Enthalpy of Formation (ΔH_f)

Calculating Enthalpy Changes of Reactions from Heat of Formation

Standard enthalpy (heat) of formation ($\triangle H_{\mathfrak{p}}^{\circ}$): change in enthalpy that accompanies the formation of \square mole of the compound in its standard state from its component <u>elements</u> their standard states.

Note: The naught, ____, simply indicates standard conditions (1 atm and 25°C).

$$\Delta H_f^{\circ}$$
 = -74.9 kJ/mol_{rxn}

$$S_{(S)} + \frac{3}{2} O_{2(g)} \rightarrow SO_3(g)$$

$$\Delta H_f^{\circ} = -396 \text{ kJ/mol}_{\text{rxn}}$$

Note: you will see <u>fractional</u> coefficients to ensure only <u>1</u> mole of compound is formed.

The ΔH_f° for <u>elements</u> (in their standard state) is always $\frac{\phi}{\phi}$ kJ/mol_{rxn}! Now you try! Write the formation reaction of NH₃: $\frac{1}{2}N_{2(g)} + \frac{3}{2}H_{2(g)} \rightarrow NH_{3(e)}$

The enthal py change for a chemical reaction, or amount of heat released or absorbed, can be determined by the following formula, which is known as Big Momma's Equation:

$$\Delta H_{rxn}^{\circ} = \Sigma \left[n \Delta H_f^{\circ}(products) \right] - \Sigma \left[n \Delta H_f^{\circ}(reactants) \right]$$
 $\left[\begin{array}{c} \circ h \\ F.C. \end{array} \right]$

$$CH_4(g) + 2 O_2(g) \rightarrow CO_2(g) + 2 H_2O(g)$$

$$\Delta H_{rxn}^{\circ} = \Delta H_{comb}^{\circ} = \left[\Delta H_{f}^{\circ} (co_{2}) + 2 \cdot \Delta H_{f}^{\circ} (H_{2}O) \right] - \left[\Delta H_{f}^{\circ} (cH_{4}) + 2(\phi) \right]$$

Enthalpy Changes of Different Types of Reactions

You will encounter a variety of Subscripts following the ΔH , however, they are simply indicating a <u>reaction</u> type of reaction or change of state.

Examples

 ΔH_{comb}° = Enthalpy of Combustion

(Heat Energy Released during Combustion Reactions)

 ΔH_{neut}° = Enthalpy of Neutralization

(Heat Energy Released during Acid-Base Neutralization Reactions)

 ΔH_{Soln}° = Enthalpy of Solution

(Heat Energy Released/Absorbed Dissolving a Solute in Water)

 ΔH_{vap}° = Enthalpy of Vaporization

(Heat Energy Absorbed to Convert from Liquid to Gas Phase)

 ΔH_{fus}° = Enthalpy of Fusion

(Heat Energy Absorbed to Convert from Solid to Liquid Phase)

Let's Practice!

1. For each reaction below, fill in the box with a subscript that specifies the type of reaction:

a.
$$NaOH(aq) + HCl(aq) \rightarrow NaCl(aq) + H_2O(l)$$

$$\Delta H_{\text{Neut}} = -55.8 \text{ kJ/mol}$$
 b. $H_2O(s) \rightarrow H_2O(l)$
$$\Delta H_{\text{LuS}} = +6.02 \text{ kJ/mol}$$

c. $CH_4(g) + 2O_2(g) \rightarrow CO_2(g) + 2H_2O(g)$

$$H_2(g)$$
, $Hg(s)$, $CO_2(g)$, $H_2O(l)$, $Br_2(l)$, $N_2(l)$ not $l \in room temp!$

3. Write a balanced molecular equation representing the enthalpy for standard heat of formation reaction of ethanol, C₂H₅OH(I).

4. Use the information provided and the balanced equation to determine ΔH_f° of carbon tetrachloride.

$$CH_4(g) + 4 \ Cl_2(g) \rightarrow CCl_4(g) + 4 \ HCl(g) \qquad \Delta H^\circ_{rxn} = -389 \ kJ/mol_{rxn}$$

Substance	ΔH°r
CH₄(g)	–75 kJ/mol _{rxn}
HCl(g)	−92 kJ/mol _{rxn}

$$\Delta H_{rxh}^{\circ} = -389 \frac{KJ}{m_{o}l_{rxh}} = \sum_{n} \Delta H_{f}^{\circ} (pr) - \sum_{n} \Delta H_{f}^{\circ} (re)$$

$$= \left[CCl_{4} + 4Hcl \right] - \left[CH_{4} + 4Cl_{2} \right]$$

$$= \left[CCl_{4} - 4(-92) \right] - \left[-75 + 4(\phi) \right]$$

$$-389 = CCl_{4} - 293 \Rightarrow \Delta H_{f}^{\circ} (CCl_{4}) = \left[-96 \frac{KJ}{mol_{rxh}} \right]$$