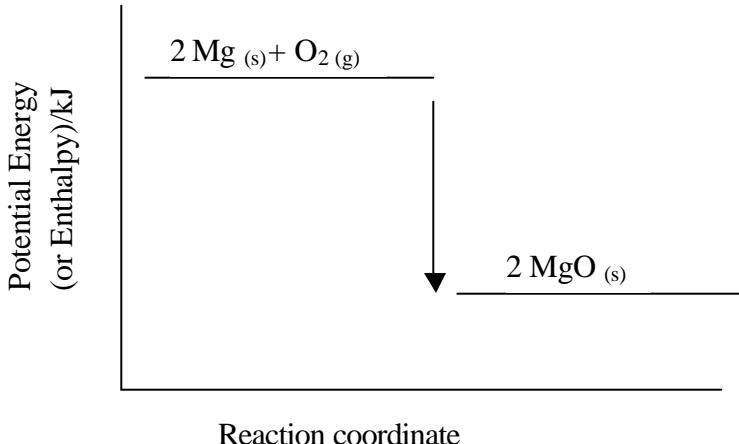


First-Year SL Chemistry

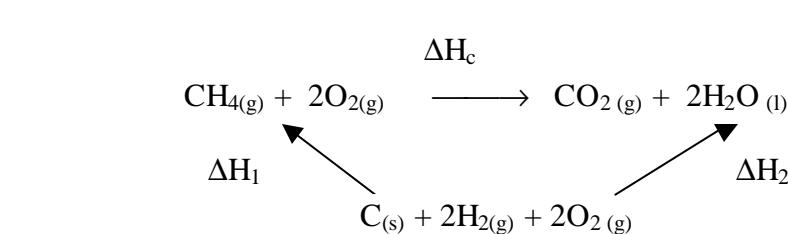
5. Energetics: Thermochemistry

Solution further problems.

- Decrease in the strength and/or the number of bonds causes energy to be released in an exothermic reaction.
- a)



- b) The reaction is exothermic.
 - c) Reactants.
- Heat gained by the calorimeter + water sample in calorimeter = - heat lost by 2nd water sample
 $\Rightarrow (\text{mass of calorimeter} \times C_p \text{ of calorimeter} \times \Delta T) + (\text{mass of water} \times C_p \text{ of water} \times \Delta T) = -(\text{mass of second water sample} \times C_p \text{ of water} \times \Delta T)$
 $\Rightarrow (\text{Heat capacity of calorimeter}) (32.6 - 24.1) \text{ K} + (0.050 \text{ kg}) (4.18 \text{ kJ kg}^{-1} \text{ K}^{-1}) (32.6 - 24.1) \text{ K}$
 $= - (0.050 \text{ kg}) (4.18 \text{ kJ kg}^{-1} \text{ K}^{-1}) (32.6 - 41.7) \text{ K}$
 $\Rightarrow (\text{Heat capacity of calorimeter}) (8.5 \text{ K}) + 1.77 \text{ kJ} = 1.90 \text{ kJ}$
 $\Rightarrow \text{Heat capacity of calorimeter} = [(1.90 - 1.77) \text{ kJ}] / 8.5 \text{ K}$
 $\Rightarrow \text{Heat capacity of calorimeter} = 0.01529 \text{ kJ K}^{-1} = 0.015 \text{ kJ K}^{-1}$
 - a) If $2 \text{NO}_{(g)} + \text{Cl}_{2(g)} \rightarrow 2 \text{NOCl}_{(g)}$ $\Delta H^\theta = -77.4 \text{ kJ}$
then $\text{NO}_{(g)} + \frac{1}{2} \text{Cl}_{2(g)} \rightarrow \text{NOCl}_{(g)}$ $\Delta H^\theta = -77.4 \text{ kJ} / 2 = -38.7 \text{ kJ}$
 - b) $6 \text{NOCl}_{(g)} \rightarrow 6 \text{NO}_{(g)} + 3 \text{Cl}_{2(g)}$ $\Delta H^\theta = -77.4 \text{ kJ} \times 3 = -232.2 \text{ kJ}$
 - $2\text{Na}_{(s)} + 2\text{S}_{(s)} + \frac{3}{2}\text{O}_{2(s)} \rightarrow \text{Na}_2\text{S}_2\text{O}_{3(s)}$ $\Delta H = \Delta H_f^q$
 - .



$$\Delta H_2 = \Delta H_1 + \Delta H_c$$

$$\Delta H_c = \Delta H_2 - \Delta H_1$$

Where, $\Delta H_1 = \Delta H_f^q [\text{CH}_4\text{(g)}] = (1 \text{ mol}) (-75 \text{ kJ mol}^{-1}) = -75 \text{ kJ}$

$$\begin{aligned}\Delta H_2 &= (2 \text{ mol}) \Delta H_f^q [\text{H}_2\text{O(l)}] + (1 \text{ mol}) \Delta H_f^q [\text{CO}_2\text{(g)}] \\ &= (2 \text{ mol}) X (-285.83 \text{ kJ mol}^{-1}) + (1 \text{ mol})(-393.51 \text{ kJ mol}^{-1}) \\ \Delta H_2 &= -956.17 \text{ kJ}\end{aligned}$$

$$\Delta H_c = -956.17 \text{ kJ} - (-75 \text{ kJ})$$

$$\Delta H_c = -890.17 \text{ kJ}$$

$$\Delta H_c = -8.90 \times 10^2 \text{ kJ mol}^{-1} (\text{mol}^{-1} \text{ obtained from the equation})$$

7. $M_R (\text{C}_5\text{H}_{10}\text{O}_5) = 155.15$

$$\begin{aligned}\text{Heat gained by the calorimeter and contents} &= \text{heat capacity} \times \Delta T \\ &= (15.8 \text{ kJ}/^\circ\text{C}) (210.54 - 20.00) ^\circ\text{C} \\ &= 3010.532 \text{ kJ}\end{aligned}$$

$\Delta H = -$ heat energy gained by calorimeter and its contents

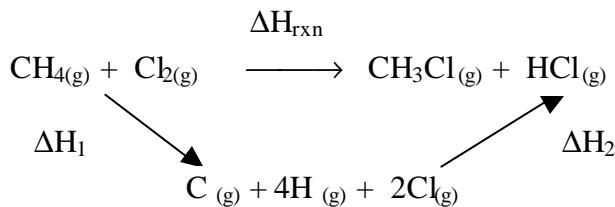
$$\Delta H = -3010 \text{ kJ}$$

$$\left(\frac{3010 \text{ kJ}}{0.548 \text{ g C}_5\text{H}_{10}\text{O}_5} \right) \left(\frac{155.15 \text{ g C}_5\text{H}_{10}\text{O}_5}{1 \text{ mol C}_5\text{H}_{10}\text{O}_5} \right) = 852343 \text{ kJ mol}^{-1}$$

$$\Delta H_c = -8.52 \times 10^5 \text{ kJ mol}^{-1}$$

8. $\text{Cl}_2\text{(g)} \rightarrow 2\text{Cl(g)}$ $\Delta H = \text{bond enthalpy}$
 $[\frac{1}{2} \text{Cl}_2\text{(g)} \rightarrow \text{Cl(g)}$ $\Delta H = \text{enthalpy of atomization}]$

9. (a)



$$\Delta H_{rxn} = \Delta H_1 + \Delta H_2$$

where, $\Delta H_1 = \text{bond breaking}$

$$\begin{aligned}&= (4 \text{ mol}) E(\text{C-H}) + (1 \text{ mol}) E(\text{Cl-Cl}) \\ &= (4 \text{ mol}) (412 \text{ kJ mol}^{-1}) + (1 \text{ mol}) (242 \text{ kJ mol}^{-1}) \\ &= 1890 \text{ kJ}\end{aligned}$$

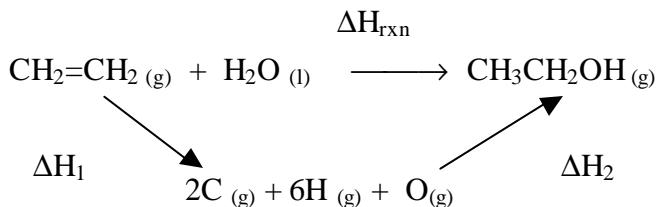
$\Delta H_2 = \text{bond making}$

$$\begin{aligned}&= (3 \text{ mol}) E(\text{C-H}) + (1 \text{ mol}) E(\text{C-Cl}) + (1 \text{ mol}) E(\text{H-Cl}) \\ &= (3 \text{ mol}) (-412 \text{ kJ mol}^{-1}) + (1 \text{ mol}) (-338 \text{ kJ mol}^{-1}) + (1 \text{ mol}) (-431 \text{ kJ mol}^{-1}) \\ &= -2005 \text{ kJ}\end{aligned}$$

$$\Delta H_{rxn} = (1890 - 2005) \text{ kJ} = -115 \text{ kJ}$$

$$\Delta H_{rxn} = -115 \text{ kJ mol}^{-1} (\text{mol}^{-1} \text{ from the equation})$$

(b)



$$\Delta H_{rxn} = \Delta H_1 + \Delta H_2$$

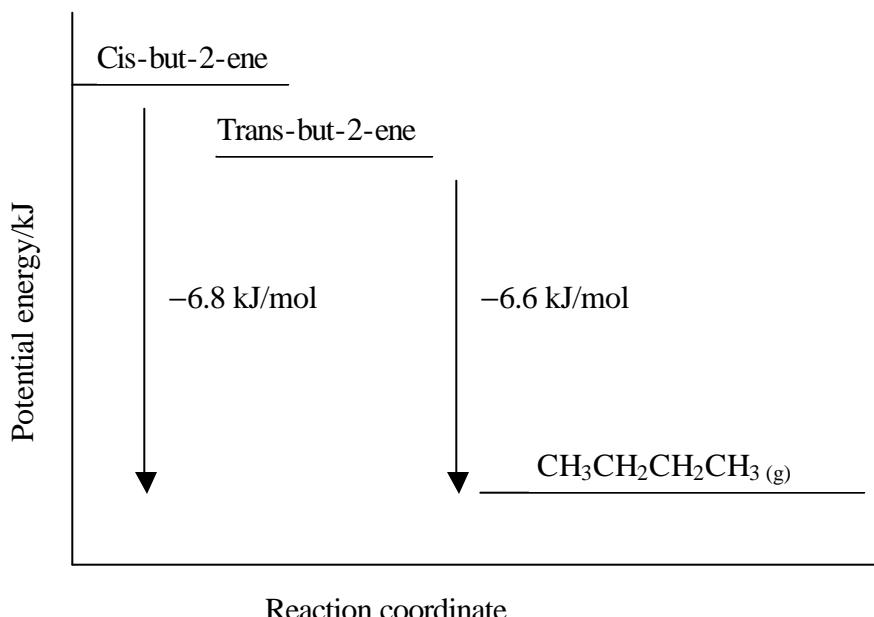
where, ΔH_1 = bond breaking

$$\begin{aligned}
 &= (4 \text{ mol}) E(\text{C-H}) + (1 \text{ mol}) E(\text{C=C}) + (2 \text{ mol}) E(\text{H-O}) \\
 &= (4 \text{ mol}) (412 \text{ kJ mol}^{-1}) + (1 \text{ mol})(612 \text{ kJ mol}^{-1}) + (2 \text{ mol}) (463 \text{ kJ mol}^{-1}) \\
 &\equiv 3186 \text{ kJ}
 \end{aligned}$$

$$\begin{aligned}\Delta H_2 = \text{bond making} &= (5 \text{ mol}) E(\text{C-H}) + (1 \text{ mol}) E(\text{C-C}) + (1 \text{ mol}) E(\text{C-O}) + (1 \text{ mol}) E(\text{O-H}) \\ &= (5 \text{ mol}) X (-412 \text{ kJ mol}^{-1}) + (1 \text{ mol})(-348 \text{ kJ mol}^{-1}) + (1 \text{ mol})(-360 \text{ kJ mol}^{-1}) + (1 \text{ mol}) \\ &\quad (-463 \text{ kJ mol}^{-1}) \\ &\equiv -3231 \text{ kJ}\end{aligned}$$

$$\Delta H_{rxn} = (3186 - 3231) \text{ kJ} = -45 \text{ kJ}$$

10.



Trans-but-2-ene is thermodynamically more stable. Because hydrogenation of trans-but-2-ene give off less heat than cis-but-2-ene, it can be said that the former is at a lower potential energy than the latter and therefore more stable as depicted above.