

2024 SUBJECT WORKBOOK Grade 12



A joint initiative between the Western Cape Education Department and Stellenbosch University.







BROADCAST SESSIONS

GRADE 12 PHYSICAL SCIENCES

Session	Date	Time	Торіс
1	10/04/2024	16h00-17h00	Organic molecules
2	08/05/2024	16h00-17h00	Reaction rates
3	15/07/2024	16h00-17h00	Acids and bases





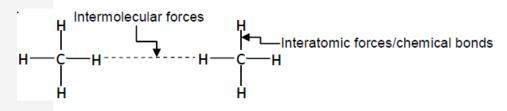


NOTE:

Intermolecular and interatomic forces (chemical bonds)

- Name and explain the different intermolecular forces (Van der Waal's forces):
 - (i) Dipole-dipole forces:
 - Forces between two polar molecules
 - (ii) Induced dipole forces or London forces:
 - Forces between non-polar molecules
 - (iii) Hydrogen bonding:
 - Forces between molecules in which hydrogen is covalently bonded to nitrogen, oxygen or fluorine a special case of dipole-dipole forces
- Describe the difference between intermolecular forces and interatomic forces (intramolecular forces) using a diagram of a group of small molecules; and in words.

Example:



- State the relationship between intermolecular forces and molecular size. For non-polar molecules, the strength of induced dipole forces increases with molecular size.
- Explain the effect of intermolecular forces on boiling point, melting point and vapour pressure.
- Boiling point:

The temperature at which the vapour pressure of a substance equals atmospheric pressure. The stronger the intermolecular forces, the higher the boiling point.

• Melting point:

The temperature at which the solid and liquid phases of a substance are at equilibrium. The stronger the intermolecular forces, the higher the melting point.

• Vapour pressure:

The pressure exerted by a vapour at equilibrium with its liquid in a closed system. The stronger the intermolecular forces, the lower the vapour pressure.

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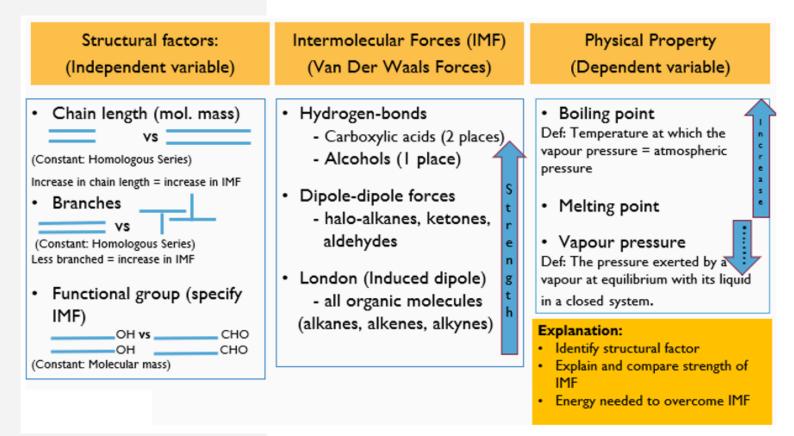
SESSION 1 | PHYSICAL PROPERTIES OF ORGANIC MOLECULES



NOTE:

- This presentation will focus on the Physical Properties of Organic molecules. To understand the concepts that will be covered in this lesson, it is important that you have a thorough knowledge of the nomenclature of all the homologous series of organic molecules.
- Examination questions on this topic will in many cases be presented as the results of an experiment in the form of a table or graph. The results of multiple experiments may be included in the same table or graph. You should be able to identify which combination of experiments may be used in a **fair test**. The skill of interpreting results in tables and graphs need practice.
- The following mind map (summary) simplifies the required knowledge to enable you to answer questions on physical properties of organic molecules. Make sure you understand the mind map.
- In some questions you will be required to identify the structural factor applicable in the experiment and explain it's influence on the strength of the IMF and then mention the effect of this on the physical property applicable. Comparing the energy required per unit time to overcome the IMF might also be required in your explanation. Read these questions carefully and let the marks awarded guide you to the expected answer for the explanation.

MIND MAP/ SUMMARY:





WORKED EXAMPLE:

Question 1

The boiling points of straight-chain alkanes and straight-chain alcohols are compared in the table.

NUMBER OF CARBON ATOMS	BOILING POINTS OF ALKANES (°C)	BOILING POINTS OF ALCOHOLS (°C)
1	- 162	64
2	- 89	78
3	- 42	98
4	- 0,5	118

1.1	Explain the increa	se in boiling points of the alkanes, as indicated in the table.	(3)
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- 1.2 Explain the difference between the boiling points of an alkane and an alcohol, each having THREE carbon atoms per molecule, by referring to the TYPE of intermolecular forces.
- 1.3 Does the vapour pressure of the alcohols INCREASE or DECREASE with an increase in the number of carbon atoms?
- 1.4 How will the boiling point of 2-methylpropane compare to that of its chain isomer? Write down HIGHER THAN, LOWER THAN or EQUAL TO. Give a reason for the

answer by referring to the structural differences between the two compounds. (2)

[10]

[10]

(4)

(1)

ANSWER:

1.1 Structure: From 1 C to 4 C atoms - The chain length/molecular size/molecular mass/ surface area increases. ✓ Intermolecular forces: Increase in strength of intermolecular forces. ✓ Energy: More energy needed to overcome intermolecular forces. \checkmark (3) 1.2 Alkanes have London forces. ✓ Alcohols have hydrogen bonding (in addition to London forces and dipole dipole forces). ✓ Hydrogen bonding are stronger intermolecular forces than London. OR More energy needed to overcome intermolecular forces in alcohols√ Alcohols have higher boiling points than alkanes. ✓ (4) 1.3 Decrease√ (1) Lower than ✓ 1.4 2-methylpropane is more branched/has a smaller surface area than butane/chain isomer. ✓ OR Butane/chain isomer is less branched /has larger surface area than 2-methylpropane (2)



REVISION:

Question 2

2.

2.

2.

The table below shows five organic compounds represented by the letters A to E.

Α	CH ₄
В	CH ₃ CH ₃
С	CH ₃ CH ₂ CH ₃
D	CH ₃ CH ₂ CH ₂ CH ₃
E	CH ₃ CH ₂ OH

2.1 Is compound B SATURATED or UNSATURATED? Give a reason for the answer. (2)

Consider the boiling points of compounds A to E given in random order below and use them, where applicable, to answer the questions that follow.

0 °C	- 162 °C	- 42 °C	- 89 °C	78 °C

2.2 Write down the boiling point of:

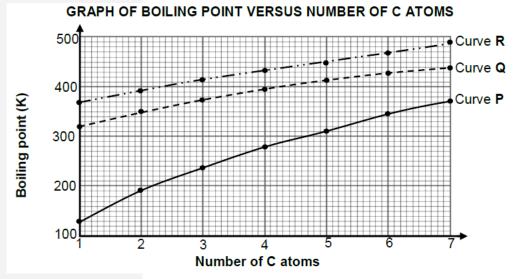
	2.2.1	Compound C	(1)
	2.2.2	Compound E	(1)
3		the difference in boiling points of compounds C and E by referring to the intermolecular forces present in EACH of these compounds.	(3)
.4		oour pressure INCREASE or DECREASE from compounds A to D? Fully the answer.	(4)
5		the vapour pressure of 2-methylpropane compare to the vapour of compound D? Write down only HIGHER THAN, LOWER THAN or O.	(1) [12]



REVISION:

Question 3

The relationship between boiling point and the number of carbon atoms in straight chain molecules of alkanes, carboxylic acids and alcohols is investigated. Curves P, Q and R are obtained.



3.1 Define the term boiling point.

(2)

3.2 For curve P, write down a conclusion that can be drawn from the above results. (2)

3.3 Identify the curve (P, Q or R) that represents each of the following:

3.3.1 Alkanes (1)

3.3.2 Carboxylic acids (1)

- 2.3 Explain the answer to QUESTION 3.3.2 by referring to the:
 - Types of intermolecular forces present in alkanes, carboxylic acids and alcohols
 - Relative strengths of these intermolecular forces
 - Energy needed

(5) **[11]**



Question 4

The boiling points of different organic compounds are given below.

	COMPOUND	BOILING POINT (°C)
Α	НСООН	101
В	CH₃COOH	118
С	CH ₃ CH ₂ COOH	141
D	CH ₃ CH ₂ CH ₂ COOH	164

4.1 Define the term boiling point.

- 4.2 Which ONE of the compounds, **A** or **B** or **C**, has the highest vapour pressure? Refer to the data in the table to give a reason for the answer. (2)
- 4.3 The boiling point of compound **B** is now compared with of compound **X**.

С	COMPOUND		BOILING POINT (°C)
E	W	CH₃COOH	118
	K	CH ₃ CH ₂ CH ₂ OH	98

4.3.1	Besides the conditions used to determine boiling points, give a reason why this is a fair comparison.	(1)
4.3.2	Is compound X a PRIMARY, SECONDARY or TERTIARY alcohol? Give a reason for the answer.	(2)
4.3.3	Fully explain the difference between the boiling points by referring to the types of intermolecular forces present in each of these compounds.	(4) [11]

(2)

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SESSION 2 | REACTION RATE



REVISION:

Question 5

- i. Name the six factors that determine the rate of a chemical reaction.
- ii. Name the two requirements according to the COLLISION THEORY for an effective collision between two molecules.

Grade 12 learners use the reaction between a sodium thiosulphate solution $(Na_2S_2O_3)$ and a hydrochloric acid solution (HCl) to investigate the factors that affect reaction rates.

The balanced equation for this reaction is :

 $Na_2S_2O_3(aq) + 2 HCI(aq) \rightarrow H_2O(I) + SO_2(g) + 2 NaCI(aq) + S(s)$

The time from the moment EQUAL VOLUMES of the two solutions are mixed untill the appearance of a certain amount of turbidity (sulphur precipitation) is taken as an indication of the reaction rate

Consider INVESTIGATION A (Experiment 1 to 3):

	Temp (°C)	[Na ₂ S ₂ O ₃] (mol.dm ⁻³)	[HCl] (mol.dm ⁻³)	Time (s)
Experiment 1	20	0,5	0,5	40
Experiment 2	20	0,9	0,5	25
Experiment 3	20	1,4	0,5	15

iii. For **INVESTIGATION A**, name the:

- a) Dependent variable
- b) Independent variable
- iv. What is the scientific conclusion that can be drawn from INVESTIGATION A?

Consider INVESTIGATION B (Experiment 4 to 6):

	Temp (°C)	[Na ₂ S ₂ O ₃] (mol.dm ⁻³)	[HCl] (mol.dm ⁻³)	Time (s)
Experiment 4	20	0,5	0,5	40
Experiment 5	30	0,5	0,5	20
Experiment 6	50	0,5	0,5	10

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SESSION 2 | REACTION RATE



REVISION:

- v. In which one of the experiments (4 to 6) in **INVESTIGATION B** is the reaction rate the fastest? Give a reason for your answer.
- vi. Explain your answer in question 6.4.1 in terms of the collision theory.

ANSWERS:

- i. Catalyst, Temperature, Gas pressure, State of division (Reaction surface), Nature of the reagents, Concentration of solutes.
- ii. Molecules will undergo effective collisions if :
 - The orientation of the collisions is correct.
 - The molecule has enough energy.
- iii. INVESTIGATION A:
 - a) Reaction Rate / Time
 - b) <u>Concentration</u> of $Na_2S_2O_3$

iv. ANY ANSWER THAT DESCRIBES THE CORRECT RELATIONSHIP : $\checkmark\checkmark$

The reaction rate increases with an increase in concentration of the $Na_2S_2O_3$

- v. Experiment 6
 - Highest temperature / Fastest time
 - Average kinetic energy the highest.
- vi. More particles have enough activation energy (Maxwell-Boltzmann curve).

More effective collisions per second



SESSION 2 | REACTION RATE



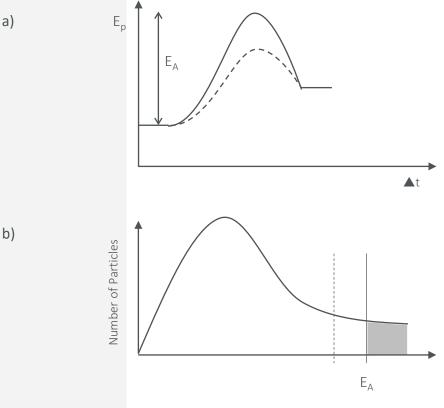
REVISION:

Question 6

- Define the term: CATALYST i.
- Draw the following graphs to indicate the effect of a catalyst on each type of graph : ii.
 - a) 3.2.1 Potential energy ▲t graph
 - b) 3.2.2 Maxwell-Boltzmann curve
 - c) 3.2.3 Reaction Rate –▲t graph
- iii. Name the catalyst used in each of the following reactions :
 - a) 3.3.1 Esterification
 - b) 3.3.2 Addition (Hydrogenation)

ANSWERS:

- CATALYST: Chemical substance that lowers the potential energy of the activated complex and i. accelerates the reaction rate without participating in the reaction itself.
- GRAPHS ii.

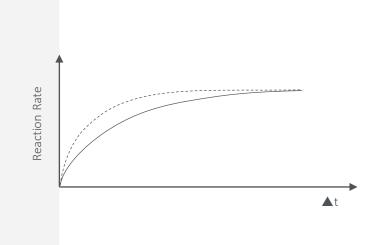


b)

SESSION 2 | REACTION RATE

ANSWERS:

c)



iii. Catalysts:

- a) Sulfuric acid (H₂SO₄)
- b) Platinum (Pt) / Nickel (Ni) / Palladium (Pd)

PHYSICAL SCIENCES

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SESSION 2 | REACTION RATE



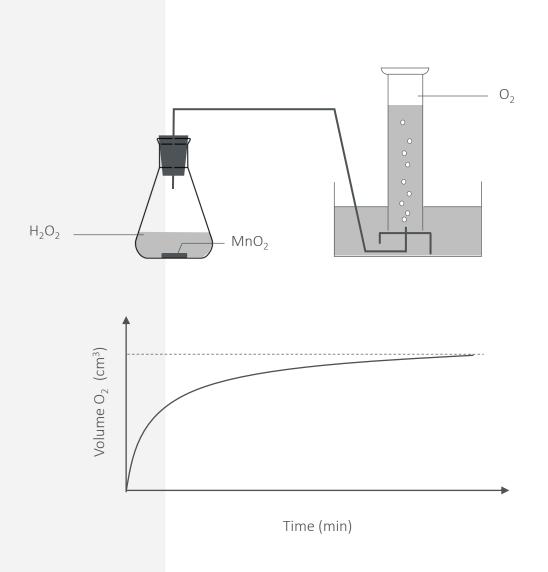
REVISION:

Question 7

Manganese dioxide (MnO_2) acts as a catalyst in the decomposition of hydrogen peroxide (H_2O_2) to form water and oxygen as products. The balanced equation is given :

 $2H_2O_2 (aq) \rightarrow 2H_2O (I) + O_2 (g)$

One gram of MnO_2 powder was added to the hydrogen peroxide, with a concentration of 2 mol.dm⁻³ at a temperature of 25°C and the volume of oxygen, which is collected by means of the downward displacement of water, was measured. The sketch below illustrates the experiment and the graph shows the rate at which the gas was collected in a gas cylinder.



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SESSION 2 | REACTION RATE



REVISION:

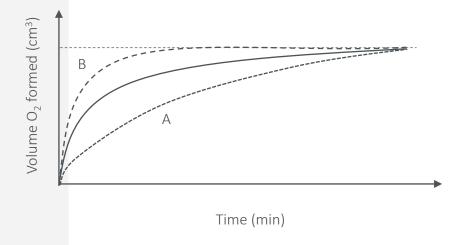
i. Define the term: CATALYST

Copy the GRAPH of time versus volume O_2 onto your answer set. The experiment is repeated twice with a change to one of the constant variables. The first experiment therefore now serves as the control experiment.

- ii. Draw the possible curves expected on the same axis system if:
 - a) No catalyst is used. (Name the new graph A)
 - b) The experiment is repeated at a higher temperature. (Name this new graph B)
- iii. What mass of MnO_2 will be left at the end of the experiment?
- iv. Define the term: REACTION RATE
- v. Name two more factors, apart from the factors already used in this question, which will influence the rate of a chemical reaction.
- vi. Refer to the collision theory to explain how the presence of a catalyst increases the rate of a reaction.

ANSWERS:

- i. Chemical substance that lowers the potential energy of the activated complex and accelerates the reaction rate without participating in the reaction itself.
- ii. GRAPHS
- a) Start and End points same as original curve. Curve BELOW the original curve.
- b) Start and End points same as original curve. Curve ABOVE the original curve.







ANSWERS:

- iii. 1g MnO₂
- iv. The change in concentration of reactants or products per unit time.
- v. Concentration, Reaction surface (state of division), Nature of the reagents
- vi. Catalyst lowers the activation energy.

More molecules possess enough energy for an effective collision.

More collisions occur in a shorter period of time.

Rate at which products are formed increases.



SESSION 2 | REACTION RATE



REVISION:

Question 8

- i. Define the following terms:
 - a) Reaction rate
 - b) Activation energy

A Grade 12 learner uses the reaction of zinc with an excess of hydrochloric acid to investigate the factors that affect the rate of a chemical reaction. The balanced equation for the reaction is :

 $Zn (s) + 2HCl (aq) \rightarrow ZnCl_2 (aq) + H_2(g)$

The learners carry out five experiments during the investigation and summarize all data in the following table. Reaction 1 is the CONTROL EXPERIMENT.

Exporimonto	Mass (Zn)	State of Division	Concentration	Temperature
Experiments		(Zn)	(HCI)	(HCI)
1	1g	Ppebbles	0,5 mol.dm ⁻³	25 °C
2	1g	Ppebbles	0,8 mol.dm ⁻³	25 °C
3	1g	Ppebbles	0,5 mol.dm ⁻³	35 °C
4	1g	Powder	0,5 mol.dm ⁻³	25 °C
5	2g	Ppebbles	0,5 mol.dm ⁻³	25 °C

- ii. Calculate the average reaction rate (in mol.s⁻¹) of reaction 1 if the reaction takes two minutes to complete.
- iii. Name the INDEPENDENT VARIABLE between the following reactions :
 - a) Experiment 1 and Experiment 2
 - b) Experiment 1 and Experiment 3
 - c) Experiment 1 and Experiment 4
 - d) Experiment 1 and Experiment 5





ANSWERS:

- i. DEFINITIONS:
 - a) The change in concentration of reactants or products per unit time.
 - b) The minimum energy required for a reaction to occur.
- ii. $n(Zn) = m \div M = 1 \div 65 = 0,015 \text{ mol}$

Average Reaction Rate = $An \div At = 0,015 \div 120$

Average Reaction Rate = $1,28 \times 10^{-4}$ mol.s⁻¹

(Answer if rounded values were used : 1,25 x 10⁻⁴ mol.s⁻¹)

- iii. VARIABLES:
 - a) Concentration of the acid
 - b) Temperature of the reaction mixture
 - c) State of division of the zinc
 - d) NONE! (Mass of the reagent is not a factor)



REVISION ACIDS & BASES

ARRHENIUS THEORY

- ACIDS donate protons (H^+) to form H_3O^+ .
- BASES donate hydroxide ions (OH⁻).

LOWRY-BRØNSTED THEORY

- Acids donates protons (H⁺).
- BASES accept protons(H⁺).

Reactions of Acids & Bases

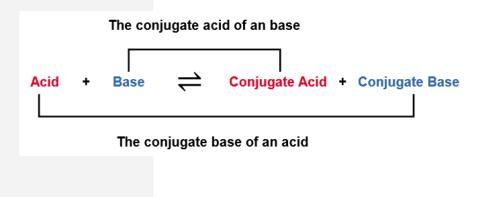
METAL OXIDE + ACID \rightarrow SALT + H₂O

METAL HYDROXIDE + ACID \rightarrow SALT + H₂O

METAL CARBONATE + ACID \rightarrow SALT + H₂O + CO₂

Conjugate Acid-Base Pairs

- When an ACID donates a proton, a CONJUGATE BASE is produced.
- When a BASE accepts a proton, a CONJUGATE ACID is produced.



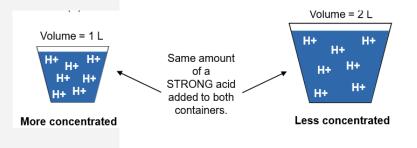


ACID-BASE STRENGTH

- Strong acids ionise completely in solution to form a high concentration of H_3O^+ ions. Examples: Hydrochloric acid (HCl), Sulfuric acid (H_2SO_4), Nitric Acid (HNO₃)
- Weak acids ionise incompletely in solution to form a low concentration of H_3O^+ ions. Examples: Ethanoic acid (CH₃COOH) Hydrofluoric acid (HF) Phosphoric acid (H₃PO₄)
- Strong bases dissociate completely in solution to form a high concentration of OHions. Examples: Sodium hydroxide (NaOH), Potassium hydroxide (KOH), Lithium hydroxide (LiOH)
- Weak bases dissociate incompletely in solution to form a low concentration of OH⁻ ions. Examples: Ammonium hydroxide (NH₄OH), Calcium hydroxide (Ca(OH)₂), Magnesium hydroxide (Mg(OH)₂)

CONCENTRATED ACIDS/BASES AND DILUTE ACIDS/BASES

- Concentrated acids/bases contain a large amount (number of moles) of acid/base in proportion to the volume of water.
- Dilute acids/bases contain a small amount (number of moles) of acid/base in proportion to the volume of water.







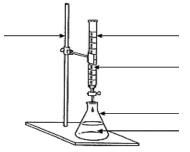
TITRATIONS

- Titrations is an experimental technique used to determine the concentration of an acid or a base using a standard solution.
- Using volumetric analysis, the **unknown concentration** of a solution (acid or base) may be determined.
- Use the equation: $C = \frac{n}{V}$
- Remember moles (n) can be calculated using mass of a substance (m) and its molar mass (M):

$$n = \frac{m}{M}$$

• Titration formula:

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







QUESTION 2

2.1	Sulphuric acid, H2SO4, ionises into two steps as follows:		
		I H2SO4(aq) + H2O(ℓ) → HSO ₄ ⁻ (aq) + H3O ⁺ (aq) Ka = 1 x 10^3	
		II $HSO_4^{-}(aq) + H2O(\ell) \rightarrow SO_4^{2-(aq)} + H3O^{+}(aq)$ Ka = 1 x 10 ⁻²	
	2.1.1	Define an acid in terms of the Lowry-Brønsted theory.	(2)
	2.1.2	Write down the NAME or FORMULA of the substance that acts as	(2)
		an ampholyte in the above equations.	
		Give a reason for the answer.	
	2.1.3	The conductivity of solutions of $HSO_4^+(aq)$ and $H_2SO_4(aq)$ are	(3)
		compared. Which solution will have a LOWER conductivity?	
		Explain the answer.	
2.2	Thop	H of a hydrochloric acid solution, HCℓ(aq), is 1,02 at 25 °C.	
2.2	ine pi	Tor a figurochione actu solution, nee(aq), is 1,02 at 25°C.	
	2.2.1	Calculate the concentration of the HC ² (aq).	(3)
		This HCℓ solution reacts with sodium carbonate, Na ₂ CO ₃ , according to the	
		following balanced equation:	
		2HCℓ(aq) + Na2CO3(aq) \rightarrow 2NaCℓ(aq) + CO2(g) + H2O(ℓ)	
		50 cm ³ of the HC ℓ solution is added to 25 cm3 of a 0,075 mol·dm ⁻³ Na ₂ CO ₃	
		solution.	
	2.2.2	Calculate the concentration of the EXCESS HC ^ℓ in the new	(8)
		solution.	



5	QUESTION 3				
The K	a values for two w	eak acids, oxalic acid	d and carbonic acid, are	e as follows:	
2 1					(2
3.1	Define the term w	/eak acid.			(2
3.2	Which acid, OXAL	IC ACID or CARBONI	C ACID, is stronger? Giv	ve a reason for the ar	nswer. (2
		NAME	FORMULA	Ka	
		Oxalic acid	(COOH) ₂	5,6 x 10 ⁻²	
		Carbonic acid	H ₂ CO ₃	4,3 x 10 ⁻⁷	
3.3	Oxalic acid ionises	s in water according	to the following balanc	ced equation:	
	(COOH)2(s) + 2H2O(ℓ) ≓ (COC	D) ₂ ²⁻ (aq) + 2H3O ⁺ (aq)		
	Write down the F	ORMULAE of the TW	/O bases in this equation	on.	(2
3.4	Learners prepare	2 dm ³ of a sodium h	ydroxide solution of co	ncentration 0,1 mol·	dm ⁻³ .
	Calculate the pH of	of the solution			(4
3.5	During a titration	of the sodium hydro	xide solution in QUEST	ION 3.4 with dilute c	oxalic acid, the
	learners find that	25,1 cm ³ of the NaO)H(aq) neutralises exac	tly 14,2 cm ³ of the (C	COOH) ₂ (aq).
	The balanced equ	ation for the reactio	n is as follows:		
	2NaOH(aq) + (CO	$OH)_2(aq) \rightarrow (COO)_2N$	a₂(aq) + 2H₂O(ℓ)		
	3.5.1 Calculate the	e concentration of th	ne oxalic acid solution.		(5
	The fellowing indi		for the tituetion.		
	The following indi	cators are available			
			PH RANGE 3,1-4,4	<u> </u>	
		B	6,0–7,6		
	3.5.2 Which ONE	of the indicators and	8,3–10,0	Inis utration?	10
		of the maleators aso	ove is most suitable for		(2
	Give a reason for				(2

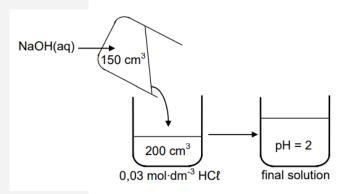
Ql	JESTION 4	
4.1	Define an acid in terms of the Lowry-Brønsted theory.	(2)
4.2	Carbonated water is an aqueous solution of carbonic acid, H_2CO_3 . $H_2CO_3(aq)$ ionises in two steps when it dissolves in water.	
	4.2.1 Write down the FORMULA of the conjugate base of $H_2CO_3(aq)$.	(1)
	4.2.2 Write down a balanced equation for the first step in the ionisation of carbonic acid.	(3)
	4.2.3 The pH of a carbonic acid solution at 25 °C is 3,4. Calculate the hydroxide ion concentration in the solution.	(5)
4.3	X is a monoprotic acid.	
	4.3.1 State the meaning of the term monoprotic.	(1)
	4.3.2 A sample of acid X is titrated with a standard sodium hydroxide solution using a suitable indicator.	
	At the endpoint it is found that 25 cm ³ of acid X is neutralised by 27,5 cm ³ of the sodium hydroxide solution of concentration 0,1 mol·dm ⁻³ .	
	Calculate the concentration of acid X.	(5)
	4.3.3 The concentration of H3O+ ions in the sample of acid X is 2,4 x 10^{-4} mol·dm ⁻³ .	
	Is acid X a WEAK or a STRONG acid? Explain the answer by referring to the answer in QUESTION 4.3.2.	(3)
	4.1 4.2	 4.2 Carbonated water is an aqueous solution of carbonic acid, H₂CO₃, H₂CO₃(aq) ionises in two steps when it dissolves in water. 4.2.1 Write down the FORMULA of the conjugate base of H₂CO₃(aq). 4.2.2 Write down a balanced equation for the first step in the ionisation of carbonic acid. 4.2.3 The pH of a carbonic acid solution at 25 °C is 3,4. Calculate the hydroxide ion concentration in the solution. 4.3 X is a monoprotic acid. 4.3.1 State the meaning of the term monoprotic. 4.3.2 A sample of acid X is titrated with a standard sodium hydroxide solution using a suitable indicator. At the endpoint it is found that 25 cm³ of acid X is neutralised by 27,5 cm³ of the sodium hydroxide solution of acid X. 4.3.3 The concentration of H3O+ ions in the sample of acid X is 2,4 x 10⁻⁴ mol·dm⁻³. Is acid X a WEAK or a STRONG acid? Explain the answer by referring to the

(3)

(7)



	QUESTION 5		
	5.1	Two acids, HX and HY, of EQUAL CONCENTRATIONS are compared.	
		The pH of HX is 2,7 and the pH of HY is 0,7.	
		E 1.1 Define en esid in terres of the Lower Defected theory	
		5.1.1 Define an acid in terms of the Lowry-Brønsted theory.	
		5.1.2 Which acid, HX or HY, is STRONGER? Give a reason for the answer.	
		5.1.3 Acid HX ionises in water according to the following equation:	
		$HX(aq) + H_2O(\ell) \rightleftharpoons H_3O^+(aq) + X^-(aq)$	
		The K_a value for the reaction is 1,8 x 10 ⁻⁵ at 25 °C.	
		Is the concentration of the hydronium ions HIGHER THAN, LOWER THAN or EQUAL	
		TO the concentration of HX? Give a reason for the answer.	
	5.2	Learners add 150 cm ³ of a sodium hydroxide solution, NaOH, of unknown concentration to 200 cm ³ of a 0,03 mol·dm ⁻³ hydrochloric acid solution, HC ² , as illustrated below. They find that the pH of the final solution is 2. Assume that the volumes are additive.	



The balanced equation for the reaction is:

 $\mathsf{HC}\ell(\mathsf{aq}) + \mathsf{NaOH}(\mathsf{aq}) \rightarrow \mathsf{NaC}\ell(\mathsf{aq}) + \mathsf{H}_2\mathsf{O}(\ell)$

Calculate the:

5.2.1 Concentration of the H_3O^+ ions in the final solution

5.2.2 Initial concentration of the NaOH(aq)