GUJARAT TECHNOLOGICAL UNIVERSITY BE - SEMESTER-III (New) EXAMINATION – WINTER 2015

Subject Code: 2133606 Subject Name: Material and Energy Balance Calculation Time: 2:30pm to 5:00pm Instructions:				Date:29/12/2015 Total Marks: 70		
2 3	 Attempt all questions. Make suitable assumptions wherever necessary. Figures to the right indicate full marks. Atomic weight: H-1,C-12,N-14,O-16,S-32,Cl-25.5, K-39, 					
				MARKS		
Q.1	1	Short Questions The rate of material system.	is zero in case of a steady state	e 1		
		(a) accumulation	(b) production			
	-	(c) input	(d) generation			
	2	A 'limiting reactant' is the one, the chemical reaction.	which decides the i	n 1		
		(a) equilibrium constant	(b) conversion			
		(c) rate constant	(d) none of these			
	3		re is less than its equilibrium vap	our 1		
	e	pressure is called the		-		
		(a) saturated	(b) superheated			
		(c) unsaturated	(d) dry gaseous			
	4		een conducted in a reactor as sho			
		can be written, are	ries around which material balance	e		
		A				
		B	L			
		_				
		(a) 1	(b) 6			
		(c) 3	(d) 4			
	5	Which of the following gravity	v scales is used exclusively for	1		
		liquids heavier than water? (a) Baumme scale	(b) Twaddel scale			
		(c) API scale	(d) none of these			
	6		f a compound containing 50% of	1		
	v	(a) AB ₃	(b) A_2B_3	*		
		(c) A_2B	(d) AB_2			
	7	Real gases approach ideal beha		1		
		(a) high pressure & high	(b) low pressure & hig	gh		
		Temptemperature.	temperature.			

8 Internal energy of a substance comprises of the			(c) high pressure & low temperature.	(d) low pressure & low temperature.	
(a) vibrational (b) rotational (c) translational (c) translational (d) all (a), (b) & (c) 9 The net heat evolved or absorbed in a chemical process, i.e. total change in the enthalpy of the system is independent of the (a) temperature & pressure. (b) number of intermediate chemical reactions (c) state of aggregation & the (d) none of these. state of combination at the beginning & the end of the reaction. 10 The heat change for the reaction, $C(s) + 2S(s) \rightarrow CS_2(l)$, is 104.2 kJ. It represents the heat of (a) formation (d) fusion 11 1 Kcal/kg. °C is equivalent to BTU/lb. °F. 1 (a) 1 (b) 2.42 (c) (4.97) (d) None of these 12 (1) Which of the following is the Claussius-Clayperon 1 (a) $PV = RT + BV + y/V^2 +$ (b) $(P + aV^2)(V-b) = RT$ $\log_{\theta} \frac{\rho}{\rho_0} = \frac{R}{R} (\frac{1}{T_0} - \frac{1}{T})$ (d) $P = \frac{RT}{V - b} - \frac{3}{TV^2}$ (c) 13 At standard conditions, 1 $N_2 + 2O_2 \rightleftharpoons 2NO_2; \Delta H^\circ = 100 \text{ kJ/mole}$ (a) 15 (b) 30 (c) 85 (d) 170 14 Heat of reaction is not influenced by (a) 15 (b) 30 (c) whether the reaction is (d) none of these. (c) whether the reaction is (d) none of these. (d) No efficiently conversion, (2) yield, (3) selectivity (f) A weight of 1.10 kg of Carbon dioxide occupie		8	Internal energy of a substance comprise	-	1
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and $b = 4.3 \times 10^{-2} \text{ m}^3/\text{kmol}$				Pal/(kmol) ²	
J/(gmol K) and T is in K. Convert that Cp is in cal/(gmol 0 F) with		(c)	The heat capacity of sulphur is $Cp = 15$		07

T in ⁰F.

OR

(c) The Antoine equation predicts the effect of temperature on vapour 07 pressure by the relation:

$$\ln P^s = A - \frac{B}{T - C}$$

Q.3	(a) (b)	Where, A, B and C are Antoine constant s. For water, the values of the constants where the vapour pressure is in kPa and temperature is in K are found to be A = 16.26205, B = 3799.887 and C = 46.854. What are these values when Vapour pressure is in mm Hg and temperature in 0 C. Define (1) dew point (2) relative humidity (3) humid heat A soap plant produced raw soap containing 50 % moisture. This is to be dried 20 % moisture before it is pressed into cakes for sale. How many 100 g soap piece can be obtained from 1000 kg of original raw soap ?	03 04
	(c)	In the BASF oil quench process to manufacture acetylene; pure oxygen and pure methane are fed to the acetylene burner. The cracked gas from the burner has the following composition: H ₂ : 56.5 %, CH ₄ : 5.2 %, C ₂ H ₄ : 0.3 %, C ₂ H ₂ : 7.5 %, C ₃ H ₆ : 0.5 %, CO: 25.8 %, CO ₂ : 4.0 %, and O ₂ :0.2 % (mole % on dry basis). Assume that formation of the other compounds, such as ammonia, is negligible. For 100 kmol cracked gas, calculate (a) methane requirement (b) oxygen requirement (c) production of water OR	07
Q.3	(a) (b)	Draw neat sketch of Recycle, bypass and purge operations A saturated solution containing 1500 kg of potassium chloride at 360 K is cooled in an open tank to 290 K. If the specific gravity 1.2, the solubility of KCl per 100 parts of water is 53.55 at 360 K and 34.5 at 290 K, calculated The capacity of the tank required	03 04
	(c)	The gaseous reaction $A = 2B + C$ takes place isothermally in a constant- pressure reactor. Starting with a mixture of 75 % A and 25 % inerts (by volume), in a specified time the volume double. Calculate the conversion achieved.	07
Q.4	(a) (b)	Define (1) Molarity, (2) Normality (3) Mole fraction Air being compressed from 100 kPa and 255 K (enthalpy 489 kJ/kg) to 1000 kPa and 278 K (enthalpy 509 kJ/kg). The exit velocity of air is 60 m/s. what is the power required (in kW) for the compressor if the load is 100 kg/hr of air ?	03 04
	(c)	Calculated the enthalpy of zinc vapour at 1200 0 C and atmosphere pressure, relative to solid at 10 0 C. Data: Melting point of Zn = 419 0 C (at 1 atm) Boiling point of Zn = 907 0 C (at 1 atm) Mean C _p of solid Zn = 0.105 kcal/kg 0 C Mean C _p of liquid Zn = 0.109 kcal/kg 0 C Heat of fusion of Zn = 1660 kcal/kgmole Heat of vaporization of Zn = 26900 kcal/kgmole Mean Cp of Zinc vapour = 4.97 kcal/kgmole 0 C Atomic weight of Zn = 65.4 kg/kgmole	07

OR

Q.4 (a) Give the statements of (1) Henry's law (2) Raoult's Law (3) 03

Dalton's law

(b) Using Watson equation, calculate latent heat of vaporization of Acetone at 313K

T1	Component	Latent	Tc	n
(boiling	_	heat of		
point		vap at T_1 ,		
temp)		Κ		
		(KJ/kmol)		
329.4	Acetone	29121	508.1	0.38
	(C_3H_6O)			

(c) Pure methane is heated from 303 K to 523 K at atmospheric 07 pressure. Calculate the heat added per kmol methane using the following[g data:

$$C_p = 19.2494 + 52.1135 \times 10^{-3}T + 11.973 \times 10^{-6}T^2 - 11.317 \times 10^{-9}T^3 \frac{KJ}{(kmol \ K)}$$

- Q.5 (a) Differentiate between sensible heat and latent heat
 - (b) For the following reaction, estimate the heat of reaction at 298 K $A + B \rightarrow C + D$ Data: $\Delta H^{0}_{f, A} = -269.8 \text{ kcal/gmol}, \Delta H^{0}_{f, B} = -195.2 \text{ kcal/gmol}, \Delta H^{0}_{f, A}$

$$c = -337.3$$
 kcal/gmol, $\Delta H^0_{f, D} = -29.05$ kcal/gmol

(c) Calculate the heat of reaction at 700 K using the following. 07

$$SO_2 + \frac{1}{2}O_2 \to SO_3$$

Comp.	$\Delta H^{0}_{f,298}(kJ/mol)$	a	b x 10 ³	c x 10 ⁶	
SO ₂	-296.81	24.77	62.95	-44.26	
O ₂	0.0	26.026	11.755	-2.3426	
SO ₃	-395.72	22.04	121.6	-91.87	
OR					

Q.5 (a) Define: (1) standard heat of formation (2) standard heat of reaction 03 (3) adiabatic flame temperature

- (b) A heat exchanger for cooling a hot hydrocarbon liquid uses 10000 kg/h of cooling water, which enters the exchanger at 294 K. The hot oil at the rate of 5000 kg/h enters at 423 K and leaves at 338 K and has an average heat capacity of 2.5 kJ/kg K. Calculate the outlet temperature of water.
- (c) Calculate the theoretical flame temperature of gas having 20 % CO and 80 % N_2 burnt with 150 % excess air. Both air and gas are being at 25 0 C.

Data: heat of formation of CO_2 = -94,052 cal /gmol , CO = -26,412 cal/ gmol at 25 $^0\!C.$

 C_{pm} : $CO_2 = 12.1$, $O_2 = 7.9$, $N_2 = 7.55 \mbox{ cal/ gmol}$ K.

 $Cp^0 = a + bT + cT^2 kJ/(kmol K)$

4

03