

1. The energy q , required to heat a substance of mass m by a temperature ΔT is given by the equation:

$$q = c \times m \times \Delta T$$

where c is the specific heat capacity – a property of the substance involved.

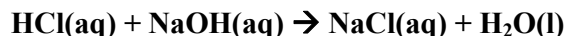
If $q = 78.2 \text{ J}$, $m = 45.6 \text{ g}$ and $\Delta T = 13.3 \text{ K}$, then

$$c = \frac{q}{m\Delta T} = \frac{78.2 \text{ J}}{(45.6 \text{ g}) \times (13.3 \text{ K})} = 0.129 \text{ J K}^{-1} \text{ g}^{-1}$$

The atomic mass of lead is 207.2 g mol^{-1} . The molar heat capacity, C , is therefore:

$$C = (0.129 \text{ J K}^{-1} \text{ g}^{-1}) \times (207.2 \text{ g mol}^{-1}) = 26.7 \text{ J K}^{-1} \text{ mol}^{-1}$$

2. The neutralization reaction is a 1:1 reaction with chemical equation:



After the reactants mix the solution has a volume of 200 mL. Assuming the solution the same density as pure water, this corresponds to a mass of:

$$\begin{aligned} \text{mass (g)} &= \text{density (g mL}^{-1}\text{)} \times \text{volume (mL)} \\ &= (0.997 \text{ g mL}^{-1}) \times (200 \text{ mL}) = 199 \text{ g} \end{aligned}$$

The temperature change, $\Delta T = (31.1 \text{ }^\circ\text{C} - 24.6 \text{ }^\circ\text{C}) = 6.7 \text{ K}$. The heat capacity of water is $4.184 \text{ J K}^{-1} \text{ g}^{-1}$. The heat change is therefore:

$$\begin{aligned} q &= c \times m \times \Delta T = (4.184 \text{ J K}^{-1} \text{ g}^{-1}) \times (199 \text{ g}) \times (6.7 \text{ K}) \\ &= 5600 \text{ J or } 5.6 \text{ kJ (heat released)} \end{aligned}$$

The number of moles of H^+ and OH^- present are both the same:

$$\begin{aligned} \text{number of moles} &= \text{concentration (M or mol L}^{-1}\text{)} \times \text{volume (L)} \\ &= (1.0 \text{ mol L}^{-1}) \times (0.100 \text{ mL}) = 0.1 \text{ mol} \end{aligned}$$

The heat change for a mole would therefore be $5.6 \text{ kJ} / 0.1 \text{ mol} = 56 \text{ kJ mol}^{-1}$. The reaction gives out heat (as the temperature rises) so it is an exothermic reaction with a negative enthalpy change: $\Delta H = -56 \text{ kJ mol}^{-1}$

3. (a) $2(\text{NH}_2)_2\text{CO(s)} + 3\text{O}_2(\text{g}) \rightarrow 4\text{H}_2\text{O(l)} + 2\text{N}_2(\text{g}) + 2\text{CO}_2(\text{g})$
 (b) The molar mass of urea, $(\text{NH}_2)_2\text{CO}$, is:

$$\text{molar mass} = 2 \times (14.01 \text{ (N)} + 2 \times 1.008 \text{ (H)}) + 12.01 \text{ (C)} + 16.00 \text{ (O)} = 60.062 \text{ g mol}^{-1}$$

The number of moles present in 6.006 g is therefore:

$$\text{number of moles} = \frac{\text{mass}}{\text{molar mass}} = \frac{6.006 \text{ g}}{60.062 \text{ g mol}^{-1}} = 0.1000 \text{ mol}$$

As this amount releases 63.4 kJ, the amount release by a mole is:

$$\Delta_{\text{comb}}H^{\circ} = -\frac{63.4 \text{ kJ}}{0.1000 \text{ mol}} = -634 \text{ kJ mol}^{-1}$$

The reaction is *exothermic* as energy is *liberated* or *released*: ΔH° is negative.

- (c) The chemical equation given in (a) corresponds to combusting *two* moles of $(\text{NH}_2)_2\text{CO}(\text{s})$. The enthalpy of the reaction is therefore:

$$\Delta_{\text{rxn}}H^{\circ} = 2 \times \Delta_{\text{comb}}H^{\circ} = 2 \times -634 = -1268 \text{ kJ mol}^{-1}$$

Using $\Delta_{\text{rxn}}H^{\circ} = \sum m\Delta_f H^{\circ}(\text{products}) - \sum n\Delta_f H^{\circ}(\text{reactants})$,

$$\begin{aligned} \Delta_{\text{rxn}}H^{\circ} = & [4\Delta_f H^{\circ}(\text{H}_2\text{O}(\text{l})) + 2\Delta_f H^{\circ}(\text{N}_2(\text{g})) + 2\Delta_f H^{\circ}(\text{CO}_2(\text{g})) \\ & - [2\Delta_f H^{\circ}((\text{NH}_2)_2\text{CO}(\text{s})) + 3\Delta_f H^{\circ}(\text{O}_2(\text{g}))] \end{aligned}$$

As $\Delta_f H^{\circ} = 0$ for an element in its standard state, this becomes:

$$\begin{aligned} \Delta_{\text{rxn}}H^{\circ} = & [(4 \times -285) + (2 \times 0) + (2 \times -393)] \\ & - [2\Delta_f H^{\circ}((\text{NH}_2)_2\text{CO}(\text{s})) + (3 \times 0)] = -1268 \text{ kJ mol}^{-1} \end{aligned}$$

Hence, $\Delta_f H^{\circ}((\text{NH}_2)_2\text{CO}(\text{s})) = -329 \text{ kJ mol}^{-1}$.

4. The temperature change, $\Delta T = 27.828 - 25.000 = 2.828 \text{ }^{\circ}\text{C}$.

As the temperature increases, heat is lost from the system and gained by the surroundings. The reaction is exothermic.

The heat change is given by:

$$q = c \times \Delta T = (96.5 \text{ kJ }^{\circ}\text{C}^{-1}) \times (2.828 \text{ }^{\circ}\text{C}) = 273 \text{ kJ}$$

This is the heat change for 0.0100 mol of propane, the internal energy change for 1 mol is therefore:

$$\Delta U = -\frac{273 \text{ kJ}}{0.0100 \text{ mol}} = -2.73 \times 10^5 \text{ kJ mol}^{-1}$$

The negative sign indicates that the system is losing energy to the surroundings. As a bomb calorimeter has a constant volume, the heat change is related to a change in internal energy, ΔU .