Patterns in the Chemical World

Is this revision??

Variation of properties from Li to Ar

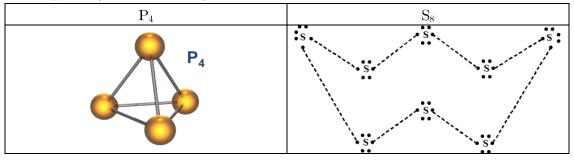
PERIODIC TABLE OF ELEMENTS

Chemical Group Block



	${f Li}$	${f Be}$	\mathbf{B}	\mathbf{C}	$\mathbf N$	O	${f F}$	Ne
Chemical	Li	Be	В	С	N_2	O_2	F_2	Ne
formula of								
elemental form								
Type	Me	etal	Semi- metal		Γ	Non-metals	5	
Nature of	Metallic bond Covalent bond Covalent bond (within molecule exc		ıle except					
bonding					Ne) + Va	an der waa	l's forces b	etween
				molecules				
	Na	Mg	Al	Si	P	\mathbf{S}	Cl	Ar
Chemical	Na	Mg	Al	Si	P_4	S_8	Cl_2	Ne
formula of								
elemental form								
\mathbf{Type}		Metal		Semi-		Non-r	metals	
				metal				
Nature of	Metallic bond		Covalent	Covalent bond (within molecule except				
bonding				bond	Ar) + Va	an der waa	l's forces b	etween
					molecules	3		

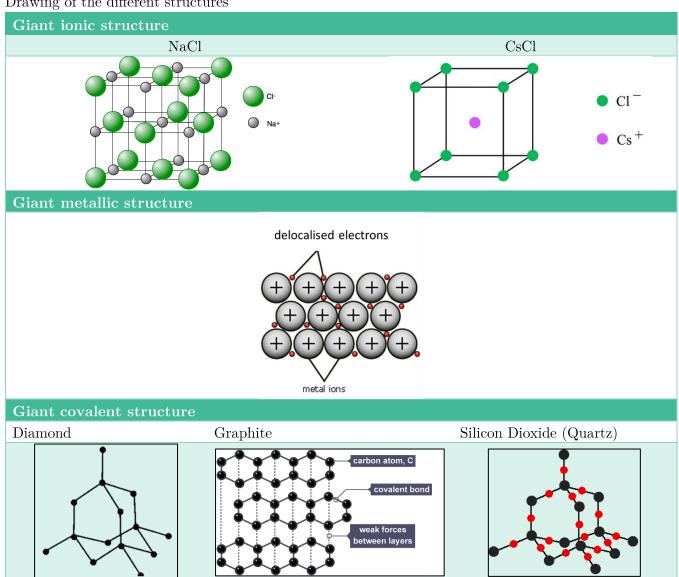
Structures of phosphorus and sulphur. In reality both elements have several allotropes, the forms P_4 and S_8 are the typical (and most stable) forms of the two elements.



Relationships between structures and electrical conductivity

Type of structure	Types of bond(s) present	Electrical conductivity	
Giant metallic structure	Metallic bond between metal	Can conduct electricity	
	ions and delocalized electrons		
Giant ionic structure	Ionic bonds between ions of	Can conduct electricity in	
	opposite polarities of charge	aqueous or molten state only	
Simple molecular structure	Covalent bonds within	Cannot conduct electricity in	
	molecules; Van der waal's forces	any state	
	between molecules		
Giant covalent structure	Covalent bonds between atoms	Cannot conduct electricity	
		(except graphite)	

Drawing of the different structures



Checkpoint

1. Explain why the melting point of diamond is higher than that of P. 2. Explain why solid sodium chloride cannot conduct electricity while solid sodium can conduct electricity. 3. Arrange Mg, Si, Cl and Ar in decreasing order of boiling point, explain your answer by making appropriate comparisons between their bonding and structures.

Properties of oxides of third period elements

	Na	Mg	Al	Si	P	S	Cl
Group	I	II	III	IV	V	VI	VII
Chemical							
formula of	$ m Na_2O$	MgO	$\mathrm{Al_2O_3}$	${ m SiO_2}$	P_4O_{10}	SO_2	OCl_2
oxide	Na ₂ O						
Nature of		т • 1	1		C 1	, 1 1	
bonding	Ionic bond			Covalent bond			
Structure of				Giant			
oxide	Giant ionic structure		covalent	Simple molecular structure			
Acid-base property	Basic Amphoteric		Acidic				

Basic oxides

Na₂O and MgO are called basic oxides because they react like a base. They react with water to form alkalis.

	$ m Na_2O$	$_{ m MgO}$
Reaction with water	$\mathrm{Na_2O}_{\mathrm{\;s\;}} + \mathrm{H_2O}_{\mathrm{\;1\;}} \rightarrow \mathrm{2NaOH}_{\mathrm{\;aq\;}}$	$\rm MgO_s \ + H_2O_1 \ \rightarrow Mg(OH)_{2 \ aq}$
Reaction with acid	${ m Na_2O}_{ m s}$ $+2{ m HCl}_{ m aq}$	${ m MgO}_{ m s}$ $+2{ m HCl}_{ m aq}$
(e.g. HCl)	$\rightarrow 2 \mathrm{NaCl}$ $_{\mathrm{aq}}$ $+ \mathrm{H_2O}$ $_{\mathrm{1}}$	$ ightarrow \mathrm{MgCl_{2~aq}} + \mathrm{H_{2}O_{1}}$

Acidic oxides

The covalent oxides are acidic as they can react with alkalis and neutralize them. Some of them can also dissolve in water to form acids.

- SiO_2
 - o Reaction with alkalis:

$$\mathrm{SiO}_{2(s)} + 2\mathrm{NaOH}_{\mathrm{\,aq\,}} \rightarrow \mathrm{Na_2SiO}_{3(\mathrm{aq})} + \mathrm{H_2O}_{\mathrm{\,1}}$$

- o SiO₂ does not dissolve in water
- P₄O₁₀
 - Reaction with alkalis:

$$\mathrm{P_4O_{10(s)} + 12NaOH}_{\mathrm{aq}} \, \rightarrow 4\mathrm{Na_3PO_{4(aq)} + 6H_2O}_{\mathrm{1}}$$

• Reaction with water:

$$\mathrm{P_4O_{10(s)}+6H_2O_{\,1}\,\rightarrow 4H_3PO_{4\,\,aq}}$$

- o Phosphoric acid is produced
- SO_2
 - o Reaction with alkalis:

$$\mathrm{SO_{2\;g}\; + 2NaOH_{\;aq}\; \rightarrow Na_{2}SO_{3\;aq}\; + H_{2}O_{\;1}}$$

• Reaction with water:

$$SO_{2(g)} + H_2O_1 \rightleftharpoons H_2SO_{3(aq)}$$

- o Sulphurous acid is produced
- OCl₂
 - o Reaction with alkali:

$$\mathrm{OCl}_{2(\mathrm{g})} + 2\mathrm{NaOH}_{\mathrm{aq}} \ \rightarrow 2\mathrm{NaOCl}_{(\mathrm{aq})} + \mathrm{H}_2\mathrm{O}_1$$

• Reaction with water

$$OCl_{2 g} + H_2O_1 \rightleftharpoons 2HOCl_{aq}$$

o Hypochlorous acid is produced

Amphoteric oxides

 Al_2O_3 is considered *amphoteric* as it has **both** acid and basic properties. In other words, it can react like an acid or a base. Al_2O_3 does not dissolve in water.

Base: 6HCl $_{\rm aq}$ + Al₂O_{3 s} $\,\rightarrow\,$ 2AlCl_{3 aq} + 3H₂O $_{\rm 1}$

Acid: 2NaOH $_{\rm aq}~+{\rm Al_2O_{3(s)}} + 3{\rm H_2O_{1}}~\rightarrow 2{\rm Na[Al~OH~_{4}]}~_{\rm aq}$

Checkpoint

- 1. Answer the following questions
 - a. Write the equation for the reaction between sodium oxide and water. State the type of bonding present in sodium oxide
 - b. With an aid of a chemical equation, explain whether P₄O₁₀ is acidic, basic or amphoteric
 - c. Relate the acid-base character of the oxides in (a) and (b) to the variation in metallic character of the elements in Period 3 of the Periodic table (sodium to argon).
 - d. Show why sulphur dioxide is an acidic gas with the aid of a chemical equation

e. Al_2O_3 is said to be amphoteric, what does it mean by being 'amphoteric', explain your answer and include relevant chemical equations.

2. DSE 2022 Q13

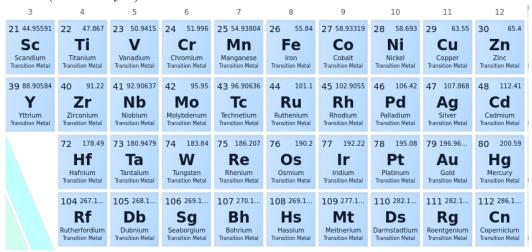
Describe the acid-base properties of the products formed (if any) when the following oxides are added to water separately. Chemical equations are NOT required. (5 marks)

Transition metals and their properties

Transition elements refer to the elements that are not a part of a main group. Most of these elements are metals, hence they are also commonly called transition metals. These elements have a few typical properties.

- Higher melting and boiling point that main group elements
- Multiple oxidation states
- Catalytic properties
- Coloured ions (when dissolved in aqueous solutions)

Most transition metals have higher boiling points than normal metals because more electrons can become delocalized and hence increase the strength of the metallic bonds between the metal ions and sea of delocalized electrons. The details of this are not covered in DSE, so the property of higher boiling point is rarely assessed. (Not seen yet)



Multiple oxidation states

Most transition metals can have multiple oxidation states because their partially filled third electron shell allows for different ways to form stable ions (i.e. different oxidation states)

Manganese

Compound	Oxidation number exhibited
MnO_2	+4
MnCl_2	+2
KMnO_4	+7

Iron

Compound	Oxidation number exhibited
FeO	+2
$\mathrm{Fe_{2}O_{3}}$	+3

Vanadium

Compound	Oxidation number exhibited
VO	+2
V_2O_3	+3
\mathbf{VC}	+4
$ m VOCl_3$	+5

Catalytic properties

This property mostly arises from the fact that transition metals can take up multiple oxidation states, they can go back and forth between the states to help catalyse reactions. Catalysts increase the rate of reaction while remaining chemically unchanged after the reaction.

Material	Process catalysed
Fe	Haber process (450°C, 200 atm, finely divided Fe) $N_2 + 3H_2 \rightleftharpoons 2NH_3$
$ m Cu/ZnO/Al_2O_3$	Methanol formation (250°C, 50-100 atm, Cu/ZnO/Al ₂ O ₃) CO + 2H ₂ \rightarrow CH ₃ OH
NiO	Steam methane reforming (700°C-1000°C, 10-20 atm, Ni) $CH_4 + H_2O \rightleftharpoons CO + 3H_2$
Pt/Rh	Catalytic converter
$\mathrm{Pt/Ni/Pd}$	Catalytic hydrogenation of alkenes

Mechanism of catalysts:

This usually will not be assessed but it can be useful to understand how transition metals catalyse reactions. Take the reaction between persulphate ions $(S_2O_8^{2-})$ and iodide ions as an example.

$${\rm S_2O_8^{2-}} + 2{\rm I^-} \rightarrow 2{\rm SO_4^{2-}} + {\rm I_2}$$

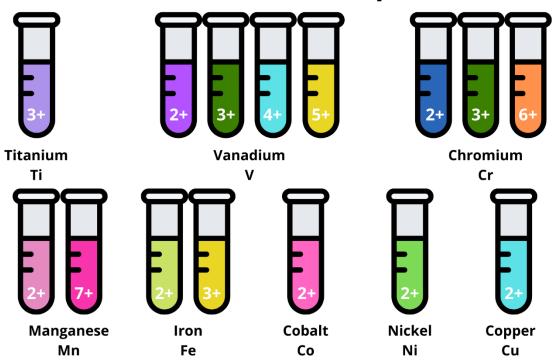
Normally the reaction proceeds quite slowly but using Fe as a catalyst can speed things up. The two ions on the reactant side are both negatively charged, a lot (kinetic) energy is required to overcome the electrostatic repulsion force.

$$\begin{split} 2\mathrm{Fe^{2+}} + \mathrm{S_2O_8^{2-}} &\to 2\mathrm{SO_4^{2-}} + 2\mathrm{Fe^{3+}} \\ 2\mathrm{Fe^{3+}} + 2\mathrm{I^-} &\to 2\mathrm{Fe^{2+}} + \mathrm{I_2} \end{split}$$

Notice how at the end of the two reactions, iron is present as Fe²⁺, meaning that it remained chemically unchanged throughout the reaction. This is a good example of how the variable oxidation states of transition metals can help to catalyse reactions (redox in this case).

Coloured ions

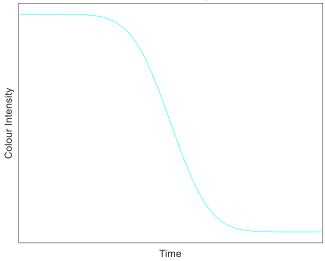
Transition Metal Ion Colors in Aqueous Solution



Checkpoint

DSE 2017 Q14

At 60° C, $MnO_4^{-}(aq)$ reacts with $C_2O_4^{2-}(aq)$ in an acidic medium to give $Mn^{2+}(aq)$, $CO_2(aq)$ and $H_2O(l)$. The graph below shows the variation of the colour intensity of the reaction mixture with time.



Based on the information above, write the chemical equation for the reaction and illustrate THREE characteristics of transition metals exhibited by manganese.

$$MnO_4^- + 8H^+ + 5e^- \rightarrow Mn^{2+} + 4H_2O \times 2$$

$$C_2O_4^{2-} \rightarrow 2CO_2 + 2e^- \ge 5$$

$$2 \rm MnO_4^- + 16 H^+ + 5 C_2 O_4^{2-} \rightarrow 2 Mn^{2+} + 8 H_2 O + 10 CO_2$$

Multiple oxidation states

Mn has an oxidation number of +7 in MnO₄, but it has an oxidation number of +2 in Mn²⁺.

Catalytic properties

The rate of reaction is proportional to the slope of the colour intensity curve against time. The slope increases as the reaction progresses. This indicates that the rate of reaction is higher after the reaction has progressed by a bit due to the presence of a catalyst, which in this case, is the product of the reaction, Mn^{2+} . Hence, Mn can catalyse chemical reactions and exhibits the catalytic properties of transitional metals.

(Mn remains chemically unchanged after the catalytic process.)

Coloured ions (when dissolved in aqueous solutions)

Colour intensity decreases with time since MnO_4^- is purple in colour while the product, Mn^{2+} is colourless (very pale pink), so manganese ions are coloured when dissolved in water.