

# thermochemistry (2)

#1  $T_f = 22.2^\circ C$

$$-q_{\text{lost}} = q_{\text{gained}}$$

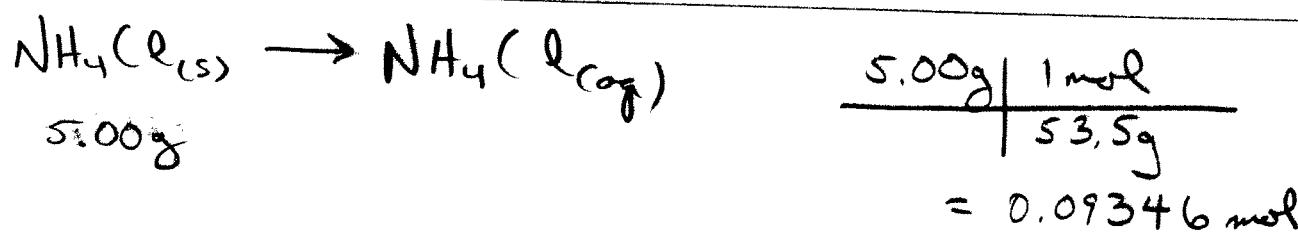
$$-q_{\text{rxn}} = q_{H_2O} + q_{\text{calorimeter}}$$

$$-(65.45 \text{ g})(0.388 \text{ J/g}^\circ C)(-58.8^\circ C) =$$

$$(125 \text{ g})(4.184 \text{ J/g}^\circ C)(2.3^\circ C) + H_c(2.3^\circ C)$$

$$1341 \text{ J} = 1203 \text{ J} + H_c(2.3^\circ C)$$

A)  $H_c = 59.9 \text{ J}^\circ C = 60. \text{ J}^\circ C$



$$-q_{\text{rxn}} = q_{\text{solt}} + q_{\text{calorimeter}}$$

$$-q_{\text{rxn}} = (155 \text{ g soln})(4.18 \text{ J/g}^\circ C)(-1.8^\circ C) + (60. \text{ J}^\circ C)(-1.8^\circ C)$$

$$-q_{\text{rxn}} = -1166 \text{ J} + -108 \text{ J} = (-1274 \text{ J})$$

B)  $q_{\text{rxn}} = 1274 \text{ J}$

C)  $\Delta H = \frac{1274 \text{ J}}{0.09346 \text{ mol}} = 13,600 \text{ J/mol} = 13.6 \text{ kJ/mol}$

A 0.712-g sample of magnesium is burned in excess oxygen in a bomb calorimeter with a heat capacity of 722 J/C° containing 350.0 g of water initially at 21.57°C. The temperature of the apparatus rises to 26.32°C

$$\frac{0.712 \text{ g Mg}}{24.31 \text{ g}} \times \frac{1 \text{ mol}}{1 \text{ mol}} = 0.02929 \text{ mol}$$

A) What is the enthalpy of combustion (in kJ/mol) of magnesium?

$$q_{rxn} = q_w + q_{calorimeter}$$

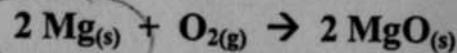
$$q_{rxn} = M_w C_w \Delta T_w + H_c \Delta T_w$$

$$q_{rxn} = (350.0 \text{ g})(4.184 \text{ J/g°C})(4.75 \text{ °C}) + (722 \text{ J/g})(4.75 \text{ °C})$$

$$q_{rxn} = 10385 \text{ J} = 10.385 \text{ kJ}$$

$$q_{rxn} = -10.385 \text{ kJ}$$

B) What is the heat of reaction in kJ for the balanced equation:



$$= -355 \text{ kJ/mol Mg} \times 2 \text{ mol Mg reacting}$$

$$\Delta H_{rxn} = -709 \text{ kJ}$$

#3

$$\frac{0.7000 \text{ L}}{1 \text{ L}} \times \frac{0.500 \text{ mol HCl}}{1 \text{ L}} = 0.350 \text{ mol HCl} \times \frac{1 \text{ mol Ba(OH)}_2}{2 \text{ mol HCl}} = 0.175 \text{ mol Ba(OH)}_2 \text{ needed}$$

$$\frac{0.3000 \text{ L}}{1 \text{ L}} \times \frac{0.500 \text{ mol}}{1 \text{ L}} = 0.150 \text{ mol Ba(OH)}_2$$

L.R.

$$1000 \text{ mL soln} = 1000 \text{ g soln}$$

$$\frac{0.150 \text{ mol Ba(OH)}_2}{1 \text{ mol Ba(OH)}_2} \times -118 \text{ kJ} = -17.7 \text{ kJ}$$

A)

(17,700 J produced)

$$q_{rxn} = -17,700 \text{ J}$$

$$q_{rxn} = q_{soln} + q_{calorimeter}$$

$$q_{rxn} = m_{soln} C_{soln} \Delta T_{soln} + H_c \Delta T_{soln}$$

$$-(-17,700 \text{ J}) = (1000 \text{ g})(4.184 \text{ J/g°C})\Delta T + (180.3 \text{ J/g})\Delta T$$

$$17700 \text{ J} = (4364.3 \text{ J})\Delta T$$

B)

$$\Delta T = 4.06 \text{ }^{\circ}\text{C} = T_f - 25.0 \text{ }^{\circ}\text{C}$$

$$T_f = 25.0 \text{ }^{\circ}\text{C} + 4.06 \text{ }^{\circ}\text{C} = 29.1 \text{ }^{\circ}\text{C}$$

$$\Delta H = \frac{q_{rxn}}{\text{mol reacting}}$$

$$\Delta H = \frac{-10.385 \text{ kJ}}{0.02929 \text{ mol}}$$

$$\Delta H = -355 \text{ kJ/mol}$$