

Topic :- THERMODYNAMICS

1 (b)
By eq. (ii) - (i),
 $C_G \rightarrow C_D; \quad \Delta H = +1.5 \text{ kJ}$

2 (a)
This is derived formula.

3 (a)
$$W = 2.303 nRT \log \frac{p_2}{p_1}$$
$$= 2.303 \times 1 \times 2 \times 300 \log \frac{10}{2} = 965.84$$
At constant temperature, $\Delta E = 0$
$$\Delta E = q + W;$$
$$q = -W = -965.84 \text{ cal}$$

4 (a)
 $H_2(g) + (1/2)O_2(g) \rightarrow H_2O(g)$
 $\Delta n = -1/2$ and thus, ΔS decreases or -ve

5 (d)
 $\Delta G = \Delta H + T \left[\frac{\partial \Delta G}{\partial T} \right]_P$ (Gibbs-Helmholtz equation)
Also, $\Delta G = \Delta H - T\Delta S$; and $-\Delta G = nF \cdot E$
 $\therefore \Delta G - \Delta H = -T\Delta S$
or $-T\Delta S = T \left[\frac{\partial \Delta G}{\partial T} \right]_P = T \left(\frac{-nF \partial E}{\partial T} \right)_P$
or $\left[\frac{\partial E}{\partial T} \right]_P = \frac{\partial S}{nF}$; similarly derive for other values.

6 (a)
Internal energy of 1 mole of gas $= \frac{3}{2}RT$

8 (c)
 $C_D \rightarrow C_G; \Delta H = -453.5 \text{ cal}$,
i.e., Energy of C_G is less and thus, more stable.

10 **(c)**
At equilibrium $\Delta G = 0$.

11 **(c)**
 $\Delta S = 16 \text{ J mol}^{-1}\text{K}^{-1}$
$$T_{\text{b.p}} = \frac{\Delta H_{\text{vapour}}}{\Delta S_{\text{vapour}}} = \frac{6 \times 1000}{16}$$
$$= 375\text{K}$$

12 **(a)**
$$\Delta S = \frac{\Delta H}{T} = \frac{1.435 \times 10^3}{273} = 5.260 \text{ cal mole}^{-1}\text{K}^{-1}$$

13 **(b)**
Endothermic reactions are those in which heat energy is absorbed.

14 **(b)**
The melting of ice at -15°C is not an spontaneous process.

15 **(c)**
Mixing of gases increases the entropy

16 **(c)**
Current flows from higher potential to lower one.

17 **(a)**
$$-W_{\text{irr.}} = P_{\text{ext.}}(V_2 - V_1)$$
$$= 1 \times (15 - 3) = 12 \text{ litre atm}$$
$$= \frac{12 \times 1.987 \times 4.184}{0.0821}$$
$$= 1.215 \times 10^3 \text{ J}$$

18 **(c)**
 ΔH for combustion of 56 litre $\text{H}_2 = \frac{-24.1 \times 56}{22.4}$
 ΔH for combustion of 56 litre $\text{CO} = \frac{-263 \times 56}{22.4}$
 \therefore Total $\Delta H = -1312 \text{ kJ}$

19 **(b)**
This is the derived formula for W_{rev} in adiabatic process.

20 **(a)**
$$\frac{1}{2} \text{H}_2 + \frac{1}{2} \text{Cl}_2 \rightarrow \text{HCl}; \quad \Delta H = -90 \text{ kJ}$$
$$\therefore \Delta H = \frac{1}{2} e_{\text{H-H}} + \frac{1}{2} e_{\text{Cl-Cl}}$$
or
$$-90 = \frac{1}{2} \times 430 + \frac{1}{2} \times 240 - e_{\text{H-Cl}}$$
$$\therefore e_{\text{H-Cl}} = 425 \text{ kJ mol}^{-1}$$

ANSWER-KEY										
Q.	1	2	3	4	5	6	7	8	9	10
A.	B	A	A	A	D	A	B	C	D	C
Q.	11	12	13	14	15	16	17	18	19	20
A.	C	A	B	B	C	C	A	C	B	A

PE