

Topic 1: Stoichiometric Relationships

1.1 Introduction to the particulate nature of matter and chemical change

Matter : Any substance (made up of one or more atoms) with space and mass.

Atom : The smallest particle of a chemical element that can exist.

Compound : Atoms of different elements chemically bond in fixed ratios to form compounds.

E.g NaCl (ratio of Na: Cl is 1:1)

Molecule : A group of atoms held together by chemical bonding, representing the smallest fundamental unit

of a chemical compound.

Why is water (H₂O) a compound and not an element? Water can be chemically broken down into constituent elements.

Difference between compounds & elements: Elemental H2 and O2 gas have different chemical & physical properties compared to the compound H₂O.

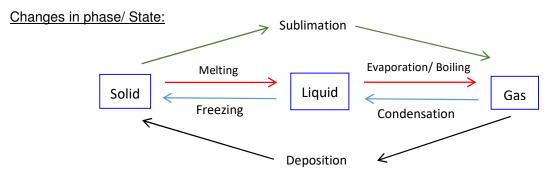
Mixtures contain 2 or more elements or compounds that are not chemically bonded together. Hence, properties of mixtures retain the individual properties of the components.

Homogenous mixture: All components are in the same physical phase and have similar properties. E.g (Salt

water, Air)

Heterogenous mixture: Components are in different physical phases and have varying properties. (Oil & Water -

subtances remain physically separate)



1.2 The mole concept

Mole:

- A unit that describes the amount of substance. Substance could be atoms, molecules, ions or electrons.
- The term "mole" describes a number 6.02 x 10²³
 One mole of substance = 6.023 x 10²³ atoms, molecules, ions or electrons.

1 mole of ¹²C atom: 6.023 x 10²³ ¹²C atoms 1 mole of NO_3 ions: 6.023×10^{23} of NO_3 ions

Relative atomic mass (A_r) : Ratio of the average mass of the isotope of atoms to $\frac{1}{42}$

mass of an atom of ¹²C atom. No units.

Relative molecular mass (M_r) : Ratio of the average mass of the molecule to $\frac{1}{12}$ mass of an

atom of ¹²C atom. No units.

Molar Mass (M): The mass of one mole of a substance. Units: g mol⁻¹



E.g. Iron (Fe) has a Molar Mass of 55.85 g mol⁻¹. This means that one mole of iron atoms $(6.023 \times 10^{23} \text{ iron atoms})$ has a mass of 55.85 g.

Sample Mass (g)

No of

Mole

(mol)

Molar

Mass

(g mol)

Determining the no. of moles of substance (or Molar Mass, Sample mass):

$$no.\,of\,\,moles = \frac{sample\,\,mass\,(g)}{molar\,\,mass\,\,(g\,\,mol^{-1})}$$

Empirical formula: Simplest ratio of the compound - E.g. Empirical formula of C₂H₄ is CH₂

Molecular formula: Actual number of atoms present in a molecule respectively.

Molecular formula: n (Empirical Formula) & Molecular Mass: n (Empirical Mass)

Structural formula: Shows the arrangement of atoms and bonds within a molecule. Used in organic chemistry.

You may be asked to find the empirical formula of a compound given its percentage composition by mass.

Example 1: A particular compound is found to be composed of 86% Carbon and 14% Hydrogen by mass, what is its empirical formula?

Step 1 – Write out the percentages in terms of grams (i.e. instead of 86% Carbon out of 100%, we have 86g of Carbon in a sample of 100g of the compound).

Step 2 – Divide by the Molar Mass of the element (determining the number of moles present in 100g of compound.

Carbon: $86 \div 12 = 7$

Hydrogen: 14 ÷ 1 = 14

Step 3 – Compare the ratio of moles present.

 $C: H \rightarrow 7: 14 = 1:2$

Therefore, the empirical formula of the compound is CH₂.

Example 2a: A compound is found to be composed of 9% Hydrogen, 55% Carbon and 36% Oxygen by mass. Find its empirical formula.

	Hydrogen	Carbon	Oxygen	
% in terms of grams	9	55	36	
Number of moles of atoms	9 ÷ 1 = 9	55 ÷ 12 = 4.58	36 ÷ 16 = 2.25	The no. of moles must be converted into
	36 ↓ ^{×4}	≈ 18.32	9	a whole number!
Ratio of moles	4 ÷9	≈ 2	1	Now we have the simplest ratio

So we have an empirical formula of C₂H₄O.

Example 2b: Given that the molar mass of the compound is 132, determine its molecular formula.

Empirical formula C_2H_4O has a molar mass of: $(2\times12) + (1\times4) + (1\times16) = 44g \text{ mol}^{-1}$

$$132 \div 44 = 3$$

This tells us that the actual molar mass is 3 times larger than the empirical mass. Therefore, the molecular formula is:

$$C_6H_{12}O_3$$

Practice Questions from IBDP HL/SL Chemistry (May 2010-2011) for 1.1 & 1.2

a) How many molecules are present in a drop of ethanol, C_2H_5OH , of mass 2.3×10^{-3} g?

$$(L = 6.0 \times 10^{23} \text{ mol}^{-1})$$

b) The relative molecular mass of a gas is 56 and its empirical formula is CH₂. What is the molecular

formula of the gas?

- c) What is the total number of hydrogen atoms in 1.0 mol of benzamide, C₆H₅CONH₂?
- d) Which is both an empirical and a molecular formula?

 $C_5 H_{12} \qquad \qquad C_5 H_{10} \qquad \qquad C_4 H_8 \qquad \qquad C_4 H_{10}$

- e) What is the mass, in g, of one molecule of ethane, C₂H₆?
- f) Which molecular formula is also an empirical formula?

 PCl_3

 C_2H_4 H_2O_2 $C_6H_{12}O_6$

- g) What is the total number of nitrogen atoms in **two** mol of NH₄NO₃?
- h) Which compound has the empirical formula with the largest mass?

 C_2H_6

 C_2H_4 C_2H_2 C_3H_6

1.3 Reacting Masses and Volume

- · Chemical Equations: Reactants react to produce substances which are known as products in a fixed relationship (i.e mole ratio).
- State symbols: (s) solid, (l) liquid (or molten), (g) gas and (aq) aqueous.
- Single arrow (→): Reaction goes to completion. All the reactants have been consumed to form products completely.
- Reversible arrows (\$\sigma\$): Reaction that has both reactant and product at the end (i.e equilibrium). The reaction proceeds in both directions, with reactants forming products and products reforming reactants simultaneously.
- Ionic equations: Used to describe the dissociation of ionic compounds in water or precipitation reactions.

For example:

NaCl (s) \rightarrow Na⁺ (aq) + Cl⁻ (aq)

Dissociation of ionic compounds

 $Ag^{+}(aq) + Cl^{-}(aq) \rightarrow AgCl(s)$

Precipitation of ionic compounds

Determining the limiting reagent / reactant:

Reactants that are added in excess to what is required.

E.g.

$$2 \text{ Ca (s)} + O_2 \text{ (g)} \rightarrow 2 \text{CaO (s)}$$

What does this chemical equation mean?

Moles: 2 moles of Ca reacts with 1 mole of O₂ to produce 2 moles of CaO.

Mass: 80.2 g of Calcium reacts with 32.0 g of Oxygen to produce 112.0 g of Calcium Oxide.

What kind of questions will appear in exams?

Any **random mass** of the reactants will appear.

Type 1: One of the reactant is in excess.

3.0 g of Calcium reacted in *excess* oxygen to produce Calcium Oxide. Find the mass of Calcium Oxide produced?

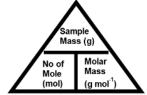
$$2 \text{ Ca (s)} + O_2 \text{ (g)} \rightarrow 2 \text{CaO (s)}$$

Answer:

No of moles or Amount of Calcium: $\frac{Sample\ Mass}{Molar\ Mass} = \frac{3.0}{40.1} = 0.0748\ mol$

Mole ratio of Ca: CaO → 1:1

Thus, amount of CaO: 0.0748 mol



Mass of Calcium oxide: No of moles x Molar Mass = 0.0748 x 56.1 = 4.196 g

Type 2: Determining the limiting agent.

3.0 g of Calcium reacted with 4.0 g of oxygen to produce Calcium Oxide. Find the mass of Calcium Oxide produced.

Amount of Calcium: 0.0748 mol Amount of Oxygen: 4.0 / 32.0 = 0.125 mol

Recall* Moles: 2 mol of Ca reacts with 1 mol of O₂

From Calcium's point of view: 0.0748 mol of Ca requires only 0.0374 mol of O₂.

→ Oxygen is in excess as we have 0.125 mol.

From Oxygen's point of view: 0.125 mol of O₂ requires 0.250 mol of Ca.

→ Ca is limited as we only have 0.0748 mol.

Only Calcium will be completely used up to form Calcium Oxide. Mole ratio of Ca:CaO is 2:2 = 1:1

Therefore, Amount of Ca = Amount of CaO = 0.0748

Mass of Calcium oxide: No of moles x Molar Mass = 0.0748 x 56.1 = 4.196 g

Balancing Equations

You will frequently be asked to balance equations in exams. Just ensure that each element has the same number of moles present in the reactant and product side of the equation.

Example 1: Na (s) + $H_2O(I) \rightarrow NaOH(aq) + H_2(g)$

In this example we have 2 moles of hydrogen atoms on the reactant side, but 3 on the product side. Therefore, the equation is not balanced. In order to balance it, we add numbers in front of each reactant/ product.

$$Na + 2H_2O \rightarrow 2NaOH + H_2$$

Now there are 4 moles hydrogen and 2 moles of Oxygen on both sides. However, the sodium is not balanced.

2Na (s) + 2H₂O (l)
$$\rightarrow$$
 2NaOH (aq) + H₂ (g)

Practice Balancing Equations

- a) $MgCO_3 + H_2SO_4 \rightarrow MgSO_4 + CO_2 + H_2O$
- b) $C_4H_8 + O_2 \rightarrow CO_2 + H_2O$
- c) $Pb(NO_3)_2 + NaCl \rightarrow PbCl_2 + NaNO_3$

Moles and Concentration

Concentration (mol
$$dm^{-3}$$
) = $\frac{No. of Moles (mol)}{Volume (dm^{-3})}$

This equation is used to calculate the number of moles of a solute (dissolved chemical) in a particular sample of solution.

Example: Find the number of moles of NaCl present in 0.300 dm⁻³ of 2.0 moldm⁻³ aqueous NaCl solution.

Moles = Conc.
$$\times$$
 Volume
= 2.0×0.300
= 0.60 mol

Calculating Moles of Gas

At room temperature (25°C / 298K) and pressure, 1 mole of gas has a volume of 24dm³.

However, at standard temperature (0°C / 273K) and pressure, 1 mole of gas has a volume of 22.7dm³.

$$Moles of Gas = \frac{Volume (dm^3)}{22.7}$$

These two equations are only to be used to calculate the moles of GAS molecules.

Example: Find the number of moles of hydrogen molecules in a container of volume 300 cm³ at:

- (a) Room temperature
- (b) Standard temperature.

Recall:
$$300 \text{cm}^3 = 0.300 \text{dm}^3$$

(a) $0.300 \div 24 = 0.0125 \text{mol}$
(b) $0.300 \div 22.7 \approx 0.0132 \text{mol}$

Ideal Gases

When working with gases, we often approximate their behaviour to an "ideal gas" to simplify any necessary calculations. There are 2 important assumptions to note about an ideal gas:

- 1. The gas molecules do not interact with one another.
- 2. The volume of each gas particle is zero.

Real gases deviate from this ideal behaviour at:

1. High pressure -

The gas particles are pushed so closely together that they occupy a larger volume collectively, to the point where the volume of each particle is not negligible.

2. Low temperature -

Each particle has very little kinetic energy, thus when they move close to other particles, they are more likely to experience some interaction due to intermolecular forces.

The ideal gas equation:

$$PV = nRT$$

Symbol	Represents	Unit
Р	Pressure	Pascals (Pa)
V	Volume	m³ (= 1000dm³)
n	no. of moles	mol
R	Gas constant = 8.31	J K ⁻¹ mol ⁻¹
Т	Temperature	К

This equation allows us to calculate the molar mass of a gas, provided its mass is given.

Example: A sample of a gas is kept in a cylinder of volume 1.00 dm³, at 100 kPa and a temperature of 18.0°C. It is found to have a mass of 1.65g. Calculate the molar mass of the gas.

Convert all given data into appropriate units:

- $1 \text{ dm}^3 = 0.00100 \text{ m}^3$
- 100 kPa = 100 000 Pa
- 18.0°C = 291 K

$$PV = nRT$$
 Rearrange to make n the subject: $n = \frac{PV}{RT}$
$$n = \frac{100000 \times 0.001}{8.31 \times 291} \approx 0.0414$$

Recall:
$$Molar\ Mass = \frac{Mass}{Moles}$$

$$Molar\ Mass = \frac{1.65}{0.414} \approx 39.9 gmol^{-1}$$

Alternatively, we can rearrange this equation to allow us to predict the response of a set amount of gas to a change in conditions.

$$\frac{PV}{T} = nR$$

For a fixed sample of gas (no. of moles does not change), the variables on the right side are both constants. This tells us that the final value of $\frac{PV}{T}$ will not change even if one of the variables involved does.

Example: A fixed sample of gas has the initial conditions of 80 kPa and a temperature of 278 K. The pressure is then increased to 110 kPa, with the volume remaining constant. Calculate the final temperature of the gas.

$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2}$$

$$T_2 = \frac{T_1 P_2 V_2}{P_1 V_1} = \frac{278 \times 110000}{80000} \approx 382K$$

1 = Initial2 = Final

Common IB Stoichiometric Relationships Questions

1.	1.01:	xic gas, A, consists of 53.8 % nitro ×10 ⁵ Pa, 1.048 g of A occupies 46 ulate the molar mass of the compo	52 cm ³ . Determine th	e empirical formula of A.		
					(Total 3 mark	(s)
2.	empi	irical formula = CN; Working must be sl	hown to get noint			
	$M_r =$	*51.9 (g mol ⁻¹);	iown io gei poini.			
	-	=C—C ==N:;			3	.01
					l	3]
_						
3.		data below is from an experiment ble of iron ore. This sample was di				
	The	resulting solution was titrated with	h a standard solution	of potassium manganate(VII),	
		nO ₄ . This procedure was carried on to form Mn ²⁺ and Fe ³⁺ and the en-			s with Fe ²⁺	
	10118	to form will and it and the cir	a point is maleated t	by a singlit pllik colour.		
		titre	1	2	3	
		initial burette reading / cm ³	1.00	23.60	10.00	
		initial burette reading / cm ³ Final burette reading / cm ³	1.00 24.60		10.00 32.50	
		Final burette reading / cm ³		46.10		
		Final burette reading / cm ³ Mass of iron ore / g	24.60	$46.10 3.682 \times 10^{-1}$		
		Final burette reading / cm ³	24.60	46.10		
		Final burette reading / cm ³ Mass of iron ore / g	24.60	$46.10 3.682 \times 10^{-1}$		
	(a)	Final burette reading / cm ³ Mass of iron ore / g concentration of KMnO ₄ solut	24.60 tion / mol dm ⁻³	$ \begin{array}{c c} 46.10 & 3 \\ 3.682 \times 10^{-1} \\ 2.152 \times 10^{-2} \end{array} $		
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		Final burette reading / cm ³ Mass of iron ore / g concentration of KMnO ₄ solut Deduce the balanced redox equa	24.60 tion / mol dm ⁻³ ation for this reaction	$ \begin{array}{c c} 46.10 & 3 \\ 3.682 \times 10^{-1} \\ 2.152 \times 10^{-2} \end{array} $	32.50	(2)
	(a) (b)	Final burette reading / cm ³ Mass of iron ore / g concentration of KMnO ₄ solut	24.60 tion / mol dm ⁻³ ation for this reaction	$ \begin{array}{c c} 46.10 & 3 \\ 3.682 \times 10^{-1} \\ 2.152 \times 10^{-2} \end{array} $	32.50	(2)

(c)	Calculate the amount, in moles, of MnO ₄ ⁻ used in the titration.	
		•••••
		(2)
(d)	Calculate the amount, in moles, of Fe present in the 3.682×10^{-1} g sample of	iron ore.
		(2)
(e)	Determine the percentage by mass of Fe present in the 3.682×10^{-1} g sample	of iron ore.
(a)	$MnO_4^-(aq) + 5Fe^{2+}(aq) + 8H^+(aq) \rightarrow Mn^{2+}(aq) + 5Fe^{3+}(aq) + 4H_2O(l)$ Award [2] if correctly balanced. Award [1] for correctly placing H^+ and H_2O . Award [1 max] for correct balanced equation but with electrons shown. Ignore state symbols.	2
(b)	Fe ²⁺ / iron(II); Do not accept iron.	1

4.

5 × 4.842 × 10 ⁻⁴ = 2.421 × 10 ⁻³ (mol); (same number of moles of Fe in the iron ore) Allow ECF from part (a) and (c) provided some mention of mole ratio is stated. 2 2 2 2 3 4 2 5 4 2 5 5 6 2 6 2 6 2 6 2 7 6 2 7 7 8 7 8 8 7 8 8 7 8 8 8	Award [II] for correct calculation. 2 d) 1 mol of MnO ₄ ⁻ reacts with 5 mol of Fe ²⁺ ; 5 × 4.842 × 10 ⁻⁴ = 2.421 × 10 ⁻³ (mol); (same number of moles of Fe in the iron ore) Allow ECF from part (a) and (c) provided some mention of mole ratio is stated. 2 e) 2.421 × 10 ⁻³ × 55.85 = 0.1352(g); 0.1352/0.3682 × 100 = 36.72 %; Allow ECF from part (d). 2 The percentage of iron(II) ions, Fe ²⁺ , in a vitamin tablet can be estimated by dissolving the ablet in dilute sulfuric acid and titrating with standard potassium manganate(VII) solution, CMnO ₄ (aq). During the process iron(II) is oxidized to iron(III) and the manganate(VII) ion is educed to the manganese(II) ion, Mn ²⁺ (aq). It was found that one tablet with a mass of 1.43 g equired 11.6 cm ³ of 2.00 × 10 ⁻² mol dm ⁻³ KMnO ₄ (aq) to reach the end-point. a) (i) State the half-equation for the oxidation of the iron(II) ions. (ii) State the half-equation for the reduction of the MnO ₄ ⁻ ions in acidic solution.	(c)		$152 \times 10^{-2} \times 2.250 \times 10^{-2}$; $\times 10^{-4}$ (mol);	
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(ii) State the half-equation for the reduction of the MnO_4^- ions in acidic solution.	(ii) State the half-equation for the reduction of the MnO_4^- ions in acidic solution.	table KM1	et in dilu nO ₄ (aq)	ite sulfuric acid and titrating with standard potassium manganate(VII). During the process iron(II) is oxidized to iron(III) and the manganate	solution, e(VII) ion is
		table KMı redu requ	et in dilunO ₄ (aq) ced to the ired 11.	the sulfuric acid and titrating with standard potassium manganate(VII). During the process iron(II) is oxidized to iron(III) and the manganate the manganese(II) ion, $\mathrm{Mn^{2+}(aq)}$. It was found that one tablet with a manganese of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ mol dm ⁻³ KMnO ₄ (aq) to reach the end-point.	solution, e(VII) ion is
(iii) Deduce the overall redox equation for the reaction.	(iii) Deduce the overall redox equation for the reaction.	table KMı redu	et in dilunO ₄ (aq) ced to the ired 11.	the sulfuric acid and titrating with standard potassium manganate(VII). During the process iron(II) is oxidized to iron(III) and the manganate the manganese(II) ion, $\mathrm{Mn^{2+}(aq)}$. It was found that one tablet with a manganese of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ mol dm ⁻³ KMnO ₄ (aq) to reach the end-point.	solution, e(VII) ion is
		table KMı redu requ	et in dilu nO ₄ (aq) ced to t ired 11.	the sulfuric acid and titrating with standard potassium manganate(VII). During the process iron(II) is oxidized to iron(III) and the manganate he manganese(II) ion, $\mathrm{Mn^{2+}(aq)}$. It was found that one tablet with a manganese of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ of $\mathrm{Cm^3}$ mol dm ⁻³ KMnO ₄ (aq) to reach the end-point. State the half-equation for the oxidation of the iron(II) ions.	solution, e(VII) ion is ass of 1.43 g
		table KMı redu requ	et in dilu nO ₄ (aq) ced to t ired 11. (i)	the sulfuric acid and titrating with standard potassium manganate(VII). During the process iron(II) is oxidized to iron(III) and the manganate he manganese(II) ion, Mn ²⁺ (aq). It was found that one tablet with a manganese of 2.00 × 10 ⁻² mol dm ⁻³ KMnO ₄ (aq) to reach the end-point. State the half-equation for the oxidation of the iron(II) ions.	solution, e(VII) ion is ass of 1.43 g

(b) (i) Calculate the amount, in moles, of MnO_4^- ions present in 11.6 cm³ of 2.00×10^{-2} mol dm⁻³ KMnO₄(aq).

(1)

(ii) Calculate the amount, in moles, of Fe²⁺ ions present in the vitamin tablet.

.....(1)

(iii) Determine the percentage by mass of Fe²⁺ ions present in the vitamin tablet.

(Total 7 marks)

- **6.** (a) (i) $Fe^{2+} \rightarrow Fe^{3+} + e^{-}$;
 - (ii) $MnO_4^- + 8H^+ + 5e^- \rightarrow Mn^{2+} + 4H_2O;$ 1
 - (iii) $MnO_4^- + 5Fe^{2+} + 8H^+ \rightarrow Mn^{2+} + 5Fe^{3+} 4H_2O;$ Accept e instead of e^- .
 - (b) (i) amount of $MnO_4^- = \frac{11.6}{1000} \times 0.0200 = 2.32 \times 10^{-4} \text{ mol};$
 - (ii) amount of Fe²⁺ = $5 \times 2.32 \times 10^{-4} = 1.16 \times 10^{-3}$ mol;

(iii) mass of Fe²⁺ = $55.85 \times 1.16 \times 10^{-3} = 6.48 \times 10^{-2}$ g;

(2)

(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample.	(e)	Dete	ermine the amount, in mol, of calcium carbonate in the sample of the eggshell	
(f) Calculate the mass and the percentage by mass of calcium carbonate in the eggshell sample. (g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark (a) (i) Copper: 0 to +2/increases by 2/+2/2+;				
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)				
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (a) (i) Copper: (b) Oto +2 / increases by 2 / +2 / 2+;				
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)				(2)
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)	(f)			ell
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)		•••••		
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)		•••••		
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)		•••••		•••
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)		•••••		
(g) Deduce one assumption made in arriving at the percentage of calcium carbonate in the eggshell sample. (Total 11 mark)				
eggshell sample. (Total 11 mark) (a) (i) Copper: 0 to +2 / increases by 2 / +2 / 2+;				(3)
(a) (i) Copper: 0 to +2 / increases by 2 / +2 / 2+;	(g)			n the
0 to $+2$ / increases by 2 / $+2$ / $2+$;		•••••	(**************************************	 (1) Fotal 11 marks)
0 to +2 / increases by 2 / +2 / 2+;				
	(a)	(i)	0 to +2 / increases by 2 / +2 / 2+;	
<i>Nitrogen:</i> +5 to +4 / decreases by 1 / -1 / 1-;				
Penalize missing + sign or incorrect notation such as 2+, 2 ⁺ or II, once only.			Penalize missing + sign or incorrect notation such as 2+, 2 ⁺	2

8.

- (ii) nitric acid/HNO₃ / NO₃ /nitrate; *Allow nitrogen from nitric acid/nitrate but not just nitrogen.* 1
- (b) (i) 0.100×0.0285 ; 2.85×10^{-3} (mol); Award [2] for correct final answer.
 - (ii) 2.85×10^{-3} (mol);
 - (iii) $(63.55 \times 2.85 \times 10^{-3}) = 0.181 \text{ g};$ Allow 63.5.
 - (iv) $\left(\frac{0.181}{0.456} \times 100 = \right) 39.7 \%$
 - (v) $\left(\frac{44.2 39.7}{44.2} \times 100 = \right) 10/10.2 \%;$ Allow 11.3 % i.e. percentage obtained in (iv) is used to divide instead of 44.2 %.
- (c) Brass has:
 delocalized electrons / sea of mobile electrons / sea of electrons
 free to move;
 No mark for just "mobile electrons".

 [10]

9. Brass is a copper containing alloy with many uses. An analysis is carried out to determine the percentage of copper present in three identical samples of brass. The reactions involved in this analysis are shown below.

Step 1:
$$\text{Cu}(s) + 2\text{HNO}_3(\text{aq}) + 2\text{H}^+(\text{aq}) \rightarrow \text{Cu}^{2+}(\text{aq}) + 2\text{NO}_2(g) + 2\text{H}_2\text{O}(l)$$

Step 2:
$$4\Gamma(aq) + 2Cu^{2+}(aq) \rightarrow 2CuI(s) + I_2(aq)$$

Step 3:
$$I_2(aq) + 2S_2O_3^{2-}(aq) \rightarrow 2\Gamma(aq) + S_4O_6^{2-}(aq)$$

		id introgen	in step 1	•
	Copper:			
	Nitrogen:			
(ii)	Identify the oxidizing agent in step 1.			
		••••••		
	ident carried out this experiment three times, with three identification obtained the following results.			
	Mass of brass = $0.456 \text{ g} \pm 0.001 \text{ g}$		1	ı
Tit	re	1	2	3
Init	re tial volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3)	0.00	0.00	0.00
Init Fin	re tial volume of 0.100 mol dm ⁻³ $S_2O_3^{2-}$ (± 0.05 cm ³) al volume of 0.100 mol dm ⁻³ $S_2O_3^{2-}$ (± 0.05 cm ³)		0.00	0.00
Init Fin	re tial volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3)	0.00	0.00	0.00
Init Fin Vol	re tial volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) al volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) tume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3) erage volume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10	0.00	0.00	0.00
Init Fin Vol	re tial volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) al volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) tume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3) erage volume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10	0.00 28.50 28.50	0.00 28.60 28.60	0.00
Init Fin Vol Ave cm	tial volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) al volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) dume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3) erage volume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3)	0.00 28.50 28.50	0.00 28.60 28.60	0.00
Init Fin Vol Ave cm	tial volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) al volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) dume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3) erage volume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3)	0.00 28.50 28.50	0.00 28.60 28.60	
Init Fin Vol Ave cm	tial volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) al volume of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.05 cm 3) dume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3) erage volume added of 0.100 mol dm $^{-3}$ S $_2$ O $_3$ ²⁻ (± 0.10 cm 3)	0.00 28.50 28.50	0.00 28.60 28.60	0.00

(2)

					(1)
		(iii)	Calculate the mass of copper in the brass.		
					(1)
		(iv)	Calculate the percentage by mass of copper in the brass.		
					(1)
		(v)	The manufacturers claim that the sample of brass contains 44.2 % copper Determine the percentage error in the result.	r by mass.	
					(1)
	(c)		reference to its metallic structure, describe how brass conducts electricity		
				 (Total 10 mar	(1) rks)
10.	(a)	n(HC	Cl)(= $0.200 \text{ mol dm}^{-3} \times 0.02720 \text{ dm}^{3}$) = $0.00544 / 5.44 \times 10^{-3} \text{ (mol)}$;	1	
	(b)		Cl) excess (= 0.100 mol dm ⁻³ × 0.02380 dm ³) = 0.00238 / × 10^{-3} (mol);		
			lize not dividing by 1000 once only in (a) and (b).	1	
	(c)	n(HC	C1) reacted (= $0.00544 - 0.00238$) = $0.00306 / 3.06 \times 10^{-3}$ (mol);	1	

Calculate the amount, in mol, of copper present in the brass.

(ii)

- (d) $2HCl(aq) + CaCO_3(s) \rightarrow CaCl_2(aq) + H_2O(l) + CO_2(g) \Box / 2H^+(aq) + CaCO_3(s) \rightarrow Ca^{2+}(aq) + H_2O(l) + CO_2(g);$ Award [1] for correct reactants and products.

 Award [1] if this equation correctly balanced.

 Award [1 max] for the following equations: $2HCl(aq) + CaCO_3(s) \rightarrow CaCl_2(aq) + H_2CO_3(aq)$ $2H^+(aq) + CaCO_3(s) \rightarrow Ca^{2+}(aq) + H_2CO_3(aq)$ Ignore state symbols.

2

2

3

(f) $M_{\rm r}({\rm CaCO_3}) \ (= 40.08 + 12.01 + 3 \times 16.00) = 100.09 / 100.1 / M = 100.09 / 100.1 \ (g \ {\rm mol}^{-1});$ Accept 100.

 $= 0.00153 / 1.53 \times 10^{-3}$ (mol);

$$m(\text{CaCO}_3)(= nM) = 0.00153 \text{ (mol)} \times 100.09 \text{ (g mol}^{-1}) = 0.153 \text{ (g)};$$

%CaCO₃ $\left(= \frac{0.153}{0.188} \times 100\right) = 81.4 \% / 81.5 \%$

Accept answers in the range 79.8 to 81.5 %. Award [3] for correct final answer.

 $n(\text{CaCO}_3) = (\frac{1}{2}n(\text{HCl})) = \frac{1}{2} \times 0.00306;$

Award [2] for correct final answer.

(e)

(g) only CaCO₃ reacts with acid / impurities are inert/non-basic / impurities do not react with the acid / nothing else in the eggshell reacts with acid / no other carbonates;

Do not accept "all calcium carbonate reacts with acid".

[11]