

NAME:.....INDEX NO.....

SIGNATURE:..... P525/3

UNNASE PRACTICAL GUIDE

P525/3

CHEMISTRY PRACTICAL

3¼ HOURS

UACE MOCK EXAMINATIONS, 2023

CHEMISTRY PRACTICAL

· 3 HOURS AND 15 MINUTES

INSTRUCTIONS TO CANDIDATES.

- ❑ *All questions are compulsory*
- ❑ *Answers to be written in the spaces provided*
- ❑ *All your work must be in blue or black ink*
- ❑ *Any work done on pencil will not be marked*
- ❑ *You are not allowed to work with the apparatus for the paper and check whether you have all the chemicals and apparatus.*
- ❑ *All working must be clearly shown.*
- ❑ *Mathematical tables and silent non-programmable scientific calculators may be used.*

(O=16, Na= 23, S=32, Mn=55)

FOR EXAMINER'S USE ONLY		
Q1	30	
Q2	34	
Q3	16	
TOTAL	<u>80</u>	

X
30

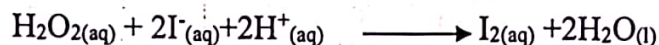
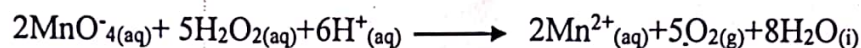
1. You are provided with the following.
- FA1: which contains 3.95g of an hydrous sodium thiosulphate, $\text{Na}_2\text{S}_2\text{O}_3$ in 500cm^3 of solution
 - FA2: Which is hydrogen peroxide solution
 - Solid T: which is a salt containing manganate (VII) ion.
 - 5% potassium iodide solution.
 - Starch solution

You are required to determine the;

- (i) Concentration of hydrogen peroxide in mol dm^{-3} of FA2
- (ii) Percentage of manganese in T.

Theory:

In acidic medium, hydrogen peroxide reacts with Manganes (VII) ions and iodide ions according to the following equations.



The iodine liberated reacts with thiosulphate ions according to the following equations.



PROCEDURES

PART A:

- (a) Using a measuring cylinder, transfer exactly 5.0cm^3 of FA2 into a 250cm^3 volumetric flask.
Make the solution up to the mark with distilled water. Label the solution FA3.
- (b) Pipette 10.0cm^3 of FA3 into a conical flask, add an equal volume of 1M sulphuric acid. Using a measuring cylinder followed by 10cm^3 of 5% potassium iodide solution. Warm the mixture to 50°C and titrate with FA1 from the burette until the solution is pale yellow.
Add starch indicator and continue the titration until the end point. Repeat the titration until you obtain consistent results.

Record your result in the table below.

Volume of pipette used $10.00 / 10.0 / 10$ \checkmark cm^3

($\frac{1}{2}$ marks)

Table I

Final burette reading (cm ³)	14.50	28.60	19.10
Initial burette reading (cm ³)	0.00	14.50	5.00
Volume of FA1 use (cm ³)	14.50	14.10	14.10

0.4 1/2

$$TR = 14.10 \pm 3$$

(4 1/2 marks)

Volumes of FA1 used for calculating average volume

14.10 and 14.10 ± 0.1 agree (1/2 marks)

Calculate the average volume of FA1 used.

(3 1/2 marks)

$$\frac{14.10 + 14.10}{2} = 14.10 \pm 0.1$$

+0.2 ✓
+0.3 ✓
+0.4 ✓
+0.5 ✓

Questions

(a) Calculate the number of moles of iodine that reacted with FA1 (2 1/2 marks)

$$R_{FA1} \text{ of } Na_2S_2O_3 = (2 \times 23) + (32 \times 2) + (3 \times 16) = 158 \checkmark$$

$$\text{Moles of } Na_2S_2O_3 = \left(\frac{3.95}{158} \right) = 0.025 \text{ moles}$$

$$500 \text{ cm}^3 \text{ of FA1 contain } 0.025 \text{ moles of } S_2O_3^{2-}$$

$$14.10 \text{ cm}^3 \text{ of FA1 contain } \left(\frac{0.025 \times 14.10}{500} \right) = 7.05 \times 10^{-4} \text{ moles}$$

$$2 \text{ moles of } S_2O_3^{2-} \text{ react with } 1 \text{ mole of } I_2$$

$$\text{Moles of } I_2 \text{ that reacted} = \frac{1}{2} \times 7.05 \times 10^{-4} = 3.525 \times 10^{-4} \text{ moles}$$

(b) Determine the concentration of FA2 in mol dm⁻³ (3 marks)

1 mole of I₂ is produced by 1 mole of H₂O₂ (FA3).

$$10 \text{ cm}^3 \text{ of FA3 contain } 3.525 \times 10^{-4} \text{ moles of } H_2O_2$$

$$250 \text{ cm}^3 \text{ of FA3 contain } \left(\frac{3.525 \times 10^{-4} \times 250}{10} \right) = 8.8125 \times 10^{-3} \text{ moles}$$

Dilution does not affect number of moles

$$5 \text{ cm}^3 \text{ of FA2 contain } 8.8125 \times 10^{-3} \text{ moles of } H_2O_2$$

$$1000 \text{ cm}^3 \text{ of FA2 contain } \left(\frac{8.8125 \times 10^{-3} \times 1000}{5} \right) \text{ moles}$$

$$= 1.7625 \text{ moles of } H_2O_2$$

PROCEDURE

PART B

(c) Weigh accurately about 0.8g of T and transfer it into a 250cm³ volumetric flask. Add about 100cm³ of 1 M sulphuric acid followed by 15cm³ of FA2. Make the solution upto the mark with distilled water and allow to stand for about five minutes. Label the solutions FA4.

(d) Pipette 10.0cm³ of FA4 into a conical flask, add an equal volume of 1M sulphuric acid using a measuring cylinder, followed by 10cm³ of 5% potassium iodide solution. Warm the mixture to 50°C and titrate with FA1 from the burette until the solution is pale yellow. Add starch indicator and continue the titration until the end point.

Repeat the titration until you obtain consistent results. Record your results in table 11 below:

RESULTS:

Mass of weighing bottle + T 2.80 ✓ g (½ mk)

Mass of empty weighing bottle 2.00 ✓ g (½ mk)

Mass of T used 0.80 ✓ g (½ mk) (2)

Volume of pipette used 10.00 / 10.10 / 10 ✓ cm³ (½ mk)

Table II

Final burette reading (cm ³)	23.70	46.20	27.50
Initial burette reading (cm ³)	1.00	23.70	5.00
Volume of FA1 used (cm ³)	22.70	22.50	22.50

TR = 22.50 ± 3 ✓ ✓ ✓

(4 ½ mks)

Volumes of FA1 used for calculating average volume

22.50 and 22.50 (agree ± 0.1) ✓ (½ mk)

∴ Calculate the average of FA1 used

$\frac{22.50 + 22.50}{2} = 22.50$

± 0.1 ✓
± 0.2 ✓
± 0.3 ✓
± 0.4 ✓
± 0.5 ✓

2 ½
(3 ½ mks)

Questions:

(a) Calculate the number of moles of

(i) Iodine that reacted with thiosulphate ions in FA1

$1\frac{1}{2}$ mark
(2½ mks)

500 cm³ of FA1 contain 0.025 moles of S₂O₃²⁻
 22.50 cm³ of FA1 contain $\left(\frac{0.025 \times 22.50}{500}\right) = 1.125 \times 10^{-3}$ mol
 2 moles of S₂O₃²⁻ react with 1 mole of I₂.
 Moles of I₂ ∴ = $\frac{1}{2} \times 1.125 \times 10^{-3} = 5.625 \times 10^{-4}$ moles of I₂

(ii) Excess hydrogen peroxide obtained in 250 cm³

$1\frac{1}{2}$ mark
(2½ mks)

1 mole of I₂ is produced by 1 mole of H₂O₂.
 10 cm³ of FA4 contain 5.625 × 10⁻⁴ moles of H₂O₂.
 ⇒ 250 cm³ of FA4 contain $\left(\frac{5.625 \times 10^{-4} \times 250}{10}\right)$
 = 0.0140625 moles of H₂O₂

(iii) Hydrogen peroxide that reacted with T

$1\frac{1}{2}$ marks
(2½ mks)

Originally
 1000 cm³ of FA2 contain 1.7625 moles of H₂O₂.
 15 cm³ of FA2 contain $\left(\frac{1.7625 \times 15}{1000}\right)$
 = 0.0264375 moles of H₂O₂
 ⇒ Moles of H₂O₂ reacted = 0.0264375 - 0.0140625
 = 0.012375 moles

(b) Determine the percentage of Manganese in T (04 mks)

5 moles of H₂O₂ reacted with 2 moles of MnO₄⁻
 0.012375 moles of H₂O₂ react with $\left(\frac{2}{5} \times 0.012375\right)$ (0.3 mark)
 = 0.00495 moles

1 mole of MnO_4^- contain 1 mole of Mn atoms
 0.00495 moles of MnO_4^- contain 0.00495 moles of Mn atoms
 Mass of Mn = $55 \times 0.00495 = 0.27225 \text{ g}$
 $\% \text{ of Mn in T} = \frac{0.2722}{0.8} \times 100 = 34.03\%$

2. You are provided with substance P which contains two cations and two anions. You are required to carry out the following tests to identify the ions present in P. Record your observations and deductions in the table below. Identify any gas(es) evolved. (32 marks)

TESTS	OBSERVATION	DEDUCTION
(a) Heat two spatula endfuls of P in a dry test tube strongly until there is no further change.	<ul style="list-style-type: none"> White solid Colourless Condensate (liquid) which turns white anhydrous CuSO_4 blue Colourless gas, turns blue litmus paper red forms white ppt with Calcium hydroxide solution Yellow residue when hot and white on cooling 	Hydrated salt or water of crystallization $\text{CO}_2 \therefore \text{CO}_3^{2-} / \text{HCO}_3^- / \text{C}_2\text{O}_4^{2-} / \text{CH}_3\text{COO}^-$ $\text{ZnO} \therefore \text{Zn}^{2+}$
(b) To two spatula endfuls of P, add 5 cm^3 of distilled water. Shake thoroughly and filter. Keep both the filtrate and residue. Divide the filtrate into three parts. (i) To the first part of the filtrate, add 4 drops of Lead (II) nitrate solution followed by dilute nitric acid.	<ul style="list-style-type: none"> Partially soluble White residue Colourless filtrate White ppt soluble in acid with effervescence 	Non transition cations in both residue and filtrate CO_3^{2-} or any one $\text{C}_2\text{O}_4^{2-}$ SO_3^{2-}
(ii) To the second part of the filtrate, add half a spatula endful of solid		

sodium hydrogen carbonate followed by 4 – 5 drops of aqueous iodine solution.	Efferescence of bubble of colourless gas turn lime water milky Brown solution turns colourless	CO_2 gas SO_3^{2-} present. (03)
(iii) Use the third part of the filtrate to carry out a test of your own choice to confirm one of the anions in P. Test: Add 2-3 drops of acidified potassium manganate (VII) solution or acidified $\text{K}_2\text{Cr}_2\text{O}_7$ solution	Purple solution turned colourless Orange solution to green	SO_3^{2-} Confirmed. (1/2)
(c) Wash the residue twice with water. Transfer it into a test and add dilute nitric acid dropwise until there is no further change (warm if necessary). Add dilute sodium hydroxide solution dropwise until in excess. Filter and keep both the filtrate and residue.	Efferescence of bubbles of a colourless gas turn lime water milky Colourless solution White ppt insoluble in excess White residue Colourless filtrate	CO_2 gas hence CO_3^{2-} Confirmed. Non transition cation Ba^{2+} , Ca^{2+} , Mg^{2+} $\text{Ba}^{2+}/\text{Ca}^{2+}/\text{Mg}^{2+}$ (0 1/2) $\text{Zn}^{2+}/\text{Al}^{3+}/\text{Pb}^{2+}/\text{Sn}^{2+}$
(d) Acidify the filtrate using dilute nitric acid. Divide the solution into three parts	White precipitate soluble in acid	$\text{Zn}^{2+}/\text{Al}^{3+}/\text{Pb}^{2+}$, $\text{Sn}^{2+}/\text{Sn}^{4+}$. (0/6)
(i) To the first part of the acidified filtrate add 3 -4 drops of dilute Sulphuric acid	No white ppt or No observable change	Pb^{2+} absent $\therefore \text{Al}^{3+}/\text{Zn}^{2+}/\text{Sn}^{2+}/\text{Sn}^{4+}$ (0/6)
(ii) To the second part of the acidified filtrate add dilute	White ppt soluble in excess to a colourless solution	Zn^{2+} present (0 1/2)

ammonia solution dropwise until in excess.		
(iii) Use the third part of the acidified filtrate to carry out a test of your own choice to confirm one of the cations in P. Test: Solid NH_4Cl ; $\text{Na}_2\text{HPO}_4(\text{aq})$ $\text{NH}_3(\text{aq})$ in excess	White ppt soluble in excess ammonia	Zn^{2+} Confirmed 02
(e) Dissolve the residue from part (c) in a minimum amount of dilute nitric acid. Divide the resultant solution into five parts.	Colourless solution	Ca^{2+} or Mg^{2+} or Ba^{2+} present 01
(i) To the first part of the acidic solution, add dilute sodium hydroxide solution dropwise until in excess.	White ppt insoluble in excess	Ca^{2+} or Mg^{2+} or Ba^{2+} present 01 1/2
(ii) To the second part of the acidic solution add ammonia solution dropwise until in excess.	White ppt insoluble	Ba^{2+} or Mg^{2+} present 01 1/2
(iii) To the third part of the acidic solution add 3 drops of potassium chromate (VI) solution followed by Ethanoic acid	No observable change or No yellow ppt	Ba^{2+} absent probably Mg^{2+} present 01 1/2

(iv) To the fourth part of the acidic solution, add excess sodium hydrogen carbonate solution and heat the mixture.	White ppt Forms on heating	Mg²⁺ <div>OK</div>
(v) Use the fifth part of the acidic solution to carry out a test of your own choice to confirm the second cation in P. Test: <div>Add $\text{NH}_4\text{Cl(s)} + \text{Na}_2\text{HPO}_4\text{(aq)}$ $+ \text{NH}_3\text{(aq)}$ in excess.</div>	<div>✓ accept</div> White ppt insoluble in excess ammonia.	Mg²⁺ Continued <div>02</div>

(f) Identify the

(i) cations in P

Zn^{2+} ✓

and

Mg^{2+} ✓

(ii) anions in P

CO_3^{2-} ✓

and

SO_3^{2-} ✓

3. You are provided with substance R which is an organic compound. You are requested to carry out the tests on R to determine the nature of R. Record your observations and deductions in the table below. X
16 (18 marks).

TESTS	OBSERVATIONS	DEDUCTIONS
(a) Burn a small amount of R on a spatula end or on a porcelain dish	Colourless liquid burns with yellow non sooty flame.	Aliphatic saturated Compound with a low Carbon Content. <div>2</div>
(b) To 1 cm ³ of R, add about 5 cm ³ of distilled water		

and shake. Divide the mixture into three parts. (i) To the first part add sodium carbonate solution	Miscible with water to a colourless solution. No effervescence	Polar compound of low molecular mass. Carboxylic acid $\frac{1}{2}$ absent
(ii) To the second part add Neutral iron (III) chloride solution.	No purple/violet colouration	Phenol absent $\frac{1}{2}$
(iii) To the third part add 5 drops of acidified Potassium dichromate (VI) solution and heat the mixture	No observable change	Non-reducing compound. Ketone or tertiary alcohol probably present $\frac{2}{2}$
(c) To 1cm ³ of R, add an equal volume of water, followed by about 4-5 drops of Brady's reagent.	Yellow ppt	Ketone present $\frac{2}{2}$
(d) To 1cm ³ of R, add tollens reagent and warm.	No silver mirror	Aldehyde absent \therefore Ketone present $\frac{2}{2}$
(e) To 1cm ³ of R, add about 3cm ³ of iodine solution followed by sodium hydroxide solution dropwise until the colour of iodine is discharged.	Pale yellow ppt	Ketone of form $\text{CH}_3\text{C}(=\text{O})\text{CH}_3$ present $\frac{1}{2}$

(f) Describe the nature of R.

Aliphatic ketone of form $\text{CH}_3\text{C}(=\text{O})\text{CH}_3$ $\frac{1}{2}$

END