

## HL Questions on Entropy and spontaneity

- 1. Given that the standard entropy of gaseous water is 189 J K<sup>-1</sup> mol<sup>-1</sup> and the standard entropy of liquid water is 69.9 J K<sup>-1</sup> mol<sup>-1</sup> calculate the standard entropy change when one mol of gaseous water condenses to the liquid state,  $H_2O(g) \rightarrow H_2O(I)$ .
- **2. (a)** Predict the sign (positive or negative) for the entropy change for the reaction:

$$2NO_2(g) \rightarrow N_2O_4(g)$$

(b) Explain why the entropy change for the above reaction is considerably more than for the reaction:

$$H_2(g) + I_2(g) \rightarrow 2HI(g)$$

**3.** Use the information below to calculate the entropy change of formation of copper(I) oxide.

| Substance          | Entropy / J K <sup>-1</sup> mol <sup>-1</sup> |
|--------------------|---|
| Cu(s)              | 33.3  |
| O <sub>2</sub> (g) | 205   |
| Cu₂O(s)            | 101   |

4. You are unlikely to use orange crystals of ammonium dichromate, (NH<sub>4</sub>)<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>(s), in a school laboratory as it is a highly toxic substance. It is also carcinogenic and can explode when heated.

When it thermally decomposes (see image on right) it is both oxidised and reduced according to the equation:

$$(NH_4)_2Cr_2O_7(s) \rightarrow Cr_2O_3(s) + N_2(g) + 4 H_2O(g)$$

(a) State what is being oxidized and what is being reduced.

Clearly the thermal decomposition of ammonium dichromate is very exothermic ( $\Delta H \ominus = -1794 \text{ kJ mol}^{-1}$ ). What can you also deduce about the change in entropy?



Thermal decomposition of ammonium dichromate (image from Flickr)



## (HL Questions on Entropy and spontaneity continued)

**5.** In a blast furnace limestone (calcium carbonate) is decomposed into calcium oxide and carbon dioxide according to the equation:

$$CaCO_3(s) \rightarrow CaO(s) + CO_2(g)$$

- (a) The standard enthalpies of formation of CaCO<sub>3</sub>(s), CaO(s) and CO<sub>2</sub>(g) are − 1207 kJ mol<sup>-1</sup>, − 635 kJ mol<sup>-1</sup> and − 394 kJ mol<sup>-1</sup> respectively.
   Calculate the standard enthalpy change, ΔH⊕, for the thermal decomposition of calcium carbonate.
- (b) The standard entropy values for CaCO<sub>3</sub>(s), CaO(s) and CO<sub>2</sub>(g) are 92.9 J K<sup>-1</sup> mol<sup>-1</sup>, 40.0 J K<sup>-1</sup> mol<sup>-1</sup> and 214 J K<sup>-1</sup> mol<sup>-1</sup> respectively.
  Calculate the change in entropy, ΔS<sup>⊕</sup>, for the thermal decomposition of calcium carbonate.
- (c) Calculate the change in free energy,  $\Delta G \ominus$ , for the thermal decomposition of calcium carbonate at 298 K.
- (d) State whether the thermal decomposition of calcium carbonate is spontaneous or non-spontaneous at 298K.
- **(e)** Explain why limestone cliffs are thermodynamically stable whereas limestone decomposes readily in a blast furnace.
- 6. Propene reacts with hydrogen at 180 °C in the presence of a nickel catalyst to give propane.

$$C_3H_6(g) + H_2(g) \rightarrow C_3H_8(g)$$

- (a) Use the information below to determine the value for the free enthalpy change for this reaction at 180 °C.
- **(b)** At 180 °C this reaction is spontaneous.

  Determine the temperature above which the reaction becomes non-spontaneous.

| Substance                         | ΔH <sub>f</sub> <sup>e</sup> / kJ mol <sup>-1</sup> | S <sup>0</sup> / J K <sup>-1</sup> mol <sup>-1</sup> |
|-----------------------------------|---|--|
| H <sub>2</sub> (g)                | 0   | 131  |
| C₃H <sub>6</sub> (g)              | + 20.4  | 267  |
| C <sub>3</sub> H <sub>8</sub> (g) | - 104   | 270  |