HW4

Enthalpy change (Δ H) – amount of chemical heat energy taken in (giving out) in a reaction. We can measure enthalpy change, but we cannot measure absolute value of enthalpy.

In an exothermic reactions heat is transferred from chemical reaction to the surrounding. Products have lower energy than reactants. Products are more stable. Δ H will be negative.

In endothermic reactions the surrounding gets colder. Products are less stable, they have higher energy than reactants. Δ H will be positive.

Lavoisier and Laplace law (1st law of thermodynamics) – the amount of energy released (or absorbed) during formation of a chemical compound is equal to the amount of energy absorbed (or released) when the same compound is destroyed. 2H₂ (g) + O₂ (g) \rightarrow 2H₂O (aq) Δ H = - 573 kJ/mol (exothermic reaction) Electrolysis of water

 $2H_2O(aq) \rightarrow 2H_2(g) + O_2(g) \Delta H = + 573 \text{ kJ/mol}$ (endothermic reaction)

Hess's law. Amount of heat given or taken in the reaction is independent of the pathway between the initial and final state.

We want to obtain Na₂SO₄. 1. 2NaOH + H₂SO₄ \rightarrow Na₂SO₄ + 2H₂O Δ H=- 131 kJ/mol 2. NaOH + H₂SO₄ \rightarrow NaHSO₄ + H₂O Δ H =- 62 kJ/mol NaHSO₄ + NaOH \rightarrow Na₂SO₄ + H₂O Δ H =-69 kJ/mol -69-62=-131

We can use Hess's law if we cannot find enthalpy change from the experiment. We can take enthalpy changes from the known chemical reaction and mathematically manipulate chemical equations. For example, we want to know the enthalpy change for synthesis of methane directly from carbon. This reaction is very difficult to perform in the lab. We can do the following manipulations:

$$C(s) + 2H_2(p) \rightarrow CM_4(p)$$

$$AH \ doe the reactions 1,2 and 3
are known from the experiments
$$I \cdot CH_4(g) + 2O_2(g) \rightarrow CO_2(g) + 2H_2O(QQ)$$

$$AH = -890 \text{ KJ/mol}$$

$$2 \cdot C(s) + O_2(g) \rightarrow CO_2(p)$$

$$AH = -394 \text{ KJ/mol}$$

$$3 \cdot 2H_2(g) + O_2(g) \rightarrow 2H_2O(QQ)$$

$$W_1 \ can \ subtract \ reactions \ 2 \ and \ 3 \ H = -572 \text{ KJ/mol}$$

$$W_1 \ can \ subtract \ reactions \ 2 \ and \ 3 \ H = -572 \text{ KJ/mol}$$

$$W_1 \ can \ subtract \ reactions \ 2 \ and \ 3 \ H = -572 \text{ KJ/mol}$$

$$H = -572 \text{ KJ/mol}$$

$$H = SMH \ reaction \ well.$$

$$h \ ave \ the \ dollowing \ mol}$$

$$Reveation:$$

$$CH_4(g) - C(s) - 2H_1(g) - 890 \text{ KJ/mol} = 394 \text{ KJ/mol}$$

$$W_2 \ veaturange \ it: \ +572 \text{ KJ/mol} = (+2M_2 \rightarrow CH_4) = 76 \text{ KJ/mol}$$$$

Questions:

1. What substance is more stable, graphite or diamond?

C (graphite) + $O_2 \rightarrow CO_2 \Delta$ H=- 393.8 kJ/mol C (diamond) + $O_2 \rightarrow CO_2 \Delta$ H=- 395.7 kJ/mol

2. Calculate enthalpy change for the following reaction $2CO(g) + O_2(g) \rightarrow 2CO_2(g)$ The enthalpy change for these reactions are known $2C(s) + O_2(g) \rightarrow 2CO(g) \Delta H=-222 \text{ kJ/mol}$ $C(s) + O_2(g) \rightarrow CO_2(g) \Delta H=-394 \text{ kJ/mol}$