

## THERMODYNAMICS

- 1) A thermodynamic state function is a quantity  
( a ) used to determine heat changes ( **b** ) **whose value is independent of path**  
( c ) used to determine pressure volume work ( d ) whose value depends on temperature only.
- 2) For the process to occur under adiabatic conditions, the correct condition is:  
( a )  $\Delta T=0$  ( b )  $\Delta P=0$  ( **c** )  **$q=0$**  ( d )  $w=0$
- 3) The enthalpies of all elements in their standard states are:  
( a ) unity ( **b** ) **zero** ( c )  $< 0$  ( d ) different for each element
- 4)  $\Delta U^\ominus$  of combustion of methane is  $-X \text{ kJ mol}^{-1}$ . The value of  $\Delta H^\ominus$  is  
( a )  $= \Delta U^\ominus$  ( b )  $> \Delta U^\ominus$  ( **c** )  $< \Delta U^\ominus$  ( d ) 0
- 5) A reaction,  $A + B \rightarrow C + D + q$  is found to have a positive entropy change. The reaction will be  
( a ) possible at high temperature ( b ) possible only at low temperature  
( c ) not possible at any temperature ( **d** ) **possible at any temperature**
- 6) As per the available data:  
( **a** )  **$\text{CH}_4(\text{g}) + 2\text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g}) + 2\text{H}_2\text{O}(\text{l}); \Delta_c H^\ominus = -890.3 \text{ KJ mol}^{-1}$**   
( b )  $\text{C}(\text{s}) + \text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g}) \Delta_c H^\ominus = -393.5 \text{ KJ mol}^{-1}$   
( c )  $\text{H}_2(\text{g}) + 1/2\text{O}_2(\text{g}) \rightarrow \text{H}_2\text{O}(\text{l}); \Delta_c H^\ominus = -285.8 \text{ KJ mol}^{-1}$
- 7) Thermodynamics is applicable to  
( **a** ) **macroscopic system only** ( b ) microscopic system only  
( c ) homogeneous system only ( d ) heterogeneous system only
- 8) An isochoric process takes place at constant  
( a ) temperature ( b ) pressure ( **c** ) **volume** ( d ) concentration
- 9) For a cyclic process, the change in internal energy of the system is  
( a ) always +ve ( **b** ) **equal to zero** ( c ) always -ve ( d ) none of the above
- 10) Which of the following properties is not a function of state?  
( **a** ) **concentration** ( b ) internal energy ( c ) enthalpy ( d ) entropy
- 11) Which of the following relation is true?  
( **a** )  **$C_p > C_u$**  ( b )  $C_u > C_p$  ( c )  $C_p = C_u$  ( d )  $C_p = C_u = 0$
- 12) Which of the following always has a negative value?  
( a ) heat of reaction ( b ) heat of solution ( **c** ) **heat of combustion**  
( d ) heat of formation
- 13) The bond energy depends upon  
( a ) size of the atom ( b ) electronegativity ( c ) bond length ( **d** ) **all of the above**
- 14) For an endothermic reaction  
( a )  $\Delta H$  is -ve ( **b** )  **$\Delta H$  is +ve** ( c )  $\Delta H$  is zero ( d ) none of these
- 15) The process depicted by the equation.  
 $\text{H}_2\text{O}(\text{s}) \rightarrow \text{H}_2\text{O}(\text{l})$   
 $\Delta H = +1.43 \text{ kcal}$  represents  
( **a** ) **fusion** ( b ) melting ( c ) evaporation ( d ) boiling
- 16) Which one is the correct unit for entropy?  
( a )  $\text{KJ mol}$  ( b )  $\text{JK}^{-1} \text{ mol}$  ( **c** )  **$\text{JK}^{-1} \text{ mol}^{-1}$**  ( d )  $\text{KJ mol}^{-1}$

17) The enthalpy of a vaporisation of  $\text{CCl}_4$  is  $30.5 \text{ kJ mol}^{-1}$ . Calculate the heat required for the vaporisation of 284g of  $\text{CCl}_4$  at constant pressure (molar mass of  $\text{CCl}_4 = 154 \text{ g mol}^{-1}$ )

**Answer :** 1 mole of  $\text{CCl}_4 = 154 \text{ g}$

Heat required for vapourising 154 g  $\text{CCl}_4 = 30.5 \text{ kJ}$

$\therefore$  Heat required for vapourising

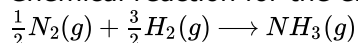
$$284 \text{ g } \text{CCl}_4 = \frac{30.5 \times 284}{154} \text{ kJ} = 56.25 \text{ kJ}$$

18) Given,  $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \longrightarrow 2\text{NH}_3(\text{g}); \Delta_r H^\circ = -92.4 \text{ kJ mol}^{-1}$

What is the standard enthalpy of formation of  $\text{NH}_3$  gas?

**Answer :** Given,  $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \longrightarrow 2\text{NH}_3(\text{g}); \Delta_r H^\circ = -92.4 \text{ kJ mol}^{-1}$

Chemical reaction for the enthalpy of formation of  $\text{NH}_3$  (g) is as follows.



Therefore,  $\Delta_f H^\circ = \frac{-92.4}{2} = -46.2 \text{ kJ mol}^{-1}$

19) Carbon monoxide is allowed to expand isothermally and reversibly from  $10\text{m}^3$  to  $20\text{m}^3$  at 300K and work obtained is 4.754kJ. Calculate the number of moles of carbon monoxide.

**Answer :**  $w = -2.303nRT \log \frac{V_2}{V_1} = 4754$

$$= -2.303 \times n \times 8.314 \times 300 \times \log \frac{20}{10}$$

On solving,  $n = 2.75 \text{ mol}$

20) Calculate the number of kJ of heat necessary to raise the temperature of 60.0g of aluminium from  $35^\circ\text{C}$  to  $55^\circ\text{C}$ . Molar heat capacity of Al is  $24 \text{ J mol}^{-1} \text{ K}^{-1}$ . Molar mass of Al =  $27 \text{ g mol}^{-1}$

**Answer :** Given, mass of Al = 60.0g

Molar mass of Al =  $27 \text{ g mol}^{-1}$

Molar heat capacity,  $C = 24 \text{ J mol}^{-1} \text{ K}^{-1}$

$$\Delta T = 55^\circ\text{C} - 35^\circ\text{C} = 20^\circ\text{C} \text{ or } 20\text{K}$$

Heat,  $q = n \cdot C \cdot \Delta T$

$$q = \frac{60}{27} \times 24 \text{ J mol}^{-1} \text{ K}^{-1} \times 20\text{K} \left( n = \frac{60}{27} \text{ mol} \right)$$

$$= 1066.66 \text{ J} = 1.067 \text{ kJ}$$

21) Although heat is path function but heats absorbed by the system under certain specific conditions is independent of path. what are those conditions? Explain when pressure remains constant.

**Answer :** At constant pressure  $q_p = \Delta U + p\Delta V$ .

But  $\Delta U + p\Delta V = \Delta H$ .

$\therefore q_p = \Delta H$ . As  $\Delta H$  is a state function, therefore,  $q_p$  is a state function.

22) Derive the relationship of work for isothermal and free expansion of an ideal gas.

**Answer :**

23) What would be the work done on or by the system if the internal energy of the system falls by 100 J even when 200 J of heat is supplied to it?

**Answer :** -300 J of work (by the system)

24) Show that for an isothermal expansion of an ideal gas  $\Delta U = 0$ .

**Answer :** For one mole of an ideal gas,  $C_V = \left( \frac{\Delta U}{\Delta T} \right)_V$

$$\text{or } \Delta U = C_V \Delta T$$

For an isothermal process, T is constant so that  $\Delta T = 0$

$$\therefore \Delta U = 0$$

25) Show that for an isothermal expansion of an ideal gas  $\Delta H = 0$ .

**Answer :**  $\Delta H = \Delta U + \Delta(pV)$

For an ideal gas,  $pV = RT$

$$\therefore \Delta H = \Delta U + \Delta(RT) = \Delta U + R\Delta T$$

Since T is constant,  $\Delta T = 0$

$$\therefore \Delta H = 0$$