## **Exercise 03**

Name:

Key

Consider the combustion of methane in air:

$$\operatorname{CH}_4(g) + 2\operatorname{O}_2(g) \to \operatorname{CO}_2(g) + 2\operatorname{H}_2\operatorname{O}(g)$$

(a) Determine the  $\Delta H_{\rm rxn}$  using the values of  $\Delta H_{\rm f}^{\rm o}$  given.

Species	ΔH <sup>o</sup> <sub>f</sub> (kJ/mol)
$CH_4(g)$	-74.6
$\mathrm{H}_{2}\mathrm{O}\left(g ight)$	-241.8
$\mathrm{CO}_{2}\left(g ight)$	-393.5

- (b) If  $1.00 \times 10^6$  J of heat were released during a combustion of methane, how many moles of CO<sub>2</sub> gas were produced?
- (a) We can determine the heat of the reaction (or heat of combustion) through the following expression. Recall that  $\Delta H_{\rm f}^{\rm o} = 0$  kJ/mol for elements in their elemental states (e.g., O<sub>2</sub> gas).

$$\Delta H_{\rm rxn} = n_{\rm CO_2} \Delta H_{\rm f,CO_2}^{\rm o} + n_{\rm O_2} \Delta H_{\rm f,O_2}^{\rm o} - n_{\rm CH_4} \Delta H_{\rm f,CH_4}^{\rm o} - n_{\rm O_2} \Delta H_{\rm f,O_2}^{\rm o}$$
  
= (1 mol CO<sub>2</sub>)  $\left(-393.5 \frac{\rm kJ}{\rm mol}\right)$  + (2 mol H<sub>2</sub>O)  $\left(-241.8 \frac{\rm kJ}{\rm mol}\right)$  - (1 mol CH<sub>4</sub>)  $\left(-74.6 \frac{\rm kJ}{\rm mol}\right)$  -  $\left(0 \frac{\rm kJ}{\rm mol}\right)$   
 $\Delta H_{\rm rxn} = -802.5 \,\rm kJ/mol$ 

(b) The heat of reaction/combustion ( $\Delta H_{rxn}$ ) tells us that for every 1 mol CH<sub>4</sub> (*g*) combusted, 802.5 kJ of heat are released. From stoichiometry, we also know that 1 mol of CH<sub>4</sub> (*g*) produces 1 mol CO<sub>2</sub> (*g*). So, if we released 1.00 × 10<sup>6</sup> J:

$$1.00 \times 10^6 \text{ J} \times \frac{1 \text{ kJ}}{1000 \text{ J}} \times \frac{1 \text{ mol } \text{CH}_4}{802.5 \text{ kJ}} \times \frac{1 \text{ mol } \text{CO}_2}{1 \text{ mol } \text{CH}_4} = 1.25 \text{ mol } \text{CO}_2$$