

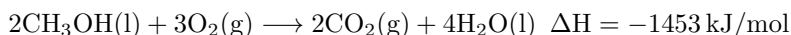
Non Sibi High School

Andover's Chem 250: Introductory/Basic Chemistry

Chapter 13, Review Quiz 1 Answers

1

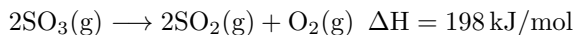
If 125 kilograms of methanol, CH_3OH , is burned according to the combustion equation below, how much heat will be released?



$$125 \text{ kg} \left(\frac{1000 \text{ g}}{1 \text{ kg}} \right) \left(\frac{1 \text{ mol CH}_3\text{OH}}{32.04 \text{ g}} \right) \left(\frac{-1453 \text{ kJ}}{2 \text{ mol CH}_3\text{OH}} \right) = -2.83 \times 10^6 \text{ kJ}$$

2

If 3.55 kJ of heat are absorbed during the decomposition reaction below, how many milliliters of sulfur trioxide gas, measured at 22°C and 712 mmHg, will decompose?



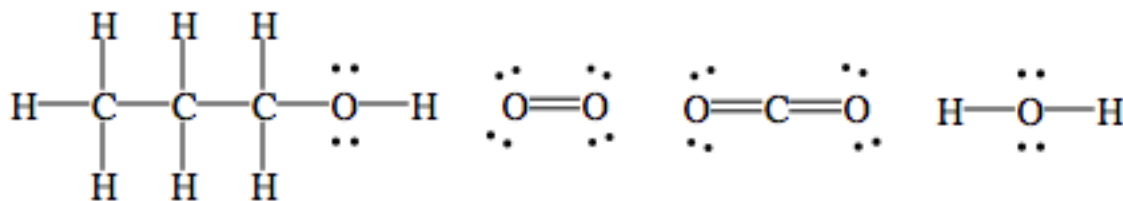
$$3.55 \text{ kJ} \left(\frac{2 \text{ mol SO}_3}{198 \text{ kJ}} \right) = 0.03586 \text{ mol SO}_3$$

$$\frac{0.03586 \text{ mol} \times 0.0821 \frac{\text{L}\cdot\text{atm}}{\text{mol}\cdot\text{K}} \times (22 + 273) \text{ K}}{\frac{712}{760} \text{ atm}} = 0.927 \text{ L} \left(\frac{1000 \text{ mL}}{1 \text{ L}} \right) = 927 \text{ mL SO}_3$$

3

For the reaction $2\text{C}_3\text{H}_7\text{OH}(g) + 9\text{O}_2(g) \longrightarrow 6\text{CO}_2(g) + 8\text{H}_2\text{O}(g)$, estimate ΔH using average bond energies.

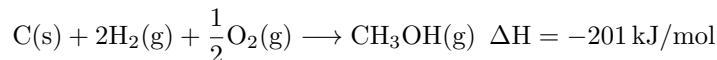
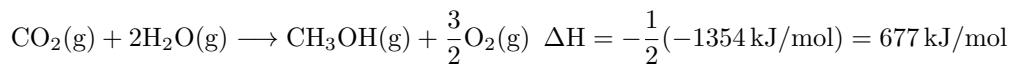
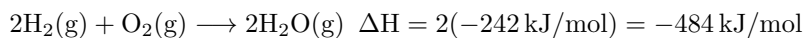
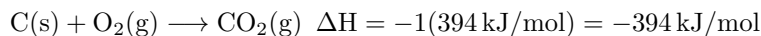
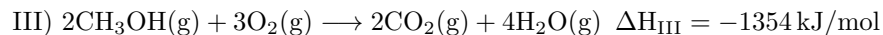
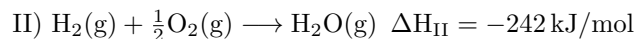
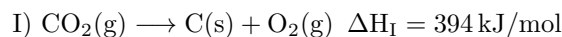
Lewis structures:



$$\begin{aligned}
 \Delta H \text{ (estimated)} &= \\
 (2 \times 7)(\text{C} - \text{H}) &+ (2 \times 2)(\text{C} - \text{C}) + (2 \times 1)(\text{C} - \text{O}) + (2 \times 1)(\text{O} - \text{H}) + (9 \times 1)(\text{O} = \text{O}) \\
 &- (6 \times 2)(\text{C} = \text{O}) - (8 \times 2)(\text{O} - \text{H}) \\
 &= 14(414) + 4(347) + 2(360) + 2(464) + 9(498) - 12(745) - 16(464) \\
 &= -3050 \text{ kJ/mol}
 \end{aligned}$$

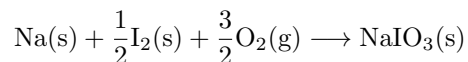
4

Calculate ΔH for the reaction $\text{C(s)} + 2\text{H}_2(\text{g}) + \frac{1}{2}\text{O}_2(\text{g}) \longrightarrow \text{CH}_3\text{OH}(\text{g})$ using the following three reactions:



5

Write the balanced formation reaction, including physical states, for solid sodium iodate, NaIO_3 .



Compound	ΔH_f° (kJ/mol)
NO(g)	90.
NO ₂ (g)	33

6

Calculate ΔH° for the reaction $2\text{NO}(\text{g}) + \text{O}_2(\text{g}) \longrightarrow 2\text{NO}_2(\text{g})$ using the following information:

$$\Delta H^\circ = 2(33) - 2(90) - 1(0) = -114 \text{ kJ/mol}$$

7

The specific heat of magnesium metal is $1.05 \text{ J/g}\cdot^\circ\text{C}$. How much heat in kilojoules is lost when a 225 gram sample of magnesium metal is cooled from 625°C to 125°C ?

$$q = 225 \text{ g} \left(\frac{1.05 \text{ J}}{\text{g}\cdot^\circ\text{C}} \right) (125 - 625)^\circ\text{C} \left(\frac{1 \text{ kJ}}{1000 \text{ J}} \right) = -118 \text{ kJ}$$

8

In an insulated calorimeter, a 475 gram piece of tin metal originally at 132°C was added to 135 grams of water originally at 19°C . The final temperature of the tin-water mixture was 36°C . Determine the specific heat of tin.

$$\begin{aligned} q_{\text{Sn lost}} &= -q_{\text{water gained}} \\ 475 \text{ g}(s_{\text{Sn}})(36 - 132)^\circ\text{C} &= -135 \text{ g} \left(\frac{4.18 \text{ J}}{\text{g}\cdot^\circ\text{C}} \right) (36 - 19)^\circ\text{C} \\ s_{\text{Sn}} &= 0.21 \text{ J/g}\cdot^\circ\text{C} \end{aligned}$$

9

The specific heat of tungsten metal is $0.13 \text{ J/g}\cdot^\circ\text{C}$. In an insulated calorimeter, a 955 gram piece of tungsten metal originally at 375°C was added to 725 grams of water originally at 18°C . Determine the final temperature of the tungsten-water mixture.

$$\begin{aligned} q_{\text{W lost}} &= -q_{\text{water gained}} \\ 955 \text{ g} \left(\frac{0.13 \text{ J}}{\text{g}\cdot^\circ\text{C}} \right) (t_{\text{final}} - 375^\circ\text{C}) &= -725 \text{ g} \left(\frac{4.18 \text{ J}}{\text{g}\cdot^\circ\text{C}} \right) (t_{\text{final}} - 18^\circ\text{C}) \\ t_{\text{final}} &= 32^\circ\text{C} \end{aligned}$$

10

In an insulated calorimeter, 18.2 grams of solid cesium hydroxide at 22.3°C was dissolved in 135.7 grams of water also at 22.3°C, after which the final temperature of the mixed solution was 36.9°C. If the specific heat of the mixed solution was 3.87 J/g·°C, determine ΔH for the dissolving process $\text{CsOH}(s) \rightarrow \text{CsOH}(aq)$ in kJ/mol CsOH.

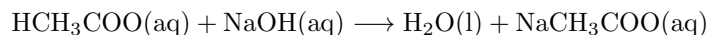
$$q_{\text{rxn lost}} = -q_{\text{soln gained}} = -(18.2 + 135.7)\text{g} \left(\frac{3.87 \text{ J}}{\text{g} \cdot ^\circ\text{C}} \right) (36.9 - 22.3)^\circ\text{C} \left(\frac{1 \text{ kJ}}{1000 \text{ J}} \right) = -8.696 \text{ kJ}$$

$$n_{\text{CsOH}} = 18.2 \text{ g} \left(\frac{1 \text{ mol}}{149.9 \text{ g}} \right) = 0.1214 \text{ mol CsOH}$$

$$\Delta H = \frac{-8.696 \text{ kJ}}{0.1214 \text{ mol CsOH}} = -71.6 \text{ kJ/mol CsOH}$$

11

In an insulated calorimeter, 55.7 mL of 1.91 M acetic acid was mixed with 62.6 mL of 1.83 M sodium hydroxide, with both solutions originally at 18.2°C. The final temperature of the mixed solutions was 30.1°C. The density of the mixed solutions was 1.03 g/mL and the specific heat of the mixed solutions was 3.96 J/g·°C. Write a balanced molecular equation, including physical states, and determine ΔH for the neutralization reaction in kJ/mol of water formed.



$$q_{\text{rxn lost}} = -q_{\text{soln gained}} = -(55.7 + 62.6)\text{mL} \left(\frac{1.03 \text{ g}}{1 \text{ mL}} \right) \left(\frac{3.96 \text{ J}}{\text{g} \cdot ^\circ\text{C}} \right) (30.1 - 18.2)^\circ\text{C} \left(\frac{1 \text{ kJ}}{1000 \text{ J}} \right) = -5.742 \text{ kJ}$$

$$55.7 \text{ mL} \left(\frac{1 \text{ L}}{1000 \text{ mL}} \right) \left(\frac{1.91 \text{ mol HCH}_3\text{COO}}{1 \text{ L}} \right) \left(\frac{1 \text{ mol H}_2\text{O}}{1 \text{ mol HCH}_3\text{COO}} \right) = 0.1064 \text{ mol H}_2\text{O}$$

$$62.6 \text{ mL} \left(\frac{1 \text{ L}}{1000 \text{ mL}} \right) \left(\frac{1.83 \text{ mol NaOH}}{1 \text{ L}} \right) \left(\frac{1 \text{ mol H}_2\text{O}}{1 \text{ mol NaOH}} \right) = 0.1147 \text{ mol H}_2\text{O}$$

HCH₃COO produces less H₂O, so HCH₃COO is the limiting reagent and 0.1064 mol H₂O is formed.

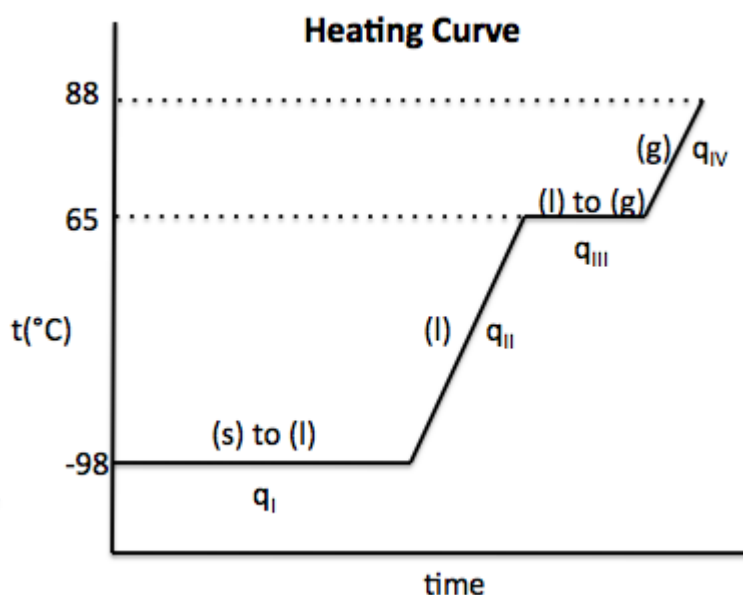
$$\Delta H = \frac{-5.742 \text{ kJ}}{0.1064 \text{ mol H}_2\text{O}} = -54.0 \text{ kJ/mol H}_2\text{O}$$

12

Consider the following data for methanol, CH₃OH:

melting point = -98°C
 boiling point = 65°C
 $\Delta H_{\text{fusion}} = 3.2 \text{ kJ/mol}$
 $\Delta H_{\text{vaporization}} = 38 \text{ kJ/mol}$
 specific heat of liquid methanol = $2.5 \text{ J/g}\cdot^{\circ}\text{C}$
 specific heat of methanol vapor = $1.7 \text{ J/g}\cdot^{\circ}\text{C}$

Sketch a heating curve that depicts solid methanol at -98°C being heated to 88°C and then calculate the total amount of heat in kilojoules absorbed when 77 grams of methanol undergoes this process.



I) solid methanol at -98°C to liquid methanol at -98°C :

$$q_{\text{I}} = 77 \text{ g} \left(\frac{1 \text{ mol}}{32.04 \text{ g}} \right) \left(\frac{3.2 \text{ kJ}}{1 \text{ mol}} \right) = 7.69 \text{ kJ}$$

II) liquid methanol at -98°C to liquid methanol at 65°C :

$$q_{\text{II}} = 77 \text{ g} \left(\frac{2.5 \text{ J}}{\text{g}\cdot^{\circ}\text{C}} \right) (65 - (-98))^{\circ}\text{C} \left(\frac{1 \text{ kJ}}{1000 \text{ J}} \right) = 31.4 \text{ kJ}$$

III) liquid methanol at 65°C to methanol vapor at 65°C :

$$q_{\text{III}} = 77 \text{ g} \left(\frac{1 \text{ mol}}{32.04 \text{ g}} \right) \left(\frac{38 \text{ kJ}}{1 \text{ mol}} \right) = 91.3 \text{ kJ}$$

IV) methanol vapor at 88°C to methanol vapor at 88°C :

$$q_{IV} = 77 \text{ g} \left(\frac{1.7 \text{ J}}{\text{g} \cdot ^\circ \text{C}} \right) (88 - 65)^\circ \text{C} \left(\frac{1 \text{ kJ}}{1000 \text{ J}} \right) = 3.01 \text{ kJ}$$

$$q_{\text{total}} = 7.69 + 31.8 + 91.3 + 3.01 = 133 \text{ kJ}$$



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