

# THE OXIDES OF PERIOD 3 ELEMENTS

### 1. Formation of oxides

All the elements in Period 3 except chlorine and argon combine directly with oxygen to form oxides.

$$4Na(s) + O_2(g) \rightarrow 2Na_2O(s)$$

Na<sub>2</sub>O is an ionic oxide.

$$2Mg(s) + O_2(g) \rightarrow 2MgO(s)$$

MgO is also an ionic oxide.

$$4Al(s) + 3O_2(g) \rightarrow 2Al_2O_3(s)$$

Al<sub>2</sub>O<sub>3</sub> is mostly ionic, but there is significant covalent character.

$$Si(s) + O_2(g) \rightarrow SiO_2(s)$$

SiO<sub>2</sub> is a giant covalent oxide.

$$P_4(s) + 5O_2(g) \rightarrow P_4O_{10}(s)$$

 $P_4O_{10}$  is a molecular covalent oxide. The oxidation number of P in this oxide is +5.

$$S(s) + O_2(g) \rightarrow SO_2(g)$$

SO<sub>2</sub> is a molecular covalent oxide.

Another oxide,  $SO_3$  is formed in a reversible process when  $SO_2$  and  $O_2$  are heated with a  $V_2O_5$  catalyst (the Contact Process)

## 2. Physical properties of oxides

The physical properties of these oxides depend on the type of bonding.

 $Na_2O$ ,  $Al_2O_3$  and MgO are ionic oxides and hence have a high melting point. MgO and  $Al_2O_3$  have a higher melting point than  $Na_2O$  since the charges are higher, resulting in a stonger attraction between the ions.

SiO<sub>2</sub> has a giant covalent structure and hence a high melting point. There are strong covalent bonds between all the atoms and thus lots of energy is required to break them.

 $P_4O_{10}$  and  $SO_3$  are molecular covalent and so only intermolecular forces exist between the molecules. The melting points are thus much lower.  $P_4O_{10}$  is a much bigger molecule than  $SO_3$  and so has a much higher melting point, as the van der Waal's forces are stronger.

| Element            | Na                | Mg    | Al              | Si                | P                     | S                     |
|--------------------|-------------------|-------|-----------------|-------------------|-----------------------|-----------------------|
| Formulae of        | Na <sub>2</sub> O | MgO   | $Al_2O_3$       | SiO <sub>2</sub>  | $P_4O_{10}$           | $SO_3$                |
| oxide              |                   |       |                 |                   |                       |                       |
| Structure of oxide | Ionic             | Ionic | Mostly<br>ionic | Giant<br>covalent | Molecular<br>covalent | Molecular<br>covalent |
|                    |                   |       |                 |                   |                       |                       |
| Melting point of   | 1275              | 2852  | 2072            | 1703              | 300                   | -10                   |
| oxide /°C          |                   |       |                 |                   |                       |                       |

### 3. Acid-base character of oxides

Ionic oxides contain the  $O^{2-}$  ion. This is a strongly basic ion which reacts with water to produce hydroxide ions:

$$O^{2}$$
- $(aq) + H_2O(l) \rightarrow 2OH$ - $(aq)$ 

Thus all ionic oxides are BASIC.

Covalent oxides do not contain ions, but have a strongly positive dipole on the atom which is not oxygen. This attracts the lone pair on water molecules, releasing H<sup>+</sup> ions:

$$MO(s) + H2O(1) \rightarrow MO(OH)^{-}(aq) + H^{+}(aq)$$

Thus all covalent oxides are ACIDIC.

Intermediate oxides can react in either of the above ways, depending on the conditions. They can thus behave as either acids or bases and are thus AMPHOTERIC.

 $Na_2O$  is a basic oxide. It dissolves in water to give an alkaline solution (pH = 14). It also reacts with acids:

$$Na_2O(s) + H_2O(l) \rightarrow 2NaOH(aq)$$
  
 $Na_2O(s) + 2H^+(aq) \rightarrow 2Na^+(aq) + H_2O(l)$ 

MgO is a basic oxide. It is only slightly soluble in water and so the solution is only slightly alkaline (pH = 9). It reacts readily with acids:

$$MgO(s) + H_2O(l) == Mg(OH)_2(s) == Mg(OH)_2(aq)$$
  
 $MgO(s) + 2H^+(aq) \rightarrow Mg^{2+}(aq) + H_2O(l)$ 

 $Al_2O_3$  is an amphoteric oxide. It is insoluble in water (pH = 7) but dissolves in both acids and alkalis:

$$Al_2O_3(s) + 6H^+(aq) \rightarrow 2Al^{3+}(aq) + 3H_2O(l)$$
  
 $Al_2O_3(s) + 3H_2O(l) + 6OH^-(aq) \rightarrow 2Al(OH)_6^{3-}(aq)$   
 $Al_2O_3(s) + 3H_2O(l) + 2OH^-(aq) \rightarrow 2Al(OH)_4^-(aq)$ 

 $SiO_2$  is an acidic oxide. It is insoluble in water (pH = 7) but dissolves in hot concentrated alkalis:

$$SiO_2(s) + 2OH^-(aq) \rightarrow SiO_3^{2-}(aq) + H_2O(l)$$

P<sub>4</sub>O<sub>10</sub> is an acidic oxide. It dissolves in water to give acidic solutions and is also soluble in alkalis:

$$P_4O_{10}(s) + 6H_2O(1) \rightarrow 4H_3PO_4(aq)$$
  $pH = 3$   
 $P_4O_{10}(s) + 12OH^-(aq) \rightarrow 4PO_4^{3-}(aq) + 6H_2O(1)$ 

SO<sub>2</sub> and SO<sub>3</sub> are acidic oxides. They dissolve in water to give acidic solutions, and also react with alkalis:

$$SO_2(g) + H_2O(l) == H_2SO_3(aq), \quad pH = 2$$
  
 $SO_3(g) + H_2O(l) \rightarrow H_2SO_4(aq), \quad pH = 1$   
 $SO_2(g) + 2OH^-(aq) \rightarrow SO_3^{2-}(aq) + H_2O(l)$   
 $SO_3(g) + 2OH^-(aq) \rightarrow SO_4^{2-}(aq) + H_2O(l)$ 

SO<sub>2</sub> is a waste gas in many industrial processes. It is harmful because it dissolves in rain water to give acid rain. It can be removed from waste gases because it dissolves in alkali and so it is passed through an alkaline solution in waste gas outlets to minimise the amount which escapes into the atmosphere.

The acid-base properties of the oxides of Period 3 can be summarised in the following table:

| Element            | Na                | Mg    | Al                             | Si               | P                              | S                  |
|--------------------|-------------------|-------|--------------------------------|------------------|--------------------------------|--------------------|
| Formulae of        | Na <sub>2</sub> O | MgO   | Al <sub>2</sub> O <sub>3</sub> | SiO <sub>2</sub> | P <sub>4</sub> O <sub>10</sub> | $SO_2$             |
| oxides             |                   |       |                                |                  |                                | $SO_3$             |
| Acid-base          | Basic             | Basic | Amphoteric                     | Acidic           | Acidic                         | Acidic             |
| character of oxide |                   |       |                                |                  |                                |                    |
| pH of solution     | 12 - 14           | 8 - 9 | 7 (insoluble)                  | 7                | 2 - 4                          | 2 - 4              |
| when dissolved in  |                   |       | (msoluble)                     | (insoluble)      |                                | (SO <sub>2</sub> ) |
| water              |                   |       |                                |                  |                                | 1 - 3              |
|                    |                   |       |                                |                  |                                | $(SO_3)$           |

The oxides therefore become more acidic on moving from left to right in the periodic table.

### THE REACTION OF PERIOD 3 ELEMENTS WITH WATER

Na, Mg, Al and Si are more electropositive than H and can reduce the water to hydrogen gas:

Na reacts vigorously with water to give the hydroxide and hydrogen:

 $2Na(s) + 2H_2O(1) \rightarrow 2NaOH(aq) + H_2(g)$ 

The resulting solution is strongly alkaline, and will have a pH of 14.

Mg reacts with steam to give the oxide and hydrogen:

 $Mg(s) + H_2O(g) \rightarrow MgO(s) + H_2(g)$ 

The resulting solution is weakly alkaline, since the oxide is slightly basic (pH = 9).

Al and Si also react with steam under certain conditions.

P, S and Cl<sub>2</sub> do not reduce water to hydrogen gas. Phosphorus and sulphur do not react with water but chlorine will disproportionate to give an acidic solution:

 $Cl_2(g) + H_2O(1) \rightarrow HClO(aq) + HCl(aq)$ 

The resulting solution contains HCl(aq) and is thus acidic (pH = 2).

The reactivity of the elements of period 3 towards water thus decreases from Na to Si, and then increases from P to Cl. The resulting solutions become increasingly acidic.