1. When 100.0 mL of 1.00 M HCl is mixed with 100.0 mL of 1.00 M NaOH, both initially at 21.1°C, are mixed in a two-cup calorimeter the temperature of the mixture rises to 27.9°C. Determine the  $\Delta H$  of neutralization for the reaction

$$\mathrm{HCl}_{(\mathrm{aq})} + \mathrm{NaOH}_{(\mathrm{aq})} \rightarrow \mathrm{NaCl}_{(\mathrm{aq})} + \mathrm{H}_2\mathrm{O}_{(\mathrm{l})}$$

By a prior experiment, the heat capacity of the calorimeter was determined to be 125 J/°C. Assume the density of the final solution is 1.0 g/mL and the specific heat of the mixture is  $4.18 \text{ J/g}^{\circ}\text{C}$ .

$$\Delta T = 27.9^{\circ}\text{C} - 21.1^{\circ}\text{C} = 6.8^{\circ}\text{C}$$

$$n_{\text{HCI}} = n_{\text{NaOH}} = 0.1000 \text{ L} \times 1.00 \text{ M} = 0.100 \text{ mol}$$

$$C_{\text{cal}} = 125 \frac{\text{J}}{\text{\circ}\text{C}}$$

$$n\Delta H + mc\Delta T + C_{\text{cal}}\Delta T = 0$$

$$\Delta H = -\frac{(mc + C_{\text{cal}})\Delta T}{n} = \frac{\left[(200.0 \text{ g})(4.18 \frac{\text{J}}{\text{g}^{\circ}\text{C}}) + 125 \frac{\text{J}}{\text{\circ}\text{C}}\right](6.8^{\circ}\text{C})}{0.100 \text{ mol}}$$

$$\Delta H = -65,350 \frac{\text{J}}{\text{mol}}$$

2. Consider the reaction

$$\mathrm{C_{12}H_{22}O_{11(s)}} \ + \ 12 \ \mathrm{O_{2(g)}} \ \rightarrow \ 12 \ \mathrm{CO_{2(g)}} \ + \ 11 \ \mathrm{H_2O_{(g)}}$$

which has a  $\Delta H$  of -5.65 x 10<sup>3</sup> kJ/mol (C<sub>12</sub>H<sub>22</sub>O<sub>11</sub>). How much heat (energy) can be produced during the complete combustion of 10.0 g of sucrose?

$$q = n\Delta H_{\text{comb}}$$

$$n = \frac{10.0 \text{ g}}{342.2 \text{ g/mol}} = 0.02922 \text{ mol}$$

$$q = 0.2922 \text{ mol}(-5.65 \times 10^3 \text{ kJ/mol}) = -165.1 \text{ kJ}$$

3. If all of the energy in question 2 were used to heat 1.0 L of water at 22.0°C, what would the final temperature of the water be? (Assume 100% energy transfer to the water.)

$$q_{\text{comb}} = -q_{\text{H}_2\text{O}}$$
  
-165.1×10<sup>3</sup> J = -1000 g(4.184  $\frac{\text{J}}{\text{g}^{\circ}\text{C}}$ )( $T_{\text{f}}$  - 22.0°C)  
 $T_{\text{f}}$  - 22.0°C = 39.46°C  
 $T_{\text{f}}$  = 61.5°C

4. Using standard enthalpies of reaction, calculate the  $\Delta H^{\circ}$  for the following reactions:

$$C_{2}H_{2(g)} + 2 H_{2(g)} \rightarrow C_{2}H_{6(g)}$$

$$C_{2}H_{2}(g) + \frac{5}{2}O_{2}(g) \rightarrow 2 CO_{2}(g) + H_{2}O(1) \qquad \Delta H^{\circ} = -1299.4 \text{ kJ}$$

$$2 C(s) + 3 H_{2}(g) \rightarrow C_{2}H_{6}(g) \qquad \Delta H^{\circ} = -84.7 \text{ kJ}$$

$$2 CO_{2}(g) \rightarrow 2 C(s) + 2 O_{2}(g) \qquad \Delta H^{\circ} = 2(393.5 \text{ kJ})$$

$$H_{2}O(1) \rightarrow H_{2}(g) + \frac{1}{2}O_{2}(g) \qquad \Delta H^{\circ} = 285.8 \text{ kJ}$$

$$C_2H_{2(g)} + 2 H_{2(g)} \rightarrow C_2H_{6(g)} \qquad \Delta H^{\circ}_{rxn} = -311.3 \text{ kJ}$$

$$2 \operatorname{CH}_{4(g)} + \frac{1}{2} \operatorname{O}_{2(g)} \rightarrow \operatorname{C}_{2} \operatorname{H}_{6(g)} + \operatorname{H}_{2} \operatorname{O}_{(l)}$$

$$2 \operatorname{CH}_{4}(g) + 4 \operatorname{O}_{2}(g) \rightarrow 2 \operatorname{CO}_{2}(g) + 4 \operatorname{H}_{2} \operatorname{O}(l) \qquad \Delta H^{\circ} = 2(-890.4 \text{ kJ})$$

$$2 \operatorname{C}(s) + 3 \operatorname{H}_{2}(g) \rightarrow \operatorname{C}_{2} \operatorname{H}_{6}(g) \qquad \Delta H^{\circ} = -84.7 \text{ kJ}$$

$$2 \operatorname{CO}_{2}(g) \rightarrow 2 \operatorname{C}(s) + 2 \operatorname{O}_{2}(g) \qquad \Delta H^{\circ} = 2(393.5 \text{ kJ})$$

$$3 \operatorname{H}_{2} \operatorname{O}(l) \rightarrow 2 \operatorname{H}_{2}(g) + \frac{3}{2} \operatorname{O}_{2}(g) \qquad \Delta H^{\circ} = 3(285.8 \text{ kJ})$$

$$2 \operatorname{CH}_{4(g)} + \frac{1}{2} \operatorname{O}_{2(g)} \rightarrow \operatorname{C}_{2} \operatorname{H}_{6(g)} + \operatorname{H}_{2} \operatorname{O}_{(l)} \qquad \Delta H^{\circ}_{rxn} = -221.1 \text{ kJ}$$