

Reg. No.

**MANIPAL INSTITUTE OF TECHNOLOGY****MANIPAL***(A constituent unit of MAHE, Manipal)***IV SEMESTER B.TECH. (CHEMICAL ENGINEERING)****ENDSEM EXAMINATIONS, JUN 2022****SUBJECT: CHEMICAL ENGINEERING THERMODYNAMICS-II [CHE 2251]****REVISED CREDIT SYSTEM****(11/06/2022)**

Time: 3 Hours

MAX. MARKS: 50

Instructions to Candidates:

- ❖ Answer **ALL** the questions.
- ❖ Missing data may be suitably assumed.

1A	Calculate the fugacity of nitrogen at 800 bar from the following data at 273 K. (Use compressibility factor method).	5												
	<table><tr><td>P, bar</td><td>50</td><td>100</td><td>200</td><td>400</td><td>800</td></tr><tr><td>PV/RT</td><td>0.9846</td><td>0.9846</td><td>1.0365</td><td>1.2557</td><td>1.7959</td></tr></table>	P , bar	50	100	200	400	800	PV/RT	0.9846	0.9846	1.0365	1.2557	1.7959	
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PV/RT	0.9846	0.9846	1.0365	1.2557	1.7959									
1B	The molar enthalpy of a binary solution at constant T and P is given by the relation $H = 120x_1 + 70x_2 + (15x_1 + 8x_2)x_1x_2$ where H is in J/mol. Calculate \bar{H}_2 as a function of x_1 .	3												
1C	Justify the following statements. (a) The concept of ideal gaseous solution is less restrictive than that of an ideal gaseous mixture. (b) Raoult's law is a simplified form of Lewis-Randall rule.	2												
2A	Check whether the given equations satisfy Gibbs Duhem equation. $\ln \gamma_1 = 500 + 140x_1 - 60x_1x_2 - 20x_2^2 + 20x_1x_2^2$ $\ln \gamma_2 = -120x_2 - 90x_1x_2 - 20x_2^2 + 20x_1x_2^2$ (Note: The final answer on both sides of the Gibbs Duhem equation should be in terms of x_1 .)	4												
2B	With the help of phase diagram, explain the effect of pressure on constant pressure equilibria.	3												
2C	Define Henry's law, Duhem theorem, Critical envelope	3												

3A	<p>Using the RK equation of state, determine the values of RK parameters for a gas mixture containing 30% propane and 70% n-pentane at 500 K and 12 bar.</p> <p>Given the properties of the components</p> <table><tr><td></td><td>Critical temperature (K)</td><td>Critical pressure (bar)</td></tr><tr><td>Propane</td><td>369.9</td><td>42.57</td></tr><tr><td>n-pentane</td><td>469.8</td><td>33.75</td></tr></table>		Critical temperature (K)	Critical pressure (bar)	Propane	369.9	42.57	n-pentane	469.8	33.75	3																		
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3B	<p>The system n-pentane (1)- n-hexane(2) n-heptane(3) forms an ideal solution. Determine the composition of liquid which is in equilibrium with vapour of composition $y_1 = 0.45$, $y_2 = 0.3$ and $y_3 = 0.25$ at 70°C. At 70°C, the vapour pressure of n-pentane, n-hexane and n-heptane are 2129.57 Torr, 785.82 Torr and 303.99 Torr respectively.</p>	3																											
3C	<p>With the help of phase diagrams, discuss the maximum boiling azeotrope.</p>	4																											
4A	<p>Methanol(1)- acetone(2) forms an azeotrope at 760 Torr with $x_1=0.2$ and $t=55.7^\circ\text{C}$. The vapour pressures at 55.7°C are $P_1^S=530.97$ Torr and $P_2^S=749.65$ Torr. Predict the P-x-y data (any two sets of data) at 55.7°C. Assume that van Laar equation is applicable to the system.</p>	4																											
4B	<p>A vapour mixture of 20 mole percent methane, 30 mole percent ethane and 50 mole percent propane is available at 30°C. Making use of the K factor chart determine the percent at which condensation begins if the mixture is isothermally compressed. Also estimate the composition of the first drop of liquid that forms.</p>	3																											
4C	<p>Vapour liquid equilibrium data for the system methanol (1)- benzene (2) at 313 K are given below. Calculate the values of $\ln(\gamma_1/\gamma_2)$.</p> <table><tr><td>x_1</td><td>0</td><td>0.141</td><td>0.304</td><td>0.468</td><td>0.643</td><td>0.702</td><td>0.878</td><td>1.00</td></tr><tr><td>y_1</td><td>0</td><td>0.507</td><td>0.531</td><td>0.543</td><td>0.566</td><td>0.578</td><td>0.670</td><td>1.000</td></tr><tr><td>P, kPa</td><td>24.46</td><td>46.52</td><td>48.32</td><td>48.73</td><td>48.81</td><td>47.61</td><td>44.52</td><td>35.12</td></tr></table>	x_1	0	0.141	0.304	0.468	0.643	0.702	0.878	1.00	y_1	0	0.507	0.531	0.543	0.566	0.578	0.670	1.000	P, kPa	24.46	46.52	48.32	48.73	48.81	47.61	44.52	35.12	3
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5A.	<p>For the following reaction the data are given at 298 K.</p> $\text{CH}_4 + 2\text{H}_2\text{O} \rightarrow \text{CO}_2 + 4\text{H}_2$ <table><tr><td></td><td>$\Delta H_{f,298}^o \times 10^{-3}, \text{J/mol}$</td><td>$\Delta G_{f,298}^o \times 10^{-3}, \text{J/mol}$</td><td>$c_p, \text{J/mol K}$</td></tr><tr><td>Methane</td><td>-74.943</td><td>-50.66</td><td>$17.449 + 60.449 \times 10^{-3}T + 1.117 \times 10^{-6}T^2$</td></tr><tr><td>Water</td><td>-241.99</td><td>-228.60</td><td>$28.850 + 12.055 \times 10^{-3}T$</td></tr><tr><td>Carbon dioxide</td><td>-393.97</td><td>-394.81</td><td>$45.369 + 8.688 \times 10^{-3}T$</td></tr><tr><td>Hydrogen</td><td></td><td></td><td>$27.012 + 3.509 \times 10^{-3}T$</td></tr></table> <p>Calculate the values of integration constants.</p>		$\Delta H_{f,298}^o \times 10^{-3}, \text{J/mol}$	$\Delta G_{f,298}^o \times 10^{-3}, \text{J/mol}$	$c_p, \text{J/mol K}$	Methane	-74.943	-50.66	$17.449 + 60.449 \times 10^{-3}T + 1.117 \times 10^{-6}T^2$	Water	-241.99	-228.60	$28.850 + 12.055 \times 10^{-3}T$	Carbon dioxide	-393.97	-394.81	$45.369 + 8.688 \times 10^{-3}T$	Hydrogen			$27.012 + 3.509 \times 10^{-3}T$	4							
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5B.	<p>Estimate the maximum conversion of ethylene to alcohol by vapour-phase hydration at 523 K and 34 bar.</p>	4																											

	$\text{C}_2\text{H}_4 (\text{g}) + \text{H}_2\text{O} (\text{g}) \rightarrow \text{C}_2\text{H}_5\text{OH} (\text{g})$ <p>The equilibrium constant varies with temperature as</p> $\ln K = \frac{4760}{T} - 1.558 \ln T + 2.22 \times 10^{-3}T - 0.29 \times 10^{-6}T^2 - 5.56$ <p>The steam to ethylene ratio in the initial mixture is 5.0. The equilibrium constant in terms of fugacity coefficient is 0.9419.</p>	
5C.	Define reaction coordinate, Lechatelier's principle	2