

Daily Tutorial Sheet-3

Level-1

- 31.(D)** $\Delta S = 10.13 = 31.2 + 51.1 - 47.3 - S_{H_2O}$
 $S_{H_2O} = 45.13 \text{ cal/k mole.}$
- 32.(A)** $\Delta G = 29.3 \times 10^3 - (2 \times 239.7 - 152.3 - 223) \times 298 = -1721.8 \text{ Joule}$
- 33.(D)** $\Delta U = 0$ [for isothermal process]
 $\Delta H = 0$
 $\therefore PV = \text{constant} \quad \Delta S = nR \ln \frac{V_2}{V_1}$
 $\Delta S > 0$ for isothermal expansion
- 34.(B)** Endothermic reaction $\Rightarrow \Delta H > 0$
 $\Delta S > 0$ as gaseous moles are increasing
- 35.(C)** In I part, entropy is decreasing as solid is formed
 In II and III, entropy is increasing as gaseous moles are increasing
- 36.(ACD)** Only heat depends on path so it is a path function
 All other options depends only on state so these are state function
- 37.(B)** $\Delta S = \frac{(\Delta H)_{\text{vap}}}{T_{B,pt}} \Rightarrow T = \frac{30000}{75} = 400 \text{ K}$
- 38.(ABD)** $P_1 V_1 = P_2 V_2$ [for isothermal]
 $\Delta U = 0$ [for isothermal]
 $\Delta H = 0$ [for isothermal]
- 39.(A)** $\Delta U = q + W$
- 40.(D)** $W = -P\Delta V = -nR\Delta T = -1 \times R \times 1 = -R$
- 41.(B)** For adiabatic reversible process;
 $PV^\gamma = K \Rightarrow P_2 = P_1 \left(\frac{V_1}{V_2} \right)^\gamma \text{ (as } V_2 > V_1)$
 \therefore So final pressure will be more for diatomic gas (less γ)
- 42.(C)** $TV^{\gamma-1} = K$
- 43.(D)** Bond formation is exothermic.
- 44.(A)** (A) At constant volume, $w = 0$
 $q < 0$ (Cooling)
 $\Delta U < 0$ (Temperature is decreasing)
 (B) $\Delta U = 0$ (isothermal)
 $w < 0$ (expansion)
 $q > 0$ ($\Delta U = q + w$)
 (C) $q = 0$ (Adiabatic)
 $w = 0$ (Vacuum)
 $\Delta U = 0$

(D) $w < 0$ (expansion)
 $q > 0$ (endothermic)
 $\Delta H = \Delta U > 0$

45.(C) (A) $q = 0$ (B) $\Delta T = 0$ (C) $\Delta H = 0$ (D) $\Delta S = 0$