

Candidate's Name : NAME

Signature: *[Signature]*

Random No.				Personal No.		
1	9	6	8	5	0	0

(Do not write your School / Centre Name or Number anywhere on this booklet.)

P525/3
CHEMISTRY
(Practical)
Nov./Dec. 2022
3¼ hours



UGANDA NATIONAL EXAMINATIONS BOARD

Uganda Advanced Certificate of Education

CHEMISTRY
(PRACTICAL)

Paper 3

3 hours 15 minutes

INSTRUCTIONS TO CANDIDATES:

Answer all questions. Use blue or black ink. Any work done in pencil will not be marked except drawings.

All your answers must be written in the spaces provided.

Mathematical tables and silent non-programmable scientific calculators may be used.

Reference books (i.e. text books, booklets on qualitative analysis etc.) should not be used.

You are not allowed to start working with the apparatus for the first 15 minutes. This time is to enable candidates read the question paper and make sure they have all the apparatus and chemicals that they may need.

For Examiners' Use Only			
Q.1	Q.2	Q.3	Total
30	30	20	80

1. You are provided with the following:

FA1, which is a solution of hydrochloric acid.

Metal X.

Substance Y, which is an oxide of X, with a formula XO.

You are required to determine the enthalpy change for the reduction of the oxide of X and comment on your answer.

PART I

Procedure

Weigh accurately about 1.2 g of X.

Using a measuring cylinder, transfer 100 cm³ of FA1 into a plastic beaker or cup. Read and record its initial temperature, in table 1.

Add at once the 1.2 g of X into FA1 in the plastic beaker or cup and at the same time, start the stop clock or watch.

Stir gently with the thermometer and record the temperature of the mixture after every half-minute in table 1, up to the fourth minute.

Results

Mass of weighing container + X = 34.7 g (1/2 mark)

Mass of weighing container alone = 33.5 g (1/2 mark)

Mass of X used = 1.2 g (1/2 mark)

Table 1

Time (minutes)	0	1/2	1	1 1/2	2	2 1/2	3	3 1/2	4
Temperature of solution (°C)	27.0	45.0	56.5	63.0	66.0	68.0	67.5	66.5	65.0

Trend [↑] Increases then decrease
Correct to 1 d.p

(4 1/2 marks)

PART II

Procedure

Weigh accurately about 2.0 g of Y.

Using a measuring cylinder, transfer 100 cm³ of FA1 into a plastic beaker or cup. Read and record its initial temperature, in table 2.

Add at once the 2.0 g of Y into FA1 in the plastic beaker or cup and at the same time, start the stop clock or watch.

Stir gently with the thermometer and record the temperature of the solution after every half-minute in table 2, up to the fourth minute.

Results

Mass of weighing container + Y = 35.7 ✓g (½ mark)

Mass of weighing container alone = 33.5 ✓g (½ mark)

Mass of Y used = 2.0 ✓g (½ mark)

Table 2

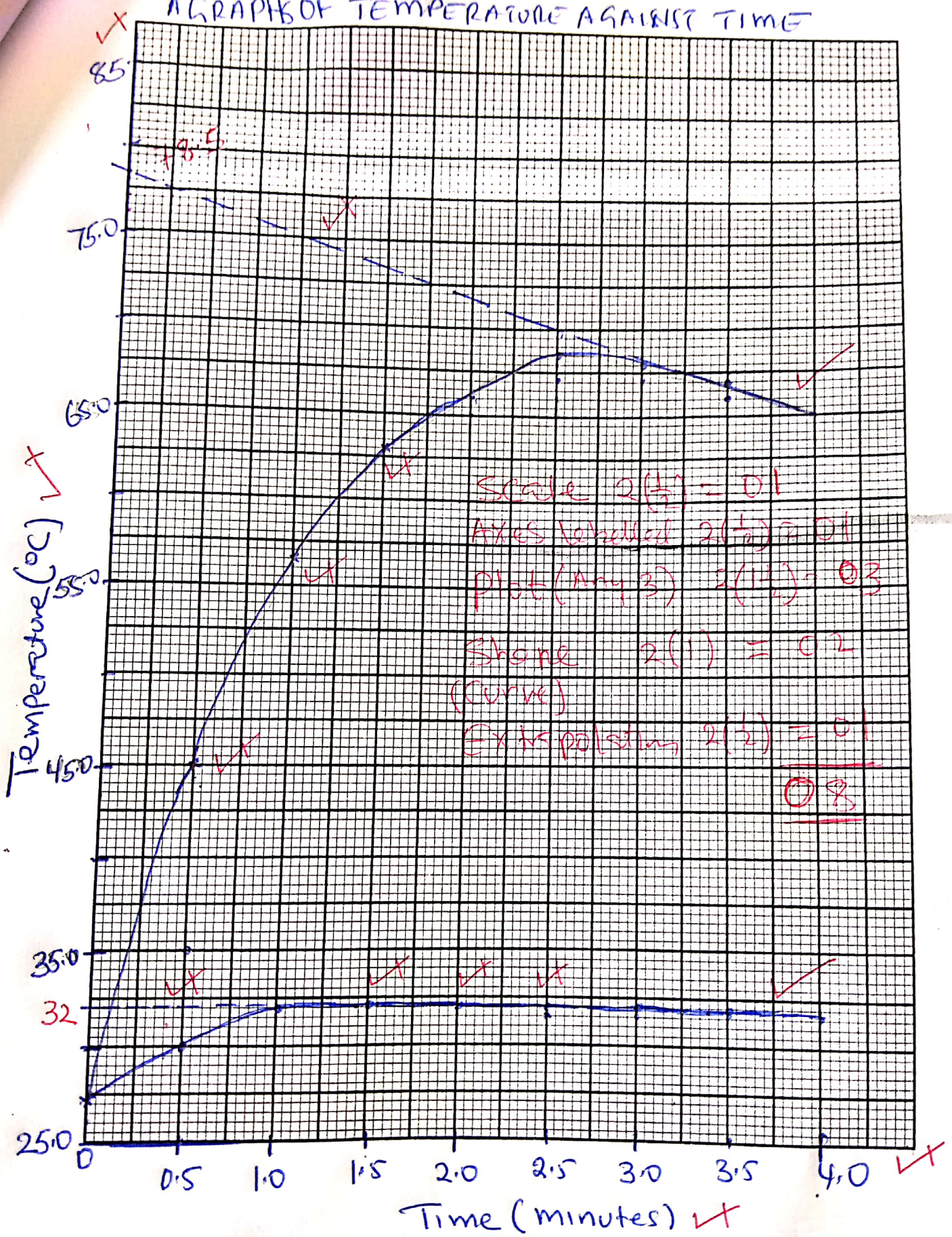
Time (minutes)	0	½	1	1½	2	2½	3	3½	4
Temperature of solution (°C)	27.0 ✓	30.0 ✓	32.0 ✓	32.0 ✓	32.0 ✓	32.0 ✓	32.0 ✓	32.0 ✓	31.5 ✓

(4½ marks)

- (a) (i) Plot on the same axes, graphs of temperature against time for results obtained in both **Part I** and **Part II**. (07 marks)
(Graph paper is provided on page 4.)

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A GRAPH OF TEMPERATURE AGAINST TIME



- (ii) From your graphs, determine the maximum temperature change for each reaction. (03 marks)

PART I

$$\text{Maximum temperature change} = 87.5 - 27 = 60.5^{\circ}\text{C}$$

PART II

$$\text{Maximum temperature change} = 32.0 - 27 = 5^{\circ}\text{C}$$

must be from the graph,
Ref if extra notation is wrong.

(02)

- (iii) Calculate the heat of reaction in Part I and Part II.

[Specific heat capacity of the solution = $4.2 \text{ J g}^{-1} \text{ K}^{-1}$ and its density = 1 g cm^{-3} in each case, equation of the reactions of X and Y are as follows:



Part I:

(01 mark)

$$\begin{aligned} \text{Heat Change} &= \text{mass} \times \text{specific heat capacity} \times \text{temperature change} \\ &= (100 \times 1) \times 4.2 \times 60.5 \end{aligned}$$

$$\begin{aligned} \text{Heat of reaction} &= -25410 \text{ Joules} \\ &= -25.41 \text{ KJ} \end{aligned}$$

Ref. of sign and unit missing

Part II:

(01 mark)

$$\begin{aligned} \text{Heat Change} &= \text{mass} \times \text{specific heat capacity} \times \text{temp change} \\ &= (100 \times 1) \times 4.2 \times 5 \end{aligned}$$

$$\begin{aligned} \text{Heat of reaction} &= -2100 \text{ Joules} \\ &= -2.1 \text{ KJ} \end{aligned}$$

Ref. of sign and unit missing

(04)

(iv) Calculate the molar heat of reactions in Part I and Part II.

(X = 24, O = 16)

(03 marks)

PART I

1.2g of X evolves 25.41 KJ ✓

24g of X evolves $\frac{25.41 \times 24}{1.2} = 508.2 \text{ KJ mol}^{-1}$ ✓ (1½)

molar heat of reaction = $-508.2 \text{ KJ mol}^{-1}$ ✓

Ref if sign and unit are missing.

PART II

molar mass of XO = $24 + 16 = 40 \text{ g}$ ✓

2.0g of XO evolves 2.1 KJ ✓

40g of XO evolves $\frac{2.1 \times 40}{2.0} = 42 \text{ KJ mol}^{-1}$ ✓ (1½)

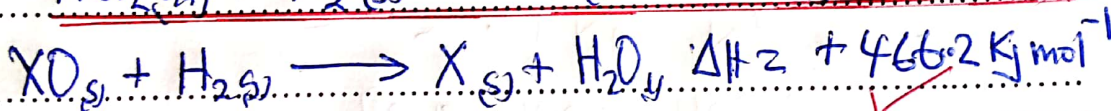
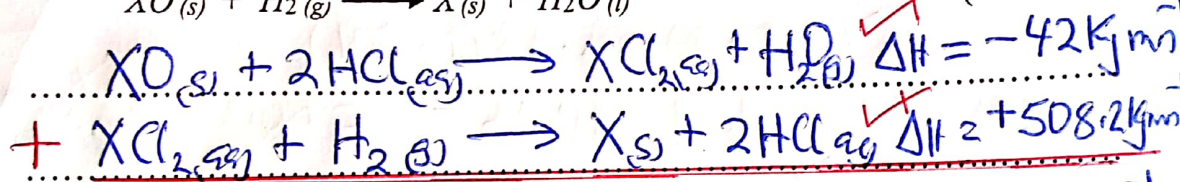
molar heat of reaction = -42 KJ mol^{-1} ✓

Ref if sign and unit are missing.

(b) Determine the heat energy change for the reaction.



(02 marks)



(02)

(c) Comment on your answer in (b).

(01 mark)

The reduction of XO to X is not feasible because the enthalpy of reaction is endothermic ✓

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T = 30

2. You are provided with substance **D** which contains **two** cations and **two** anions. You are required to carry out the tests in table 3 to identify the cations and anions in **D**. Identify any gas(es) evolved. Record your observations and deductions in the table.

Table 3

(30 marks)

TESTS	OBSERVATIONS	DEDUCTIONS
(a) To two spatula end-fuls of D in a test tube, add dilute nitric acid until there is no further change.	Green powdered solid dissolves to form a green solution with effervescence of a colourless gas that turns lime water milky, blue litmus red	Cu^{2+} , Ni^{2+} ✓ Fe^{2+} or Cr^{3+} ✓ Suspected ✓ CO_2 evolved ✓ CO_3^{2-} or HCO_3^- ✓ Present ✓
(b) To two spatula end-fuls of D in a test tube, add about 5 cm^3 of distilled water, shake well and filter. Keep both the filtrate and the residue.	Partly soluble Green residue Colourless filtrate	Cu^{2+} , Ni^{2+} , Fe^{2+} ✓ Cr^{3+} suspected ✓ non coloured metal ions present e.g. Al^{3+} , Pb^{2+} , Zn^{2+} ✓
(c) Divide the filtrate into three parts. (i) To the first part, add silver nitrate solution followed by dilute nitric acid.	Yellow precipitate soluble in dilute nitric acid	PO_4^{3-} ✓ Suspected ✓
(ii) To the second part, add aqueous iron(II) chloride solution.	Pale yellow precipitate	PO_4^{3-} ✓ suspected ✓

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TESTS	OBSERVATIONS	DEDUCTIONS
(iii) Use the third part of the filtrate to carry out a test of your own choice to confirm one of the anions in D. Test: To the third part concentrated nitric acid was added followed by Ammonium molybdate and dilute aqueous ammonia solution	Yellow precipitate ✓ Soluble in aqueous ammonia ✓	PO_4^{3-} ✓ Confirmed present $2\frac{1}{2}$
(d) Wash the residue twice with water. Transfer it into a test tube, add dilute nitric acid and warm. Add dilute sodium hydroxide solution drop-wise until in excess. Filter and keep both the filtrate and the residue.	Dissolves to form a green solution with effervescence of colourless gas that turns blue litmus red and turns lime water milky. Green ppt insoluble in excess Green residue Colourless filtrate	Cu^{2+}, Ni^{2+}, Fe^{2+} or Cr^{3+} suspected CO_2 evolved CO_3^{2-} Confirmed Ni^{2+} or Fe^{2+} (05) Ni^{2+} or Fe^{2+} Al^{3+}, Pb^{2+}, Zn^{2+}, Sn^{2+}
(e) To the filtrate from part (d), add dilute nitric acid drop-wise until the solution is just acidic. Divide the solution into three parts.	White precipitate ✓ Soluble in dilute nitric acid	Al^{3+}, Pb^{2+} ✓ Zn^{2+}, Sn^{2+} (01) Sn^{4+} suspected

$8\frac{1}{2}$

TESTS	OBSERVATIONS	DEDUCTIONS
(i) To the first part of the acidified filtrate, add dilute sodium hydroxide solution drop-wise until in excess.	White precipitate ✓ Soluble ✓	Al^{3+} , Pb^{2+} , Zn^{2+} ✓ Sn^{2+} or Sn^{4+} ✓ Suspected ^{Any one} (1½)
(ii) To the second part of the acidified filtrate, add dilute ammonia solution drop-wise until in excess.	White precipitate ✓ Soluble ✓	Zn^{2+} ✓ Present (1½)
(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 D. Test: To the third part solid Ammonium chloride was added followed by disodium hydrogen phosphate solution and excess ammonia solution ✓	White precipitate ✓ Soluble ✓ excess ammonia	Zn^{2+} ✓ Confirmed (2½)
(f) Dissolve the residue in part (d) in a minimum amount of dilute nitric acid. Divide the resultant solution into three parts.	Green residue dissolves to form a green solution ✓	Ni^{2+} or Fe^{2+} ✓ Suspected present (01)

6½

TESTS	OBSERVATIONS	DEDUCTIONS
(i) To the first part, add dilute sodium hydroxide solution drop-wise until in excess.	Green precipitate Insoluble in excess	Ni^{2+} ✓ Fe^{2+} suspected present (1½)
(ii) To the second part, add dilute ammonia solution drop-wise until in excess.	Green precipitate Soluble to form a pale blue solution	Ni^{2+} ✓ present (1½)
(iii) Use the third part to carry out a test of your own choice to confirm the second cation in D. Test: excess To the third Ammonia solution was added followed by dimethylglyoxime solution ✓	Red / bright pink precipitate	Ni^{2+} ✓ Confirmed present (02)

(g) (i) The cations in D are Zn^{2+} (e (iii)) and Ni^{2+} (f (iii)) (02)
(ii) The anions in D are CO_3^{2-} (d) and PO_4^{3-} c (iii)

(07)

T = 30

3. You are provided with substance L, which is an organic compound. You are required to carry out the tests in table 4 to determine the nature of L. Record your observations and deductions in the table.

Table 4

(20 marks)

TESTS	OBSERVATIONS	DEDUCTIONS	
(a) Burn a small amount of L on a spatula-end or in porcelain dish.	A colourless liquid burns with a yellow sooty flame ✓	Aromatic compound ✓ Aliphatic compound with high carbon content ✓	02
(b) To about 1 cm ³ of L, add about 2 cm ³ of distilled water. Shake the mixture and test with litmus. Divide the mixture into four parts. (i) To the first part, add 2-3 drops of 2, 4 - dinitrophenyl hydrazine solution.	Miscible to form a colourless solution ✓ No effect on both blue and red litmus paper ✓	Polar ✓ Aliphatic compound ✓ Neutral compound present ✓	2½
(ii) To the second part, add a half a spatula endful of solid sodium hydrogencarbonate.	no observable change / no yellow precipitate formed ✓	Carbonyl compound absent ✓	02
(iii) To the third part, add 2-3 drops of neutral iron(III) chloride solution.	no observable change / no purple colouration formed ✓	Carboxylic acid ✓ absent ✓	02
(iv) To the fourth part, add 2-3 drops of acidified potassium dichromate solution and heat.	no observable change / orange solution persisted on heating ✓	Phenol ✓ absent ✓ Reducing agent absent ✓ Primary alcohol ✓ Secondary alcohol absent ✓	02

TESTS	OBSERVATIONS	DEDUCTIONS
(c) To 1 cm ³ of L add about an equal volume of ethanoic acid, followed by about 2-3 drops of concentrated sulphuric acid and heat the mixture. Pour it into a beaker of cold water and allow to stand.	Sweet fruity Smell ✓	ester ✓ formed Tertiary ✓ alcohol ✓ present 02
(d) To 1 cm ³ of L, add 2-3 drops of concentrated sulphuric acid and heat. Pass the vapour evolving into a test tube containing acidified potassium manganate(VII) solution.	White fumes / vapour turns acidified potassium manganate (VII) solution colourless ✓	Tertiary ✓ alcohol ✓ dehydrated to form an alkene ✓ 02
(e) To 1 cm ³ of L, add 4-5 drops of Lucas reagent.	Immediate cloudy solution formed ✓	Tertiary ✓ alcohol ✓ Confirmed present 02

(f) Describe the nature of L.

Aliphatic tertiary alcohol with high Carbon Content (1½)

7½

T = 20