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Manipal Institute of Technology, Manipal

(A Constituent Institute of Manipal University)



IV SEMESTER B.TECH (CHEMICAL ENGINEERING) END SEMESTER MAKEUP EXAMINATIONS, JUN/JUL 2016

SUBJECT: **CHEMICAL ENGINEERING THERMODYNAMICS-II [CHE 2201]**

REVISED CREDIT SYSTEM

Time: 3 Hours

MAX. MARKS: 100

Instructions to Candidates:

- ❖ Answer **ALL** questions.
- ❖ Missing data, if any, may be suitably assumed.

1A.	Calculate the fugacity of propane at 12 bar and 310 K using the following data.							08
	P, bar	1.7	3.4	6.8	10.2	11.7	13.6	
	V, m ³ /kg	0.3313	0.1609	0.0754	0.0468	0.0382	0.021	
	Given the molecular weight of propane is 44.							
1B.	The molar enthalpy of a binary solution at constant T and P is given by the relation $H = 400x_1 + 600x_2 + (40x_1 + 20x_2)x_1x_2$ where H is in J/mol. Determine \bar{H}_1 and \bar{H}_2 as functions of x_1 and the numerical values of the pure component enthalpies H_1 and H_2 .							08
1C.	Discuss the tangent-intercept method for determining partial molar property.							04
2A.	The activity coefficient of thallium in amalgams at 298 K are given below.							10
	x_2	0	0.00326	0.01675	0.04856	0.0986	0.168	
	γ_2	1.0	1.042	1.231	1.776	2.811	4.321	
	Determine the activity coefficient of mercury (component 1) in the solution.							
2B.	Prove that if Henry's law is obeyed by component 1 in a binary solution over certain concentration range, Lewis-Randall rule (Raoult's law) will be obeyed by component 2 over the same concentration range.							06
2C.	Derive the expression for the effect of temperature and pressure on activity.							04
3A.	Discuss with a sketch, maximum and minimum boiling azeotrope.							06

3B.	Mixtures of n-Heptane (A) and n-Octane (B) are expected to behave ideally. The total pressure over the system is 101.3 kPa. Using the vapour pressure data given below draw the boiling point diagram.	07																														
	<table><tr><td>T, k</td><td>371.4</td><td>378</td><td>383</td><td>388</td><td>393</td><td>398.6</td></tr><tr><td>P_A, kPa</td><td>101.3</td><td>125.3</td><td>140.0</td><td>160.0</td><td>179.9</td><td>205.5</td></tr><tr><td>P_B, kPa</td><td>44.4</td><td>55.6</td><td>64.5</td><td>74.8</td><td>86.6</td><td>101.3</td></tr></table>	T, k	371.4	378	383	388	393	398.6	P_A, kPa	101.3	125.3	140.0	160.0	179.9	205.5	P_B, kPa	44.4	55.6	64.5	74.8	86.6	101.3										
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3C.	2-propanol and 1-propanol forms an ideal solution. Prepare P-x-y diagram at 90 ⁰ C. The Antoine constants are	07																														
	<table><tr><td></td><td>A</td><td>B</td><td>C</td></tr><tr><td>2-propanol (1)</td><td>8.87829</td><td>2010.330</td><td>252.636</td></tr><tr><td>1-propanol (2)</td><td>8.37895</td><td>1788.020</td><td>227.438</td></tr></table>		A	B	C	2-propanol (1)	8.87829	2010.330	252.636	1-propanol (2)	8.37895	1788.020	227.438																			
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4A.	A mixture contains 45% (mol) methanol (A), 30% (mol) ethanol (B) and the rest n-propanol (C). Liquid solution may be assumed to be ideal and perfect gas law is valid for the vapour phase. Calculate at a total pressure of 101.3 kPa, the bubble point and the vapour composition. The vapour pressures of the pure liquids are given below.	10																														
	<table><tr><td>Temperature, K</td><td>333</td><td>343</td><td>353</td><td>363</td></tr><tr><td>P_A, kPa</td><td>81.97</td><td>133.29</td><td>186.61</td><td>266.58</td></tr><tr><td>P_B, kPa</td><td>49.32</td><td>73.31</td><td>106.63</td><td>166.61</td></tr><tr><td>P_C, kPa</td><td>39.32</td><td>62.65</td><td>93.30</td><td>133.29</td></tr></table>	Temperature, K	333	343	353	363	P_A, kPa	81.97	133.29	186.61	266.58	P_B, kPa	49.32	73.31	106.63	166.61	P_C, kPa	39.32	62.65	93.30	133.29											
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4B.	The following VLE data were obtained for ethanol – water system at 298 K. Test whether the data are thermodynamically consistent by slope of ln γ method.	10																														
	<table><tr><td>X₁</td><td>0.122</td><td>0.163</td><td>0.226</td><td>0.320</td><td>0.337</td><td>0.437</td><td>0.440</td><td>0.579</td><td>0.830</td></tr><tr><td>Y₁</td><td>0.474</td><td>0.531</td><td>0.562</td><td>0.582</td><td>0.589</td><td>0.620</td><td>0.619</td><td>0.685</td><td>0.849</td></tr><tr><td>P,kPa</td><td>5.57</td><td>6.02</td><td>6.38</td><td>6.76</td><td>6.80</td><td>7.02</td><td>7.04</td><td>7.30</td><td>7.78</td></tr></table>	X ₁	0.122	0.163	0.226	0.320	0.337	0.437	0.440	0.579	0.830	Y ₁	0.474	0.531	0.562	0.582	0.589	0.620	0.619	0.685	0.849	P,kPa	5.57	6.02	6.38	6.76	6.80	7.02	7.04	7.30	7.78	
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P,kPa	5.57	6.02	6.38	6.76	6.80	7.02	7.04	7.30	7.78																							
	The vapour pressures of ethanol and water are, respectively, 7.86 kPa and 3.17 kPa.																															
5A.	Derive van't Hoff equation for showing the effect of temperature on equilibrium constant.	08																														
5B.	For the vapour phase hydration of ethylene to ethanol according to $C_2H_4 + H_2O \rightarrow C_2H_5OH$ The equilibrium constants were measured at temperature 420 K and 600 K. They are 6.8×10^{-2} and 1.9×10^{-3} respectively. The specific heat data is as follows.	12																														
	<table><tr><td></td><td>$c_p, J/mol K$</td></tr><tr><td>Ethylene</td><td>$11.886 + 120.12 \times 10^{-3}T - 36.649 \times 10^{-6}T^2$</td></tr><tr><td>Water</td><td>$30.475 + 9.652 \times 10^{-3}T + 1.189 \times 10^{-6}T^2$</td></tr><tr><td>Ethanol</td><td>$29.358 + 166.9 \times 10^{-3}T - 50.09 \times 10^{-6}T^2$</td></tr></table>		$c_p, J/mol K$	Ethylene	$11.886 + 120.12 \times 10^{-3}T - 36.649 \times 10^{-6}T^2$	Water	$30.475 + 9.652 \times 10^{-3}T + 1.189 \times 10^{-6}T^2$	Ethanol	$29.358 + 166.9 \times 10^{-3}T - 50.09 \times 10^{-6}T^2$																							
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	Develop general expressions for the equilibrium constant and standard free energy change as functions of temperature.																															