184  kmol = amount of A r	eacted	
Kmol A reacted		
Conversion of A =	* 100	
Total kmol of A feed		
59.184		
Conversion of A = * 100 = <b>98.64</b> %	<b>Ans</b>	
60		
6-f Sensible Heat :		4
Sensible heat is the heat that must be trans	sferred to raise or lower the 2	
temperature of a substance or mixture of subst	ance.	
Latent Heat: When matter undergoes a phase	e change, the enthalpy change 2	
associated with unit amount of matter at const	ant temperature and pressure is	
known as Latent Heat of phase change.		