

**GUJARAT TECHNOLOGICAL UNIVERSITY****BE - SEMESTER-III (NEW) EXAMINATION – WINTER 2021****Subject Code:3130507****Date:21-02-2022****Subject Name:Chemical Engineering Thermodynamics I****Time:10:30 AM TO 01:00 PM****Total Marks:70****Instructions:**

1. Attempt all questions.
2. Make suitable assumptions wherever necessary.
3. Figures to the right indicate full marks.
4. Simple and non-programmable scientific calculators are allowed.

		Marks
<b>Q.1</b>	(a) State Zeroth law and Third law of thermodynamics	<b>03</b>
	(b) Discuss in brief about choice of refrigerant for refrigeration system.	<b>04</b>
	(c) Explain vapour compression refrigerant cycle with neat flow diagram and T-S diagram.	<b>07</b>
<b>Q.2</b>	(a) Discuss strength and limitations of thermodynamics in chemical engineering.	<b>03</b>
	(b) Select whether the following properties are extensive or intensive: (a) temperature, (b) volume, (c) specific volume (d) heat capacity, (e) potential energy, (f) pressure (g) Internal energy (h) Enthalpy.	<b>04</b>
	(c) Starting from fundamentals, Derive a mathematical expression of the first law of thermodynamics for a steady state flow process.	<b>07</b>
	<b>OR</b>	
(c)	Using Maxwell's equation prove that : $dH = C_p dT + V(1 - \beta T) dP$ $dS = C_p dT/T - \beta V dP$ . Where $\beta =$ Volume expansivity.	<b>07</b>
<b>Q.3</b>	(a) How many degrees of freedom have each of the following system? (1) Liquid water in equilibrium with its vapor. (2) Liquid water in equilibrium with a mixture of water vapor and nitrogen. (3) A liquid solution of alcohol in water in equilibrium with its vapor.	<b>03</b>
	(b) From the First Principle $dU = dQ - dW$ prove the following $\left(\frac{\partial T}{\partial V}\right)_S = -\left(\frac{\partial P}{\partial S}\right)_V$	<b>04</b>
	(c) Discuss Cluasius Inequality in detail	<b>07</b>
<b>OR</b>		
<b>Q.3</b>	(a) Give answer with Justification: Two reversible heat engines are operated between absolute temperature $T_1$ , $T_2$ & $T_3$ as shown in <b>figure (i)</b> . If the work done per cycle by two engine are same then the temperature $T_2$ must be: a. Arithmetic mean of $T_1$ & $T_3$ b. Geometric mean of $T_1$ & $T_3$ c. Logarithmic mean of $T_1$ & $T_3$ d. Harmonic mean of $T_1$ & $T_3$	<b>03</b>
	(b) Explain concept of entropy in brief.	<b>04</b>
	(c) Discuss the Thermodynamic temperature scale in detail	<b>07</b>
	<b>Q.4</b>	(a) Explain physical significance of Virial coefficients.
	(b) Give significance of Compressibility factor and volume expansivity	<b>04</b>

- (c) Discuss the PVT- behavior of pure liquids with PV and PT diagram for a pure material. **07**

**OR**

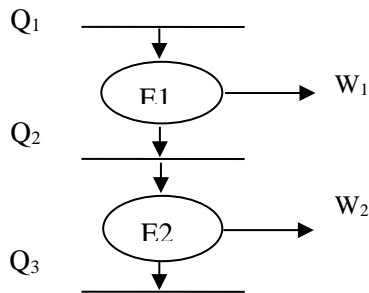
- Q.4** (a) Explain the principle of corresponding states and discuss the generalized compressibility chart. **03**  
 (b) Define: i) standard heat of formation ii) standard heat of combustion iii) standard heat of reaction iv) latent heat **04**  
 (c) Calculate the volume occupied by one mole of oxygen at 300 K and 100 bar using **07**  
 (a) The ideal gas law  
 (b) The van der Waals equation.  
 $a = 0.1378 \text{ N m}^4/\text{mol}^2$  and  $b = 3.18 \times 10^{-5} \text{ m}^3/\text{mol}$

- Q.5** (a) Define: sonic velocity, nozzle and Mach no. **03**  
 (b) Write a Short note on Ejector **04**  
 (c) The standard heat of reaction at 298K is -42.433 kJ for the reaction  $\text{C}_2\text{H}_4(\text{g}) + \text{H}_2\text{O}(\text{g}) \rightarrow \text{C}_2\text{H}_5\text{OH}(\text{g})$ . Calculate the heat of reaction at 400 K. The constants in the heat capacity equation  $C_p = a + bT + cT^2$  are as given below: ( $C_p$  is in J/mol.K and T in K) **07**

Component	a	b	c
$\text{C}_2\text{H}_4$	11.85	$119.75 \times 10^{-3}$	$-36.53 \times 10^{-6}$
$\text{H}_2\text{O}$	30.38	$9.62 \times 10^{-3}$	$1.19 \times 10^{-6}$
$\text{C}_2\text{H}_5\text{OH}$	29.27	$166.39 \times 10^{-3}$	$-49.93 \times 10^{-6}$

**OR**

- Q.5** (a) Explain working principle of Linde liquefaction process in brief **03**  
 (b) Explain isenthalpic process in detail **04**  
 (c) Using Hess's law, calculate the heat of formation for chloroform ( $\text{CHCl}_3$ ) **07**  
 with the following data:  
 (a)  $\text{CHCl}_3(\text{g}) + 0.5\text{O}_2(\text{g}) + \text{H}_2\text{O} \rightarrow \text{CO}_2(\text{g}) + 3\text{HCl}(\text{g})$ ;  $\Delta H^\circ_{298} = -509.93 \text{ kJ}$   
 (b)  $\text{H}_2(\text{g}) + 0.5\text{O}_2(\text{g}) \rightarrow \text{H}_2\text{O}(\text{l})$ ;  $\Delta H^\circ_{298} = -296.03 \text{ kJ}$   
 (c)  $\text{C}(\text{s}) + \text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g})$ ;  $\Delta H^\circ_{298} = -393.78 \text{ kJ}$   
 (d)  $0.5\text{H}_2(\text{g}) + 0.5\text{Cl}_2(\text{g}) \rightarrow \text{HCl}(\text{g})$ ;  $\Delta H^\circ_{298} = -167.57 \text{ kJ}$



**Figure (i)**