- Thermo chemistry is the study of heat changes that accompany chemical reactions and phase changes.
- In chemical reactions energy is either absorbed or released.
   According to this there are two types of reactions;
   endothermic and exothermic.
  - a. Endothermic Reactions
    - Energy is absorbed by reactants and total potential

energy of reactants is smaller than that of products.

87.9 kJ + C(s) + 2S(s)  $\longrightarrow$  CS<sub>2</sub> (l) The reaction above is an example for endothermic reactions.



R C (Reaction Coordinate)

- b. Exothermic Reactions
  - Energy is released by reactants and total potential

energy of reactants is greater than that of products.







- Enthalpy (H) is the heat content of a substance at constant pressure.
- The change in enthalpy for a reaction is called the enthalpy of reaction (ΔH).

ΔH = ΣH<sub>products</sub> - ΣH<sub>reactants</sub>
If ΣH<sub>products</sub> > ΣH<sub>reactants</sub>, then ΔH > 0 so the reaction is endothermic. Similarly,
If ΣH<sub>products</sub> < ΣH<sub>reactants</sub>, then ΔH < 0 so the reaction is exothermic.</li>



Enthalpy Changes
for Exothermic and
<b>Endothermic Reactions</b>

Type of reaction	Sign of ∆H <sub>rxn</sub>
Exothermic	Negative
Endothermic	Positive





R C (Reaction Coordinate)

#### Standard Heat of Formation ( $\Delta H^{\circ}_{\phi}$ )

- The heat change when 1 mole compound is produced from its elements in their most stable states (under 1 atm pressure and at 25°C is called as standard heat of formation, and shown by  $\Delta H^{o}_{f}$ .
- $\Delta H_{f}^{o}$  of the free atoms (K, Fe, Na, S, P, Cu...etc) and free simple molecules (O<sub>2</sub>, N<sub>2</sub>, Cl<sub>2</sub>, P<sub>4</sub>, ...etc) are accepted as zero.

Heat of a reaction,  $\Delta H^{\circ}$  can be calculated by using  $\Delta H^{\circ}_{f}$  values.  $\Delta H^{o} = \Sigma H^{o}_{\text{f(products)}} - \Sigma H^{o}_{\text{f(reactants)}}$ Example 3 Find the heat of the reaction ( $\Delta H^{\circ}$ )  $SO_2(g) + \frac{1}{2}O_2(g) \rightarrow SO_3(g)$ by using  $\Delta H_{f}^{o}$  of the compounds given  $\Delta H^{o}_{f(SO2)} = -297 \text{ kJ/mol}, \Delta H^{o}_{f(SO3)} = -396 \text{ kJ/mol}$  $\Delta H^{o}_{f(O2)} = 0$  kJ/mol

Solution

 $\Delta H^{o} = \Sigma H^{o}_{f(products)} - \Sigma H^{o}_{f(reactants)}$   $\Delta H^{o} = \Delta H^{o}_{f(SO3)} - [\Delta H^{o}_{f(SO2)} + \frac{1}{2} \Delta H^{o}_{f(O2)}]$   $\Delta H^{o} = (-396) - [(-297) + \frac{1}{2} \times (0)]$   $\Delta H^{o} = -99 \text{ kJ}$ *Example 4* 

When 2.4 g of graphite burnt with  $O_2$  completely, 78.70 kJ heat is released. What is the molar enthalpy of the formation of  $CO_2$ ?



**Solution** 

 $C_{(graphite)} + O_{2}(g) \rightarrow CO_{2}(g)$ 2.4 g C releases 78.70 kJ <u>12 g (1mol) C releases x</u> x = 393.5 kJ

 $\Delta H^{o} = \Sigma H^{o}_{f(\text{products})} - \Sigma H^{o}_{f(\text{reactants})}$  $\Delta H^{o} = \Delta H^{o}_{f(\text{CO2})} - [\Delta H^{o}_{f(\text{C})} + \Delta H^{o}_{f(\text{O2})}]$  $-393.5 = \Delta H^{o}_{f(\text{CO2})} - [(0) + (0)]$  $\Delta H^{o}_{f(\text{CO2})} = -393.5 \text{ kJ}$ 

The combustion reaction of propane is  $C_3H_8(g) + 5O_2(g) \rightarrow 3CO_2(g) + 4H_2O(I) \Delta H^0 = -2220.3 \text{ kJ}$ If  $\Delta H^{o}_{f(CO2)}$  and  $\Delta H^{o}_{f(H2O)}$  values are -393.5 kJ/mol and -286 kJ/mol respectively find  $\Delta H^{o}_{f(C3H8)}$ ? Example 6  $Fe_2O_3(s) + 3CO(g) \rightarrow 2Fe(s) + 3CO_2(g)$ If  $\Delta H^{o}_{f(Fe2O3)}$ ,  $\Delta H^{o}_{f(CO)}$  and  $\Delta H^{o}_{f(CO2)}$  values are -826 kJ/mol, -110.5 kJ/mol and -393.5 kJ/mol respectively find  $\Delta H^{\circ}$  for the reaction.