



**Western Cape
Government**

FOR YOU

Eden & Central Karoo Education District
West Coast Education District

PHYSICAL SCIENCES

CHEMISTRY REVISION

8 November 2024

NOTHING IN LIFE IS TO BE FEARED, IT IS ONLY TO BE UNDERSTOOD.
NOW IS THE TIME TO UNDERSTAND MORE, SO THAT WE MAY FEAR LESS.



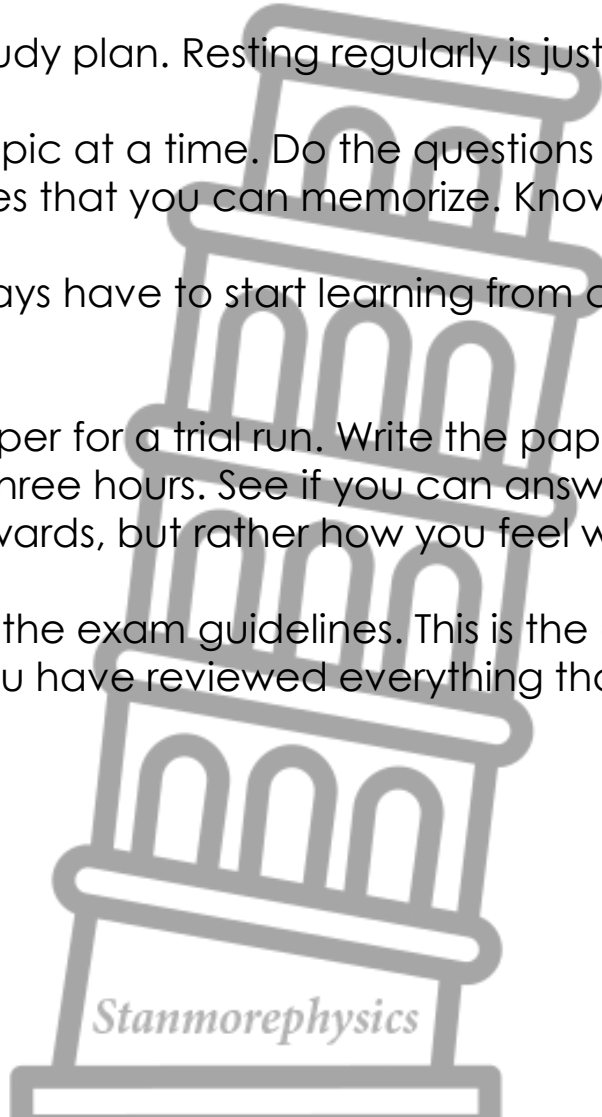
MARIE CURIE





STUDY TIPS :

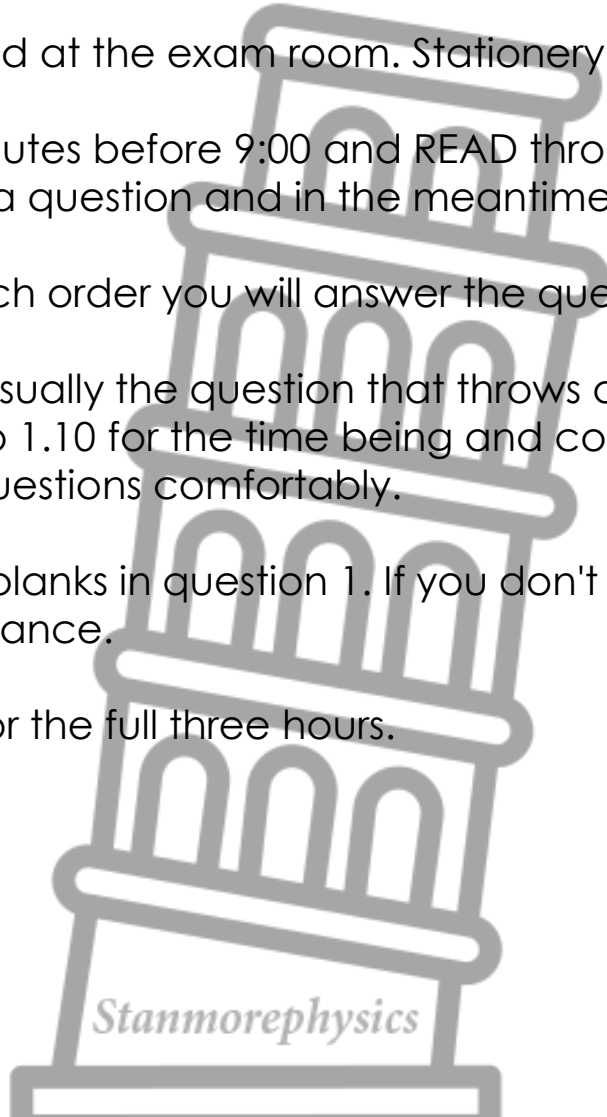
1. Work with a study plan. Resting regularly is just as important as spending hours with your books!
2. Review one topic at a time. Do the questions in previous papers which are only about that specific topic alone. Make key notes that you can memorize. Know your laws & definitions.
3. You don't always have to start learning from chapter 1. Start with the last topic or the topic with which you still have challenges.
4. Leave one paper for a trial run. Write the paper on Sunday as if you were sitting in the exam room. You don't have to sit through it for three hours. See if you can answer the paper from start to finish. The idea is not that you have to mark the answers afterwards, but rather how you feel when you're done.
5. Read through the exam guidelines. This is the examiner's "demarcation". Check that you have reviewed everything that the examiner considers important.





EXAM TIPS FOR MONDAY

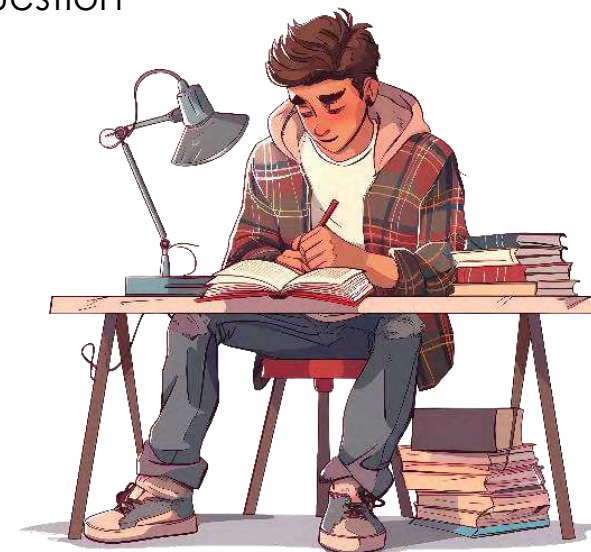
1. Arrive prepared at the exam room. Stationery, calculator, admission letter, ID.
2. Use the 10 minutes before 9:00 and READ through the question paper. You may not write anything yet, but you can read through a question and in the meantime identify the questions that are easy and the routine questions.
3. Decide in which order you will answer the question paper.
4. Question 1 is usually the question that throws off your time management. Leave the question until last. Write the numbers 1.1 to 1.10 for the time being and come back later when you are sure you have enough time and can work through the questions comfortably.
5. Do not leave blanks in question 1. If you don't know and you guess an answer you have at least a 25% chance.
6. Sit and work for the full three hours.

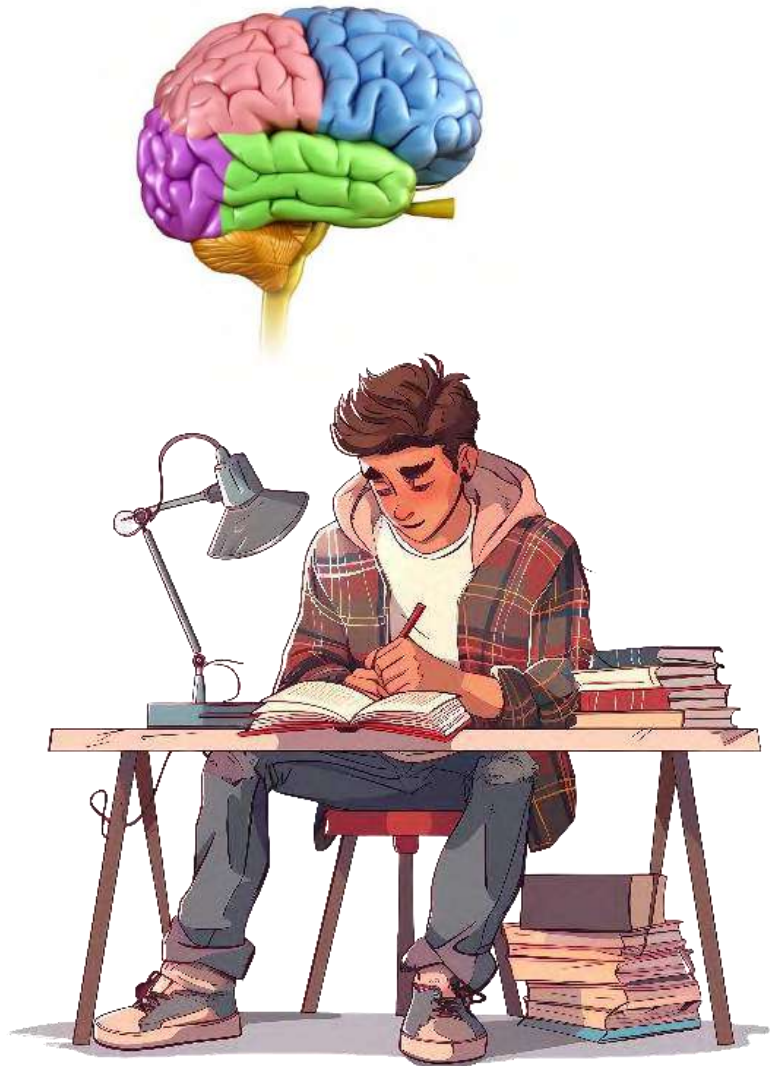




EXAM TIPS TO GET EXTRA MARKS

1. Know your laws & definitions WORD-FOR-WORD. The examiner sometimes doubts whether you know your work if you start to explain things in your own words.
2. Write the formulas exactly as they appear on the formula sheet. Do not try to change the subject of the formula in your mind.
3. You will only get a mark for the formula if you have at least tried some form of substitution. Even if the substitution is completely wrong, you still get the formula mark!
4. The examiner marks along with the error. If you are therefore not sure whether you have done a calculation correctly, do not just stop and leave the question open. Substitute your answer in the following sub question if that is what the question requires. The examiner will take a calculator and mark your answer along with your error.
5. Add all sketches, graphs, diagrams as neatly as possible with labels.
6. If you have to explain something, do it step by step and logically. Like what you learned to do in class.
For example:
 - ❶ Compare the boiling points of two compounds.
 - ❷ Predict the reaction that is favored on the basis of Le Chatelier's principle.





- * Laws
- * Definitions
- * Theory
- * Captions
- * Recognize
- * Name

15%



- * Simple routine Graphs
- * Draw Free Body Diagrams
- * Recognize data in a different format
- * Simple SI Conversions
- * Simple application of theory

40%



- * Calculations
- * Algebraic applications
- * Draw Graphs (new situations & interpretations)
- * Compare situations (Before & After)
- * Interpret diagrams
- * Quantitative chemistry & Stoichiometry

35%



- * Design steps and strategies to solve complex problems.
- * Calculations (Two separate systems that have common factors)
- * Appreciation for applying theory in new situation.

10%





basic education

Department:
Basic Education
REPUBLIC OF SOUTH AFRICA

**NATIONAL
SENIOR CERTIFICATE**

GRADE 12

**PHYSICAL SCIENCES: CHEMISTRY (P2)
NOVEMBER 2023**

Marks: 150

TIME: 3 hours

This question paper consists of 18 pages and 3 data sheets.

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Please turn over.

- Question 1 – Multiple choice questions
- Question 2 – Organic Molecules
- Question 3 – Organic IMK
- Question 4 – Organic Reactions
- Question 5 – Reaction rate
- Question 6 – Chemical Equilibrium
- Question 7 – Acid-Base Reactions
- Question 8 – Galvanic Cells
- Question 9 – Electrolytic Cells





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INSTRUCTIONS AND INFORMATION

1. Write down your examination number and center number in the appropriate spaces on the ANSWER BOOK.
2. This paper consists of 10 questions. Answer ALL the questions in the ANSWER BOOK.
3. Start EACH question on a NEW page in the ANSWER BOOK.
4. Number the answers correctly according to the numbering system used in this paper.
5. Leave ONE line open between two sub-questions, e.g. between QUESTION 2.1 and QUESTION 2.2.
6. You may use a non-programmable calculator.
7. You may use appropriate mathematical tools.
8. Show ALL formulas and substitutions in ALL calculations.
9. Round your final numerical answers to a minimum of TWO decimal places.
10. Give short (concise) motivations, discussions, etc. where necessary.
11. You are advised to use the attached DETAIL SHEETS.
12. Write neatly and legibly.



QUESTION 2

The letters A to H in the table below represent eight organic compounds.

A	Heptanoic acid	B	$\text{CH}_3(\text{CH}_2)_3\text{COOCH}_3$
C	4-ethyl-3,3-difluorohexane	D	Hexanoic acid
E	$\begin{array}{c} \text{CH}_3-\text{CH}-\overset{\text{CH}_2}{\parallel}{\text{C}}-\text{CH}_3 \\ \\ \text{CH}_3 \end{array}$	F	$\begin{array}{c} \text{O} \\ \\ \text{CH}_3-\text{CH}-\text{C}-\text{CH}_2-\text{CH}_3 \\ \\ \text{CH}_3 \end{array}$
G	$\begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_3-\text{C}-\text{CH}_2-\text{CH}_3 \\ \\ \text{C}=\text{O} \\ \\ \text{H}-\text{O} \end{array}$	H	$\begin{array}{cccc} \text{H} & \text{H} & \text{O} & \text{H} \\ & & & \\ \text{H}-\text{C} & -\text{C} & -\text{C} & -\text{C}-\text{H} \\ & & & \\ \text{H} & \text{H} & & \text{H} \end{array}$

2.1 Define the term organic compound.

2.2 Write down the IUPAC name of compound: 2.2.1 E ; 2.2.2 H

2.3 Write down the:

2.3.1 STRUCTURAL formula of compound B

2.3.2 STRUCTURAL formula of compound C

2.3.3 General formula of the homologous series to which compound E belongs

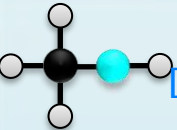
2.3.4 STRUCTURAL formula of the FUNCTIONAL group of compound F

2.3.5 IUPAC name of the alcohol needed to produce compound B

2.4 Write down the letter(s) of the compound(s) that:

2.4.1 Is a FUNCTIONAL isomer of compound G

2.4.2 Are CHAIN isomers of each other



ORGANIC MOLECULES

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Make sure you know the correct wording of all definitions!

TYPES OF FORMULAS:

Molecular Formula
Structure Formula
Condensed Formula

IUPAC**SUBSTITUENTS**

x2 di-

x3 tri-

x4 tetra-

FUNCTIONAL GROUPS:

Hydroxyl group
Carbonyl group
Carboxyl group

HYDROCARBONS

ALKANES $\text{C}_n\text{H}_{2n+2}$

ALKENES C_nH_{2n}

ALKYNES $\text{C}_n\text{H}_{2n-2}$

ISOMERS:

Structural Isomers
Positional Isomers
Functional Isomers

HALOGEN COMPOUNDS

HALOALKANES

HALOALKENES

HALOALKYNES

OXYGEN COMPOUNDS

ALCOHOLS

CARBOXYLIC ACIDS

ALDEHYDES

KETONES

PREFIXES

x1 C Meth-

x2 C Eth-

x3 C Prop-

x4 C But-


x5 C Pent-

x6 C Hex-

x7 C Hept-

x8 C Oct-



1. The **SUFFIXES** (-on ; -ene ; -yn ; -anol ; anoic acid ; -anal ; -anone) indicate the **HOMOLOGOUS SERIES** to which the molecule belongs.
 2. The **PREFIXES** (met-; et-; prop-; but-; pent-; hex-; hept- and oct-) indicate the **NUMBER OF CARBONS** in the parent chain.
 4. The **POSITION OF THE FUNCTIONAL GROUP** must be indicated in the middle of the name of the parent chain.
 5. Carbons in the parent chain are numbered so that the **SMALLEST NUMBER** is used in the IUPAC name.
 6. The **FUNCTIONAL GROUP GETS PREFERRED** and the carbon it is attached to is given the smallest possible numerical value first.
 7. The **POSITION** of the **FUNCTIONAL GROUP** is always indicated in the IUPAC name if two or more positional isomers exist. Eg: pantan-2-one.
 8. When naming **HALOALKANES**, neither the branching nor the halogens take priority over the other. The IUPAC rule regarding the smallest numbering of carbons and alphabetical order is in question.
 9. **BRANCHES** from the parent chain are indicated with the suffix (-yl). The specific carbon in the parent chain where the branch is attached must be indicated in front of the IUPAC name.
 10. **MORE THAN ONE SUBSTITUENT OF THE SAME KIND** is indicated by the prefixes (di-, tri- or tetra-). The specific carbons in the parent chain to which each substituent is attached must be indicated in front of the IUPAC name.
 11. The names of branches are indicated in **ALPHABETICAL ORDER** in the IUPAC name. Ignore the prefixes in this rule.
 12. There **MUST** be a **COMMA** between any two numbers in the molecule's name.
 13. There **MUST** be a **HYPHEN** between any word and a number.
 14. The carbons are numbered in such a way that the IUPAC name always indicates the **SMALLEST POSSIBLE COMBINATION OF NUMBERS**.
- 

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C	4-ethyl-3,3-difluorohexane	D	Hexanoic acid
E	$\begin{array}{c} \text{CH}_3-\text{CH}-\overset{\text{CH}_2}{\parallel}{\text{C}}-\text{CH}_3 \\ \\ \text{CH}_3 \end{array}$	F	$\begin{array}{c} \text{O} \\ \\ \text{CH}_3-\text{CH}-\text{C}-\text{CH}_2-\text{CH}_3 \\ \\ \text{CH}_3 \end{array}$
G	$\begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_3-\text{C}-\text{CH}_2-\text{CH}_3 \\ \\ \text{C}=\text{O} \\ \\ \text{H}-\text{O} \end{array}$	H	$\begin{array}{cccc} \text{H} & \text{H} & \text{O} & \text{H} \\ & & & \\ \text{H}-\text{C} & -\text{C} & -\text{C} & -\text{C}-\text{H} \\ & & & \\ \text{H} & \text{H} & & \text{H} \end{array}$

2.1 Define the term organic compound.

ORGANIC COMPOUND: Molecule/compound containing carbon(atoms).

ORGANIC MOLECULES

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TYPES OF FORMULAS:

Molecular Formula
Structure Formula
Condensed Formula

I U P A C**SUBSTITUENTS**

x2	di-
----	-----

x3	tri-
----	------

x4	tetra-
----	--------

HYDROCARBONS

ALKANES	$\text{C}_n\text{H}_{2n+2}$
---------	-----------------------------

ALKENES	C_nH_{2n}
---------	---------------------------

ALKYNES	$\text{C}_n\text{H}_{2n-2}$
---------	-----------------------------

ISOMERS:

Structural Isomers
Positional Isomers
Functional Isomers

HALOGEN COMPOUNDS

HALOALKANES	x1 C	Meth-
-------------	------	-------

HALOALKENES	x2 C	Eth-
-------------	------	------

HALOALKYNES	x3 C	Prop-
-------------	------	-------

OXYGEN COMPOUNDS

ALCOHOLS	x4 C	But-
----------	------	------

CARBOXYLIC ACIDS	x5 C	Pent-
------------------	------	-------

ALDEHYDES	x6 C	Hex-
-----------	------	------

KETONES	x7 C	Hept-
---------	------	-------

	x8 C	Oct-
--	------	------



QUESTION 2

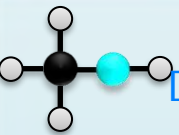
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G	$\begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_3-\text{C}-\text{CH}_2-\text{CH}_3 \\ \\ \text{C}=\text{O} \\ \\ \text{H}-\text{O} \end{array}$	H	$\begin{array}{cccc} & \text{H} & \text{H} & \text{O} & \text{H} \\ & & & & \\ \text{H}- & \text{C} & -\text{C} & -\text{C} & -\text{C}-\text{H} \\ & & & & \\ & \text{H} & \text{H} & & \text{H} \end{array}$

2.2 Write down the IUPAC name of compound: 2.2.1 E ; 2.2.2 H

2.2.1 2,3-dimethylbut-1-one

2.2.2 Butane-2-one / 2-butanone / butanone



ORGANIC MOLECULES

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TYPES OF FORMULAS:

Molecular Formula
Structure Formula
Condensed Formula

IUPAC**SUBSTITUENTS**

x2	di-
----	-----

x3	tri-
----	------

x4	tetra-
----	--------

HYDROCARBONS

ALKANES	$\text{C}_n\text{H}_{2n+2}$
---------	-----------------------------

ALKENES	C_nH_{2n}
---------	---------------------------

ALKYNES	$\text{C}_n\text{H}_{2n-2}$
---------	-----------------------------

HALOGEN COMPOUNDS

HALOALKANES

HALOALKENES

HALOALKYNES

OXYGEN COMPOUNDS

ALCOHOLS

CARBOXYLIC ACIDS

ALDEHYDES

KETONES

PREFIXES

x1 C	Meth-
------	-------

x2 C	Eth-
------	------

x3 C	Prop-
------	-------

x4 C	But-
------	------

x5 C	Pent-
------	-------

x6 C	Hex-
------	------

x7 C	Hept-
------	-------

x8 C	Oct-
------	------

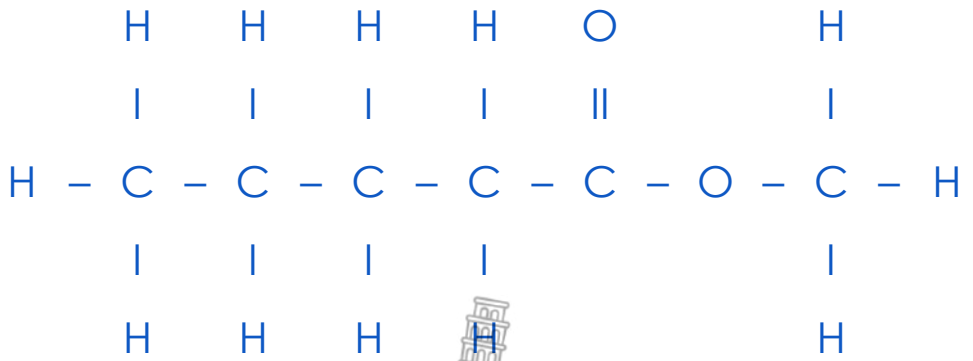


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2.3 Write down the:
2.3.1 STRUCTURAL formula of compound B



ORGANIC MOLECULES

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TYPES OF FORMULAS:
Molecular Formula
Structure Formula
Condensed Formula

I U P A C

SUBSTITUENTS	
x2	di-
x3	tri-
x4	tetra-

FUNCTIONAL GROUPS:
Hydroxyl group
Carbonyl group
Carboxyl group

HYDROCARBONS	
ALKANES	$\text{C}_n\text{H}_{2n+2}$
ALKENES	C_nH_{2n}
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ISOMERS:
Structural Isomers
Positional Isomers
Functional Isomers

HALOGEN COMPOUNDS	
HALOALKANES	
HALOALKENES	
HALOALKYNES	

PREFIXES	
x1 C	Meth-
x2 C	Eth-
x3 C	Prop-
x4 C	But-
x5 C	Pent-
x6 C	Hex-
x7 C	Hept-
x8 C	Oct-

OXYGEN COMPOUNDS	
ALCOHOLS	
CARBOXYLIC ACIDS	
ALDEHYDES	
KETONES	

QUESTION 2

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2.3 Write down the:

2.3.3 General formula of the homologous series to which compound E belongs

2.3.3 C_nH_{2n}

ORGANIC MOLECULES

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TYPES OF FORMULAS:

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FUNCTIONAL GROUPS:

Hydroxyl group
Carbonyl group
Carboxyl group

ISOMERS:

Structural Isomers
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HALOGEN COMPOUNDS

HALOALKANES

HALOALKENES

HALOALKYNES

OXYGEN COMPOUNDS

ALCOHOLS

CARBOXYLIC ACIDS

ALDEHYDES

KETONES

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x2 C Eth-

x3 C Prop-

x4 C But-

x5 C Pent-

x6 C Hex-

x7 C Hept-

x8 C Oct-



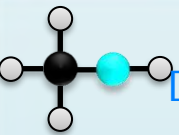
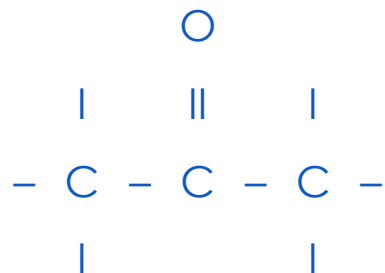
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2.3 Write down the:

2.3.4 STRUCTURAL formula of the FUNCTIONAL group of compound F



ORGANIC MOLECULES

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TYPES OF FORMULAS:

Molecular Formula
Structure Formula
Condensed Formula

I U P A C**SUBSTITUENTS**

x2	di-
----	-----

x3	tri-
----	------

x4	tetra-
----	--------

HYDROCARBONS

ALKANES	$\text{C}_n\text{H}_{2n+2}$
---------	-----------------------------

ALKENES	C_nH_{2n}
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ALKYNES	$\text{C}_n\text{H}_{2n-2}$
---------	-----------------------------

ISOMERS:

Structural Isomers
Positional Isomers
Functional Isomers

HALOGEN COMPOUNDS

HALOALKANES	x1 C	Meth-
-------------	------	-------

HALOALKENES	x2 C	Eth-
-------------	------	------

HALOALKYNES	x3 C	Prop-
-------------	------	-------

OXYGEN COMPOUNDS

ALCOHOLS	x4 C	But-
----------	------	------

CARBOXYLIC ACIDS	x5 C	Pent-
------------------	------	-------

ALDEHYDES	x6 C	Hex-
-----------	------	------

KETONES	x7 C	Hept-
---------	------	-------

	x8 C	Oct-
--	------	------



Alkanes	Alkenes	Alkynes
$\begin{array}{c} & \\ -\text{C}- & \text{C}- \\ & \end{array}$	$\begin{array}{c} & \\ -\text{C}=\text{C}- \\ & \end{array}$	$-\text{C}\equiv\text{C}-$

Haloalkanes	Haloalkenes	Haloalkynes
$\begin{array}{c} & \\ \text{X}-\text{C}- & \text{C}- \\ & \end{array}$	$\begin{array}{c} & \\ \text{X}-\text{C}=\text{C}- \\ & \end{array}$	$\text{X}-\text{C}\equiv\text{C}-$

Alcohols	Carboxylic acids	Aldehydes	Ketones	Esters
Hydroxyl group	Carboxyl group	Formyl group	Carbonyl group	
$\begin{array}{c} \\ -\text{C}-\text{O}-\text{H} \\ \end{array}$	$\begin{array}{c} \text{O} \\ \\ -\text{C}-\text{O}-\text{H} \end{array}$	$\begin{array}{c} \text{O} \\ \\ -\text{C}-\text{H} \end{array}$	$\begin{array}{c} & \text{O} & \\ & & \\ & -\text{C}- & \text{C}- \\ & & \end{array}$	$\begin{array}{c} & & \text{O} \\ & & \\ & -\text{O}- & \text{C}- \\ & & \end{array}$



QUESTION 2

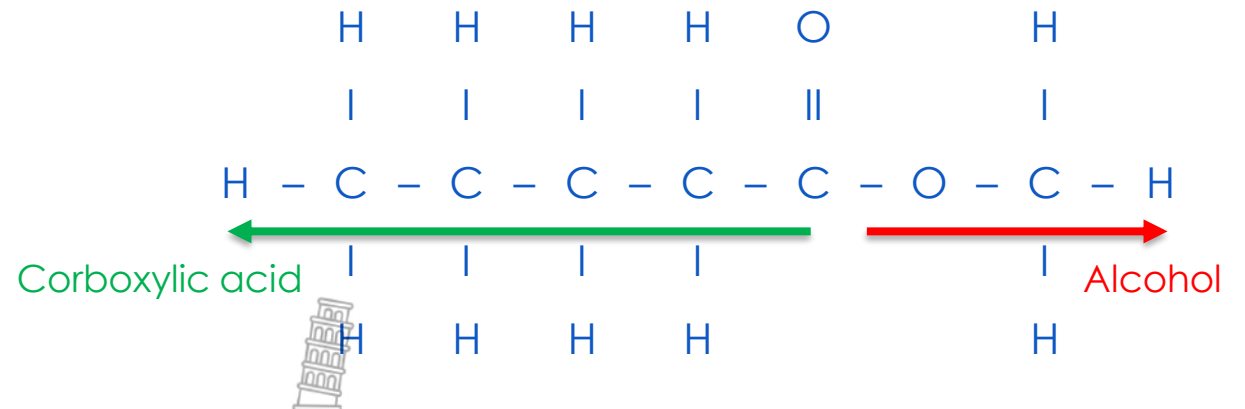
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C	4-ethyl-3,3-difluorohexane	D	Hexanoic acid
E	$\begin{array}{c} \text{CH}_2 \\ \\ \text{CH}_3 - \text{CH} - \text{C} - \text{CH}_3 \\ \\ \text{CH}_3 \end{array}$	F	$\begin{array}{c} \text{O} \\ \\ \text{CH}_3 - \text{CH} - \text{C} - \text{CH}_2 - \text{CH}_3 \\ \\ \text{CH}_3 \end{array}$
G	$\begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_3 - \text{C} - \text{CH}_2 - \text{CH}_3 \\ \\ \text{C} = \text{O} \\ \\ \text{H} - \text{O} \end{array}$	H	$\begin{array}{ccccccc} & \text{H} & \text{H} & \text{O} & \text{H} & & \\ & & & & & & \\ \text{H} & - \text{C} & - \text{C} & - \text{C} & - \text{C} & - \text{H} & \\ & & & & & & \\ & \text{H} & \text{H} & & \text{H} & & \end{array}$

2.3 Write down the:

2.3.5 IUPAC name of the alcohol needed to produce compound B

2.3.5 Methanol



ORGANIC MOLECULES

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Make sure you know the correct wording of all definitions!

TYPES OF FORMULAS:

Molecular Formula
Structure Formula
Condensed Formula

I U P A C**SUBSTITUENTS**

x2 di-

x3 tri-

x4 tetra-

HYDROCARBONS

ALKANES $\text{C}_n\text{H}_{2n+2}$

ALKENES C_nH_{2n}

ALKYNES $\text{C}_n\text{H}_{2n-2}$

FUNCTIONAL GROUPS:

Hydroxyl group
Carbonyl group
Carboxyl group

ISOMERS:

Structural Isomers
Positional Isomers
Functional Isomers

HALOGEN COMPOUNDS

HALOALKANES

HALOALKENES

HALOALKYNES

PREFIXES

x1 C Meth-

x2 C Eth-

x3 C Prop-

x4 C But-

x5 C Pent-

x6 C Hex-

x7 C Hept-

x8 C Oct-

OXYGEN COMPOUNDS

ALCOHOLS

CARBOXYLIC ACIDS

ALDEHYDES

KETONES

QUESTION 2

The letters A to H in the table below represent eight organic compounds.

A	Heptanoic acid $C_7H_{14}O_2$	B	$CH_3(CH_2)_3COOCH_3$ $C_6H_{12}O_2$ Ester
C	4-ethyl-3,3-difluorohexane	D	Hexanoic acid $C_6H_{12}O_2$ Carboxylic acid
E	$ \begin{array}{c} \text{CH}_2 \\ \\ \text{CH}_3-\text{CH}-\text{C}-\text{CH}_3 \\ \\ \text{CH}_3 \end{array} $ $C_8H_{16}F_2$ C_6H_{12}	F	$ \begin{array}{c} \text{O} \\ \\ \text{CH}_3-\text{CH}-\text{C}-\text{CH}_2-\text{CH}_3 \\ \\ \text{CH}_3 \end{array} $ $C_6H_{12}O$
G	$ \begin{array}{c} \text{CH}_3 \\ \\ \text{CH}_3-\text{C}-\text{CH}_2-\text{CH}_3 \\ \\ \text{C}=\text{O} \\ \\ \text{H}-\text{O} \end{array} $ $C_6H_{12}O_2$ Carboxylic acid	H	$ \begin{array}{cccc} \text{H} & \text{H} & \text{O} & \text{H} \\ & & & \\ \text{H}-\text{C} & -\text{C} & -\text{C} & -\text{C}-\text{H} \\ & & & \\ \text{H} & \text{H} & & \text{H} \end{array} $ C_4H_8O

2.4 Write down the letter(s) of the compound(s) that:

2.4.1 Is a FUNCTIONAL isomer of compound G

2.4.2 Are CHAIN isomers of each other

2.4.1 B

2.4.2 D & G

ORGANIC MOLECULES

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TYPES OF FORMULAS:

Molecular Formula
Structure Formula
Condensed Formula

I U P A C**SUBSTITUENTS**

x2 di-

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x4 tetra-

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ALKANES C_nH_{2n+2}

ALKENES C_nH_{2n}

ALKYNES C_nH_{2n-2}

FUNCTIONAL GROUPS:

Hydroxyl group
Carbonyl group
Carboxyl group

ISOMERS:

Structural Isomers
Positional Isomers
Functional Isomers

HALOGEN COMPOUNDS

HALOALKANES

HALOALKENES

HALOALKYNES

OXYGEN COMPOUNDS

ALCOHOLS

CARBOXYLIC ACIDS

ALDEHYDES

KETONES

PREFIXES

x1 C Meth-

x2 C Eth-

x3 C Prop-

x4 C But-

x5 C Pent-

x6 C Hex-

x7 C Hept-

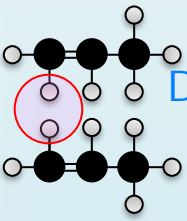
x8 C Oct-



QUESTION 3

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INTERMOLECULAR FORCES



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VAPOR PRESSURE: This is the pressure that an enclosed vapor in equilibrium with its liquid exerts on the surface of the liquid.

TYPES of IMF

London (Dispersion) forces
Dipole-Dipole forces
Hydrogen bonds

COMPARE COMPOUNDS

1. ID 2 IMK
2. Strongest IMK
3. Most ENERGY
4. Declare PROPERTIES

Melting Point & Boiling Point

DIRECTLY PROPORTIONAL TO THE STRENGTH OF THE IMF

Vapor pressure

INVERSE PROPORTIONAL TO THE STRENGTH OF THE IMF

The relationship between boiling point and the molecular mass of aldehydes, carboxylic acids and primary alcohols is investigated. Curves P, R and S are obtained. All compounds used are straight chain molecules.

3.1 Define the term boiling point.

3.2 Write down the conclusion that can be made for curve P.

3.3 Explain the answer to QUESTION 3.2 in terms of the structures of the compounds.

3.4 Curve R represents the alcohols.

3.4.1 Which homologous series is represented by curve S?

3.4.2 Explain the answer to QUESTION 3.4.1 by referring to the strength of intermolecular forces.

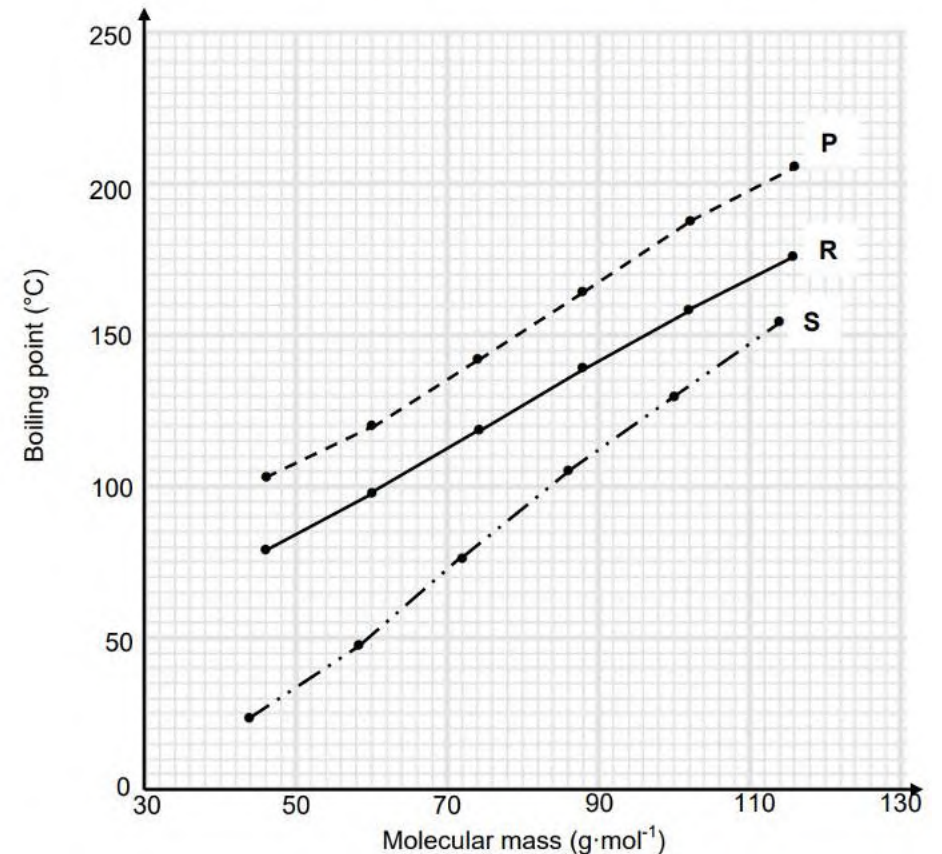
3.5 For curve R, write down the:

3.5.1 Molecular mass of the compound with a boiling point of 97 °C

3.5.2 IUPAC name of the compound in QUESTION 3.5.1

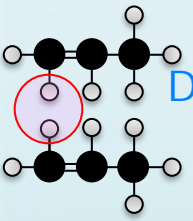
3.6 Two compounds, A and B, used in this investigation have a molecular mass of 74 g.mol⁻¹. A has a boiling point of 118 °C and B a boiling point of 142 °C. Explain the difference in these boiling points by referring to the structures of these compounds.

GRAPH OF BOILING POINT VERSUS MOLECULAR MASS



QUESTION 3

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INTERMOLECULAR FORCES

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DIRECTLY PROPORTIONAL TO THE STRENGTH OF THE IMF

Vapor pressure

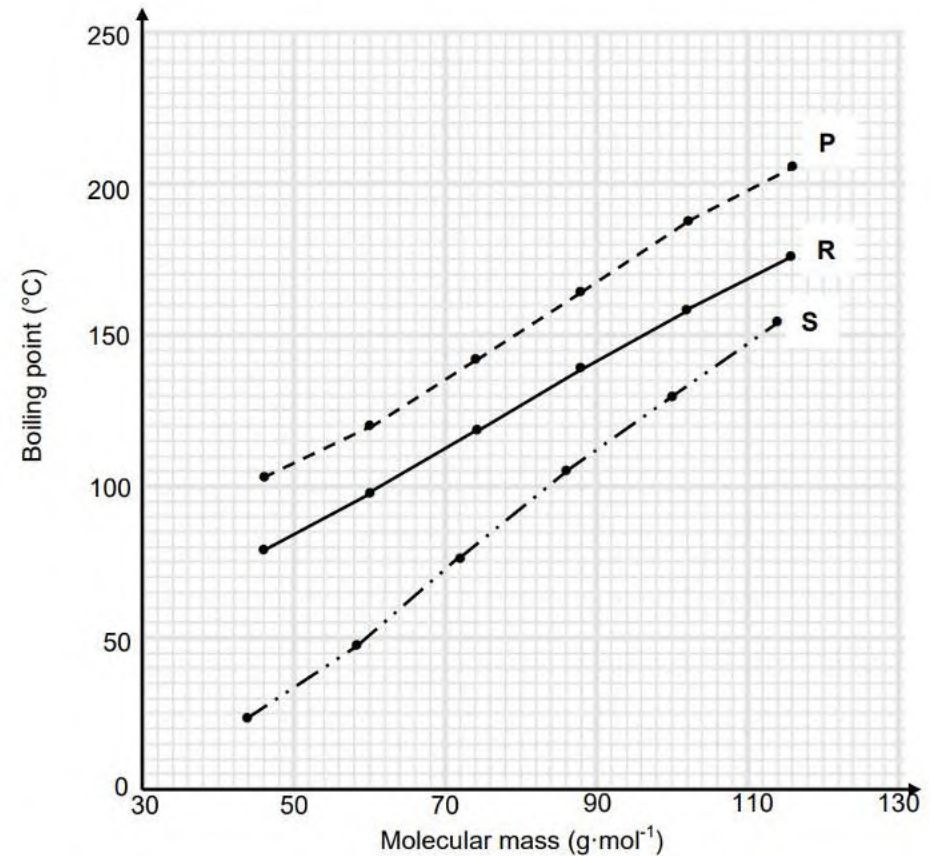
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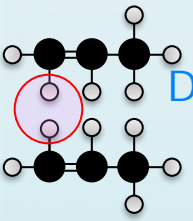
BOILING POINT: The temperature at which the vapour pressure of a LIQUID equals the atmospheric pressure.

GRAPH OF BOILING POINT VERSUS MOLECULAR MASS



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3.2 Write down the conclusion that can be made for curve P.

The higher the molecular mass, the higher the boiling point.

OR

As the molecular mass increases, so does the boiling point.

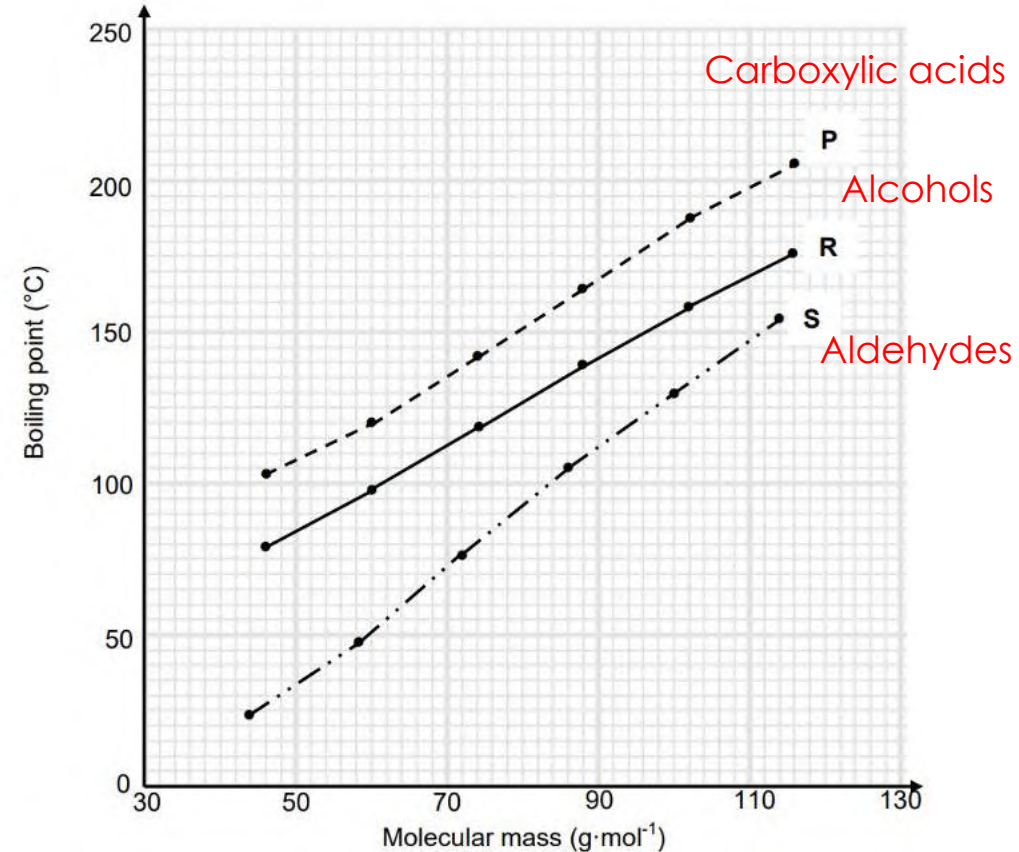
OR

The longer the C-chain, the higher the boiling point. The longer the C-chain, the higher the boiling point.

OR

The boiling point and the molecular mass are proportional.

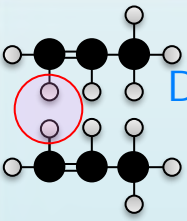
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3.3 Explain the answer to QUESTION 3.2 in terms of the structures of the compounds.

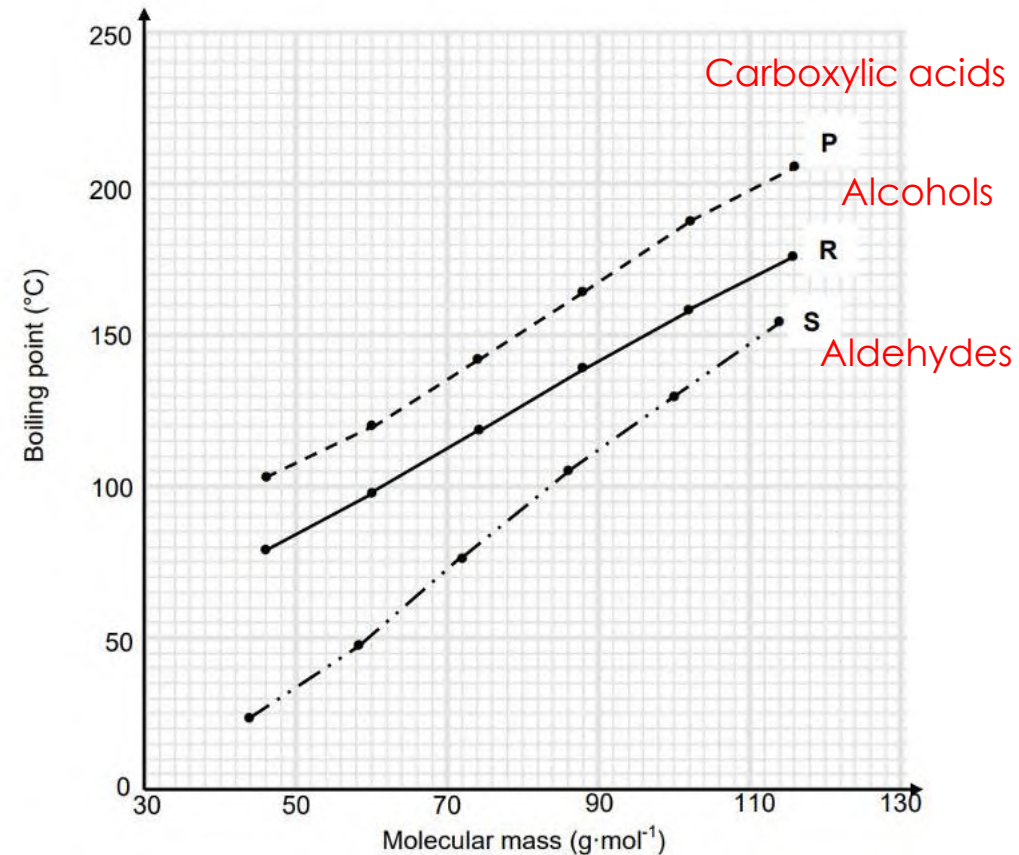
Strength of the intermolecular forces increases with increase In:

- molar mass
- Chain length
- contact surface.

More sites for London forces.

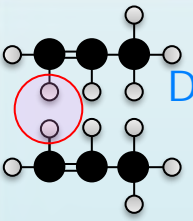
More energy required to overcome intermolecular forces.

GRAPH OF BOILING POINT VERSUS MOLECULAR MASS



QUESTION 3

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INTERMOLECULAR FORCES

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3.4.2 Explain the answer to QUESTION 3.4.1 by referring to the strength of intermolecular forces.

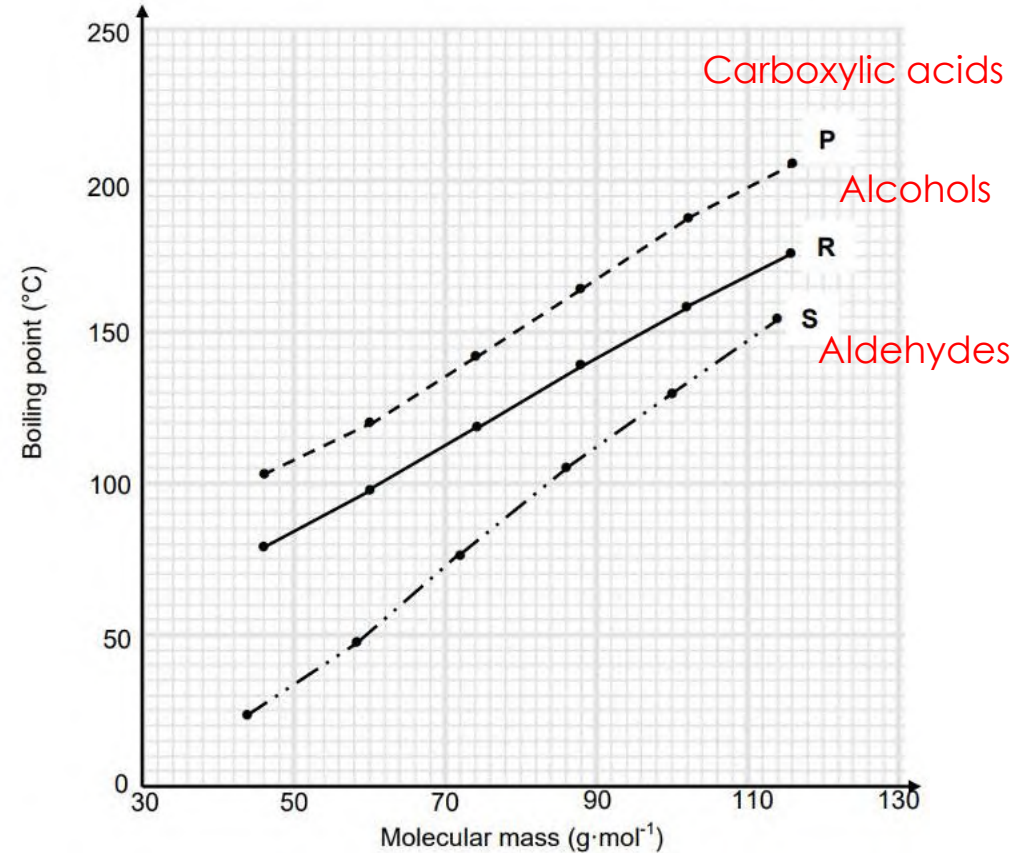
3.4.1 Aldehydes

3.4.2 Aldehydes have the weakest intermolecular forces.

Thus the aldehydes have the lowest boiling points.

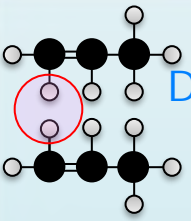
The least energy needed to overcome the intermolecular forces.

GRAPH OF BOILING POINT VERSUS MOLECULAR MASS



QUESTION 3

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INTERMOLECULAR FORCES

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The relationship between boiling point and the molecular mass of aldehydes, carboxylic acids and primary alcohols is investigated. Curves P, R and S are obtained. All compounds used are straight chain molecules.

3.5 For curve R, write down the:

3.5.1 Molecular mass of the compound with a boiling point of 97 °C

3.5.2 IUPAC name of the compound in QUE!

3.5.1 Alcohol with a molar mass of 60 g.mol⁻¹

3.5.2 Propan-1-ol

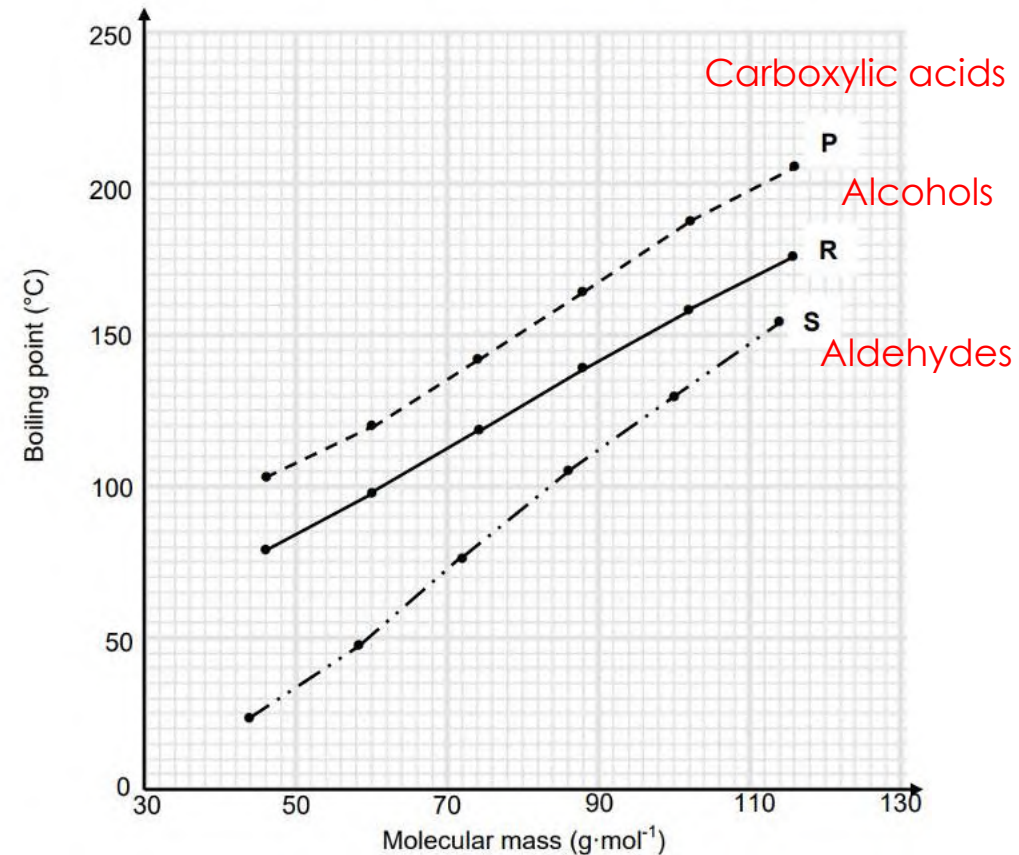


$$C_1H_4O = 1(12) + 4(1) + 1(16) = 32 \text{ g.mol}^{-1}$$

$$C_2H_6O = 2(12) + 6(1) + 1(16) = 46 \text{ g.mol}^{-1}$$

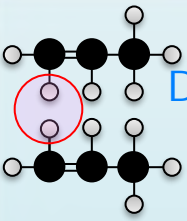
$$C_3H_8O = 3(12) + 8(1) + 1(16) = 60 \text{ g.mol}^{-1}$$

GRAPH OF BOILING POINT VERSUS MOLECULAR MASS



QUESTION 3

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The relationship between boiling point and the molecular mass of aldehydes, carboxylic acids and primary alcohols is investigated. Curves P, R and S are obtained. All compounds used are straight chain molecules.

3.6 Two compounds, A and B, used in this investigation have a molecular mass of $74 \text{ g}\cdot\text{mol}^{-1}$. A has a boiling point of $118 \text{ }^\circ\text{C}$ and B a boiling point of $142 \text{ }^\circ\text{C}$. Explain the difference in these boiling points by referring to the structures of these compounds.

Both compounds has strong hydrogen bonds.

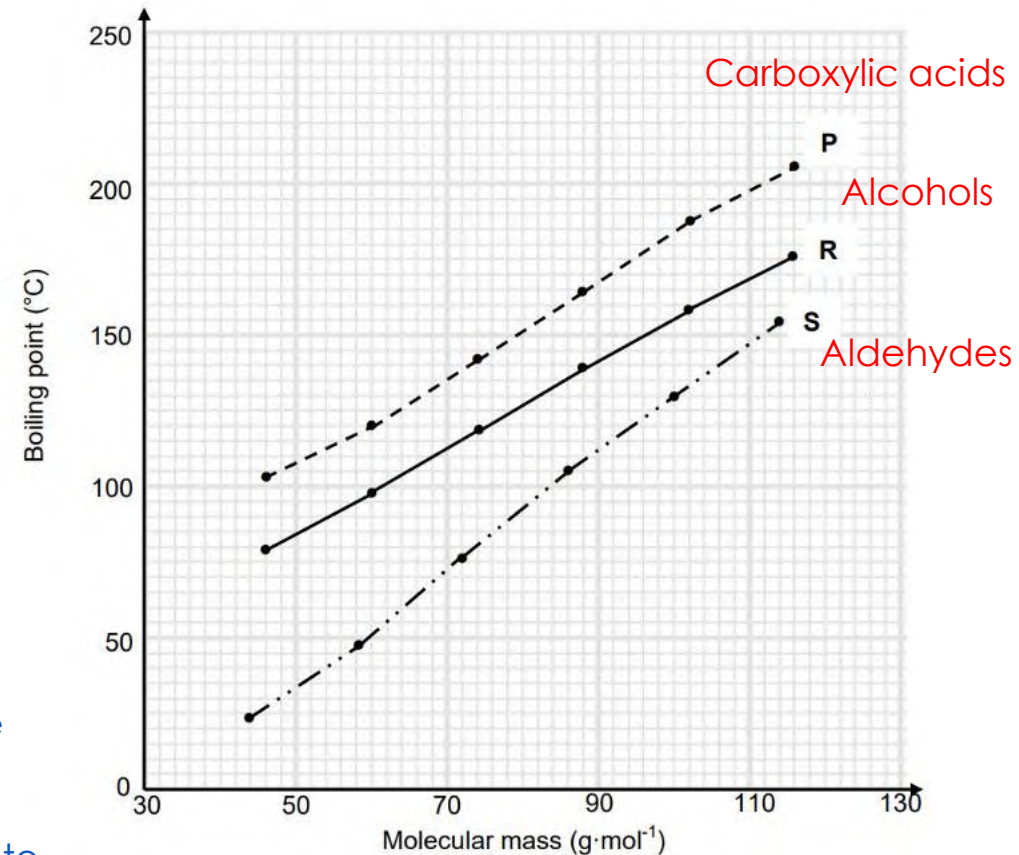
Carboxylic acids have two sites for hydrogen bonds while Alcohols have only one site for a hydrogen bond.

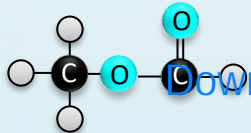
Thus, the intermolecular forces of Carboxylic acids are stronger than the intermolecular forces of Alcohols.

Carboxylic acids require more energy to overcome the intermolecular forces.

Carboxylic acids therefore have higher boiling points.

GRAPH OF BOILING POINT VERSUS MOLECULAR MASS





ORGANIC REACTIONS

QUESTION 4

NSC 2023



Combustion, Esterification, Cracking

ADDITION REACTIONS

HIDROGENATIONS	ALKENE + $H_2 \rightarrow$ ALKANE (CATALYST: Pt/Ni/Pd & Non-polar solvent, Heat)
HALOGENATION	ALKENE + $X_2 \rightarrow$ HALOALKANE (NO H_2O)
HIDROHALOGENATION	ALKENE + $HX \rightarrow$ HALOALKANE (NO H_2O)
HIDRATATION	ALKENE + $H_2O \rightarrow$ ALCOHOL (H_2SO_4 (dil) / H_3PO_4 (dil) & Heat in the form of steam)

ELIMINATION REACTIONS

DEHYDROGENATION	ALKANE - $H_2 \rightarrow$ ALKENE (CATALYST: Pt / Ni / Pd)
DEHALOGENATION	HALOALKANE - $X_2 \rightarrow$ ALKENE (Unreactive solvent)
DEHYDROHALOGENATION	HALOALKANE - $HX \rightarrow$ ALKENE (NaOH(c)/KOH(c), Ethanol, Heat ... Reflux)
DEHIDRATATION	ALCOHOL - $H_2O \rightarrow$ ALKENE (CATALYST: H_2SO_4 (c) / H_3PO_4 (c) & Heat)

SUBSTITUTION REACTIONS




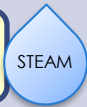


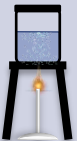



HALOGENATION	ALKANE + $X_2 \rightarrow$ HALOALKANE (IN THE PRESENCE OF SUNLIGHT)
HYDROLYSIS	HALOALKANE + $OH^- \rightarrow$ ALCOHOL (NaOH(aq) / KOH(aq) & Ethanol & Low Heat)
HIDROHALOGENATION	ALCOHOL + $HX \rightarrow$ HALOALKANE (CATALYST: H_2SO_4 (c) / H_3PO_4 (c) & Heat)

MARKOVNIKOV
ADDITION REACTIONS

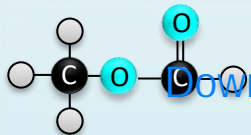
ZAITEV
ELIMINATION REACTIONS





Hydrogenation	Halogenation	Hydrohalogenation	Hydration
 <p>Pt / Ni / Pd Non-polar</p>			<p>STRONG H_2SO_4 DILUTE H_3PO_4</p> 
Dehydrogenation	Dehalogenation	Dehydrohalogenation	Dehydration
<p>Pt / Ni / Pd</p>	 <p>Unreactive</p>	<p>STRONG NaOH (c) KOH</p>  <p>Etanol</p>	<p>STRONG H_2SO_4 (c) H_3PO_4</p> 
Halogenation	Hydrolysis		Hydrohalogenation
	<p>STRONG NaOH DILUTE KOH</p>  <p>Etanol</p>	<p>STRONG H_2SO_4 (c) H_3PO_4</p> 	





ORGANIC REACTIONS

Combustion, Esterification, Cracking

ADDITION REACTIONS

HIDROGENATIONS	ALKENE + $H_2 \rightarrow$ ALKANE (CATALYST: Pt/Ni/Pd & Non-polar solvent, Heat)
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DEHIDRATATION	ALCOHOL - $H_2O \rightarrow$ ALKENE (CATALYST: H_2SO_4 (c) / H_3PO_4 (c) & Heat)

SUBSTITUTION REACTIONS

HALOGENATION	ALKANE + $X_2 \rightarrow$ HALOALKANE (IN THE PRESENCE OF SUNLIGHT)
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MARKOVNIKOV
ADDITION REACTIONS

ZAITEV
ELIMINATION REACTIONS

QUESTION 4

NSC 2023

4.1 Consider the cracking reaction below. $C_{16}H_{34} \rightarrow C_6H_{14} + C_6H_x + 2C_yH_z$

4.1.1 Define cracking.

4.1.2 Write down the values represented by x, y and z in the equation above.

Compound C_6H_{14} undergoes complete combustion.

4.1.3 Using MOLECULAR FORMULAE, write down the balanced equation for this reaction.

4.2 Consider the equations for reactions I to III below. A and B represent organic compounds that are POSITIONAL ISOMERS. X is an inorganic product.

I	$CH_3CH_2CHCHCH_3 + HCl \rightarrow A + B$
II	$A \xrightarrow[\Delta]{H_2O} CH_3CH_2CH_2CH(OH)CH_3 + X$
III	$CH_3CH_2CH_2CH(OH)CH_3 \longrightarrow CH_3CH_2CHCHCH_3 + H_2O$

Write down the:

4.2.1 Definition of positional isomers

4.2.2 Type of reaction represented by reaction I

4.2.3 STRUCTURAL formula of compound B

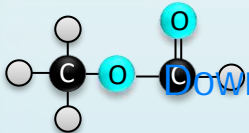
4.2.4 Formula of X

4.2.5 Inorganic reagent for reaction III

Compound A can be converted directly to the organic product of reaction III.

4.2.6 Besides heat, write down the reaction condition needed for this conversion.

4.2.7 Write down TWO terms that describe this type of reaction.



ORGANIC REACTIONS

QUESTION 4

NSC 2023

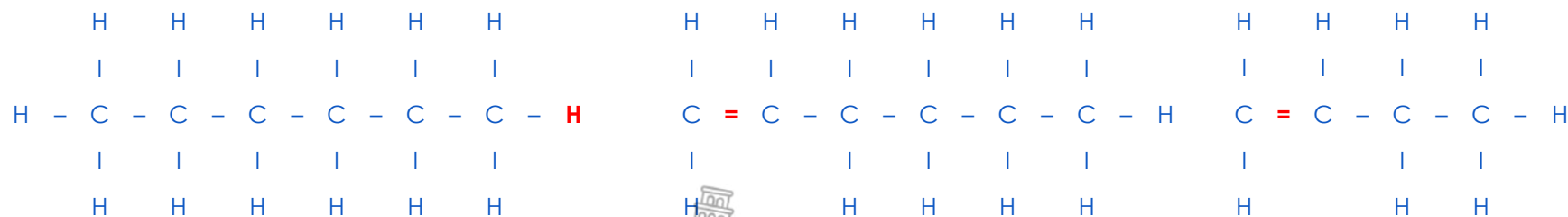
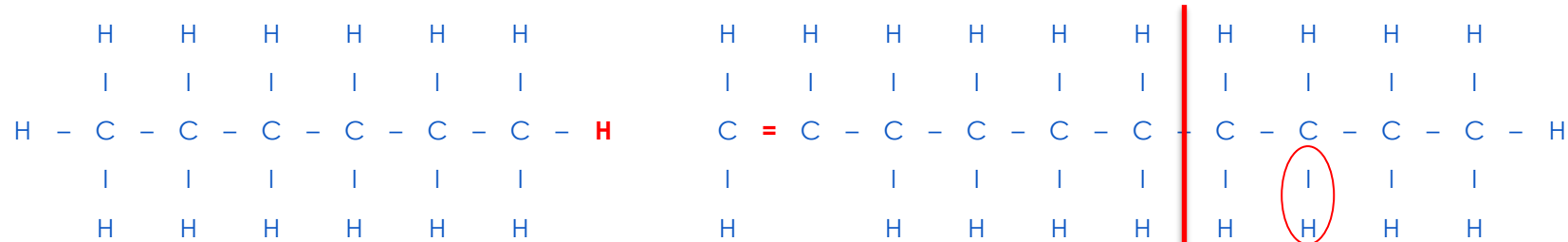
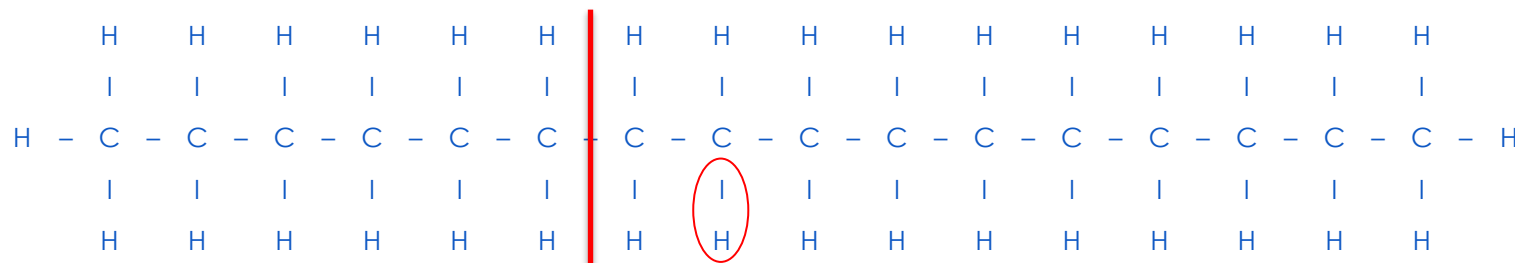
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(Alkane) (Alkane) (Alkene) (Alkene)

4.1.1 Define cracking.

4.1.2 Write down the values represented by x, y and z in the equation above.

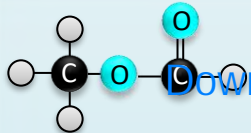
The chemical process/reaction in which longer chain hydrocarbon/alkane molecule is broken down into shorter (more useful) molecules.



MARKOVNIKOV
ADDITION REACTIONS

ZAITEV
ELIMINATION REACTIONS





ORGANIC REACTIONS

Combustion, Esterification, Cracking

ADDITION REACTIONS

HIDROGENATIONS	ALKENE + H ₂ → ALKANE (CATALYST: Pt/Ni/Pd & Non-polar solvent, Heat)
HALOGENATION	ALKENE + X ₂ → HALOALKANE (NO H ₂ O)
HIDROHALOGENATION	ALKENE + HX → HALOALKANE (NO H ₂ O)
HIDRATATION	ALKENE + H ₂ O → ALCOHOL (H ₂ SO ₄ (dil) / H ₃ PO ₄ (dil) & Heat in the form of steam)

ELIMINATION REACTIONS

DEHYDROGENATION	ALKANE - H ₂ → ALKENE (CATALYST: Pt / Ni / Pd)
DEHALOGENATION	HALOALKANE - X ₂ → ALKENE (Unreactive solvent)
DEHYDROHALOGENATION	HALOALKANE - HX → ALKENE (NaOH(c)/KOH(c), Ethanol, Heat ... Reflux)
DEHIDRATATION	ALCOHOL - H ₂ O → ALKENE (CATALYST: H ₂ SO ₄ (c) / H ₃ PO ₄ (c) & Heat)

SUBSTITUTION REACTIONS

HALOGENATION	ALKANE + X ₂ → HALOALKANE (IN THE PRESENCE OF SUNLIGHT)
HYDROLYSIS	HALOALKANE + OH ⁻ → ALCOHOL (NaOH(aq) / KOH(aq) & Ethanol & Low Heat)
HIDROHALOGENATION	ALCOHOL + HX → HALOALKANE (CATALYST: H ₂ SO ₄ (c) / H ₃ PO ₄ (c) & Heat)

MARKOVNIKOV
ADDITION REACTIONS

ZAITEV
ELIMINATION REACTIONS

QUESTION 4

Compound C₆H₁₄ undergoes complete combustion.

4.1.3 Using MOLECULAR FORMULAE, write down the balanced equation for this reaction.

EXCESS OXYGEN : Compound + O₂ → CO₂ + H₂O

Reactants & Products:



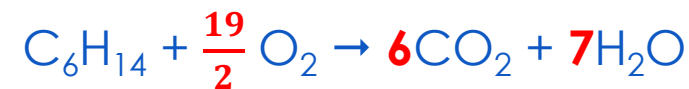
Balance C:



Balance H:

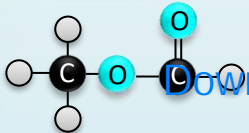


Balance O:



Empirical :



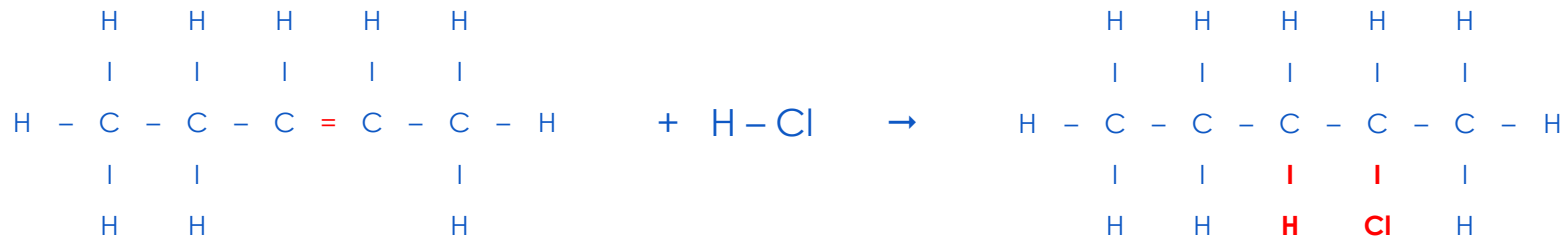


ORGANIC REACTIONS

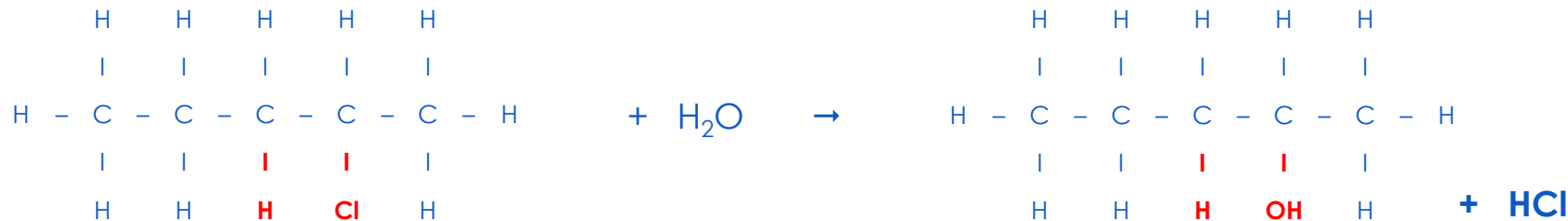
QUESTION 4

NSC 2023

4.2 Consider the equations for reactions I to III below. A and B represent organic compounds that are POSITIONAL ISOMERS. X is an inorganic product.



I	$\text{CH}_3\text{CH}_2\text{CHCHCH}_3 + \text{HCl} \rightarrow \text{A} + \text{B}$
II	$\text{A} \xrightarrow[\Delta]{\text{H}_2\text{O}} \text{CH}_3\text{CH}_2\text{CH}_2\text{CH}(\text{OH})\text{CH}_3 + \text{X}$
III	$\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}(\text{OH})\text{CH}_3 \longrightarrow \text{CH}_3\text{CH}_2\text{CHCHCH}_3 + \text{H}_2\text{O}$



Write down the:
4.2.3 STRUCTURAL formula of compound B

SEE STRUCTURE DRAWN ABOVE.



Combustion, Esterification, Cracking

ADDITION REACTIONS

HIDROGENATIONS	ALKENE + $\text{H}_2 \rightarrow$ ALKANE (CATALYST: Pt/Ni/Pd & Non-polar solvent, Heat)
HALOGENATION	ALKENE + $\text{X}_2 \rightarrow$ HALOALKANE (NO H_2O)
HIDROHALOGENATION	ALKENE + $\text{HX} \rightarrow$ HALOALKANE (NO H_2O)
HIDRATATION	ALKENE + $\text{H}_2\text{O} \rightarrow$ ALCOHOL (H_2SO_4 (dil) / H_3PO_4 (dil) & Heat in the form of steam)

ELIMINATION REACTIONS

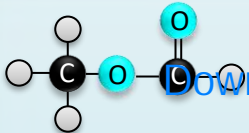
DEHYDROGENATION	ALKANE - $\text{H}_2 \rightarrow$ ALKENE (CATALYST: Pt / Ni / Pd)
DEHALOGENATION	HALOALKANE - $\text{X}_2 \rightarrow$ ALKENE (Unreactive solvent)
DEHYDROHALOGENATION	HALOALKANE - $\text{HX} \rightarrow$ ALKENE (NaOH(c)/KOH(c), Ethanol, Heat ... Reflux)
DEHYDRATION	ALCOHOL - $\text{H}_2\text{O} \rightarrow$ ALKENE (CATALYST: H_2SO_4 (c) / H_3PO_4 (c) & Heat)

SUBSTITUTION REACTIONS

HALOGENATION	ALKANE + $\text{X}_2 \rightarrow$ HALOALKANE (IN THE PRESENCE OF SUNLIGHT)
HYDROLYSIS	HALOALKANE + $\text{OH}^- \rightarrow$ ALCOHOL (NaOH(aq) / KOH(aq) & Ethanol & Low Heat)
HIDROHALOGENATION	ALCOHOL + $\text{HX} \rightarrow$ HALOALKANE (CATALYST: H_2SO_4 (c) / H_3PO_4 (c) & Heat)

MARKOVNIKOV
ADDITION REACTIONS

ZAITEV
ELIMINATION REACTIONS



ORGANIC REACTIONS

Combustion, Esterification, Cracking

ADDITION REACTIONS

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HALOGENATION	ALKENE + X ₂ → HALOALKANE (NO H ₂ O)
HIDROHALOGENATION	ALKENE + HX → HALOALKANE (NO H ₂ O)
HIDRATATION	ALKENE + H ₂ O → ALCOHOL (H ₂ SO ₄ (dil) / H ₃ PO ₄ (dil) & Heat in the form of steam)

ELIMINATION REACTIONS

DEHYDROGENATION	ALKANE - H ₂ → ALKENE (CATALYST: Pt / Ni / Pd)
DEHALOGENATION	HALOALKANE - X ₂ → ALKENE (Unreactive solvent)
DEHYDROHALOGENATION	HALOALKANE - HX → ALKENE (NaOH(c)/KOH(c), Ethanol, Heat ... Reflux)

DEHYDRATION: ALCOHOL - H₂O → ALKENE
(CATALYST: H₂SO₄(c) / H₃PO₄(c) & Heat)

SUBSTITUTION REACTIONS

HALOGENATION	ALKANE + X ₂ → HALOALKANE (IN THE PRESENCE OF SUNLIGHT)
HYDROLYSIS	HALOALKANE + OH ⁻ → ALCOHOL (NaOH(aq) / KOH(aq) & Ethanol & Low Heat)
HIDROHALOGENATION	ALCOHOL + HX → HALOALKANE (CATALYST: H ₂ SO ₄ (c) / H ₃ PO ₄ (c) & Heat)

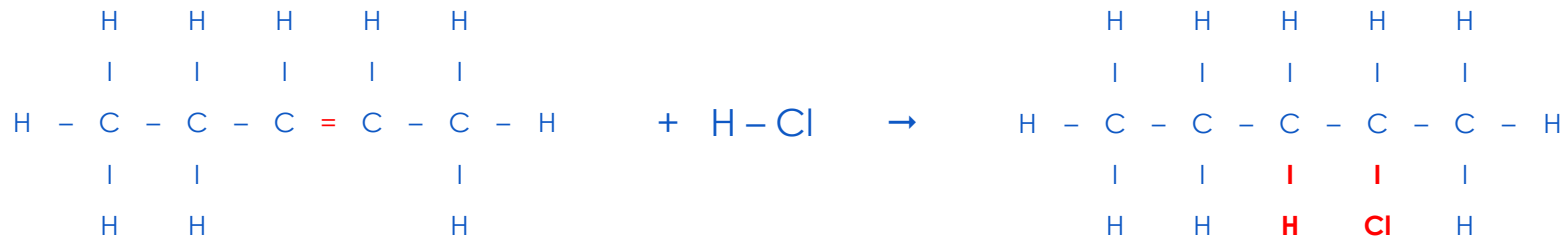
MARKOVNIKOV
ADDITION REACTIONS

ZAITEV
ELIMINATION REACTIONS

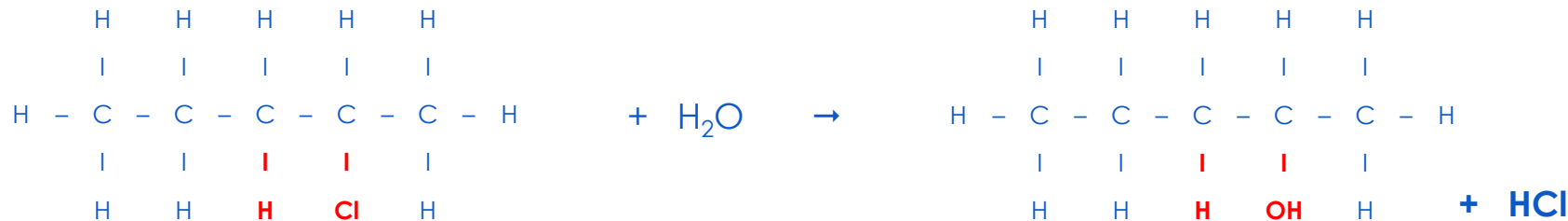
QUESTION 4

NSC 2023

4.2 Consider the equations for reactions I to III below. A and B represent organic compounds that are POSITIONAL ISOMERS. X is an inorganic product.



I	CH ₃ CH ₂ CHCHCH ₃ + HCl → A + B
II	A $\xrightarrow[\Delta]{\text{H}_2\text{O}}$ CH ₃ CH ₂ CH ₂ CH(OH)CH ₃ + X
III	CH ₃ CH ₂ CH ₂ CH(OH)CH ₃ → CH ₃ CH ₂ CHCHCH ₃ + H ₂ O

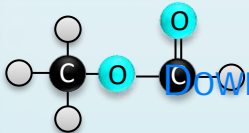


Write down the:

4.2.5 Inorganic reagent for reaction III

Concentrated sulfuric acid / H₂SO₄ (c) / Concentrated phosphoric acid / H₃PO₄ (c)



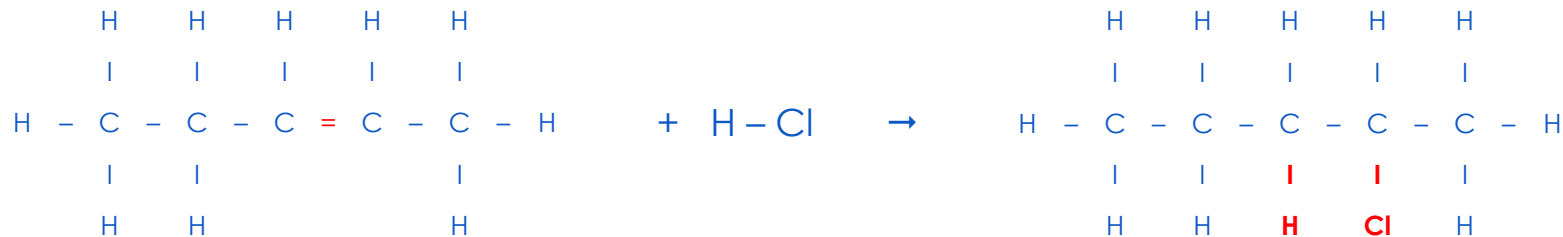


ORGANIC REACTIONS

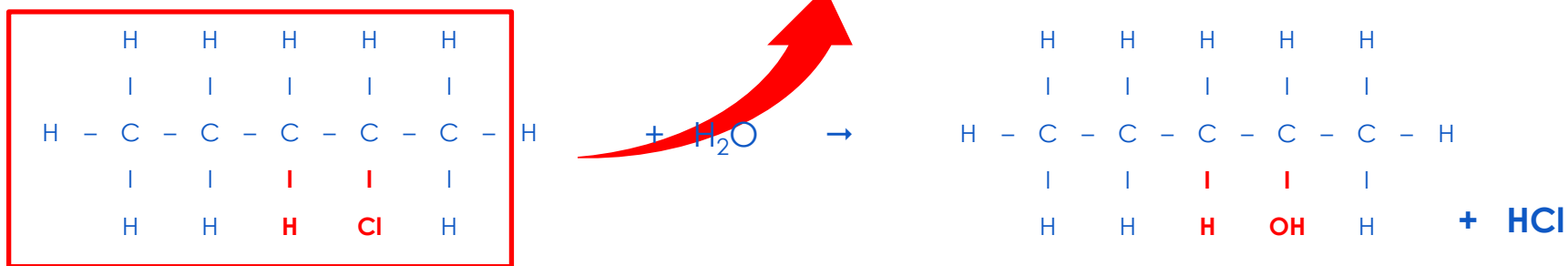
QUESTION 4

NSC 2023

4.2 Consider the equations for reactions I to III below. A and B represent organic compounds that are POSITIONAL ISOMERS. X is an inorganic product.



I	$\text{CH}_3\text{CH}_2\text{CHCHCH}_3 + \text{HCl} \rightarrow \text{A} + \text{B}$
II	$\text{A} \xrightarrow[\Delta]{\text{H}_2\text{O}} \text{CH}_3\text{CH}_2\text{CH}_2\text{CH}(\text{OH})\text{CH}_3 + \text{X}$
III	$\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}(\text{OH})\text{CH}_3 \longrightarrow \text{CH}_3\text{CH}_2\text{CHCHCH}_3 + \text{H}_2\text{O}$



Compound A can be converted directly to the organic product of reaction III.
4.2.6 Besides heat, write down the reaction condition needed for this conversion

Strong base/NaOH /KOH/ LiOH/sodium hydroxide/potassium hydroxide/dissolved in ethanol

ADDITION REACTIONS

HIDROGENATIONS	ALKENE + $\text{H}_2 \rightarrow$ ALKANE (CATALYST: Pt/Ni/Pd & Non-polar solvent, Heat)
HALOGENATION	ALKENE + $\text{X}_2 \rightarrow$ HALOALKANE (NO H_2O)
HIDROHALOGENATION	ALKENE + $\text{HX} \rightarrow$ HALOALKANE (NO H_2O)
HIDRATATION	ALKENE + $\text{H}_2\text{O} \rightarrow$ ALCOHOL (H_2SO_4 (dil) / H_3PO_4 (dil) & Heat in the form of steam)

ELIMINATION REACTIONS

DEHYDROGENATION	ALKANE - $\text{H}_2 \rightarrow$ ALKENE (CATALYST: Pt / Ni / Pd)
DEHALOGENATION	HALOALKANE - $\text{X}_2 \rightarrow$ ALKENE (Unreactive solvent)
DEHYDROHALOGENATION	HALOALKANE - $\text{HX} \rightarrow$ ALKENE (NaOH(c)/KOH(c), Ethanol, Heat ... Reflux)
DEHYDRATION	ALCOHOL - $\text{H}_2\text{O} \rightarrow$ ALKENE (CATALYST: H_2SO_4 (c) / H_3PO_4 (c) & Heat)

SUBSTITUTION REACTIONS

HALOGENATION	ALKANE + $\text{X}_2 \rightarrow$ HALOALKANE (IN THE PRESENCE OF SUNLIGHT)
HYDROLYSIS	HALOALKANE + $\text{OH}^- \rightarrow$ ALCOHOL (NaOH(aq) / KOH(aq) & Ethanol & Low Heat)
HIDROHALOGENATION	ALCOHOL + $\text{HX} \rightarrow$ HALOALKANE (CATALYST: H_2SO_4 (c) / H_3PO_4 (c) & Heat)

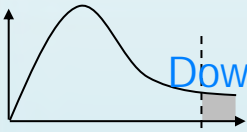
MARKOVNIKOV
ADDITION REACTIONS

ZAITEV
ELIMINATION REACTIONS

QUESTION 5

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REACTION RATE



Make sure you know the correct wording of all definitions!

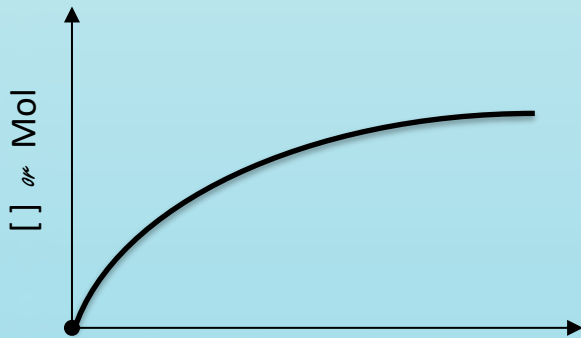
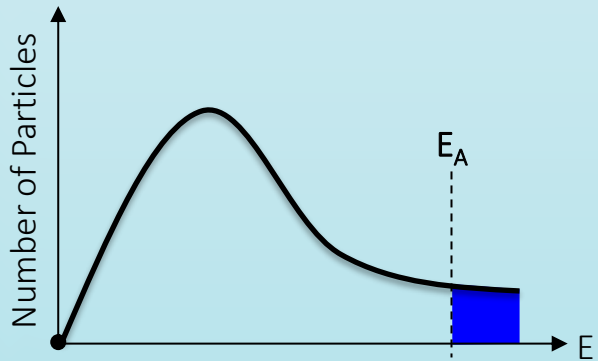
FACTORS:

Temperature / Concentration
Catalyst / Pressure
Nature of the reactants
State of division

$$RT = \pm \frac{\Delta[\quad]}{\Delta\text{Time}}$$

COLLISION THEORY

1. Identify the factor that differs.
2. Difference between particles.
3. Number of effective collisions per unit time.
4. Effect on reaction rate.

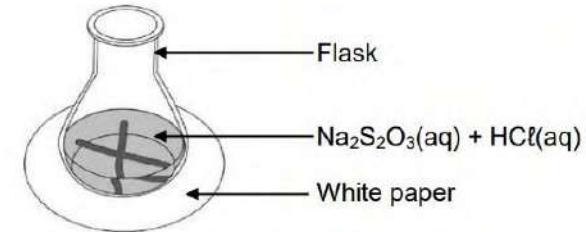


The reaction between EXCESS dilute hydrochloric acid and sodium thiosulphate is used to investigate factors that influence reaction rate. $\text{Na}_2\text{S}_2\text{O}_3(\text{aq}) + 2\text{HCl}(\text{aq}) \rightarrow 2\text{NaCl}(\text{aq}) + \text{S}(\text{s}) + \text{H}_2\text{O}(\text{l}) + \text{SO}_2(\text{g})$

The concentration of $\text{HCl}(\text{aq})$ used is $1 \text{ mol}\cdot\text{dm}^{-3}$. The same volume of $\text{HCl}(\text{aq})$ is used in each run. The time taken for the cross on the paper under the flask to become invisible is measured.

The table below summarises the reaction conditions and results of the experiment.

RUN	VOLUME $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ (cm^3)	VOLUME $\text{H}_2\text{O}(\text{l})$ ADDED (cm^3)	CONCENTRATION $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ ($\text{mol}\cdot\text{dm}^{-3}$)	TIME (s)
1	50	0	0,13	20,4
2	40	10	0,10	26,7
3	30	20	P	33,3



5.1 Define reaction rate.

5.2 Write down the independent variable for this investigation.

5.3 Calculate the value of P in the table.

5.4 When 0,21 g of sulphur has formed in Run 1, the cross becomes invisible. Calculate the average reaction rate with respect to sodium thiosulphate, $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$, in $\text{g}\cdot\text{s}^{-1}$.

Another investigation is performed at different temperatures.

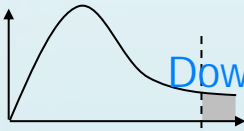
5.5 Sketch the Maxwell-Boltzmann distribution curve for the reaction at 20°C . Label this curve as A. On the same set of axis, draw the curve that will be obtained at 35°C and label it as B.

5.6 Explain the effect of temperature on reaction rate in terms of the collision theory.

QUESTION 5

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REACTION RATE



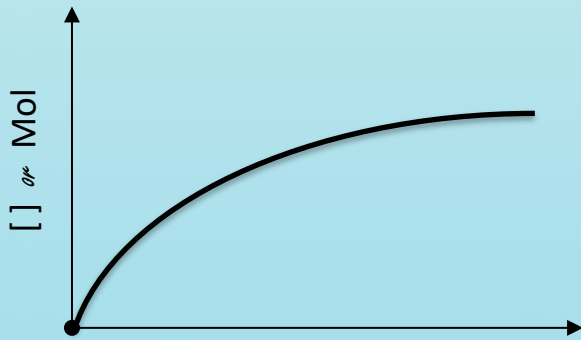
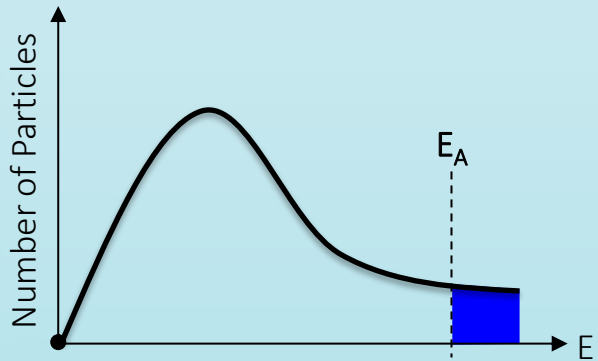
Make sure you know the correct wording of all definitions!

- FACTORS:**
 Temperature / Concentration
 Catalyst / Pressure
 Nature of the reactants
 State of division

$$RT = \pm \frac{\Delta[]}{\Delta \text{Time}}$$

COLLISION THEORY

1. Identify the factor that differs.
2. Difference between particles.
3. Number of effective collisions per unit time.
4. Effect on reaction rate.

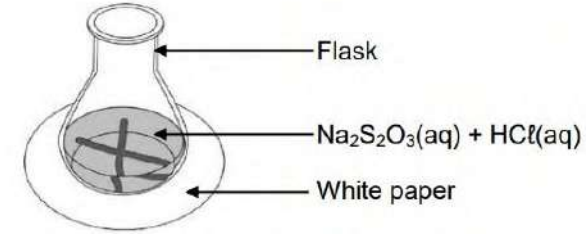


The reaction between EXCESS dilute hydrochloric acid and sodium thiosulphate is used to investigate factors that influence reaction rate. $\text{Na}_2\text{S}_2\text{O}_3(\text{aq}) + 2\text{HCl}(\text{aq}) \rightarrow 2\text{NaCl}(\text{aq}) + \text{S}(\text{s}) + \text{H}_2\text{O}(\text{l}) + \text{SO}_2(\text{g})$

The concentration of $\text{HCl}(\text{aq})$ used is $1 \text{ mol}\cdot\text{dm}^{-3}$. The same volume of $\text{HCl}(\text{aq})$ is used in each run. The time taken for the cross on the paper under the flask to become invisible is measured.

The table below summarises the reaction conditions and results of the experiment.

RUN	VOLUME $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ (cm^3)	VOLUME $\text{H}_2\text{O}(\text{l})$ ADDED (cm^3)	CONCENTRATION $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ ($\text{mol}\cdot\text{dm}^{-3}$)	TIME (s)
1	50	0	0,13	20,4
2	40	10	0,10	26,7
3	30	20	P	33,3



5.1 Define reaction rate.

REACTION RATE: The change in concentration of reactants or products per unit time.



QUESTION 5



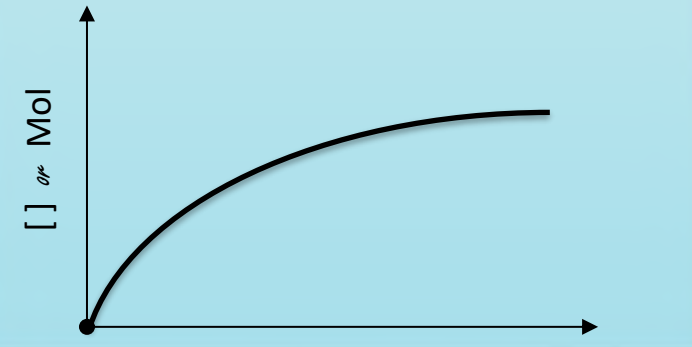
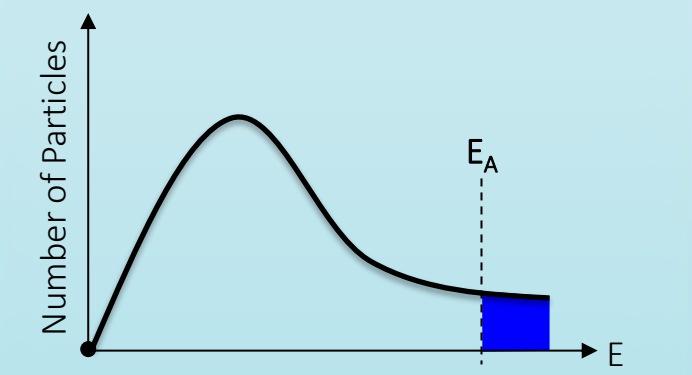
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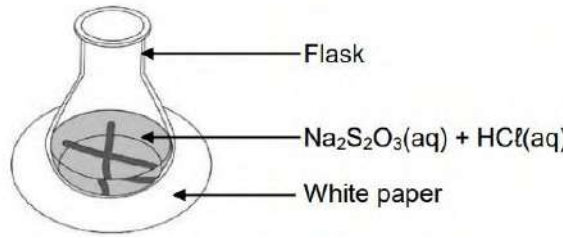
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1	50	0	0,13	20,4
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3	30	20	P	33,3

50 cm^3



5.2 Write down the independent variable for this investigation.

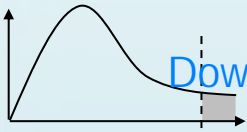
Concentration of the sodium thiosulphate ($\text{Na}_2\text{S}_2\text{O}_3$)



QUESTION 5

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REACTION RATE



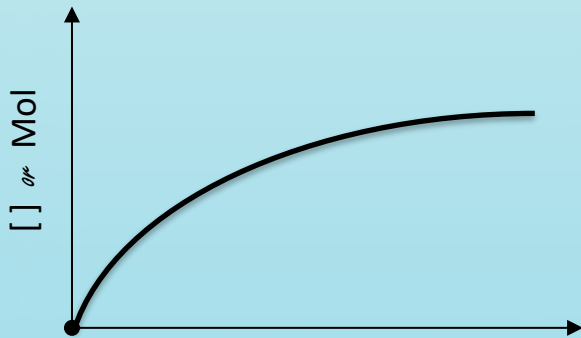
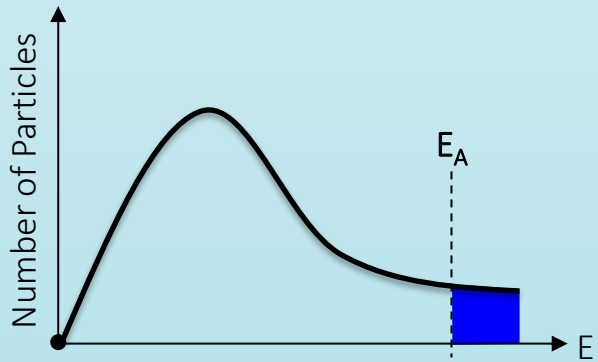
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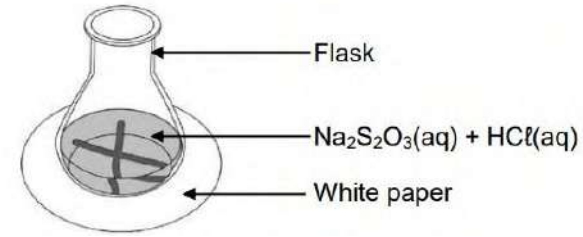


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2	40	10	0,10	26,7
3	30	20	P	33,3



50 cm³

5.3 Calculate the value of P in the table.



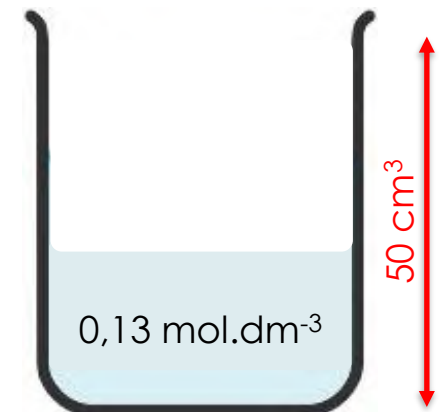
$$n = cV = (0,13)(0,05)$$

$$n = 0,0065 \text{ mol}$$



$$n = cV = (0,13)(0,04)$$

$$n = 0,0052 \text{ mol}$$

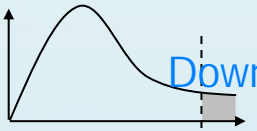


$$n = cV = (0,13)(0,03)$$

$$n = 0,0039 \text{ mol}$$

QUESTION 5

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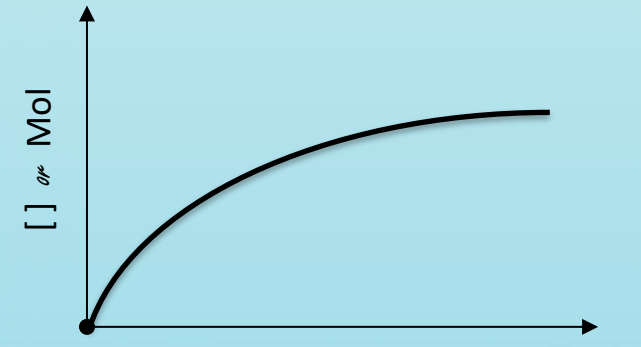
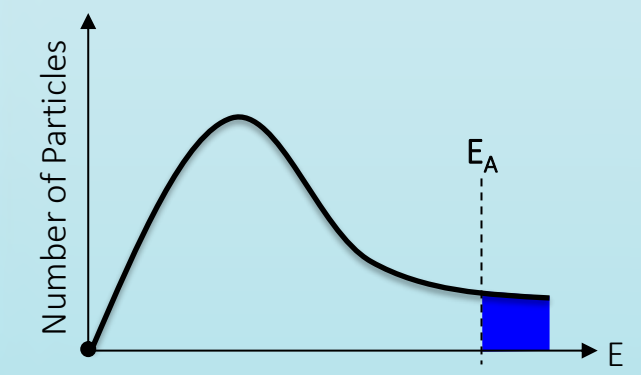
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 State of division

$$RT = \pm \frac{\Delta[\quad]}{\Delta\text{Time}}$$

COLLISION THEORY

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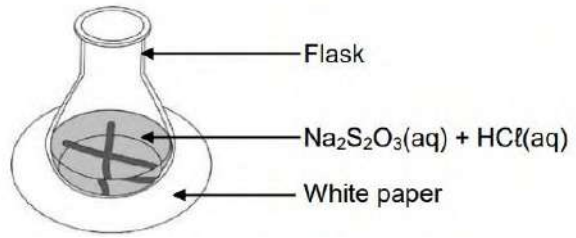


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The concentration of $\text{HCl}(\text{aq})$ used is $1 \text{ mol}\cdot\text{dm}^{-3}$. The same volume of $\text{HCl}(\text{aq})$ is used in each run. The time taken for the cross on the paper under the flask to become invisible is measured.

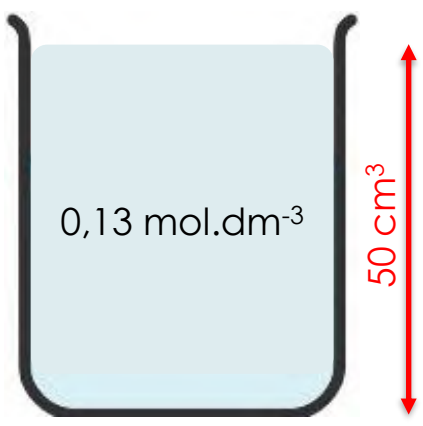
The table below summarises the reaction conditions and results of the experiment.

RUN	VOLUME $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ (cm^3)	VOLUME $\text{H}_2\text{O}(\text{l})$ ADDED (cm^3)	CONCENTRATION $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ ($\text{mol}\cdot\text{dm}^{-3}$)	TIME (s)
1	50	0	0,13	20,4
2	40	10	0,10	26,7
3	30	20	P	33,3



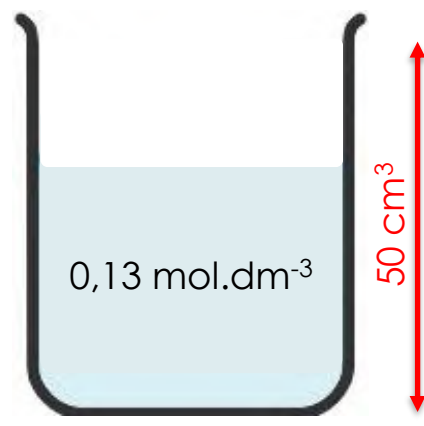
50 cm³

5.3 Calculate the value of P in the table.

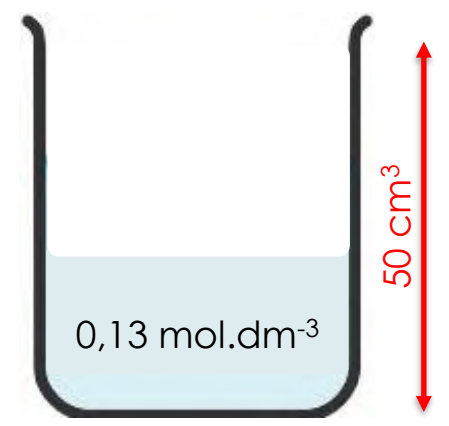


$$n = cV = (0,13)(0,05)$$

$$n = 0,0065 \text{ mol}$$



$$c = \frac{0,0052}{0,05} = 0,104 \text{ mol}\cdot\text{dm}^{-3}$$

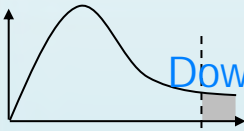


$$c = \frac{0,0039}{0,05} = 0,078 \text{ mol}\cdot\text{dm}^{-3}$$

QUESTION 5

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REACTION RATE



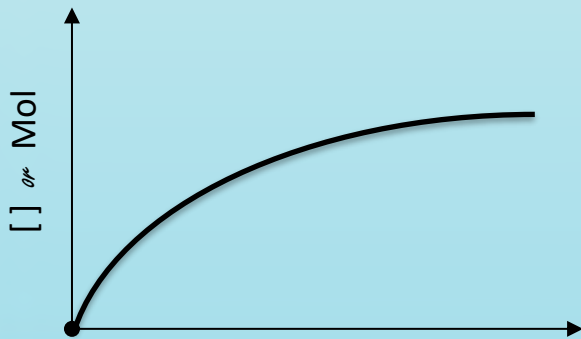
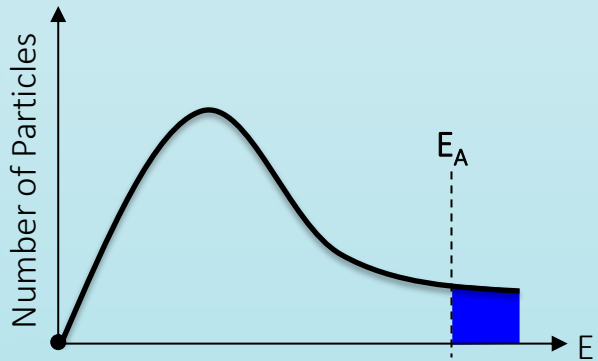
Make sure you know the correct wording of all definitions!

- FACTORS:**
 Temperature / Concentration
 Catalyst / Pressure
 Nature of the reactants
 State of division

$$RT = \pm \frac{\Delta[]}{\Delta \text{Time}}$$

COLLISION THEORY

1. Identify the factor that differs.
2. Difference between particles.
3. Number of effective collisions per unit time.
4. Effect on reaction rate.



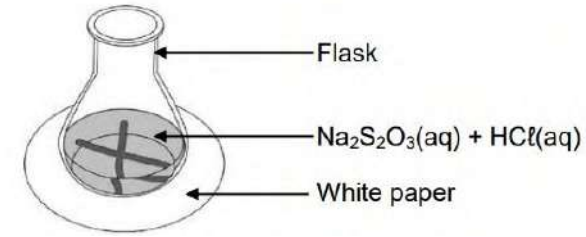
The reaction between EXCESS dilute hydrochloric acid and sodium thiosulphate is used to investigate factors that influence reaction rate. $\text{Na}_2\text{S}_2\text{O}_3(\text{aq}) + 2\text{HCl}(\text{aq}) \rightarrow 2\text{NaCl}(\text{aq}) + \text{S}(\text{s}) + \text{H}_2\text{O}(\text{l}) + \text{SO}_2(\text{g})$

The concentration of $\text{HCl}(\text{aq})$ used is $1 \text{ mol}\cdot\text{dm}^{-3}$. The same volume of $\text{HCl}(\text{aq})$ is used in each run. The time taken for the cross on the paper under the flask to become invisible is measured.

The table below summarises the reaction conditions and results of the experiment.

RUN	VOLUME $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ (cm^3)	VOLUME $\text{H}_2\text{O}(\text{l})$ ADDED (cm^3)	CONCENTRATION $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ ($\text{mol}\cdot\text{dm}^{-3}$)	TIME (s)
1	50	0	0,13	20,4
2	40	10	0,10	26,7
3	30	20	P	33,3

50 cm³



5.4 When 0,21 g of sulphur has formed in Run 1, the cross becomes invisible. Calculate the average reaction rate with respect to sodium thiosulphate, $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$, in $\text{g}\cdot\text{s}^{-1}$.

	$\text{Na}_2\text{S}_2\text{O}_3$	2HCl	2NaCl	S	H_2O	SO_2
n_i						
Δn	0,0065625			0,0065625		
n_f						
	$0,0065625 = \frac{\Delta m}{158}$			$\Delta n = \frac{\Delta m}{M}$		
	$\Delta m = 1,036875$			$\Delta n = \frac{0,21}{32}$		
				$\Delta n = 0,0065625$		

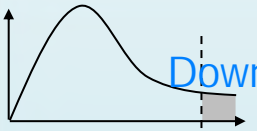
$$\text{Rate} = \frac{\Delta m}{\Delta t}$$

$$\text{Rate} = \frac{1,036875}{20,4}$$

$$\text{Rate} = 0,051 \text{ g}\cdot\text{s}^{-1}$$

QUESTION 5

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REACTION RATE

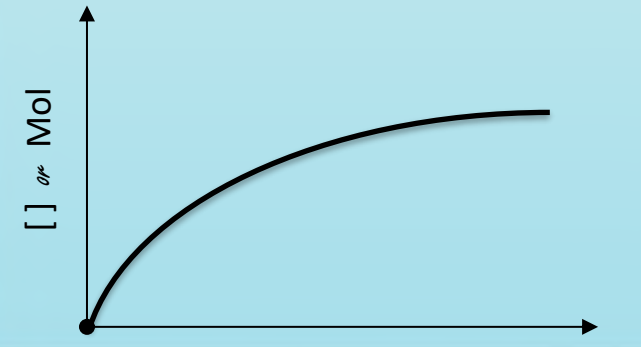
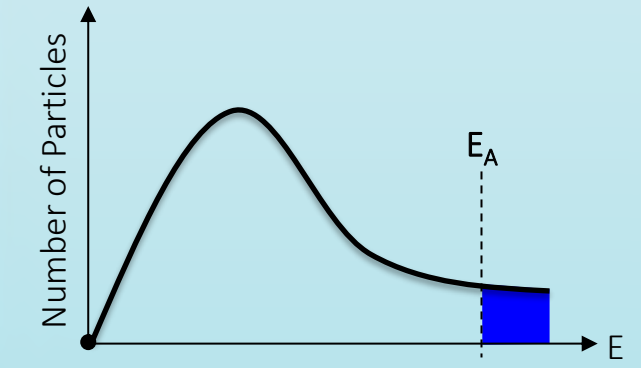
Make sure you know the correct wording of all definitions!

- FACTORS:**
 Temperature / Concentration
 Catalyst / Pressure
 Nature of the reactants
 State of division

$$RT = \pm \frac{\Delta[]}{\Delta \text{Time}}$$

COLLISION THEORY

1. Identify the factor that differs.
2. Difference between particles.
3. Number of effective collisions per unit time.
4. Effect on reaction rate.



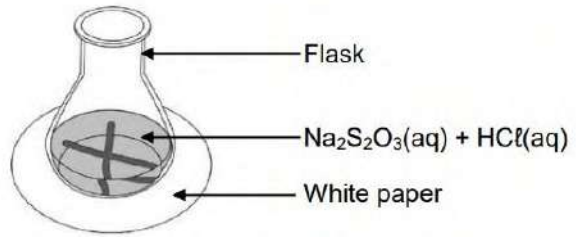
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The concentration of $\text{HCl}(\text{aq})$ used is $1 \text{ mol}\cdot\text{dm}^{-3}$. The same volume of $\text{HCl}(\text{aq})$ is used in each run. The time taken for the cross on the paper under the flask to become invisible is measured.

The table below summarises the reaction conditions and results of the experiment.

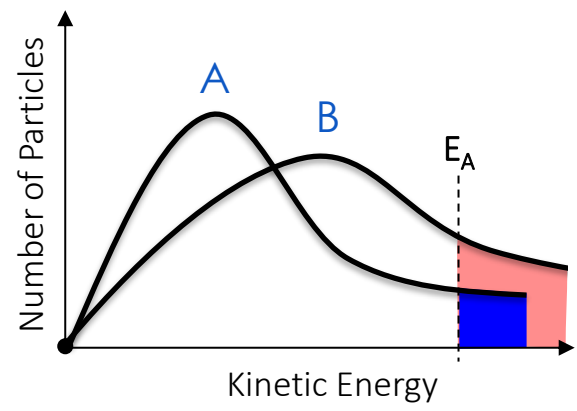
RUN	VOLUME $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ (cm^3)	VOLUME $\text{H}_2\text{O}(\text{l})$ ADDED (cm^3)	CONCENTRATION $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ ($\text{mol}\cdot\text{dm}^{-3}$)	TIME (s)
1	50	0	0,13	20,4
2	40	10	0,10	26,7
3	30	20	P	33,3

50 cm³



Another investigation is performed at different temperatures.

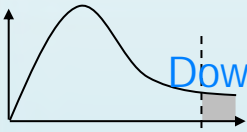
5.5 Sketch the Maxwell-Boltzmann distribution curve for the reaction at 20 °C. Label this curve as A. On the same set of axis, draw the curve that will be obtained at 35 °C and label it as B.



QUESTION 5

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REACTION RATE



Make sure you know the correct wording of all definitions!

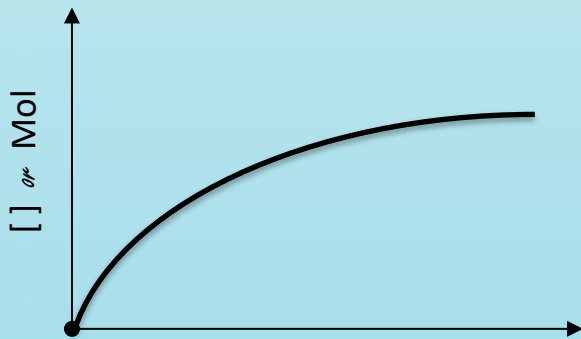
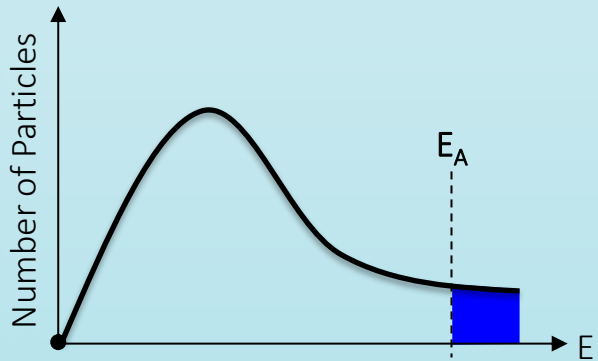
FACTORS:

Temperature / Concentration
Catalyst / Pressure
Nature of the reactants
State of division

$$RT = \pm \frac{\Delta[\quad]}{\Delta\text{Time}}$$

COLLISION THEORY

1. Identify the factor that differs.
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3. Number of effective collisions per unit time.
4. Effect on reaction rate.



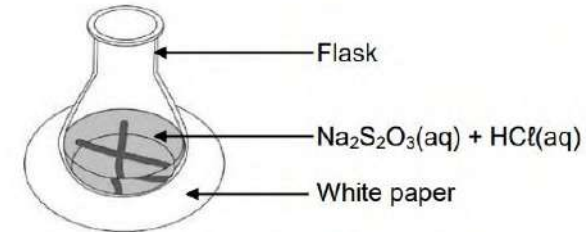
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The concentration of $\text{HCl}(\text{aq})$ used is $1 \text{ mol}\cdot\text{dm}^{-3}$. The same volume of $\text{HCl}(\text{aq})$ is used in each run. The time taken for the cross on the paper under the flask to become invisible is measured.

The table below summarises the reaction conditions and results of the experiment.

RUN	VOLUME $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ (cm^3)	VOLUME $\text{H}_2\text{O}(\text{l})$ ADDED (cm^3)	CONCENTRATION $\text{Na}_2\text{S}_2\text{O}_3(\text{aq})$ ($\text{mol}\cdot\text{dm}^{-3}$)	TIME (s)
1	50	0	0,13	20,4
2	40	10	0,10	26,7
3	30	20	P	33,3

50 cm^3



5.6 Explain the effect of temperature on reaction rate in terms of the collision theory.

- At a higher temperature, the particles have more kinetic energy.
- More molecules have kinetic energy equal to or greater than the activation energy.
- More effective collisions per unit time.
- Reaction rate increases.





Complete the following Google form if you would like the video link and other resources to be sent to you:

<https://bit.ly/4dIYf3i>

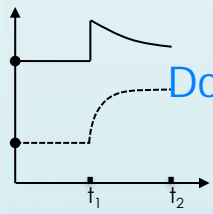


5 minutes
OXYGEN BREAK



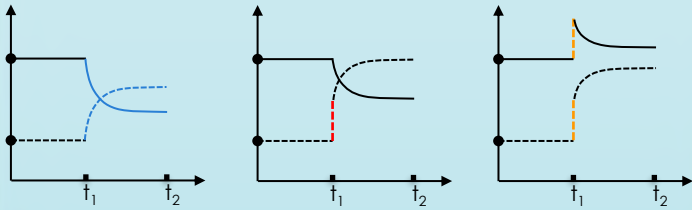
QUESTION 6

CHEMICAL EQUILIBRIUM
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Make sure you know the correct wording of all definitions!

FACTORS		
ENDOTHERMIC	↑ T ↓	EXOTHERMIC
AWAY	↑ [] ↓	TOWARDS
LEAST MOL	↑ P ↓	MOST MOL



- ID Disturbance.
- Le Chatelier
- Reaction favoured.
- Result.

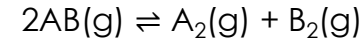
Is the reaction **Exo**- or **Endo**thermic?

Temperature ↑ or ↓
Le Chatelier
Reaction favoured
K_c value ↑ or ↓

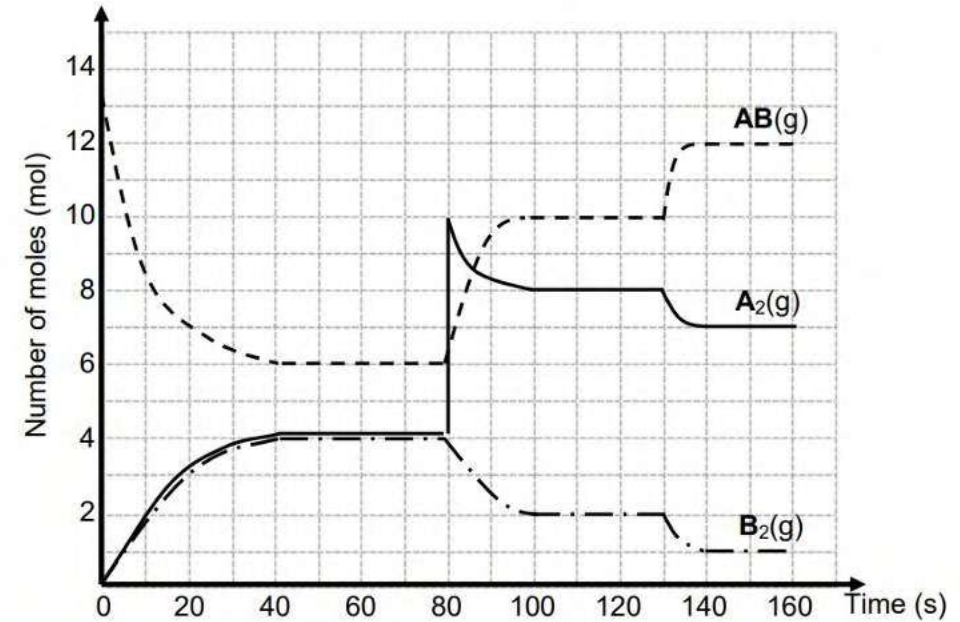
$$K_c = \frac{[A]^a[B]^b}{[C]^c[D]^d}$$

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n _i			0	0
▲n				
n _f				
c = $\frac{n}{V}$				

Consider the following hypothetical reaction reaching equilibrium in a 4 dm³ closed container at 150 °C.



The graph below shows the changes in the amounts of reactants and products over time.



6.1 Write down the meaning of the term reversible reaction.

6.2 State Le Chatelier's principle.

6.3 A change was made to the equilibrium mixture at t = 80 s.

6.3.1 Write down the change made at t = 80 s.

6.3.2 Use Le Chatelier's principle to explain how the system reacts to this change.

6.4 Calculate the equilibrium constant, K_c, at t = 120 s.

6.5 At t = 130 s the temperature of the system is decreased to 100 °C.

6.5.1 Draw a potential energy diagram for this reaction.

6.5.2 Will the equilibrium constant, K_c, at 100 °C be GREATER THAN, LESS THAN or EQUAL TO the K_c at 150 °C? Explain the answer.

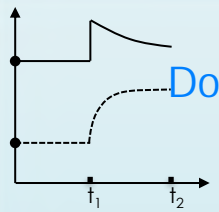
6.6 The initial reaction now takes place in the presence of a catalyst at 150 °C. Describe the changes that will be observed on the graph between t = 0 s and t = 60 s.



QUESTION 6

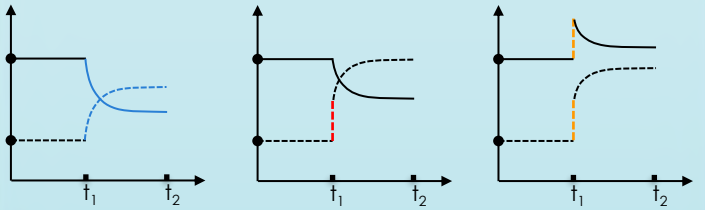
CHEMICAL EQUILIBRIUM

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Make sure you know the correct wording of all definitions!

	FACTORS		
ENDOTHERMIC	↑ T ↓	EXOTHERMIC	
AWAY	↑ [] ↓	TOWARDS	
LEAST MOL	↑ P ↓	MOST MOL	



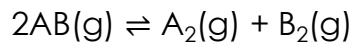
- ID Disturbance.
- Le Chatelier
- Reaction favoured.
- Result.

Is the reaction **Exo-** or **Endo**thermic?
 Temperature ↑ or ↓
 Le Chatelier
 Reaction favoured
 K_c value ↑ or ↓

$$K_c = \frac{[A]^a[B]^b}{[C]^c[D]^d}$$

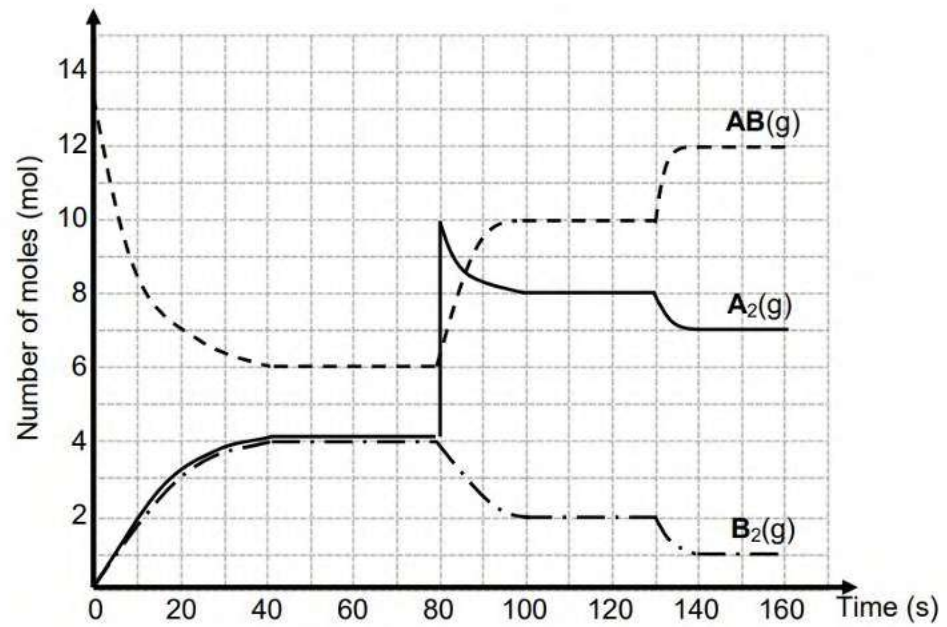
	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n_i			0	0
Δn				
n_f				
$c = \frac{n}{v}$				

Consider the following hypothetical reaction reaching equilibrium in a 4 dm³ closed container at 150 °C.



The graph below shows the changes in the amounts of reactants and products over time.

6.1 Write down the meaning of the term reversible reaction.



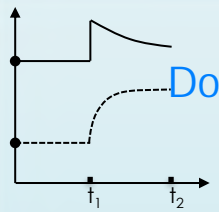
A reaction in which products can be converted back to reactants, and (vice versa).



QUESTION 6


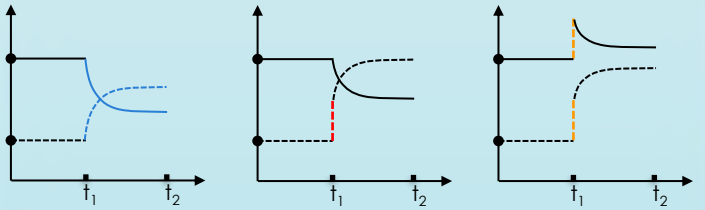
CHEMICAL EQUILIBRIUM

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Make sure you know the correct wording of all definitions!

	FACTORS		
ENDOTHERMIC	↑ T ↓	EXOTHERMIC	
AWAY	↑ [] ↓	TOWARDS	
LEAST MOL	↑ P ↓	MOST MOL	

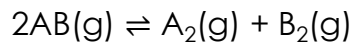
- ID Disturbance.
- Le Chatelier
- Reaction favoured.
- Result.

Is the reaction **Exo**- or **Endo**thermic?
 Temperature ↑ or ↓
 Le Chatelier
 Reaction favoured
 K_c value ↑ or ↓

$$K_c = \frac{[A]^a[B]^b}{[C]^c[D]^d}$$

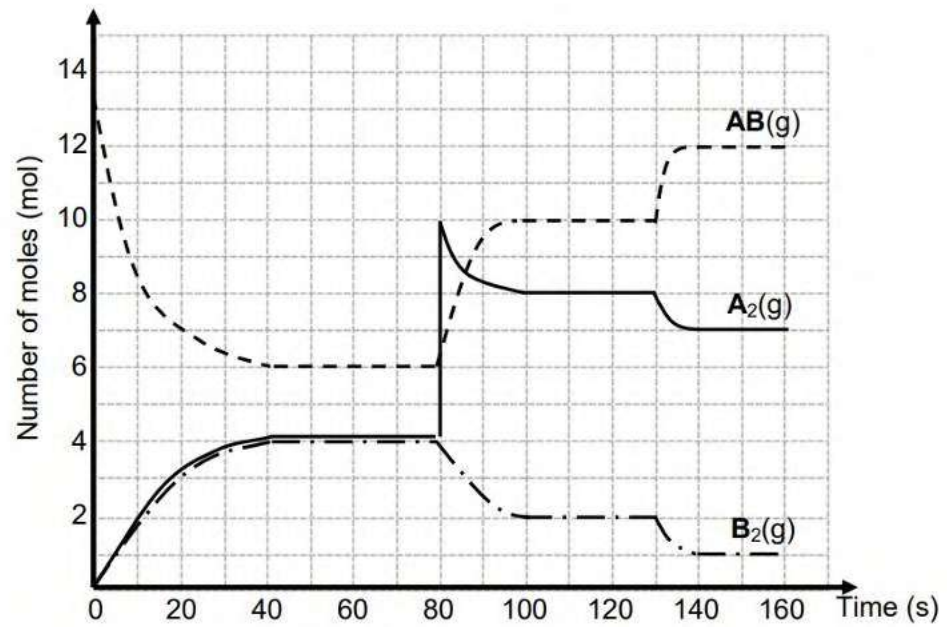
	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n_i			0	0
Δn				
n_f				
$c = \frac{n}{v}$				

Consider the following hypothetical reaction reaching equilibrium in a 4 dm³ closed container at 150 °C.



The graph below shows the changes in the amounts of reactants and products over time.

6.2 State Le Chatelier's principle.



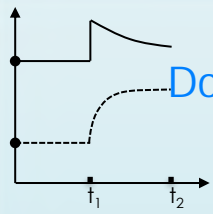
LE CHATELIER'S PRINCIPLE:

When the equilibrium in a closed system is disturbed, the system will re-instate a new equilibrium by favouring the reaction that will oppose the disturbance.



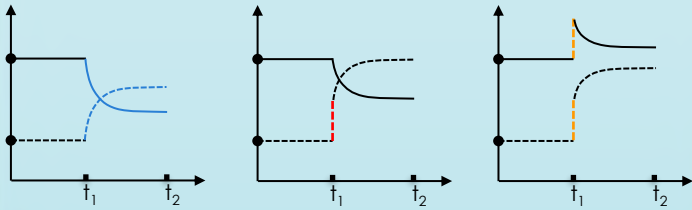
QUESTION 6

CHEMICAL EQUILIBRIUM
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FACTORS		
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AWAY	↑ [] ↓	TOWARDS
LEAST MOL	↑ P ↓	MOST MOL



- ID Disturbance.
- Le Chatelier
- Reaction favoured.
- Result.

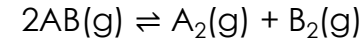
Is the reaction **Exo-** or **Endo**thermic?

Temperature ↑ or ↓
Le Chatelier
Reaction favoured
K_c value ↑ or ↓

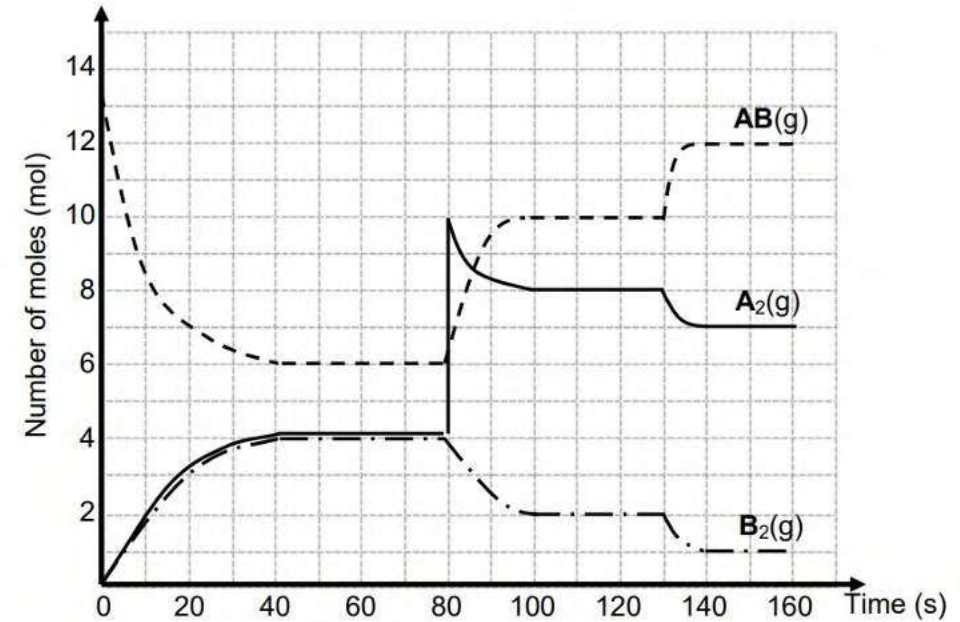
$$K_c = \frac{[A]^a[B]^b}{[C]^c[D]^d}$$

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n _i			0	0
▲n				
n _f				
c = $\frac{n}{V}$				

Consider the following hypothetical reaction reaching equilibrium in a 4 dm³ closed container at 150 °C.



The graph below shows the changes in the amounts of reactants and products over time.



6.3 A change was made to the equilibrium mixture at t = 80 s.

6.3.1 Write down the change made at t = 80 s.

6.3.2 Use Le Chatelier's principle to explain how the system reacts to this change.

6.3.1 The concentration A₂(g) was increased

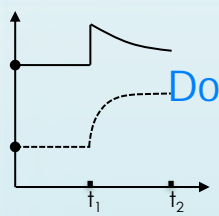
6.3.2 An increase in concentration of A₂ favors the reaction that decreases the concentration (away from the increase).

The reverse reaction is favored.
The amount of moles of the reactants increases.



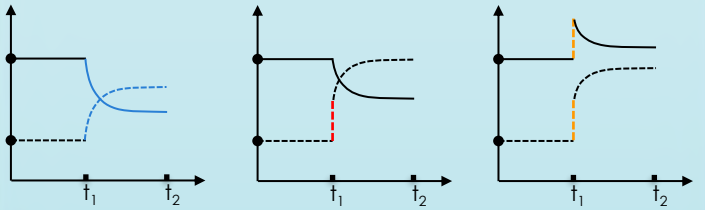
QUESTION 6

CHEMICAL EQUILIBRIUM
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	FACTORS		
ENDOTHERMIC	↑ T ↓	EXOTHERMIC	
AWAY	↑ [] ↓	TOWARDS	
LEAST MOL	↑ P ↓	MOST MOL	



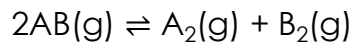
- ID Disturbance.
- Le Chatelier
- Reaction favoured.
- Result.

Is the reaction **Exo**- or **Endo**thermic?
Temperature ↑ or ↓
Le Chatelier
Reaction favoured
K_c value ↑ or ↓

$$K_c = \frac{[A]^a[B]^b}{[C]^c[D]^d}$$

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n _i			0	0
Δn				
n _f				
c = $\frac{n}{V}$				

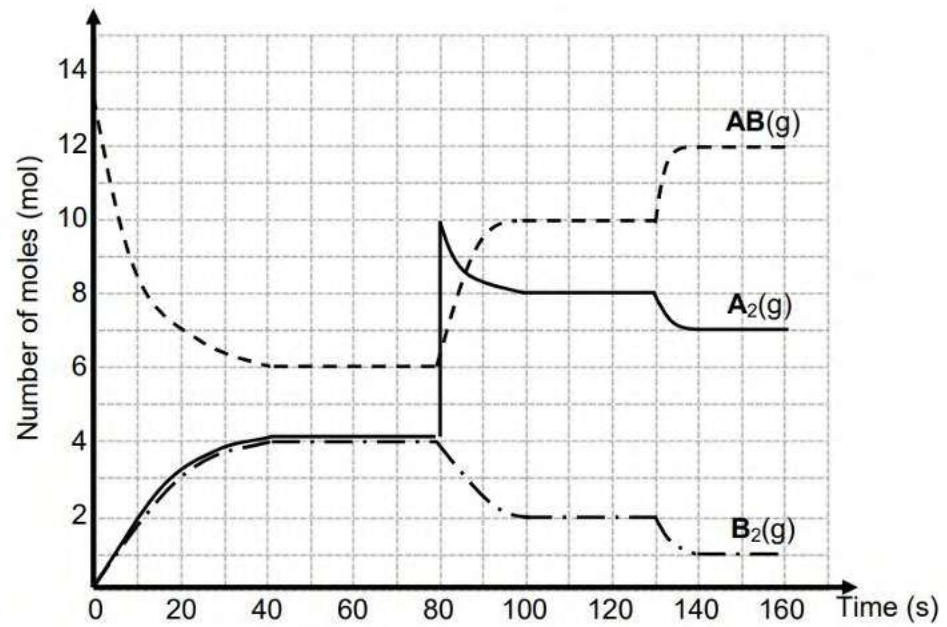
Consider the following hypothetical reaction reaching equilibrium in a 4 dm³ closed container at 150 °C.



The graph below shows the changes in the amounts of reactants and products over time.

6.4 Calculate the equilibrium constant, K_c, at t = 120 s.

	2AB	A ₂	B ₂
n _i	14	0	0
Δn	-8	+4	+4
n _f	6	4	4
c = $\frac{n}{V}$	c = $\frac{6}{4}$	c = $\frac{4}{4}$	c = $\frac{4}{4}$



$$K_c = \frac{[A_2][B_2]}{[AB]^2}$$

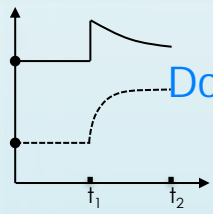
$$K_c = \frac{(1)(1)}{(1,5)^2}$$

$$K_c = 0,44$$



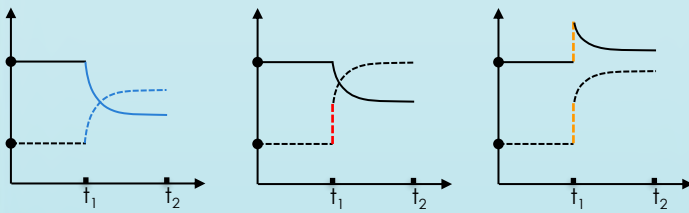
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- ID Disturbance.
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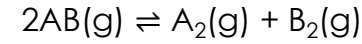
Is the reaction **Exo-** or **Endo**thermic?

Temperature ↑ or ↓
Le Chatelier
Reaction favoured
 K_c value ↑ or ↓

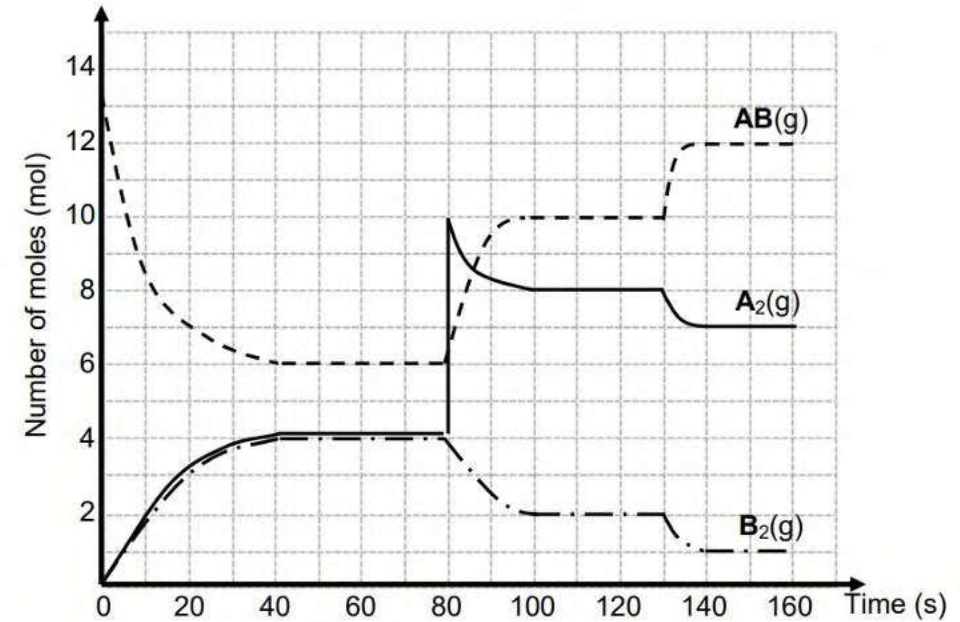
$$K_c = \frac{[A]^a[B]^b}{[C]^c[D]^d}$$

	H_2SO_4	$2NaOH$	Na_2SO_4	$2H_2O$
n_i			0	0
Δn				
n_f				
$c = \frac{n}{v}$				

Consider the following hypothetical reaction reaching equilibrium in a 4 dm^3 closed container at $150 \text{ }^\circ\text{C}$.



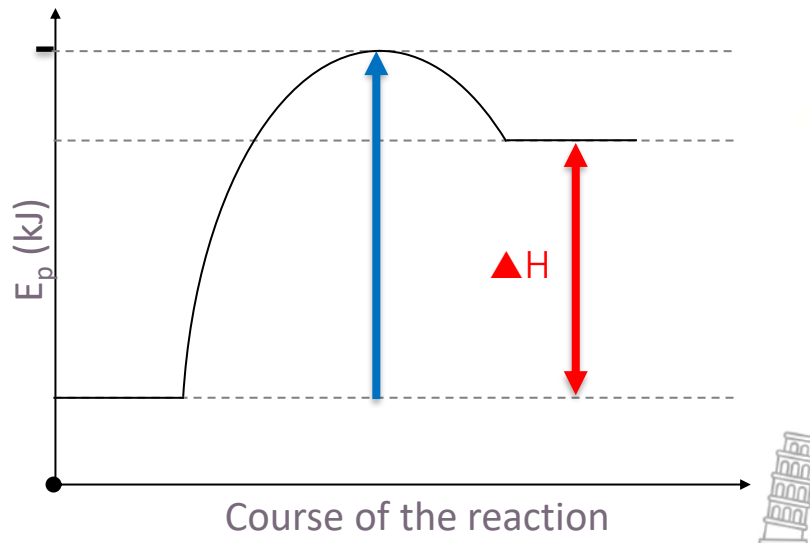
The graph below shows the changes in the amounts of reactants and products over time.



6.5 At $t = 130 \text{ s}$ the temperature of the system is decreased to $100 \text{ }^\circ\text{C}$.

6.5.1 Draw a potential energy diagram for this reaction.

6.5.2 Will the equilibrium constant, K_c , at $100 \text{ }^\circ\text{C}$ be GREATER THAN, LESS THAN or EQUAL TO the K_c at $150 \text{ }^\circ\text{C}$? Explain the answer.



Temperature decreases. ↑

$[AB]$ increases. ↓

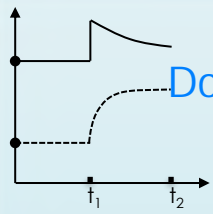
∴ Reverse reaction is exothermic.

∴ Forward reaction is endothermic.

K_c decreases with a decrease in temperature.

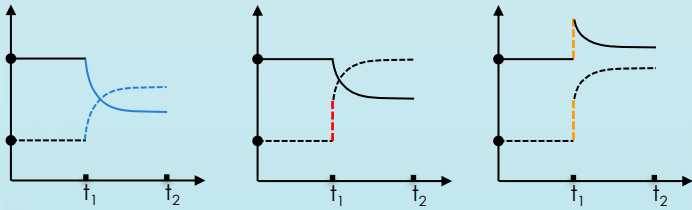
QUESTION 6

CHEMICAL EQUILIBRIUM
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Make sure you know the correct wording of all definitions!

FACTORS		
ENDOTHERMIC	↑ T ↓	EXOTHERMIC
AWAY	↑ [] ↓	TOWARDS
LEAST MOL	↑ P ↓	MOST MOL



- ID Disturbance.
- Le Chatelier
- Reaction favoured.
- Result.

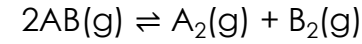
Is the reaction **Exo-** or **Endo**thermic?

Temperature ↑ or ↓
Le Chatelier
Reaction favoured
 K_c value ↑ or ↓

$$K_c = \frac{[A]^a[B]^b}{[C]^c[D]^d}$$

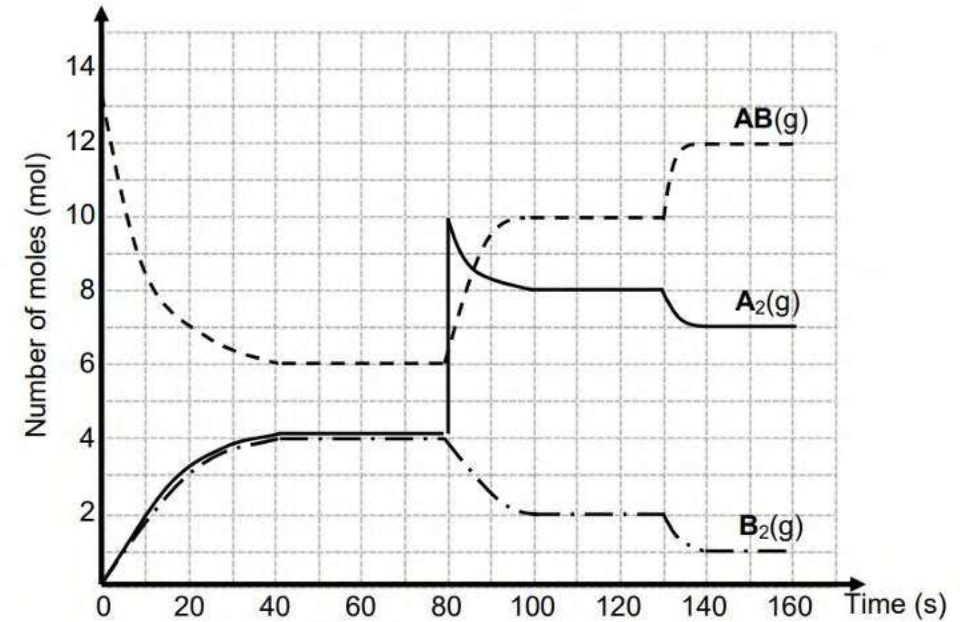
	H_2SO_4	$2NaOH$	Na_2SO_4	$2H_2O$
n_i			0	0
Δn				
n_f				
$c = \frac{n}{v}$				

Consider the following hypothetical reaction reaching equilibrium in a 4 dm^3 closed container at $150 \text{ }^\circ\text{C}$.



The graph below shows the changes in the amounts of reactants and products over time.

6.6 The initial reaction now takes place in the presence of a catalyst at $150 \text{ }^\circ\text{C}$. Describe the changes that will be observed on the graph between $t = 0 \text{ s}$ and $t = 60 \text{ s}$.



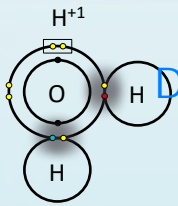
CATALYST increases the forward AND the reverse reaction rates.

All three curves will start and end at the same place, but the gradients of all three curves are steeper and reach equilibrium faster.



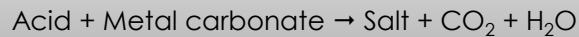
QUESTION 7

ACID-BASE REACTIONS



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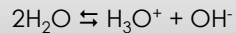
INDICATORS

Methyl Orange (Red/Yellow)
Bromothymol Blue (Yellow/Blue)
Phenolphthalein (Colorless/Pink)

TITRATIONS

$$\frac{c_a v_a}{c_b v_b} = \frac{n_a}{n_b}$$

pH-CALCULATIONS



$$K_w = [\text{H}_3\text{O}^+][\text{OH}^-] = 1 \times 10^{-14}$$

$$\text{pH} = -\log[\text{H}_3\text{O}^+]$$

Hydrolysis of
Salts

QUANTITATIVE CHEMISTRY

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n : n	1	2	1	2
n _i				
▲n				
n _f				

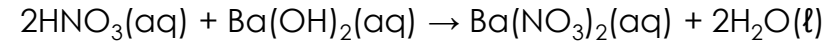
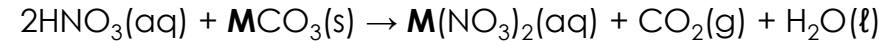
To identify metal **M** in an unknown metal carbonate, **MCO**₃, the following procedure is carried out:

Step 1: 0,198 g of IMPURE **MCO**₃ is reacted with 25 cm³ of 0,4 mol.dm⁻³ nitric acid, HNO₃(aq).

Step 2: The EXCESS HNO₃(aq) is then neutralised with 20 cm³ of 0,15 mol.dm⁻³ barium hydroxide, Ba(OH)₂(aq).

Assume that the volumes are additive.

The following reactions take place:



7.1 Define the term strong base.

7.2 Calculate the:

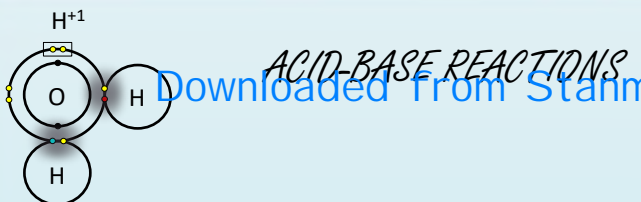
7.2.1 Number of moles of Ba(OH)₂(aq) that reacted with the excess HNO₃(aq)

7.2.2 pH of the solution after Step 1

7.3 The percentage purity of the **MCO**₃(s) in the sample is 85%. Identify metal **M**.

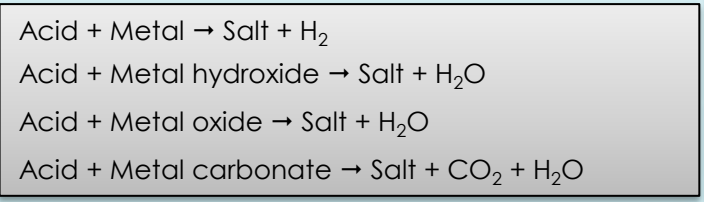


QUESTION 7



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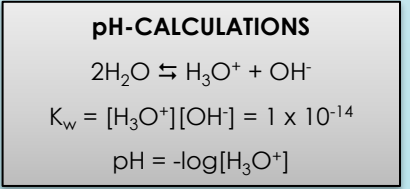
Make sure you know the correct wording of all definitions!



INDICATORS
 Methyl Orange (Red/Yellow)
 Bromothymol Blue (Yellow/Blue)
 Phenolphthalein (Colorless/Pink)

TITRATIONS

$$\frac{c_a V_a}{c_b V_b} = \frac{n_a}{n_b}$$



Hydrolysis of
Salts

QUANTITATIVE CHEMISTRY

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n : n	1	2	1	2
n _i				
▲n				
n _f				

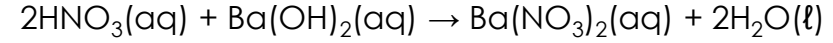
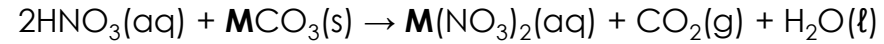
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Assume that the volumes are additive.

The following reactions take place:



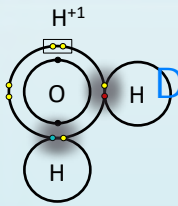
7.1 Define the term strong base.

STRONG BASE: Base that completely dissociates in water to form a high concentration of hydroxyde ions.



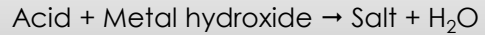
QUESTION 7

ACID-BASE REACTIONS



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Make sure you know the correct wording of all definitions!



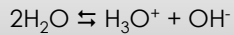
INDICATORS

Methyl Orange (Red/Yellow)
Bromothymol Blue (Yellow/Blue)
Phenolphthalein (Colorless/Pink)

TITRATIONS

$$\frac{c_a v_a}{c_b v_b} = \frac{n_a}{n_b}$$

pH-CALCULATIONS



$$K_w = [\text{H}_3\text{O}^+][\text{OH}^-] = 1 \times 10^{-14}$$

$$\text{pH} = -\log[\text{H}_3\text{O}^+]$$

Hydrolysis of
Salts

QUANTITATIVE CHEMISTRY

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n : n	1	2	1	2
n _i				
▲n				
n _f				

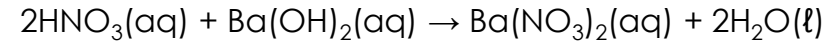
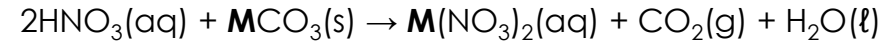
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Step 2: The EXCESS HNO₃(aq) is then neutralised with 20 cm³ of 0,15 mol.dm⁻³ barium hydroxide, Ba(OH)₂(aq).

Assume that the volumes are additive.

The following reactions take place:



7.2 Calculate the:

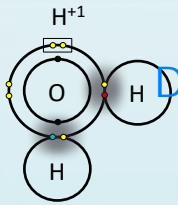
7.2.1 Number of moles of Ba(OH)₂(aq) that reacted with the excess HNO₃(aq)

7.2.2 pH of the solution after Step 1



QUESTION 7

ACID-BASE REACTIONS



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Make sure you know the correct wording of all definitions!

- Acid + Metal → Salt + H₂
- Acid + Metal hydroxide → Salt + H₂O
- Acid + Metal oxide → Salt + H₂O
- Acid + Metal carbonate → Salt + CO₂ + H₂O

INDICATORS

- Methyl Orange (Red/Yellow)
- Bromothymol Blue (Yellow/Blue)
- Phenolphthalein (Colorless/Pink)

TITRATIONS

$$\frac{c_a V_a}{c_b V_b} = \frac{n_a}{n_b}$$

pH-CALCULATIONS

$$2H_2O \rightleftharpoons H_3O^+ + OH^-$$

$$K_w = [H_3O^+][OH^-] = 1 \times 10^{-14}$$

$$pH = -\log[H_3O^+]$$

Hydrolysis of Salts

QUANTITATIVE CHEMISTRY

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n : n	1	2	1	2
n _i				
▲n				
n _f				

To identify metal **M** in an unknown metal carbonate, **MCO₃**, the following procedure is carried out:

Step 1: 0,198 g of IMPURE **MCO₃** is reacted with 25 cm³ of 0,4 mol.dm⁻³ nitric acid, HNO₃(aq).

Step 2: The EXCESS HNO₃(aq) is then neutralised with 20 cm³ of 0,15 mol.dm⁻³ barium hydroxide, Ba(OH)₂(aq).

Assume that the volumes are additive.

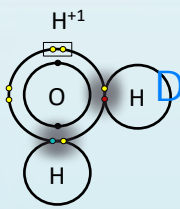
The following reactions take place:



	2HNO ₃	MCO ₃	M(NO ₃) ₂	CO ₂	H ₂ O
ni			0	0	0
Δn					
nf	0,006				
	c = 0,4 mol.dm ⁻³ V = 0,025 dm ³	m = 0,198 g			

	2HNO ₃	Ba(OH) ₂	Ba(NO ₃) ₂	H ₂ O
ni	0,006	0,003	0	0
Δn	0,006	0,003		
nf	0	0		
		c = 0,15 mol.dm ⁻³ V = 0,02 dm ³ n = cV = (0,15)(0,02)		





ACID-BASE REACTIONS

Make sure you know the correct wording of all definitions!

- Acid + Metal → Salt + H₂
- Acid + Metal hydroxide → Salt + H₂O
- Acid + Metal oxide → Salt + H₂O
- Acid + Metal carbonate → Salt + CO₂ + H₂O

- ### INDICATORS
- Methyl Orange (Red/Yellow)
 - Bromothymol Blue (Yellow/Blue)
 - Phenolphthalein (Colorless/Pink)

TITRATIONS

$$\frac{c_a V_a}{c_b V_b} = \frac{n_a}{n_b}$$

pH-CALCULATIONS

$$2\text{H}_2\text{O} \rightleftharpoons \text{H}_3\text{O}^+ + \text{OH}^-$$

$$K_w = [\text{H}_3\text{O}^+][\text{OH}^-] = 1 \times 10^{-14}$$

$$\text{pH} = -\log[\text{H}_3\text{O}^+]$$

Hydrolysis of Salts

QUANTITATIVE CHEMISTRY

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n : n	1	2	1	2
n _i				
Δn				
n _f				

	2HNO ₃	MCO ₃	M(NO ₃) ₂	CO ₂	H ₂ O
n _i			0	0	0
Δn					
n _f	0,006				
	c = 0,4 mol.dm ⁻³ V = 0,025 dm ³	m = 0,198 g			

Assume that the volumes are additive

	2HNO ₃	Ba(OH) ₂	Ba(NO ₃) ₂	H ₂ O
n _i	0,006	0,003	0	0
Δn	0,006	0,003		
n _f	0	0		
		c = 0,15 mol.dm ⁻³ V = 0,02 dm ³ n = cV = (0,15)(0,02)		

7.2.1 Number of moles of Ba(OH)₂(aq) that reacted with the excess HNO₃(aq)

7.2.2 pH of the solution after Step 1

7.2.1 ① $n = cV = (0,15)(0,02) = 0,003 \text{ mol}$

② $\Delta n_{\text{Ba(OH)}_2} = 0,003 \text{ mol}$

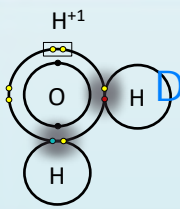
7.2.2 ③ $\Delta n_{\text{HNO}_3} : \Delta n_{\text{Ba(OH)}_2} = 2 : 1$

∴ $\Delta n_{\text{HNO}_3} = 0,006 \text{ mol}$

④ $c_f(\text{HNO}_3) = \frac{n}{V} = \frac{0,006}{0,025} = 0,24 \text{ mol.dm}^{-3}$

$\text{pH} = -\log[\text{H}_3\text{O}^+]$

$\text{pH} = -\log(0,24) = 0,62$



ACID-BASE REACTIONS

Make sure you know the correct wording of all definitions!

- Acid + Metal → Salt + H₂
- Acid + Metal hydroxide → Salt + H₂O
- Acid + Metal oxide → Salt + H₂O
- Acid + Metal carbonate → Salt + CO₂ + H₂O

- ### INDICATORS
- Methyl Orange (Red/Yellow)
 - Bromothymol Blue (Yellow/Blue)
 - Phenolphthalein (Colorless/Pink)

TITRATIONS

$$\frac{c_a V_a}{c_b V_b} = \frac{n_a}{n_b}$$

pH-CALCULATIONS

$$2H_2O \rightleftharpoons H_3O^+ + OH^-$$

$$K_w = [H_3O^+][OH^-] = 1 \times 10^{-14}$$

$$pH = -\log[H_3O^+]$$

Hydrolysis of Salts

QUANTITATIVE CHEMISTRY

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n : n	1	2	1	2
n _i				
Δn				
n _f				

	2HNO ₃	MCO ₃	M(NO ₃) ₂	CO ₂	H ₂ O
n _i	0,01 5	0,00235294117 8	0	0	0
Δn	0,004 6	0,002 7			
nf	0,006 4				
	c = 0,4 mol.dm ⁻³ V = 0,025 dm ³ n = cV = (0,4)(0,025)				

Assume that the volumes are additive

	2HNO ₃	Ba(OH) ₂	Ba(NO ₃) ₂	H ₂ O
n _i	0,006 4	0,003 1	0	0
Δn	0,006 3	0,003 2		
nf	0	0		
		c = 0,15 mol.dm ⁻³ V = 0,02 dm ³ n = cV = (0,15)(0,02)		

7.3 The percentage purity of the MCO₃(s) in the sample is 85%. Identify metal **M**.

5 $n_{(HNO_3)} = cV = (0,4)(0,025) = 0,01 \text{ mol}$

6 $\Delta n_{(HNO_3)} = 0,01 - 0,006 = 0,004 \text{ mol}$

7 $\Delta n_{(HNO_3)} : \Delta n_{(MCO_3)} = 2 : 1$

$\therefore \Delta n_{(MCO_3)} = 0,002 \text{ mol}$

8 85% van $n_i(MCO_3) = 0,002 \text{ mol}$

$\therefore n_i(MCO_3) = 0,00235294117 \text{ mol}$

$\therefore n_i(MCO_3) = \frac{m}{M}$

$0,00235294117 = \frac{0,198}{M+12+3(16)}$

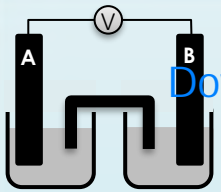
$\therefore M = 24,15 \text{ g.mol}^{-1}$ en M is Mg



QUESTION 8

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GALVANIC CELLS



Make sure you know the correct wording of all definitions!

CELL NOTATION: Anode // Cathode
 Element / Ion // Ion / Element
 $Zn / Zn^{2+} // Cu^{2+} / Cu$

STANDARD CONDITIONS

Temp. 25 °C (298 K)
 Concentration 1 mol·dm⁻³
 Pressure 101,3 kPa.

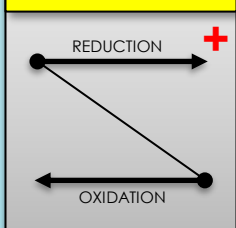
ELECTRODES

Metal
 Gas

$$E^{\theta}_{cell} = E^{\theta}_{cathode} - E^{\theta}_{anode}$$

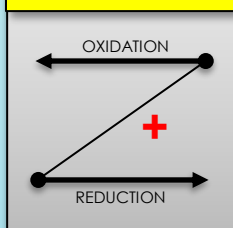
$$E^{\theta}_{cell} = E^{\theta}_{reduction} - E^{\theta}_{oxidation}$$

TABLE 4A

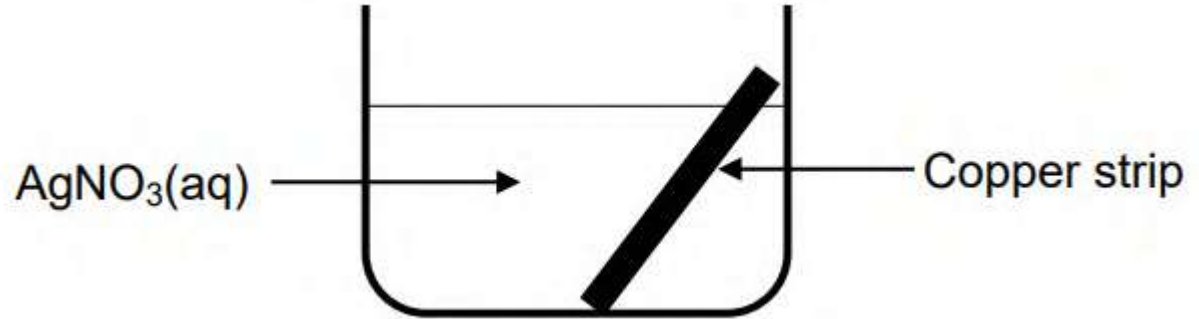


Spontaneous Reaction

TABLE 4B



A cleaned pure copper strip, Cu(s), is placed in a beaker containing a colourless silver nitrate solution, AgNO₃(aq), at 25 °C, as shown below.



After a while, it is observed that the solution in the beaker becomes blue.

8.1 Write down:

8.1.1 ONE other OBSERVABLE change, besides the solution turning blue

8.1.2 The NAME or FORMULA of the oxidising agent

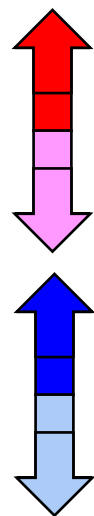
8.2 Explain the answer to QUESTION 8.1.1 by referring to the relative strengths of the oxidising agents or reducing agents.



TABLE 4A: REDOX HALF REACTIONS
E⁰ (V)

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$F_2(g) + 2e^- \rightleftharpoons 2F^-$	+ 2,87
$Co^{3+} + e^- \rightleftharpoons Co^{2+}$	+ 1,81
$H_2O_2 + 2H^+ + 2e^- \rightleftharpoons 2H_2O$	+1,77
$MnO_4^- + 8H^+ + 5e^- \rightleftharpoons Mn^{2+} + 4H_2O$	+ 1,51
$Cl_2(g) + 2e^- \rightleftharpoons 2Cl^-$	+ 1,36
$Cr_2O_7^{2-} + 14H^+ + 6e^- \rightleftharpoons 2Cr^{3+} + 7H_2O$	+ 1,33
$O_2(g) + 4H^+ + 4e^- \rightleftharpoons 2H_2O$	+ 1,23
$MnO_2 + 4H^+ + 2e^- \rightleftharpoons Mn^{2+} + 2H_2O$	+ 1,23
$Pt^{2+} + 2e^- \rightleftharpoons Pt$	+ 1,20
$Br_2(l) + 2e^- \rightleftharpoons 2Br^-$	+ 1,07
$NO_3^- + 4H^+ + 3e^- \rightleftharpoons NO(g) + 2H_2O$	+ 0,96
$Hg^{2+} + 2e^- \rightleftharpoons Hg(l)$	+ 0,85
$Ag^+ + e^- \rightleftharpoons Ag$	+ 0,80
$NO_3^- + 2H^+ + e^- \rightleftharpoons NO_2(g) + H_2O$	+ 0,80
$Fe^{3+} + e^- \rightleftharpoons Fe^{2+}$	+ 0,77
$O_2(g) + 2H^+ + 2e^- \rightleftharpoons H_2O_2$	+ 0,68
$I_2 + 2e^- \rightleftharpoons 2I^-$	+ 0,54
$Cu^+ + e^- \rightleftharpoons Cu$	+ 0,52
$SO_2 + 4H^+ + 4e^- \rightleftharpoons S + 2H_2O$	+ 0,45
$2H_2O + O_2 + 4e^- \rightleftharpoons 4OH^-$	+ 0,40
$Cu^{2+} + 2e^- \rightleftharpoons Cu$	+ 0,34
$SO_4^{2-} + 4H^+ + e^- \rightleftharpoons SO_2(g) + 2H_2O$	+ 0,17


TABLE 4B: REDOX HALF REACTIONS
E⁰ (V)

$Li^+ + e^- \rightleftharpoons Li$	- 3,05
$K^+ + e^- \rightleftharpoons K$	- 2,93
$Cs^+ + e^- \rightleftharpoons Cs$	- 2,92
$Ba^{2+} + 2e^- \rightleftharpoons Ba$	- 2,90
$Sr^{2+} + 2e^- \rightleftharpoons Sr$	- 2,89
$Ca^{2+} + 2e^- \rightleftharpoons Ca$	- 2,87
$Na^+ + e^- \rightleftharpoons Na$	- 2,71
$Mg^{2+} + 2e^- \rightleftharpoons Mg$	- 2,36
$Al^{3+} + 3e^- \rightleftharpoons Al$	- 1,66
$Mn^{2+} + 2e^- \rightleftharpoons Mn$	- 1,18
$Cr^{2+} + 2e^- \rightleftharpoons Cr$	- 0,91
$2H_2O + 2e^- \rightleftharpoons H_2(g) + 2OH^-$	- 0,83
$Zn^{2+} + 2e^- \rightleftharpoons Zn$	- 0,76
$Cr^{3+} + 3e^- \rightleftharpoons Cr$	- 0,74
$Fe^{2+} + 2e^- \rightleftharpoons Fe$	- 0,44
$Cr^{3+} + e^- \rightleftharpoons Cr^{2+}$	- 0,41
$Cd^{2+} + 2e^- \rightleftharpoons Cd$	- 0,40
$Co^{2+} + 2e^- \rightleftharpoons Co$	- 0,28
$Ni^{2+} + 2e^- \rightleftharpoons Ni$	- 0,27
$Sn^{2+} + 2e^- \rightleftharpoons Sn$	- 0,14
$Pb^{2+} + 2e^- \rightleftharpoons Pb$	- 0,13
$Fe^{3+} + 3e^- \rightleftharpoons Fe$	- 0,06

QUESTION 8



GALVANIC CELLS

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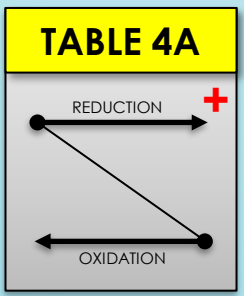
CELL NOTATION: Anode // Cathode
 Element / Ion // Ion / Element
 Zn / Zn²⁺ // Cu²⁺ / Cu

STANDARD CONDITIONS
 Temp. 25 °C (298 K)
 Concentration 1 mol·dm⁻³
 Pressure 101,3 kPa.

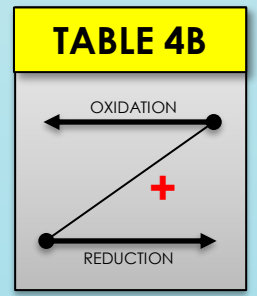
ELECTRODES
 Metal
 Gas

$$E^{\theta}_{cell} = E^{\theta}_{cathode} - E^{\theta}_{anode}$$

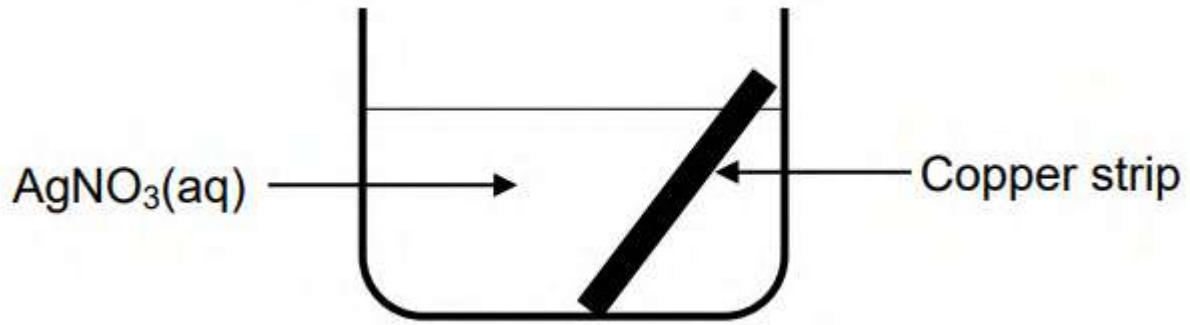
$$E^{\theta}_{cell} = E^{\theta}_{reduction} - E^{\theta}_{oxidation}$$



Spontaneous Reaction



A cleaned pure copper strip, Cu(s), is placed in a beaker containing a colourless silver nitrate solution, AgNO₃(aq), at 25 °C, as shown below.



After a while, it is observed that the solution in the beaker becomes blue.

*Copper strip corrodes / gets eaten away / loses mass.
 Silver settles down.*

8.1 Write down:

8.1.1 ONE other OBSERVABLE change, besides the solution turning blue

8.1.2 The NAME or FORMULA of the oxidising agent

Ag⁺ -ion / Silver ion / AgNO₃

TABLE 4A

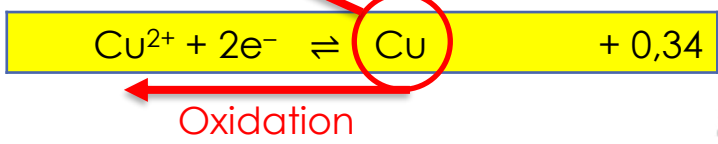
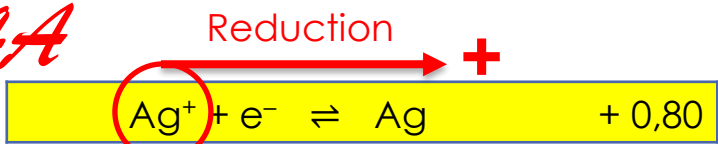
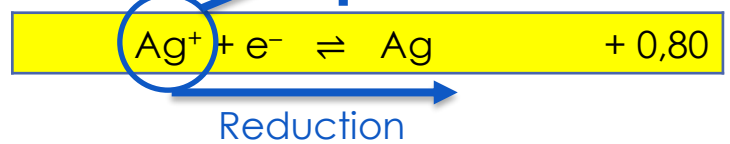
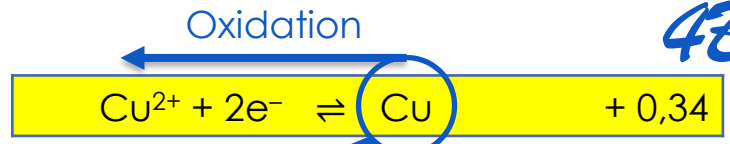


TABLE 4B



QUESTION 8



GALVANIC CELLS

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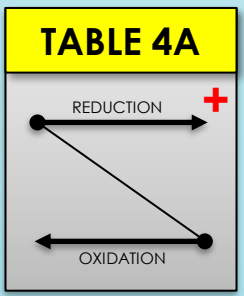
CELL NOTATION: Anode // Cathode
 Element / Ion // Ion / Element
 Zn / Zn²⁺ // Cu²⁺ / Cu

STANDARD CONDITIONS
 Temp. 25 °C (298 K)
 Concentration 1 mol·dm⁻³
 Pressure 101,3 kPa.

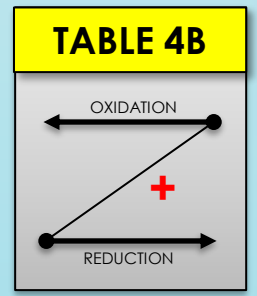
ELECTRODES
 Metal
 Gas

$$E^{\theta}_{\text{cell}} = E^{\theta}_{\text{cathode}} - E^{\theta}_{\text{anode}}$$

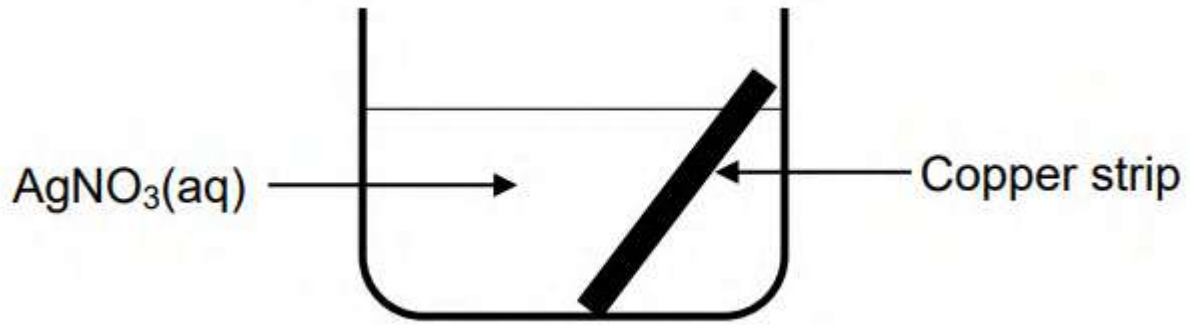
$$E^{\theta}_{\text{cell}} = E^{\theta}_{\text{reduction}} - E^{\theta}_{\text{oxidation}}$$



Spontaneous Reaction



A cleaned pure copper strip, Cu(s), is placed in a beaker containing a colourless silver nitrate solution, AgNO₃(aq), at 25 °C, as shown below.



After a while, it is observed that the solution in the beaker becomes blue.

8.2 Explain the answer to QUESTION 8.1.1 by referring to the relative strengths of the oxidising agents or reducing agents.

TABLE 4A

Reduction

$$\text{Ag}^+ + e^- \rightleftharpoons \text{Ag} \quad +0,80$$

$$\text{Cu}^{2+} + 2e^- \rightleftharpoons \text{Cu} \quad +0,34$$

Oxidation

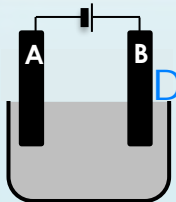
TABLE 4B

Oxidation

$$\text{Cu}^{2+} + 2e^- \rightleftharpoons \text{Cu} \quad +0,34$$

$$\text{Ag}^+ + e^- \rightleftharpoons \text{Ag} \quad +0,80$$

Reduction



ELECTROLYTIC CELLS

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EXPLAIN STRENGTH OF **REDUCING AGENTS** AND **OXIDIZING AGENTS** IN REDOX REACTIONS.

TABLE 4A

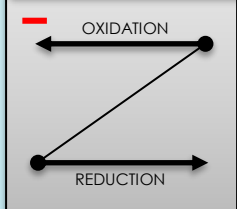
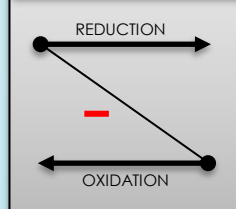


TABLE 4B



Non-Spontaneous Reaction

CARBON

TYPE OF ELECTROLYTE

- Molten Salt
- Dissolved Salt

METAL

TYPE OF ELECTROLYTE

- Electroplating
- Electrefining

ELECTRODES

TABLE 4A

Half-reactions/Halfreactions	E ⁰ (V)
F ₂ (g) + 2e ⁻ = 2F ⁻	+2,87
Co ³⁺ + e ⁻ = Co ²⁺	+1,81
H ₂ O ₂ + 2H ⁺ + 2e ⁻ = 2H ₂ O	+1,77
MnO ₄ ⁻ + 8H ⁺ + 5e ⁻ = Mn ²⁺ + 4H ₂ O	+1,51
Cl ₂ (g) + 2e ⁻ = 2Cl ⁻	+1,36
Cr ₂ O ₇ ²⁻ + 14H ⁺ + 6e ⁻ = 2Cr ³⁺ + 7H ₂ O	+1,33
Reduction	
Br ₂ (l) + 2e ⁻ = 2Br ⁻	+1,07
NO ₃ ⁻ + 4H ⁺ + 3e ⁻ = NO(g) + 2H ₂ O	+0,96
Hg ²⁺ + 2e ⁻ = Hg(l)	+0,85
Ag ⁺ + e ⁻ = Ag	+0,80
NO ₃ ⁻ + 2H ⁺ + e ⁻ = NO ₂ (g) + H ₂ O	+0,80
Fe ³⁺ + e ⁻ = Fe ²⁺	+0,77
O ₂ (g) + 2H ⁺ + 2e ⁻ = H ₂ O ₂	+0,68
I ₂ + 2e ⁻ = 2I ⁻	+0,54
Cu ⁺ + e ⁻ = Cu	+0,52
SO ₂ + 4H ⁺ + 4e ⁻ = S + 2H ₂ O	+0,45
2H ₂ O + O ₂ + 4e ⁻ = 4OH ⁻	+0,40
Cu ²⁺ + 2e ⁻ = Cu	+0,34
SO ₄ ²⁻ + 4H ⁺ + 2e ⁻ = SO ₂ (g) + 2H ₂ O	+0,17
Cu ²⁺ + e ⁻ = Cu ⁺	+0,16
Sn ⁴⁺ + 2e ⁻ = Sn ²⁺	+0,15
Oxidation	
Fe ²⁺ + 2e ⁻ = Fe	-0,08
Pb ²⁺ + 2e ⁻ = Pb	-0,13
Sn ²⁺ + 2e ⁻ = Sn	-0,14
Ni ²⁺ + 2e ⁻ = Ni	-0,27
Co ³⁺ + 2e ⁻ = Co	-0,28
Cd ²⁺ + 2e ⁻ = Cd	-0,40
Cr ³⁺ + e ⁻ = Cr ²⁺	-0,41
Fe ²⁺ + 2e ⁻ = Fe	-0,44
Cr ³⁺ + 3e ⁻ = Cr	-0,74
Zn ²⁺ + 2e ⁻ = Zn	-0,76
2H ₂ O + 2e ⁻ = H ₂ (g) + 2OH ⁻	-0,83
Cr ²⁺ + 2e ⁻ = Cr	-0,91
Mn ²⁺ + 2e ⁻ = Mn	-1,18
Al ³⁺ + 3e ⁻ = Al	-1,66
Mg ²⁺ + 2e ⁻ = Mg	-2,36
Na ⁺ + e ⁻ = Na	-2,71
Ca ²⁺ + 2e ⁻ = Ca	-2,87
Sr ²⁺ + 2e ⁻ = Sr	-2,89
Ba ²⁺ + 2e ⁻ = Ba	-2,90
Cs ⁺ + e ⁻ = Cs	-2,92
K ⁺ + e ⁻ = K	-2,93
Li ⁺ + e ⁻ = Li	-3,05

↑ INCREASING STRENGTH OF OXIDIZING AGENTS

↓ INCREASING STRENGTH OF REDUCING AGENTS

TABLE 4B

Half-reactions/Halfreactions	E ⁰ (V)
Li ⁺ + e ⁻ = Li	-3,05
K ⁺ + e ⁻ = K	-2,93
Cs ⁺ + e ⁻ = Cs	-2,92
Ba ²⁺ + 2e ⁻ = Ba	-2,90
Sr ²⁺ + 2e ⁻ = Sr	-2,89
Ca ²⁺ + 2e ⁻ = Ca	-2,87
Na ⁺ + e ⁻ = Na	-2,71
Mg ²⁺ + 2e ⁻ = Mg	-2,36
Al ³⁺ + 3e ⁻ = Al	-1,66
Mn ²⁺ + 2e ⁻ = Mn	-1,18
Cr ²⁺ + 2e ⁻ = Cr	-0,91
2H ₂ O + 2e ⁻ = H ₂ (g) + 2OH ⁻	-0,83
Zn ²⁺ + 2e ⁻ = Zn	-0,76
Cr ³⁺ + 3e ⁻ = Cr	-0,74
Fe ²⁺ + 2e ⁻ = Fe	-0,44
Cr ³⁺ + e ⁻ = Cr ²⁺	-0,41
Cd ²⁺ + 2e ⁻ = Cd	-0,40
Co ²⁺ + 2e ⁻ = Co	-0,28
Ni ²⁺ + 2e ⁻ = Ni	-0,27
Sn ²⁺ + 2e ⁻ = Sn	-0,14
Pb ²⁺ + 2e ⁻ = Pb	-0,13
Fe ³⁺ + 3e ⁻ = Fe	-0,06
Oxidation	
Sn ²⁺ + 2e ⁻ = Sn	+0,15
Cu ²⁺ + e ⁻ = Cu ⁺	+0,16
SO ₄ ²⁻ + 4H ⁺ + 2e ⁻ = SO ₂ (g) + 2H ₂ O	+0,17
Cu ⁺ + e ⁻ = Cu	+0,52
2H ₂ O + O ₂ + 4e ⁻ = 4OH ⁻	+0,40
SO ₂ + 4H ⁺ + 4e ⁻ = S + 2H ₂ O	+0,45
Cu ⁺ + e ⁻ = Cu	+0,52
I ₂ + 2e ⁻ = 2I ⁻	+0,54
O ₂ (g) + 2H ⁺ + 2e ⁻ = H ₂ O ₂	+0,68
Fe ³⁺ + e ⁻ = Fe ²⁺	+0,77
NO ₃ ⁻ + 2H ⁺ + e ⁻ = NO ₂ (g) + H ₂ O	+0,80
Ag ⁺ + e ⁻ = Ag	+0,80
Hg ²⁺ + 2e ⁻ = Hg(l)	+0,85
Reduction	
Cr ₂ O ₇ ²⁻ + 14H ⁺ + 6e ⁻ = 2Cr ³⁺ + 7H ₂ O	+1,33
Cl ₂ (g) + 2e ⁻ = 2Cl ⁻	+1,36
MnO ₄ ⁻ + 8H ⁺ + 5e ⁻ = Mn ²⁺ + 4H ₂ O	+1,51
H ₂ O ₂ + 2H ⁺ + 2e ⁻ = 2H ₂ O	+1,77
Co ³⁺ + e ⁻ = Co ²⁺	+1,81
F ₂ (g) + 2e ⁻ = 2F ⁻	+2,87

↑ INCREASING STRENGTH OF REDUCING AGENTS

QUESTION 8



GALVANIC CELLS

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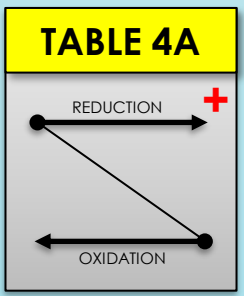
CELL NOTATION: Anode // Cathode
 Element / Ion // Ion / Element
 Zn / Zn²⁺ // Cu²⁺ / Cu

STANDARD CONDITIONS
 Temp. 25 °C (298 K)
 Concentration 1 mol·dm⁻³
 Pressure 101,3 kPa.

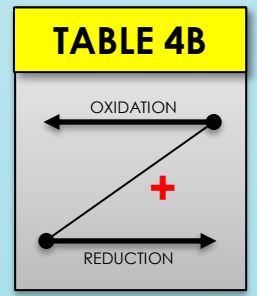
ELECTRODES
 Metal
 Gas

$$E^{\theta}_{cell} = E^{\theta}_{cathode} - E^{\theta}_{anode}$$

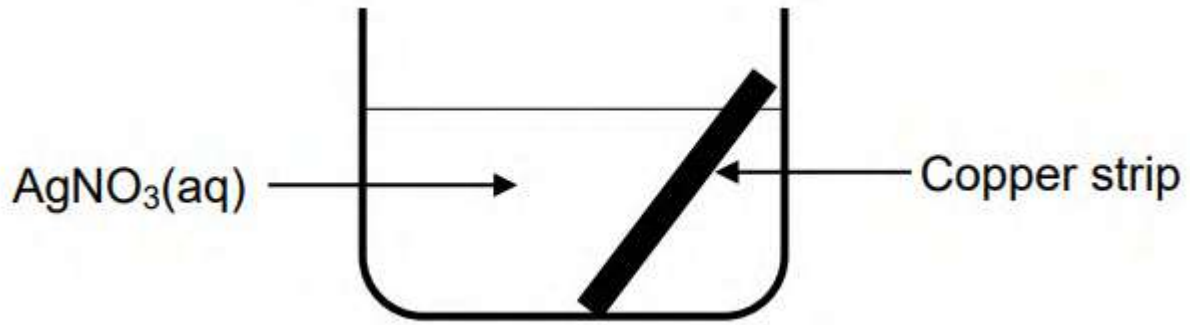
$$E^{\theta}_{cell} = E^{\theta}_{reduction} - E^{\theta}_{oxidation}$$



Spontaneous Reaction



A cleaned pure copper strip, Cu(s), is placed in a beaker containing a colourless silver nitrate solution, AgNO₃(aq), at 25 °C, as shown below.



After a while, it is observed that the solution in the beaker becomes blue.

8.2 Explain the answer to QUESTION 8.1.1 by referring to the relative strengths of the oxidising agents or reducing agents.

Cu is a stronger reducing agent than Ag and will therefore reduce Ag to Ag⁺.
 Ag⁺ is a stronger oxidizing agent than Cu²⁺ and will therefore oxidize Cu²⁺ to Cu.

TABLE 4A

Reduction → +

$$Ag^{+} + e^{-} \rightleftharpoons Ag \quad + 0,80$$

$$Cu^{2+} + 2e^{-} \rightleftharpoons Cu \quad + 0,34$$

Oxidation ←

TABLE 4B

Oxidation ←

$$Cu^{2+} + 2e^{-} \rightleftharpoons Cu \quad + 0,34$$

$$Ag^{+} + e^{-} \rightleftharpoons Ag \quad + 0,80$$

Reduction →

QUESTION 8



GALVANIC CELLS

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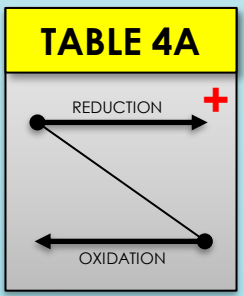
CELL NOTATION: Anode // Cathode
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STANDARD CONDITIONS
 Temp. 25 °C (298 K)
 Concentration 1 mol·dm⁻³
 Pressure 101,3 kPa.

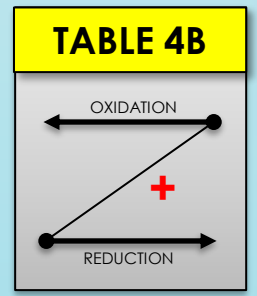
ELECTRODES
 Metal
 Gas

$$E^{\theta}_{\text{cell}} = E^{\theta}_{\text{cathode}} - E^{\theta}_{\text{anode}}$$

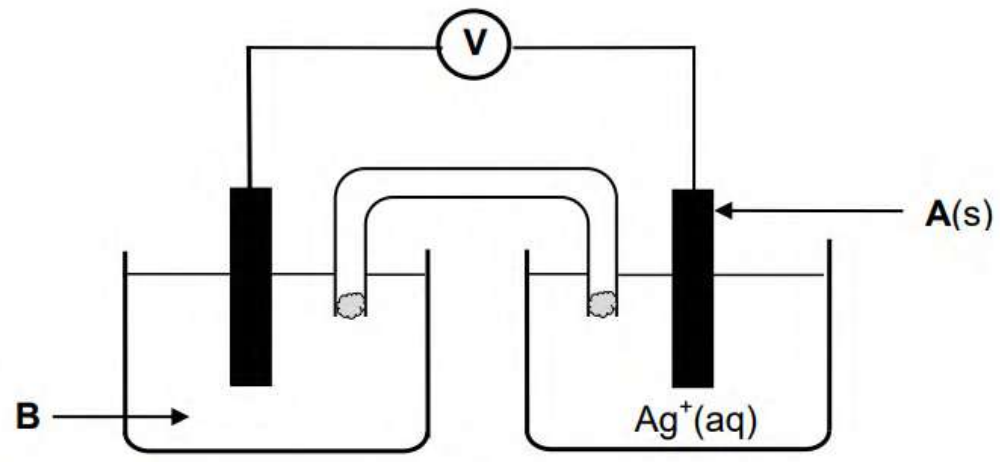
$$E^{\theta}_{\text{cell}} = E^{\theta}_{\text{reduction}} - E^{\theta}_{\text{oxidation}}$$



Spontaneous Reaction



A galvanic cell is now set up using Cu and Ag strips as electrodes. A simplified diagram of the cell is shown below.



8.3 Write down the:

8.3.1 NAME or FORMULA of electrode A

8.3.2 NAME or FORMULA of solution B

8.3.3 Overall (net) balanced equation for the cell reaction

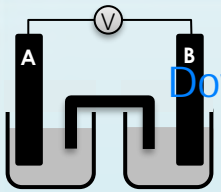
8.4 The salt bridge contains potassium nitrate, KNO₃(aq). Write down the FORMULA of the ion in the salt bridge that will move into the silver ion solution. Choose from K⁺ (aq) or NO₃⁻ (aq). Give a reason for the answer.



QUESTION 8

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GALVANIC CELLS



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CELL NOTATION: Anode // Cathode
 Element / Ion // Ion / Element
 Zn / Zn²⁺ // Cu²⁺ / Cu

STANDARD CONDITIONS

Temp. 25 °C (298 K)
 Concentration 1 mol·dm⁻³
 Pressure 101,3 kPa.

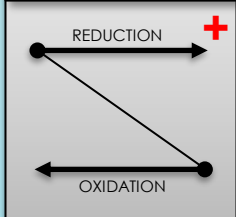
ELECTRODES

Metal
 Gas

$$E^{\theta}_{\text{cell}} = E^{\theta}_{\text{cathode}} - E^{\theta}_{\text{anode}}$$

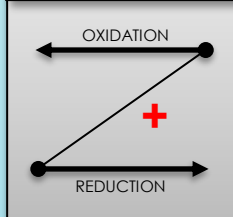
$$E^{\theta}_{\text{cell}} = E^{\theta}_{\text{reduction}} - E^{\theta}_{\text{oxidation}}$$

TABLE 4A

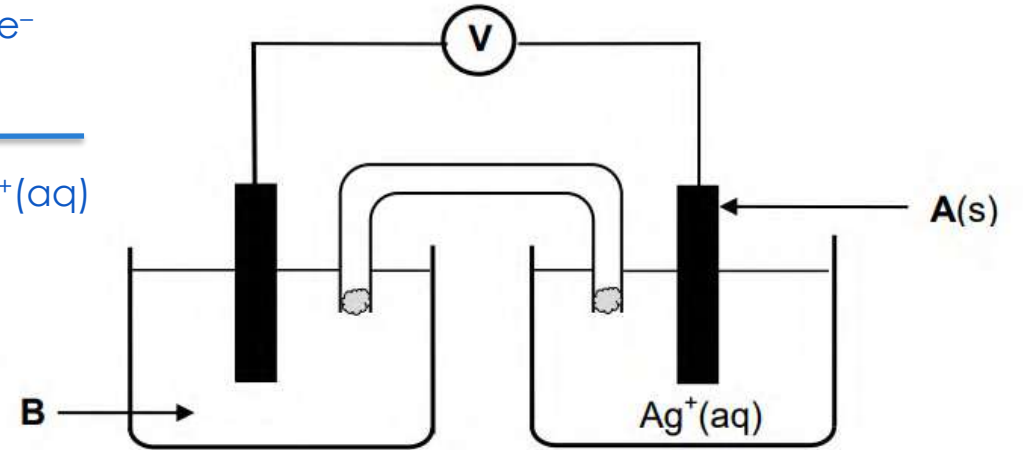
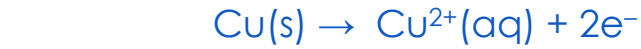


Spontaneous Reaction

TABLE 4B



A galvanic cell is now set up using Cu and Ag strips as electrodes. A simplified diagram of the cell is shown below.



8.3 Write down the:

8.3.1 NAME or FORMULA of electrode A Silver / Ag

8.3.2 NAME or FORMULA of solution B Copper nitrate / Cu(NO₃)₂

8.3.3 Overall (net) balanced equation for the cell reaction

TABLE 4A

Reduction	+
$\text{Ag}^{+} + \text{e}^{-} \rightleftharpoons \text{Ag}$	+ 0,80
$\text{Cu}^{2+} + 2\text{e}^{-} \rightleftharpoons \text{Cu}$	+ 0,34
Oxidation	

TABLE 4B

Oxidation	
$\text{Cu}^{2+} + 2\text{e}^{-} \rightleftharpoons \text{Cu}$	+ 0,34
$\text{Ag}^{+} + \text{e}^{-} \rightleftharpoons \text{Ag}$	+ 0,80
Reduction	

QUESTION 8



GALVANIC CELLS

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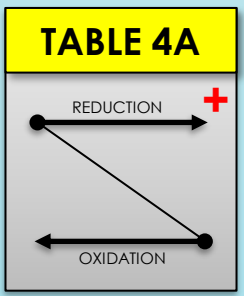
CELL NOTATION: Anode // Cathode
 Element / Ion // Ion / Element
 Zn / Zn²⁺ // Cu²⁺ / Cu

STANDARD CONDITIONS
 Temp. 25 °C (298 K)
 Concentration 1 mol·dm⁻³
 Pressure 101,3 kPa.

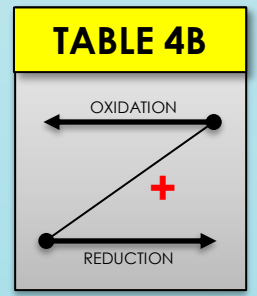
ELECTRODES
 Metal
 Gas

$$E^\theta_{\text{cell}} = E^\theta_{\text{cathode}} - E^\theta_{\text{anode}}$$

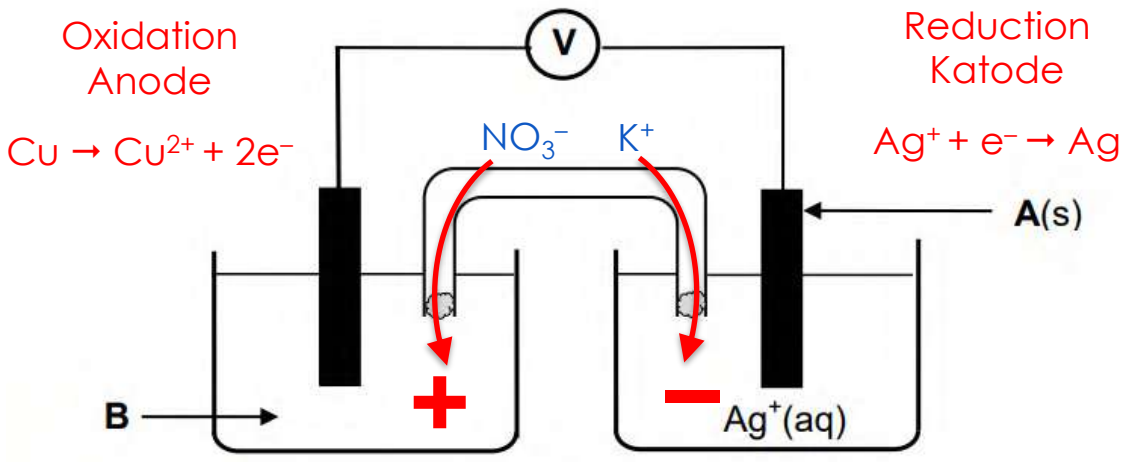
$$E^\theta_{\text{cell}} = E^\theta_{\text{reduction}} - E^\theta_{\text{oxidation}}$$



Spontaneous Reaction



A galvanic cell is now set up using Cu and Ag strips as electrodes. A simplified diagram of the cell is shown below.



8.4 The salt bridge contains potassium nitrate, KNO₃(aq). Write down the FORMULA of the ion in the salt bridge that will move into the silver ion solution. Choose from K⁺ (aq) or NO₃⁻ (aq). Give a reason for the answer.

Answer: K⁺

TABLE 4A

Reduction

$$\text{Ag}^+ + e^- \rightleftharpoons \text{Ag} \quad + 0,80$$

$$\text{Cu}^{2+} + 2e^- \rightleftharpoons \text{Cu} \quad + 0,34$$

Oxidation

TABLE 4B

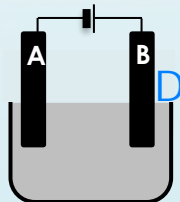
Oxidation

$$\text{Cu}^{2+} + 2e^- \rightleftharpoons \text{Cu} \quad + 0,34$$

$$\text{Ag}^+ + e^- \rightleftharpoons \text{Ag} \quad + 0,80$$

Reduction

QUESTION 9

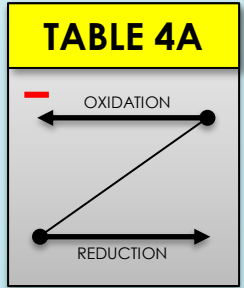


ELECTROLYTIC CELLS

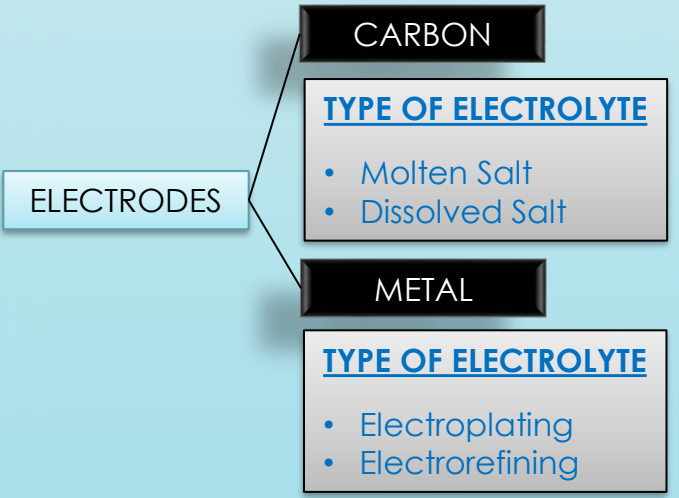
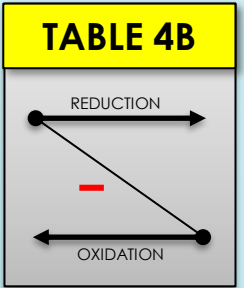
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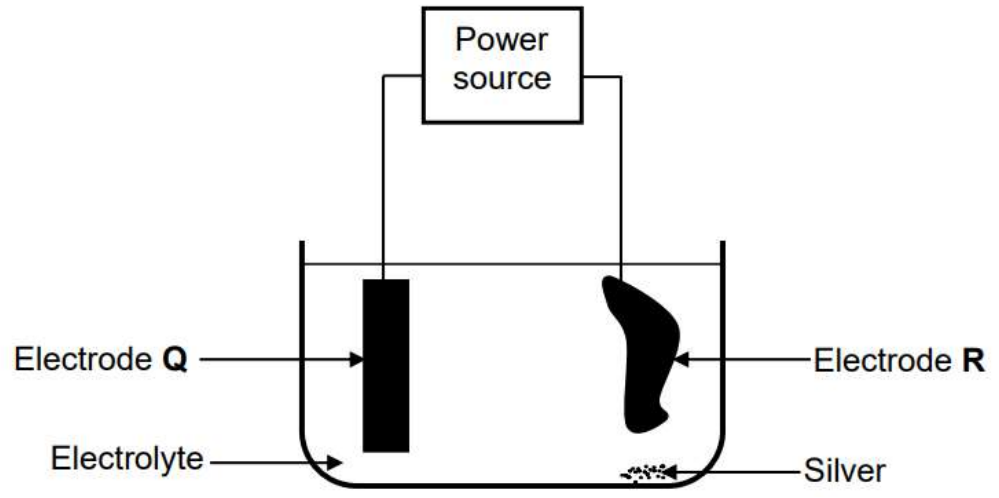
EXPLAIN STRENGTH OF REDUCING AGENTS AND OXIDIZING AGENTS IN REDOX REACTIONS.



Non-Spontaneous Reaction



An electrolytic cell is set up to purify a piece of copper that contains silver and zinc as impurities. A simplified diagram of the cell is shown below. Electrode R is impure copper.



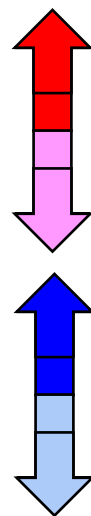
- 9.1 Define the term electrolysis.
- 9.2 Write down the reaction taking place at electrode Q.
- 9.3 In which direction do the electrons flow in the external circuit? Choose from Q to R or R to Q.
- 9.4 Calculate the current needed to form 16 g of copper when the cell operates for five hours.
- 9.5 During this electrolysis, only copper and zinc are oxidised. Give a reason why the silver is not oxidised.



TABLE 4A: REDOX HALF REACTIONS
E⁰ (V)

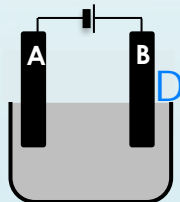
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$F_2(g) + 2e^- \rightleftharpoons 2F^-$	+ 2,87
$Co^{3+} + e^- \rightleftharpoons Co^{2+}$	+ 1,81
$H_2O_2 + 2H^+ + 2e^- \rightleftharpoons 2H_2O$	+1,77
$MnO_4^- + 8H^+ + 5e^- \rightleftharpoons Mn^{2+} + 4H_2O$	+ 1,51
$Cl_2(g) + 2e^- \rightleftharpoons 2Cl^-$	+ 1,36
$Cr_2O_7^{2-} + 14H^+ + 6e^- \rightleftharpoons 2Cr^{3+} + 7H_2O$	+ 1,33
$O_2(g) + 4H^+ + 4e^- \rightleftharpoons 2H_2O$	+ 1,23
$MnO_2 + 4H^+ + 2e^- \rightleftharpoons Mn^{2+} + 2H_2O$	+ 1,23
$Pt^{2+} + 2e^- \rightleftharpoons Pt$	+ 1,20
$Br_2(l) + 2e^- \rightleftharpoons 2Br^-$	+ 1,07
$NO_3^- + 4H^+ + 3e^- \rightleftharpoons NO(g) + 2H_2O$	+ 0,96
$Hg^{2+} + 2e^- \rightleftharpoons Hg(l)$	+ 0,85
$Ag^+ + e^- \rightleftharpoons Ag$	+ 0,80
$NO_3^- + 2H^+ + e^- \rightleftharpoons NO_2(g) + H_2O$	+ 0,80
$Fe^{3+} + e^- \rightleftharpoons Fe^{2+}$	+ 0,77
$O_2(g) + 2H^+ + 2e^- \rightleftharpoons H_2O_2$	+ 0,68
$I_2 + 2e^- \rightleftharpoons 2I^-$	+ 0,54
$Cu^+ + e^- \rightleftharpoons Cu$	+ 0,52
$SO_2 + 4H^+ + 4e^- \rightleftharpoons S + 2H_2O$	+ 0,45
$2H_2O + O_2 + 4e^- \rightleftharpoons 4OH^-$	+ 0,40
$Cu^{2+} + 2e^- \rightleftharpoons Cu$	+ 0,34
$SO_4^{2-} + 4H^+ + e^- \rightleftharpoons SO_2(g) + 2H_2O$	+ 0,17


TABLE 4B: REDOX HALF REACTIONS
E⁰ (V)

$Li^+ + e^- \rightleftharpoons Li$	- 3,05
$K^+ + e^- \rightleftharpoons K$	- 2,93
$Cs^+ + e^- \rightleftharpoons Cs$	- 2,92
$Ba^{2+} + 2e^- \rightleftharpoons Ba$	- 2,90
$Sr^{2+} + 2e^- \rightleftharpoons Sr$	- 2,89
$Ca^{2+} + 2e^- \rightleftharpoons Ca$	- 2,87
$Na^+ + e^- \rightleftharpoons Na$	- 2,71
$Mg^{2+} + 2e^- \rightleftharpoons Mg$	- 2,36
$Al^{3+} + 3e^- \rightleftharpoons Al$	- 1,66
$Mn^{2+} + 2e^- \rightleftharpoons Mn$	- 1,18
$Cr^{2+} + 2e^- \rightleftharpoons Cr$	- 0,91
$2H_2O + 2e^- \rightleftharpoons H_2(g) + 2OH^-$	- 0,83
$Zn^{2+} + 2e^- \rightleftharpoons Zn$	- 0,76
$Cr^{3+} + 3e^- \rightleftharpoons Cr$	- 0,74
$Fe^{2+} + 2e^- \rightleftharpoons Fe$	- 0,44
$Cr^{3+} + e^- \rightleftharpoons Cr^{2+}$	- 0,41
$Cd^{2+} + 2e^- \rightleftharpoons Cd$	- 0,40
$Co^{2+} + 2e^- \rightleftharpoons Co$	- 0,28
$Ni^{2+} + 2e^- \rightleftharpoons Ni$	- 0,27
$Sn^{2+} + 2e^- \rightleftharpoons Sn$	- 0,14
$Pb^{2+} + 2e^- \rightleftharpoons Pb$	- 0,13
$Fe^{3+} + 3e^- \rightleftharpoons Fe$	- 0,06

QUESTION 9

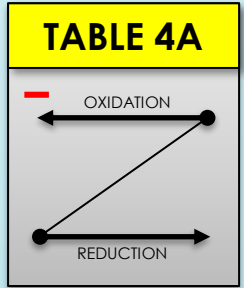


ELECTROLYTIC CELLS

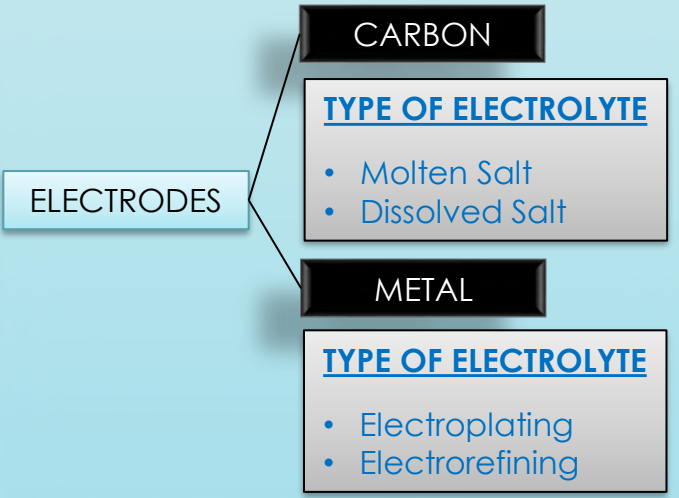
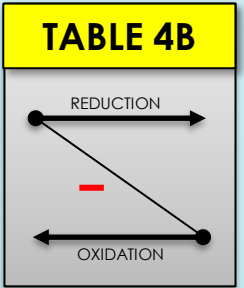
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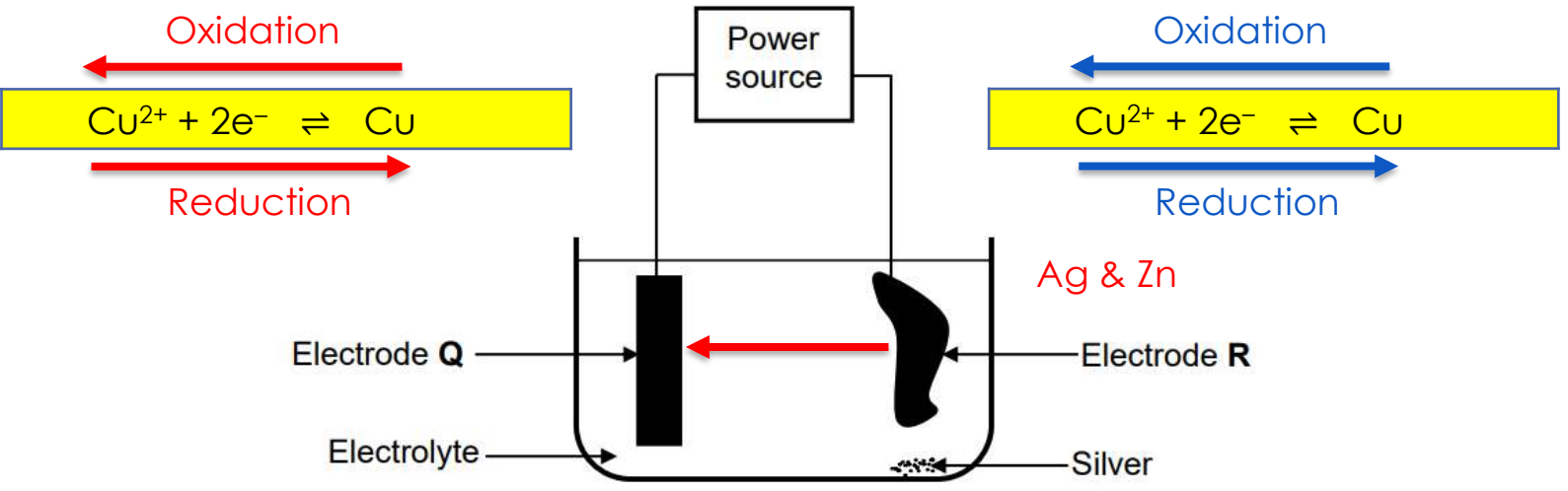
EXPLAIN STRENGTH OF REDUCING AGENTS AND OXIDIZING AGENTS IN REDOX REACTIONS.



Non-Spontaneous Reaction



An electrolytic cell is set up to purify a piece of copper that contains silver and zinc as impurities. A simplified diagram of the cell is shown below. Electrode R is impure copper.

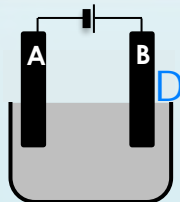


9.1 Define the term electrolysis.

ELECTROLYTIC: A chemical process in which electrical energy is converted into chemical energy.



QUESTION 9

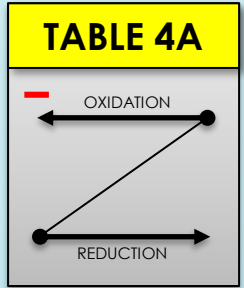


ELECTROLYTIC CELLS

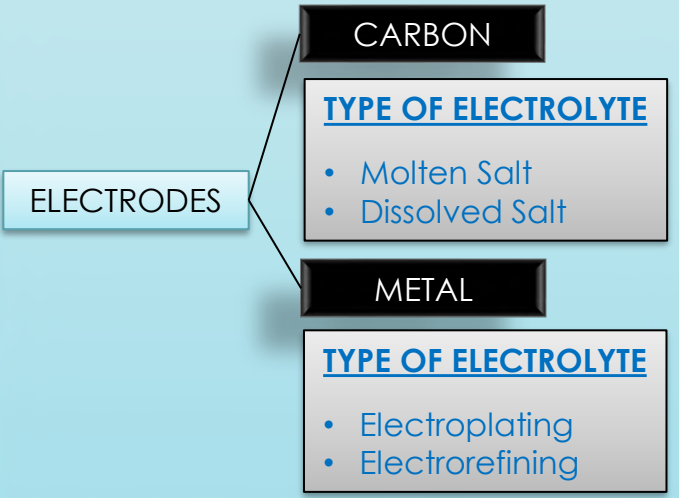
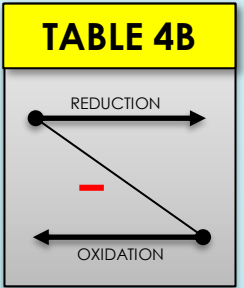
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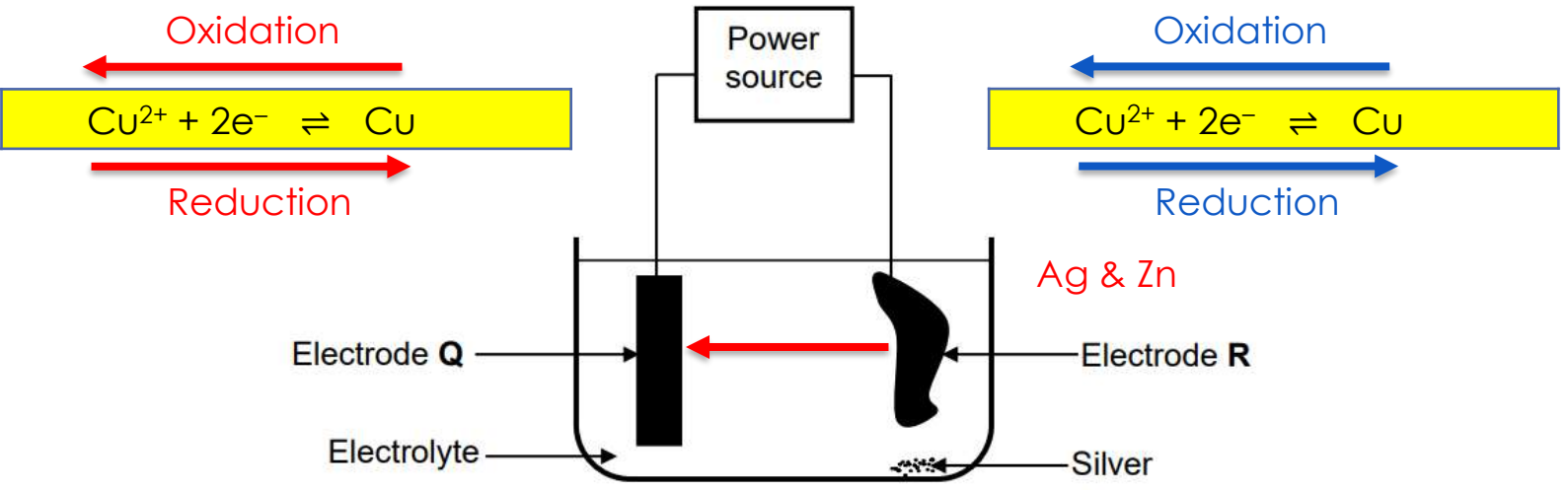
EXPLAIN STRENGTH OF REDUCING AGENTS AND OXIDIZING AGENTS IN REDOX REACTIONS.



Non-Spontaneous Reaction



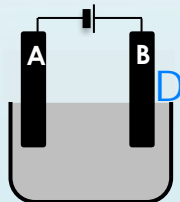
An electrolytic cell is set up to purify a piece of copper that contains silver and zinc as impurities. A simplified diagram of the cell is shown below. Electrode R is impure copper.



9.2 Write down the reaction taking place at electrode Q.



QUESTION 9

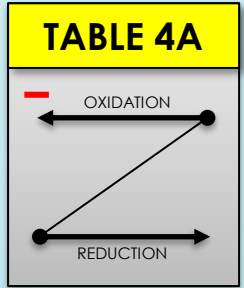


ELECTROLYTIC CELLS

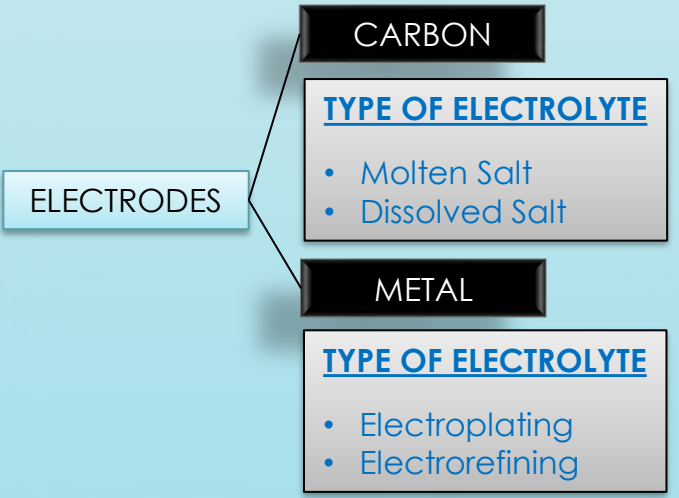
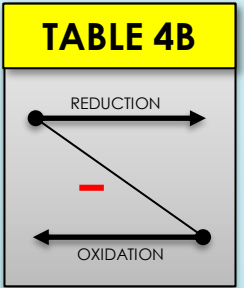
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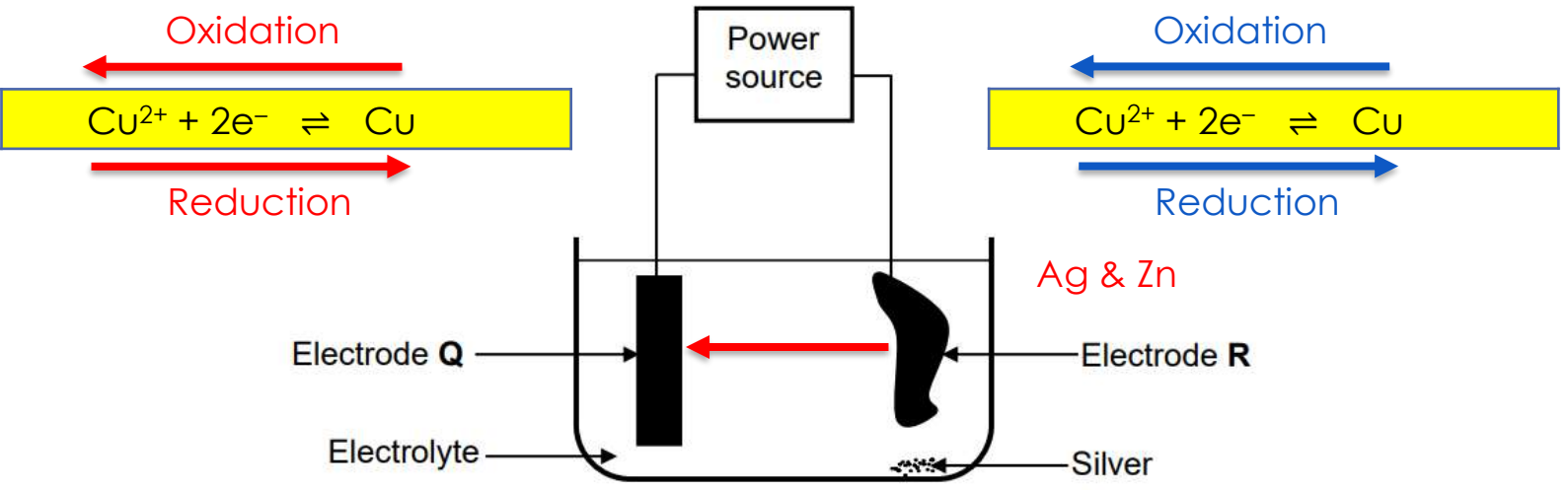
EXPLAIN STRENGTH OF REDUCING AGENTS AND OXIDIZING AGENTS IN REDOX REACTIONS.



Non-Spontaneous Reaction



An electrolytic cell is set up to purify a piece of copper that contains silver and zinc as impurities. A simplified diagram of the cell is shown below. Electrode R is impure copper.



9.3 In which direction do the electrons flow in the external circuit? Choose from Q to R or R to Q.

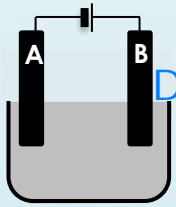
R to Q



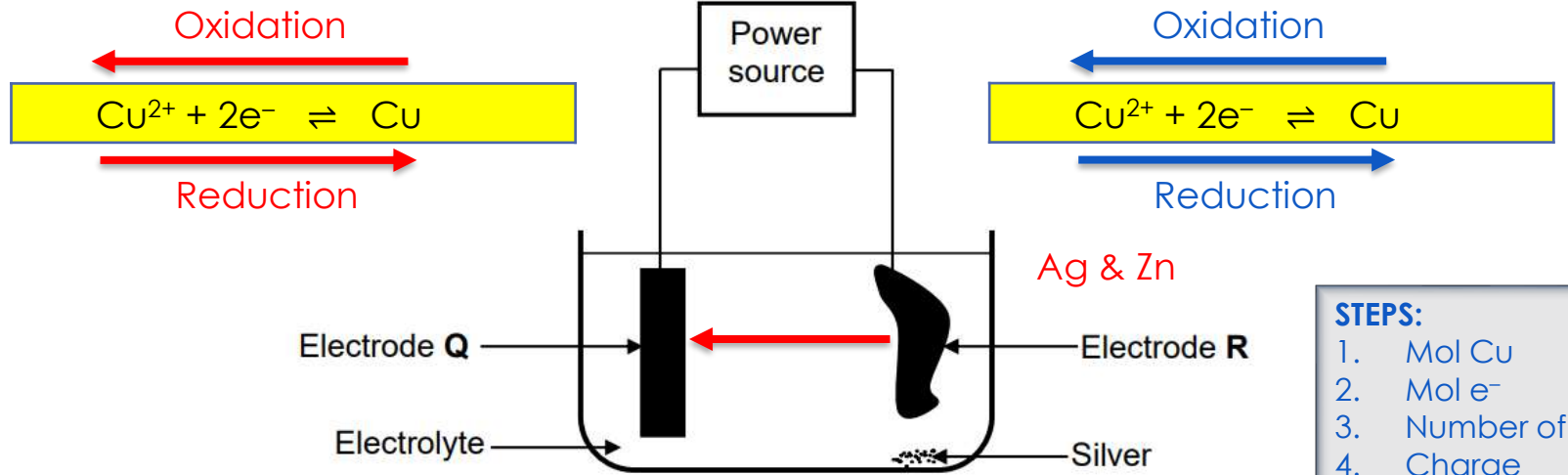
QUESTION 9

ELECTROLYTIC CELLS

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An electrolytic cell is set up to purify a piece of copper that contains silver and zinc as impurities. A simplified diagram of the cell is shown below. Electrode R is impure copper.



- STEPS:**
1. Mol Cu
 2. Mol e⁻
 3. Number of e⁻
 4. Charge
 5. Current

9.4 Calculate the current needed to form 16 g of copper when the cell operates for five hours.

1 $n(\text{Cu}) = \frac{m}{M} = \frac{16}{63,5} = 0,25 \text{ mol}$

4 $n = \frac{Q}{q_e} \therefore Q = (3,034 \times 10^{23})(1,6 \times 10^{-19}) = 48539,21 \text{ C}$

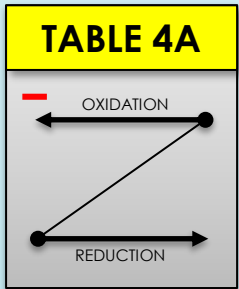
2 $n(e^-) = 2 \times 0,25 = 0,5 \text{ mol}$

3 $n = \frac{N}{N_A} \therefore N = n \cdot N_A = (0,5)(6,02 \times 10^{23}) = 3,034 \times 10^{23} e^-$

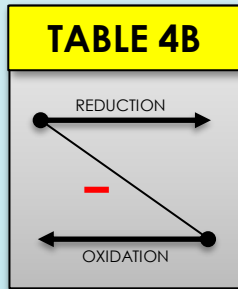
5 $I = \frac{Q}{\Delta t} = \frac{48539,21}{18000} = 2,68 \text{ A}$

Make sure you know the correct wording of all definitions!

EXPLAIN STRENGTH OF REDUCING AGENTS AND OXIDIZING AGENTS IN REDOX REACTIONS.



Non-Spontaneous Reaction



ELECTRODES

CARBON

TYPE OF ELECTROLYTE

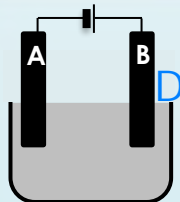
- Molten Salt
- Dissolved Salt

METAL

TYPE OF ELECTROLYTE

- Electroplating
- Electrefining

QUESTION 9

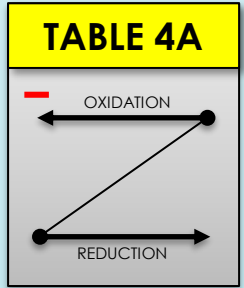


ELECTROLYTIC CELLS

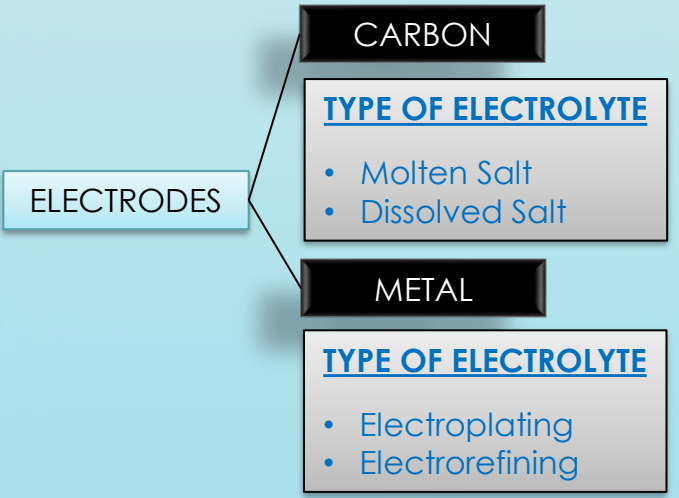
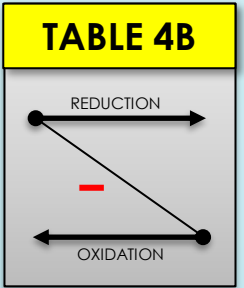
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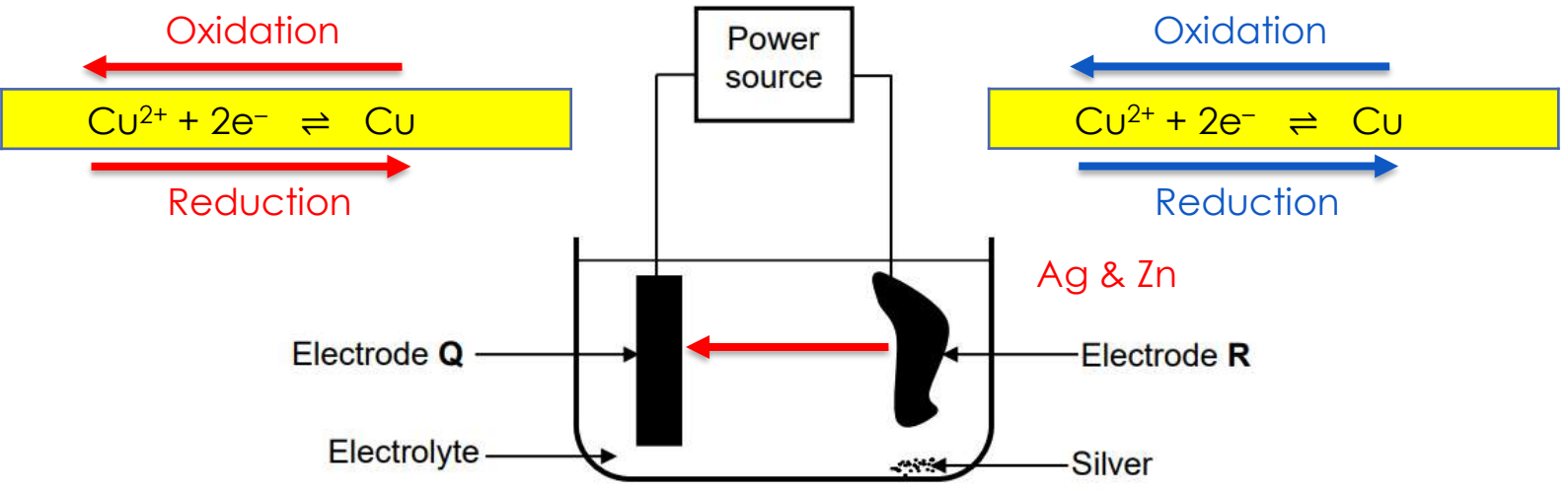
EXPLAIN STRENGTH OF REDUCING AGENTS AND OXIDIZING AGENTS IN REDOX REACTIONS.



Non-Spontaneous Reaction

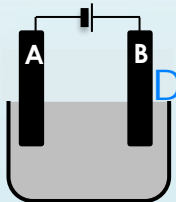


An electrolytic cell is set up to purify a piece of copper that contains silver and zinc as impurities. A simplified diagram of the cell is shown below. Electrode R is impure copper.



9.5 During this electrolysis, only copper and zinc are oxidised. Give a reason why the silver is not oxidised.





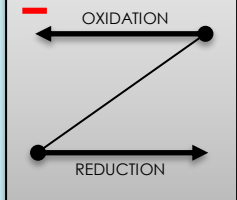
ELECTROLYTIC CELLS

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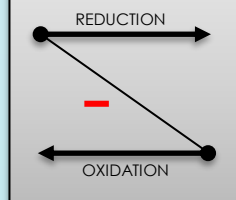
EXPLAIN STRENGTH OF **REDUCING AGENTS** AND **OXIDIZING AGENTS** IN REDOX REACTIONS.

TABLE 4A



Non-Spontaneous Reaction

TABLE 4B



ELECTRODES

CARBON

TYPE OF ELECTROLYTE

- Molten Salt
- Dissolved Salt

METAL

TYPE OF ELECTROLYTE

- Electroplating
- Electrefining

TABLE

4A

Half-reactions/Halfreactions	E ⁰ (V)
F ₂ (g) + 2e ⁻ = 2F ⁻	+2,87
Oxidation	
MnO ₄ ⁻ + 8H ⁺ + 5e ⁻ = Mn ²⁺ + 4H ₂ O	+1,51
Cl ₂ (g) + 2e ⁻ = 2Cl ⁻	+1,36
Cr ₂ O ₇ ²⁻ + 14H ⁺ + 6e ⁻ = 2Cr ³⁺ + 7H ₂ O	+1,33
O ₂ (g) + 4H ⁺ + 4e ⁻ = 2H ₂ O	+1,23
MnO ₂ + 4H ⁺ + 2e ⁻ = Mn ²⁺ + 2H ₂ O	+1,23
Pt ²⁺ + 2e ⁻ = Pt	+1,20
Br ₂ (l) + 2e ⁻ = 2Br ⁻	+1,07
NO ₃ ⁻ + 4H ⁺ + 3e ⁻ = NO(g) + 2H ₂ O	+0,96
Hg ²⁺ + 2e ⁻ = Hg(l)	+0,85
Ag ⁺ + e ⁻ = Ag	+0,80
NO ₃ ⁻ + 2H ⁺ + e ⁻ = NO ₂ (g) + H ₂ O	+0,80
Fe ³⁺ + e ⁻ = Fe ²⁺	+0,77
O ₂ (g) + 2H ⁺ + 2e ⁻ = H ₂ O ₂	+0,68
I ₂ + 2e ⁻ = 2I ⁻	+0,54
Cu ⁺ + e ⁻ = Cu	+0,52
SO ₂ + 4H ⁺ + 4e ⁻ = S + 2H ₂ O	+0,45
2H ₂ O + O ₂ + 4e ⁻ = 4OH ⁻	+0,40
Cu ²⁺ + 2e ⁻ = Cu	+0,34
SO ₄ ²⁻ + 4H ⁺ + 2e ⁻ = SO ₂ (g) + 2H ₂ O	+0,17
Cu ²⁺ + e ⁻ = Cu ⁺	+0,16
Sn ⁴⁺ + 2e ⁻ = Sn ²⁺	+0,15
S + 2H ⁺ + 2e ⁻ = H ₂ S(g)	+0,14
2H ⁺ + 2e ⁻ = H ₂ (g)	0,00
Fe ²⁺ + 3e ⁻ = Fe	-0,06
Pb ²⁺ + 2e ⁻ = Pb	-0,13
Sn ²⁺ + 2e ⁻ = Sn	-0,14
Ni ²⁺ + 2e ⁻ = Ni	-0,27
Co ³⁺ + 2e ⁻ = Co	-0,28
Cd ²⁺ + 2e ⁻ = Cd	-0,40
Cr ³⁺ + e ⁻ = Cr ²⁺	-0,41
Fe ²⁺ + 2e ⁻ = Fe	-0,44
Cr ³⁺ + 3e ⁻ = Cr	-0,74
Zn ²⁺ + 2e ⁻ = Zn	-0,76
2H ₂ O + 2e ⁻ = H ₂ (g) + 2OH ⁻	-0,83
Cr ²⁺ + 2e ⁻ = Cr	-0,91
Mn ²⁺ + 2e ⁻ = Mn	-1,18
Al ³⁺ + 3e ⁻ = Al	-1,66
Mg ²⁺ + 2e ⁻ = Mg	-2,36
Na ⁺ + e ⁻ = Na	-2,71
Ca ²⁺ + 2e ⁻ = Ca	-2,87
Sr ²⁺ + 2e ⁻ = Sr	-2,89
Ba ²⁺ + 2e ⁻ = Ba	-2,90
Cs ⁺ + e ⁻ = Cs	-2,92
K ⁺ + e ⁻ = K	-2,93
Li ⁺ + e ⁻ = Li	-3,05

INCREASING STRENGTH OF OXIDIZING AGENTS

INCREASING STRENGTH OF REDUCING AGENTS

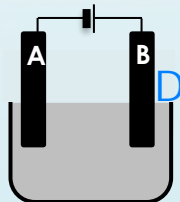
TABLE

4B

Half-reactions/Halfreactions	E ⁰ (V)
Li ⁺ + e ⁻ = Li	-3,05
Oxidation	
Sr + 2e ⁻ = Sr ²⁺	-2,89
Ca ²⁺ + 2e ⁻ = Ca	-2,87
Na ⁺ + e ⁻ = Na	-2,71
Mg ²⁺ + 2e ⁻ = Mg	-2,36
Al ³⁺ + 3e ⁻ = Al	-1,66
Mn ²⁺ + 2e ⁻ = Mn	-1,18
Cr ²⁺ + 2e ⁻ = Cr	-0,91
2H ₂ O + 2e ⁻ = H ₂ (g) + 2OH ⁻	-0,83
Zn ²⁺ + 2e ⁻ = Zn	-0,76
Cr ³⁺ + 3e ⁻ = Cr	-0,74
Fe ²⁺ + 2e ⁻ = Fe	-0,44
Cr ³⁺ + e ⁻ = Cr ²⁺	-0,41
Cd ²⁺ + 2e ⁻ = Cd	-0,40
Co ²⁺ + 2e ⁻ = Co	-0,28
Ni ²⁺ + 2e ⁻ = Ni	-0,27
Sn ²⁺ + 2e ⁻ = Sn	-0,14
Pb ²⁺ + 2e ⁻ = Pb	-0,13
Fe ³⁺ + 3e ⁻ = Fe	-0,06
2H ⁺ + 2e ⁻ = H ₂ (g)	0,00
S + 2H ⁺ + 2e ⁻ = H ₂ S(g)	+0,14
Sn ⁴⁺ + 2e ⁻ = Sn ²⁺	+0,15
Cu ²⁺ + e ⁻ = Cu ⁺	+0,16
SO ₄ ²⁻ + 4H ⁺ + 2e ⁻ = SO ₂ (g) + 2H ₂ O	+0,17
Cu ²⁺ + 2e ⁻ = Cu	+0,34
2H ₂ O + O ₂ + 4e ⁻ = 4OH ⁻	+0,40
SO ₂ + 4H ⁺ + 4e ⁻ = S + 2H ₂ O	+0,45
Cu ⁺ + e ⁻ = Cu	+0,52
I ₂ + 2e ⁻ = 2I ⁻	+0,54
O ₂ (g) + 2H ⁺ + 2e ⁻ = H ₂ O ₂	+0,68
Fe ³⁺ + e ⁻ = Fe ²⁺	+0,77
NO ₃ ⁻ + 2H ⁺ + e ⁻ = NO ₂ (g) + H ₂ O	+0,80
Ag ⁺ + e ⁻ = Ag	+0,80
Hg ²⁺ + 2e ⁻ = Hg(l)	+0,85
NO ₃ ⁻ + 4H ⁺ + 3e ⁻ = NO(g) + 2H ₂ O	+0,96
Br ₂ (l) + 2e ⁻ = 2Br ⁻	+1,07
Pt ²⁺ + 2e ⁻ = Pt	+1,20
MnO ₂ + 4H ⁺ + 2e ⁻ = Mn ²⁺ + 2H ₂ O	+1,23
O ₂ (g) + 4H ⁺ + 4e ⁻ = 2H ₂ O	+1,23
Cr ₂ O ₇ ²⁻ + 14H ⁺ + 6e ⁻ = 2Cr ³⁺ + 7H ₂ O	+1,33
Cl ₂ (g) + 2e ⁻ = 2Cl ⁻	+1,36
MnO ₄ ⁻ + 8H ⁺ + 5e ⁻ = Mn ²⁺ + 4H ₂ O	+1,51
H ₂ O ₂ + 2H ⁺ + 2e ⁻ = 2H ₂ O	+1,77
Co ³⁺ + e ⁻ = Co ²⁺	+1,81
F ₂ (g) + 2e ⁻ = 2F ⁻	+2,87

INCREASING STRENGTH OF REDUCING AGENTS

QUESTION 9

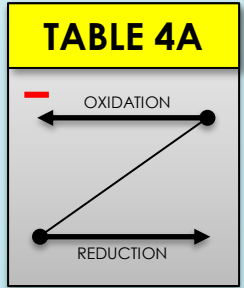


ELECTROLYTIC CELLS

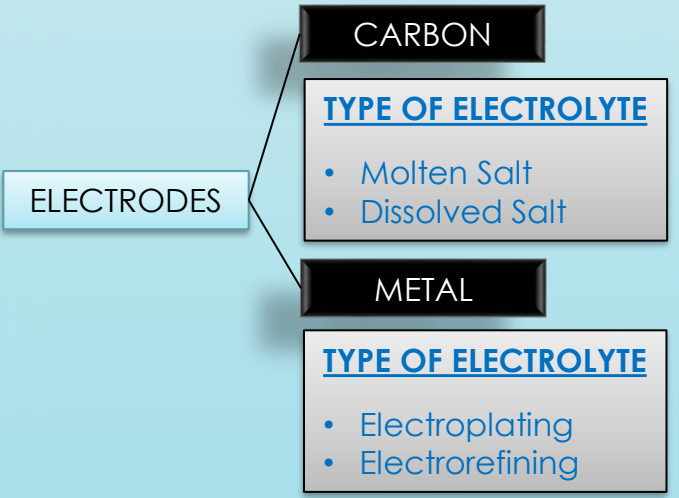
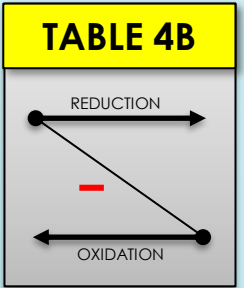
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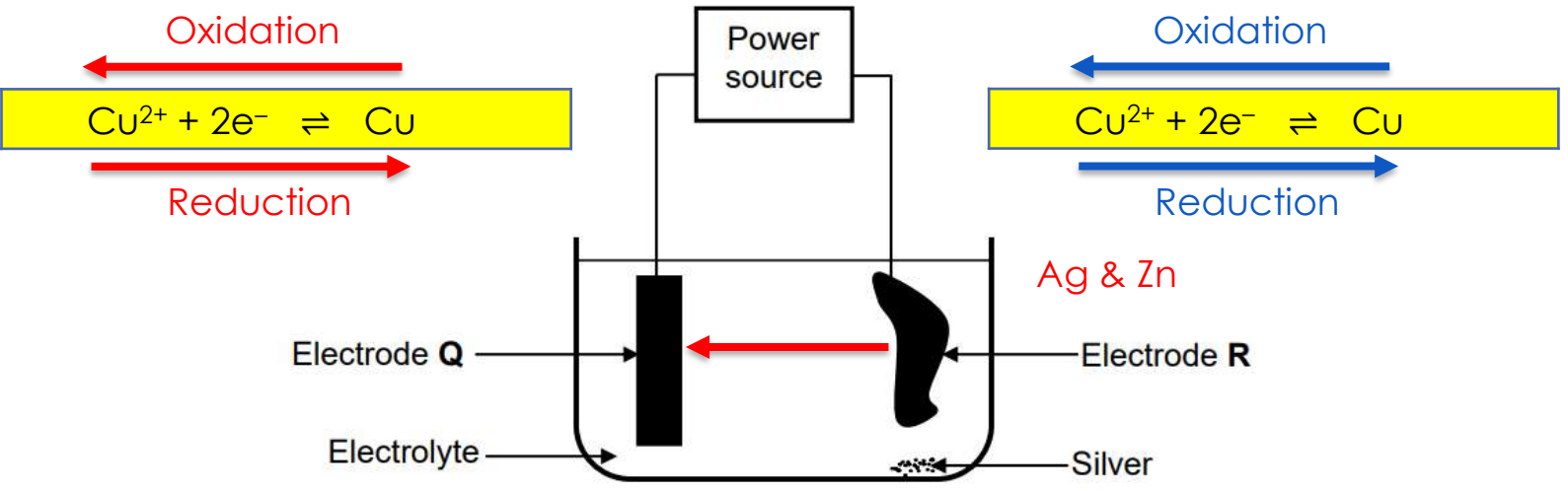
EXPLAIN STRENGTH OF REDUCING AGENTS AND OXIDIZING AGENTS IN REDOX REACTIONS.



Non-Spontaneous Reaction



An electrolytic cell is set up to purify a piece of copper that contains silver and zinc as impurities. A simplified diagram of the cell is shown below. Electrode R is impure copper.



9.5 During this electrolysis, only copper and zinc are oxidised. Give a reason why the silver is not oxidised.

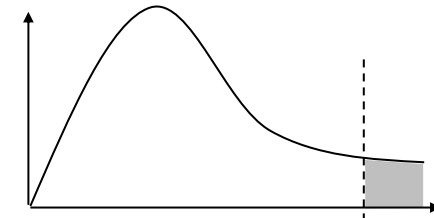
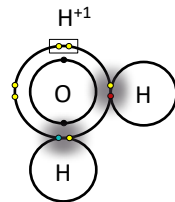
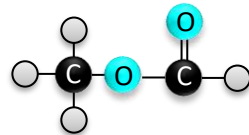
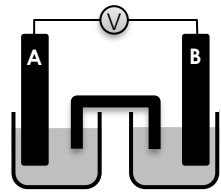
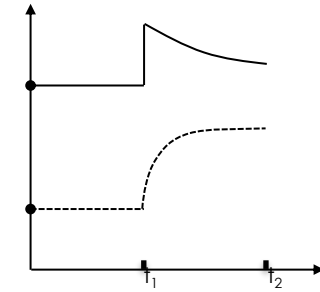
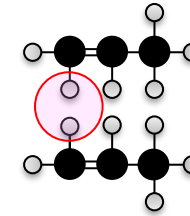
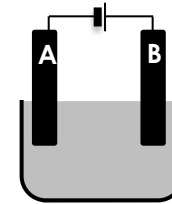
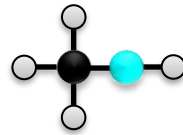
Silver is a **WEAKER REDUCING AGENT** than Cu and will therefore not be oxidized.





Good luck with the Chemistry Exam!

	H ₂ SO ₄	2NaOH	Na ₂ SO ₄	2H ₂ O
n : n	1	2	1	2
n _i				
▲n				
n _f				



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