

Two Ways to Find ΔH° for a Reaction

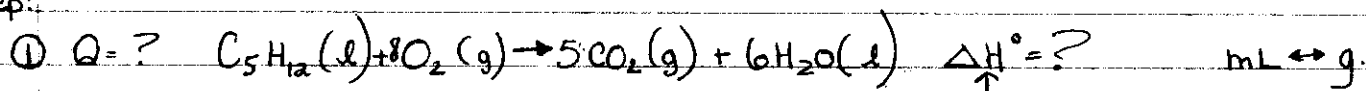
Nov. 9th

1. Experimental Data
2. Previously Determined ΔH° values (Heat's of Formation)

ΔH° ← as composed to
25°C, 1 atmosphere pressure (SATP)
change in enthalpy (change in chemical potential energy)

Eg. 2 g of C_5H_{12} is reacted with oxygen in a combustion reaction.
Determine ΔH° for this reaction given that heat produced by the
combustion warms 500 mL water from 20.000°C to 46.457°C (p. 299)

Step:



$m = 500 \text{ mL} \rightarrow 500 \text{ g}$

$c = 4.184 \text{ J/g}^\circ\text{C}$

$\Delta T = 46.457^\circ\text{C} - 20.000^\circ\text{C} = 26.457^\circ\text{C}$

Written per mol
of reactant.

$Q = mc\Delta T$

$Q = 500 \text{ g} \times 4.184 \text{ J/g}^\circ\text{C} \times 26.457^\circ\text{C}$

$Q = 55348 \text{ J} \rightarrow 55.348 \text{ kJ}$

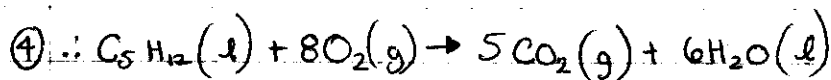
② Conversions.

$55.348 \text{ kJ} \times \frac{72.17 \text{ g}}{1 \text{ mol } C_5H_{12}} = 1997.23 \text{ kJ}$

$2.00 \text{ g } C_5H_{12} \quad 1 \text{ mol } C_5H_{12} \quad 1 \text{ mol } C_5H_{12}$

③ $\Delta H = -Q$

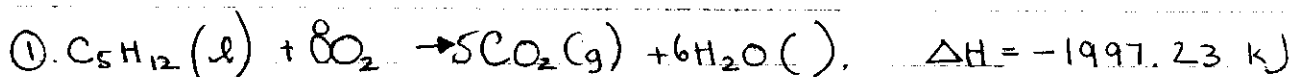
$\Delta H = -1997.23 \text{ kJ} / \text{mol } C_5H_{12}$



$\Delta H^\circ = -1997.23 \text{ kJ}$

Given that $\Delta H^\circ = -1997.23 \text{ kJ}$ for the combustion of C_5H_{12} .

Determine the final temperature of 500 mL of water at $20,000^\circ\text{C}$ if warmed by the combination of 200 g of pentane.



② $Q = -\Delta H$

$Q = 1997.23 \text{ kJ/mol C}_5\text{H}_{12}$

③ Conversions (scale \rightarrow size)

$$200 \text{ g C}_5\text{H}_{12} \times \frac{1 \text{ mol C}_5\text{H}_{12}}{72.17 \text{ g C}_5\text{H}_{12}} \times \frac{1997.23 \text{ kJ}}{1 \text{ mol C}_5\text{H}_{12}} = 55.348 \text{ kJ}$$

④ $Q = 55.348 \text{ kJ} \rightarrow 55348 \text{ J}$ $\Delta T = \frac{Q}{mC}$

$m = 500 \text{ mL} \rightarrow 500 \text{ g}$

$C = 4.184 \text{ J/g}^\circ\text{C}$

$\Delta T = ?$

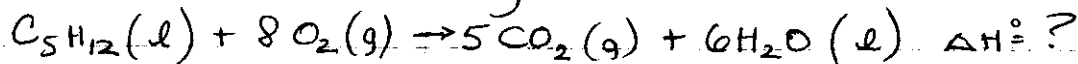
$\Delta T = \frac{55348 \text{ J}}{500 \text{ g} \times 4.184 \text{ J/g}^\circ\text{C}}$

$T_F = 26.457^\circ\text{C} + 20000^\circ\text{C}$
 $T_F = 46.457^\circ\text{C}$

$\Delta T = 26.457^\circ\text{C}$

* Schlenker's mistake $\Delta H \neq -1997.23$ for the combustion of pentane.

Find ΔH° for a reaction using heats of formation.



$$\Delta H^\circ = \left[\underbrace{5 \Delta H^\circ_{\text{CO}_2}}_{\text{pot. energy of CO}_2} + \underbrace{6 \Delta H^\circ_{\text{H}_2\text{O}(\text{l})}}_{\text{pot. energy of H}_2\text{O}} \right] - \left[\underbrace{\Delta H^\circ_{\text{C}_5\text{H}_{12}}}_{\text{pot. energy C}_5\text{H}_{12}} + \underbrace{8 \Delta H^\circ_{\text{O}_2}}_{\text{pot. energy of O}_2^*} \right] \quad (\text{p. 799})$$

$$\Delta H^\circ = \left[5(-393.5 \text{ kJ}) + 6(-285.8 \text{ kJ}) \right] - \left[(-173.5 \text{ kJ}) + 8(0^*) \right]$$

$\Delta H^\circ = -3508.8 \text{ kJ}$

* all elements in their natural state at SATP have been assigned $\Delta H^\circ = 0$