



Rewarding Learning

ADVANCED
General Certificate of Education
2010

Centre Number

71

Candidate Number

Chemistry
Assessment Unit A2 2
assessing

Analytical, Transition Metals, Electrochemistry
and Further Organic Chemistry
[AC222]



AC222

TUESDAY 1 JUNE, AFTERNOON

TIME

2 hours.

INSTRUCTIONS TO CANDIDATES

Write your Centre Number and Candidate Number in the spaces provided at the top of this page.

Answer **all seventeen** questions.

Answer **all ten** questions in **Section A**. Record your answers by marking the appropriate letter on the answer sheet provided. Use only the spaces numbered 1 to 10. Keep in sequence when answering.

Answer **all seven** questions in **Section B**. Write your answers in the spaces provided in this question paper.

INFORMATION FOR CANDIDATES

The total mark for this paper is 120.

Quality of written communication will be assessed in question **16(b)**. In Section A all questions carry equal marks, i.e. **two** marks for each question.

In Section B the figures in brackets printed down the right-hand side of pages indicate the marks awarded to each question or part question.

A Periodic Table of Elements (including some data) is provided.

For Examiner's
use only

Question Number	Marks
Section A	
1–10	
Section B	
11	
12	
13	
14	
15	
16	
17	

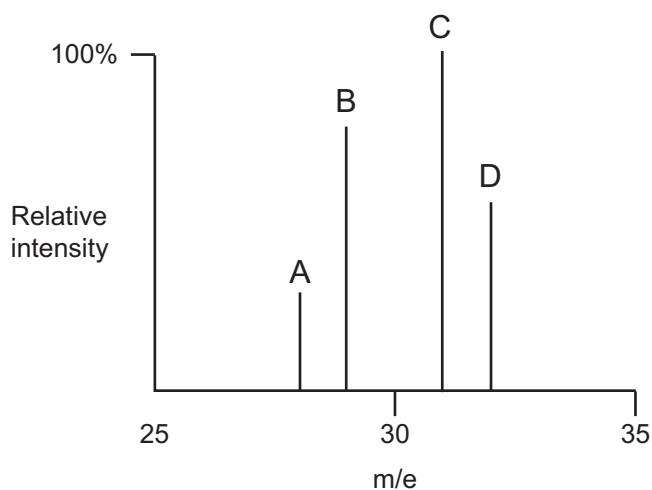
Total
Marks

Section A

For each of the following questions only **one** of the lettered responses (A–D) is correct.

Select the correct response in each case and mark its code letter by connecting the dots as illustrated on the answer sheet.

- 1 The mass spectrum of methanol is shown below.
Which one of the following is the base peak?



- 2 Which one of the following represents the electronic configuration for the Fe^{3+} ion?

- A $1s^2 2s^2 2p^6 3s^2 3p^6 3d^5$
- B $1s^2 2s^2 2p^6 3s^2 3p^6 3d^3 4s^2$
- C $1s^2 2s^2 2p^6 3s^2 3p^6 3d^4 4s^1$
- D $1s^2 2s^2 2p^6 3s^2 3p^6 3d^6 4s^2$

- 3 Which one of the following formulae **cannot** be determined by colorimetry?

- A $[\text{Cu}(\text{NH}_3)_4(\text{H}_2\text{O})_2]^{2+}$
- B $[\text{Fe}(\text{SCN})(\text{H}_2\text{O})_5]^{2+}$
- C $[\text{Ni}(\text{NH}_3)_4(\text{H}_2\text{O})_2]^{2+}$
- D $[\text{Zn}(\text{NH}_3)_4(\text{H}_2\text{O})_2]^{2+}$

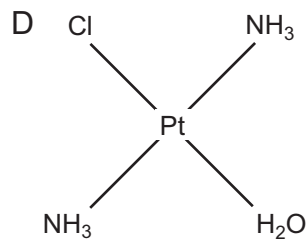
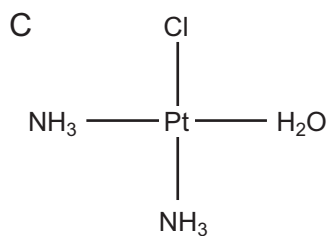
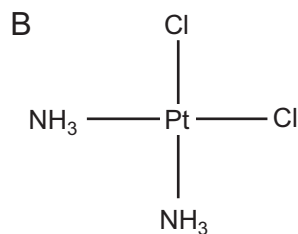
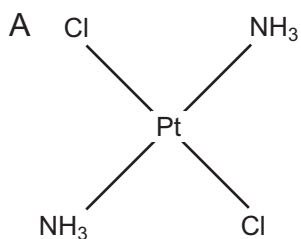
4 The standard electrode potentials for a series of redox equations are listed below.

	$E^{\ominus}/V$
$Mn^{2+}(aq) + 2e^{-} \rightleftharpoons Mn(s)$	-1.19
$Fe^{2+}(aq) + 2e^{-} \rightleftharpoons Fe(s)$	-0.44
$Ni^{2+}(aq) + 2e^{-} \rightleftharpoons Ni(s)$	-0.25
$I_2(s) + 2e^{-} \rightleftharpoons 2I^{-}(aq)$	+0.54
$Fe^{3+}(aq) + e^{-} \rightleftharpoons Fe^{2+}(aq)$	+0.77
$Ag^{+}(aq) + e^{-} \rightleftharpoons Ag(s)$	+0.80

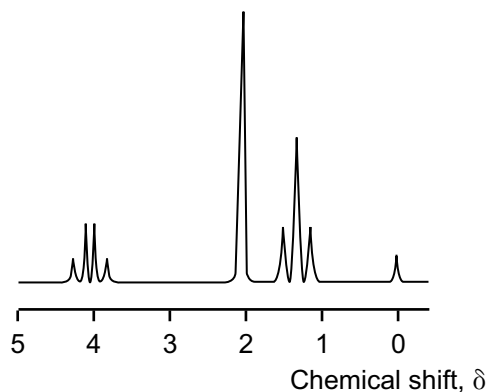
Which one of the elements listed will reduce $Fe^{3+}(aq)$ to $Fe^{2+}(aq)$, but not to $Fe(s)$?

- A Iodine
- B Manganese
- C Nickel
- D Silver

5 Which one of the following represents the structure of cisplatin?



6 The nmr spectrum of a compound **X** is shown below.



Which one of the following is **X**?

- A $\text{CH}_3\text{CH}_2\text{CH}_2\text{COOH}$
- B $\text{CH}_3\text{CH}_2\text{COOCH}_2\text{CH}_3$
- C $\text{CH}_3\text{COOCH}_2\text{CH}_3$
- D $\text{CH}_3\text{CH}_2\text{COOH}$

7 A green solid was dissolved in water and the resulting solution divided into two portions. Sodium hydroxide solution was added to one of the portions and ammonia solution to the other. The results are summarised in the table below.

Solution added	Few drops of solution	Excess solution
Sodium hydroxide solution	Green precipitate	No effect
Ammonia solution	Green precipitate	Dissolves to form a blue solution

Which one of the following ions was present in the green solid?

- A Cr^{3+}
- B Cu^{2+}
- C Fe^{2+}
- D Ni^{2+}

8 Which one of the following is the colour of chrome alum crystals?

- A Black
- B Green
- C Orange
- D Violet

- 9 25.0 cm³ of hydrogen peroxide solution were added to excess acidified potassium iodide solution and the resulting solution made up to 500 cm³.
25.0 cm³ of the diluted solution reacted with 36.4 cm³ of sodium thiosulphate solution of concentration 0.10 mol dm⁻³.
Which one of the following is the concentration of the undiluted hydrogen peroxide?

- A 0.07 mol dm⁻³
- B 0.15 mol dm⁻³
- C 1.46 mol dm⁻³
- D 2.91 mol dm⁻³

- 10 Which one of the following pairs of monomers will **not** combine to form a polymer?

- A HOOC(CH₂)₄COOH and H₂N(CH₂)₆NH₂
- B H₂N(CH₂)₆NH₂ and H₂N(CH₂)₆NH₂
- C CH₂CH₂ and CH₂CH₂
- D HOOC(CH₂)₄COOH and HO(CH₂)₂OH

Section B

Answer **all seven** questions in this section

- 11 (a)** Complete the table below by giving the name of the indicator and the colour change at the end point.

Titration	Indicator	Colour change	
		from	to
Edta added to magnesium ions			
Thiosulphate ions added to iodine			

[6]

- (b)** Some iron tablets, used to treat anaemia, contain iron(II) fumarate ($\text{FeC}_4\text{H}_2\text{O}_4$).

Five of these iron tablets were dissolved in dilute sulphuric acid and the solution made up to 250 cm^3 with distilled water. On titration 25.0 cm^3 of this solution reacted with 18.7 cm^3 of 0.01 mol dm^{-3} acidified potassium manganate(VII) solution.

- (i)** Write the equation for the reaction of iron(II) ions with acidified manganate(VII) ions.

_____ [2]

- (ii)** What is the colour change at the end point of the titration?

From _____ to _____ [2]

- (iii)** Calculate the mass of iron(II) fumarate in each tablet.

 _____ [4]

Examiner Only

Marks Remark

12 (a) Polythene is a typical addition polymer.

(i) State the conditions necessary for the formation of HD polythene.

Temperature: _____

Pressure: _____

Catalyst: _____ [3]

(ii) Making reference to their structures, explain the difference in flexibility between HD and LD polythene.

[2]

(b) Polyethylene terephthalate is a polyester.

(i) Draw the repeating unit for polyethylene terephthalate.

[2]

(ii) Give **one** use for polyethylene terephthalate.

[1]

Examiner Only	
Marks	Remarks

- (c) Explain in terms of entropy why dimethylglyoxime displaces the water ligands in the hydrated nickel(II) ion to form the nickel(II) dimethylglyoxime complex.

[2]

Examiner Only	
Marks	Remark

(d) The mechanism for nitrating toluene is similar to that for nitrating benzene.

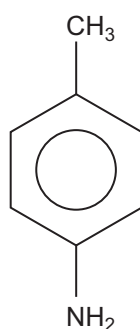
(i) What name is given to this mechanism?

 [2]

(ii) Draw the mechanism for the mononitration of toluene.

[2]

(e) 4-nitrotoluene can be converted to toluidine.



Toluidine

(i) Name the reagents which could be used to convert 4-nitrotoluene to toluidine.

[2]

(ii) A salt of toluidine is formed during the reduction. How can toluidine be liberated from this salt?

[1]

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(Questions continue overleaf)

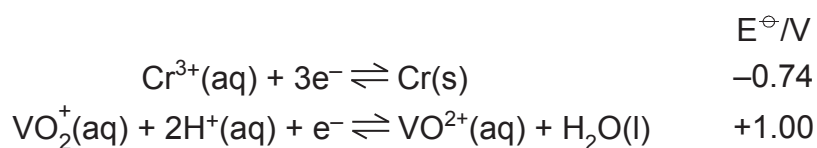
(c) An acidified solution of ammonium vanadate(V) undergoes a series of reductions when it is stirred with zinc.

(i) Complete the table below, giving the colour of $\text{VO}_2^+(\text{aq})$ and $\text{V}^{3+}(\text{aq})$ formed at the different stages of the reduction.

Ion	Colour
$\text{VO}_2^+(\text{aq})$	Yellow
$\text{VO}^{2+}(\text{aq})$	
$\text{V}^{3+}(\text{aq})$	
$\text{V}^{2+}(\text{aq})$	Violet

[2]

(ii) Chromium will also reduce ammonium vanadate(V).



Write an equation for the reaction between chromium and $\text{VO}_2^+(\text{aq})$ ions and calculate the e.m.f. of the reaction.

e.m.f. _____ [3]

Examiner Only

Marks Remark

17 Proteins are formed from amino acids.

- (ii) Write an equation for the reaction between aspartic acid and asparagine to form a dipeptide.

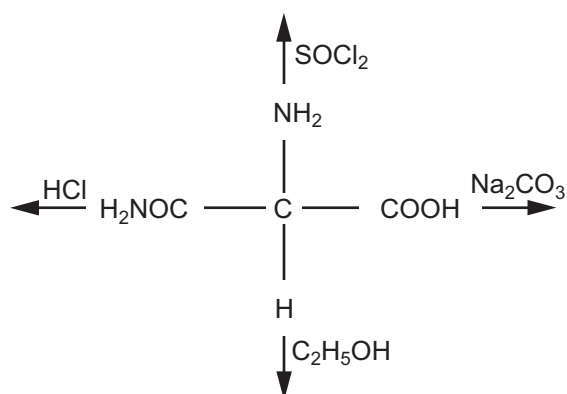
_____ [2]

- (iii) Amino acids dissolve in water to form a dipolar ion (zwitterion).

Draw the structure of the dipolar ion formed by asparagine.

[1]

- (iv) Complete the following flow diagram to show the structure of the organic product formed.



[4]

Examiner Only	
Marks	Remark

**Published Mark Schemes for
GCE A2 Chemistry**

Summer 2010

NORTHERN IRELAND GENERAL CERTIFICATE OF SECONDARY EDUCATION (GCSE) AND NORTHERN IRELAND GENERAL CERTIFICATE OF EDUCATION (GCE)

MARK SCHEMES (2010)

Foreword

Introduction

Mark Schemes are published to assist teachers and students in their preparation for examinations. Through the mark schemes teachers and students will be able to see what examiners are looking for in response to questions and exactly where the marks have been awarded. The publishing of the mark schemes may help to show that examiners are not concerned about finding out what a student does not know but rather with rewarding students for what they do know.

The Purpose of Mark Schemes

Examination papers are set and revised by teams of examiners and revisers appointed by the Council. The teams of examiners and revisers include experienced teachers who are familiar with the level and standards expected of 16- and 18-year-old students in schools and colleges. The job of the examiners is to set the questions and the mark schemes; and the job of the revisers is to review the questions and mark schemes commenting on a large range of issues about which they must be satisfied before the question papers and mark schemes are finalised.

The questions and the mark schemes are developed in association with each other so that the issues of differentiation and positive achievement can be addressed right from the start. Mark schemes therefore are regarded as a part of an integral process which begins with the setting of questions and ends with the marking of the examination.

The main purpose of the mark scheme is to provide a uniform basis for the marking process so that all the markers are following exactly the same instructions and making the same judgements in so far as this is possible. Before marking begins a standardising meeting is held where all the markers are briefed using the mark scheme and samples of the students' work in the form of scripts. Consideration is also given at this stage to any comments on the operational papers received from teachers and their organisations. During this meeting, and up to and including the end of the marking, there is provision for amendments to be made to the mark scheme. What is published represents this final form of the mark scheme.

It is important to recognise that in some cases there may well be other correct responses which are equally acceptable to those published: the mark scheme can only cover those responses which emerged in the examination. There may also be instances where certain judgements may have to be left to the experience of the examiner, for example, where there is no absolute correct response – all teachers will be familiar with making such judgements.

The Council hopes that the mark schemes will be viewed and used in a constructive way as a further support to the teaching and learning processes.

CONTENTS

	Page
A2 1	1
A2 2	11
A2 3 Practical Examination 1	19
A2 3 Practical Examination 2	25

New
Specification



Rewarding Learning

ADVANCED
General Certificate of Education
2010

Chemistry

Assessment Unit A2 1

assessing

Periodic Trends and Further Organic,
Physical and Inorganic Chemistry

[AC212]

FRIDAY 21 MAY, AFTERNOON

**MARK
SCHEME**

- Section A**
- 1** C
 - 2** B
 - 3** C
 - 4** C
 - 5** A
 - 6** B
 - 7** D
 - 8** C
 - 9** C
 - 10** B

[2] for each correct answer

[20]

Section A

**AVAILABLE
MARKS**

20

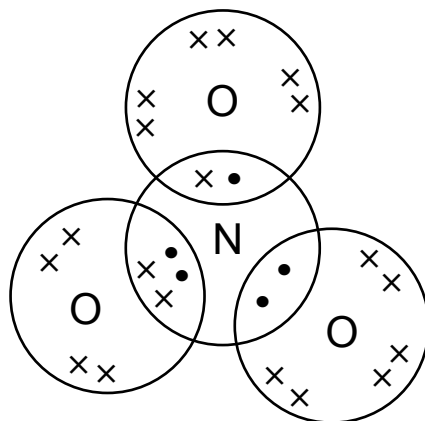
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Section B

11 (a) (i) $\text{NH}_3 + \text{HNO}_3 \rightarrow \text{NH}_4\text{NO}_3$ [1]

(ii) acid [1]
salt of strong acid and weak base [1] [2]

(iii)



each error [−1] [2]

(b) (i) $\text{NH}_4\text{NO}_3 \rightarrow \text{N}_2\text{O} + 2\text{H}_2\text{O}$ [1]

(ii) ΔG always negative [1]

(c) (i) volume / smell / storage / composition [1]

(ii) mass $\text{NH}_4\text{NO}_3 = 80$

$$\% \text{N} = (2 \times 14 / 80) \times 100$$

$$= 35\%$$

award [2] directly for correct answer, each error [−1] [2]

(d) (i) leaching of fertilisers / detergents [1]

(ii) $= 0.05 \times 10^{-3} \text{g per cm}^3$

$$= 0.05 \text{ g per dm}^3$$

$$= 0.05/62$$

$$= 8.06 \times 10^{-4} \text{ M}$$

each error [−1], carry error through [3]

(iii) iron +2 to +3 oxidation [1]
nitrogen reduction +3 to +2 [1] [2]

12 (a) $pK_a = 4.87$

$$K_a = 1.35 \times 10^{-5} \text{ mol dm}^{-3}$$

$$K_a = [\text{CH}_3\text{CH}_2\text{COO}^-][\text{H}^+] / [\text{CH}_3\text{CH}_2\text{COOH}]$$

$$1.35 \times 10^{-5} = [\text{H}^+]^2 / 0.05$$

$$[\text{H}^+]^2 = 6.74 \times 10^{-7}$$

$$[\text{H}^+] = 8.2 \times 10^{-4}$$

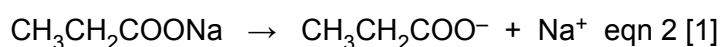
$$\text{pH} = -\log(8.2 \times 10^{-4})$$

$$\text{pH} = 3.09 \text{ or } 3.085 \text{ or } 3.1$$

each error [-1], carry error through

[4]

(b) $\text{CH}_3\text{CH}_2\text{COOH} \rightleftharpoons \text{CH}_3\text{CH}_2\text{COO}^- + \text{H}^+$ eqn 1 [1]



add acid

pushes equilibrium 1 to lhs / H^+ react with $\text{CH}_3\text{CH}_2\text{COO}^-$ [1]

salt acts as a source of $\text{CH}_3\text{CH}_2\text{COO}^-$ [1]

to a maximum of 4

[4]

(c) (i) $\text{CH}_3\text{CH}_2\text{COOH} + \text{CH}_3\text{OH} \rightleftharpoons \text{CH}_3\text{CH}_2\text{COOCH}_3 + \text{H}_2\text{O}$

[1]

(ii) catalyst / increases yield / pushes eqn to rhs / absorbs water [1]

[1]

(iii) propanoic acid has hydrogen bonding (between O-H)

the ester has no hydrogen bonding [1]

comment on relative strengths of bonding [1]

[2]

(iv) higher yield / not reversible [1]

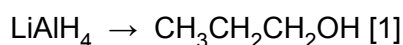
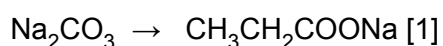
faster [1]

other product gaseous [1]

any **two**

[2]

(d) $\text{NH}_3 \rightarrow \text{C}_2\text{H}_5\text{COONH}_4$ [1]

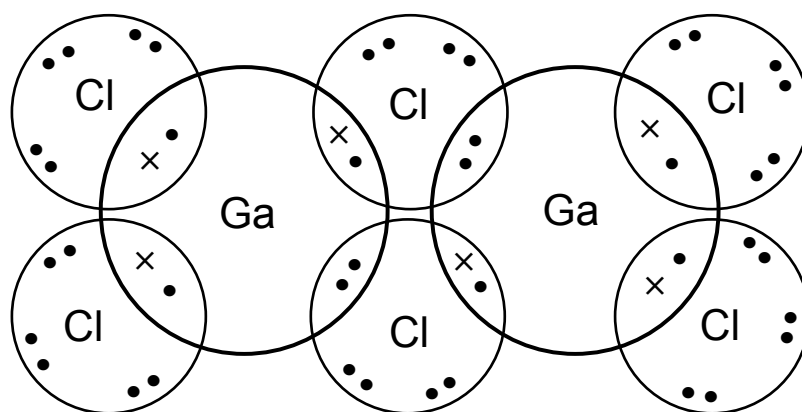


[3]

17

- 13 (a) (i) sulphur [1]
- (ii) magnesium oxide [1]
- (iii) PCl_5 or $\text{PCl}_4^+\text{PCl}_6^-$ [1]
- (iv) basic [1]
acidic [1]
acidic [1] [3]
- (b) (i) $\text{Ga}_2\text{O}_3 + 6\text{HCl} \rightarrow 2\text{GaCl}_3 + 3\text{H}_2\text{O}$ [2]
unbalanced [-1]

(ii)



[2]

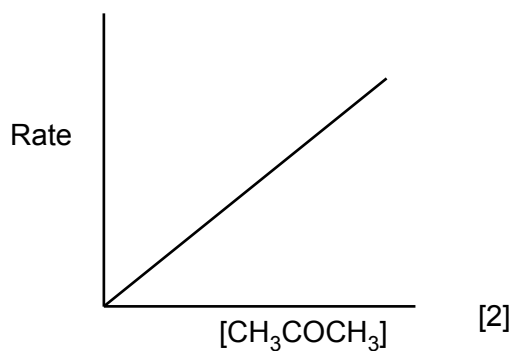
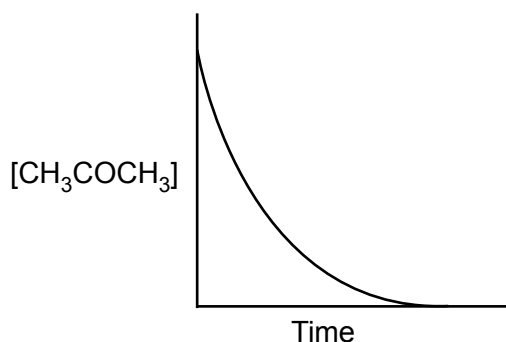
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14 (a) $2\text{KI} + 3\text{H}_2\text{SO}_4 + \text{MnO}_2 \rightarrow \text{I}_2 + 2\text{KHSO}_4 + \text{MnSO}_4 + 2\text{H}_2\text{O}$ [2]
unbalanced [-1]

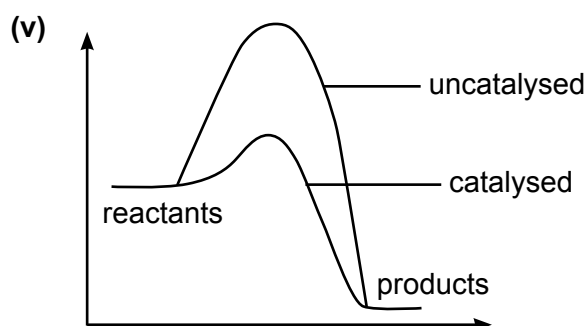
(b) (i) colorimeter [1] [1]

(ii) rate constant [1]
 $\text{dm}^3\text{mol}^{-1}\text{s}^{-1}$ [1] [2]

(iii)



(iv) $\times 4$ [1]



missing label [-1] [2]

(vi) increases k (as E_{act} is smaller) [1]

- (c) (i) known mass of oil [1]
 add Wij's solution and place in dark and add KI(aq) [1]
 prepare blank [1]
 titrate with standard sodium thiosulphate [1]
 starch indicator [1] [5]

Quality of written communication: [2]

2 marks The candidate expresses ideas clearly and fluently through well-linked sentences and paragraphs. Arguments are generally relevant and well-structured. There are few errors of grammar, punctuation and spelling.

1 mark The candidate expresses ideas clearly, if not always fluently. Arguments may sometimes stray from the point. There may be some errors of grammar, punctuation and spelling, but not such as to suggest a weakness in these areas.

0 marks The candidate expresses ideas satisfactorily, but without precision. Arguments may be of doubtful relevance or obscurely presented. Errors in grammar, punctuation and spelling are sufficiently intrusive to disrupt the understanding of the passage.

- (ii) saturated / few C=C double bonds [1]

- (d) (i) $\text{H}_2 + \text{I}_2 \rightarrow 2\text{HI}$
 1 1 0
 0.25 0.25 1.5
 $K_c = [\text{HI}]^2 / [\text{H}_2] [\text{I}_2]$
 $= 1.5^2 / 0.25 \times 0.25$
 $= 36$ [2]
 no units [1] [3]

- (ii) volumes would cancel [1]

- (iii) 0.0277 or $\frac{1}{36}$ (carry error through from part (i)) [1]

- (e) $\text{Pb}(\text{NO}_3)_2 + 2\text{KI} \rightarrow \text{PbI}_2 + 2\text{KNO}_3$
 moles KI = $50 \times 0.4 / 1000 = 0.02$ gives 0.01 mol PbI_2
 moles PbI_2 formed = $3.8 / 461 = 0.00824$
 $\% \text{ yield} = (0.00824 / 0.01) \times 100$
 $= 82.4\%$
 award [3] for correct answer
 each error [-1], carry error through [3]

- 15 (a)** +7 [1]
+3 [1] [2]
- (b) (i)** $1s^2 2s^2 2p^6$ [1]
 $1s^2 2s^2 2p^6 3s^2 3p^6$ [1] [2]
- (ii)** X = first ionisation energy of Mg [1]
Y = (twice) electron affinity for chlorine [1]
Z = (standard) enthalpy of formation of $MgCl_2$ [1] [3]
- (iii)** $+148 + 738 + 1451 + 242 - 696 = -642 + \Delta H_{latt}$
 $\Delta H_{latt} = +2525 \text{ (kJ mol}^{-1}\text{)}$
each error [-1] [2]
- (c) (i)** $CH_3CH_2CH_2COOH + SOCl_2 \rightarrow CH_3CH_2CH_2COCl + SO_2 + HCl$ [1]
 $CH_3CH_2CH_2COOH + PCl_5 \rightarrow CH_3CH_2CH_2COCl + POCl_3 + HCl$ [1] [2]
- (ii)** more pure [1]
other products are gaseous [1] [2]
- (d) (i)** (increase dissociation)
increase temperature moves eqn to rhs [1]
to absorb thermal energy / endothermic direction [1] [2]
- (ii)** (decreased dissociation)
eqn moves to lhs [1]
to side with fewer molecules / reduce pressure [1] [2]
- (iii)** $SO_2Cl_2 \rightarrow SO_2 + Cl_2$
2 0 0
0.5 1.5 1.5
partial pressure $SO_2 = 1.5 / 3.5 \times 150 = 64.285$
partial pressure $Cl_2 = 64.285$
partial pressure $SO_2Cl_2 = 0.5 / 3.5 \times 150 = 21.43$
 $K_p = PP(SO_2) \times PP(Cl_2) / PP(SO_2Cl_2)$
 $= (64.285)^2 / 21.43$
 $= 192.85$ [3] kPa [1]
each error [-1], carry error through [4]

			AVAILABLE MARKS
(e) $\text{SO}_2\text{Cl}_2 + 2\text{H}_2\text{O} \rightarrow \text{H}_2\text{SO}_4 + 2\text{HCl}$ 1 mole $\text{SO}_2\text{Cl}_2 \rightarrow 4$ moles H^+ 135 g = 1 mole $[\text{H}^+] = 4\text{M}$ $\text{pH} = -\log 4$ $= -0.6$ each error $[-1]$, carry error through			24
16 (a)	$\text{C}_2\text{H}_5\text{OH} \rightarrow \text{CH}_3\text{CHO} + \text{H}_2$	[1]	
(b) (i)	rotate (the plane) [1] of plane polarised light [1] (plane must be mentioned at least once)	[2]	
(ii)		[2]	
(iii)	$\text{CH}_3-\text{CH}=\text{CH}-\text{CHO}$	[1]	6
Section B			100
Total			120



Rewarding Learning

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General Certificate of Education
2010

Chemistry

Assessment Unit A2 2

assessing

Analytical, Transition Metals, Electrochemistry
and Further Organic Chemistry

[AC222]

TUESDAY 1 JUNE, AFTERNOON

**MARK
SCHEME**

- Section A**
- 1** C
 - 2** A
 - 3** D
 - 4** C
 - 5** B
 - 6** C
 - 7** D
 - 8** D
 - 9** C
 - 10** B

[2] for each correct answer

[20]

Section A

AVAILABLE MARKS	
	20
	20

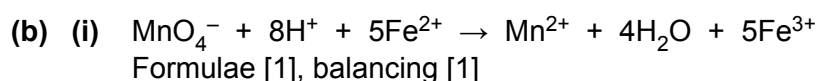
Section B

11 (a)	Titration	Indicator	Colour change	
			from	to
	Magnesium/Edta	Eriochrome black T	Red	Blue
	Iodine/Thiosulphate	Starch	Blue/Black	Colourless

[1] for the correct indicator

[1] for the correct colour change

[6]



[2]

(ii) Colourless [1] to pink [1]

[2]

(iii) Moles of $\text{MnO}_4^- = (18.7 \times 0.01)/1000 = 1.87 \times 10^{-4}$

Moles of $\text{FeC}_4\text{H}_2\text{O}_4$ in $25 \text{ cm}^3 = (1.87 \times 10^{-4}) \times 5 = 9.35 \times 10^{-4}$

Moles of $\text{FeC}_4\text{H}_2\text{O}_4$ in $250 \text{ cm}^3 = 9.35 \times 10^{-3}$

Mass of $\text{FeC}_4\text{H}_2\text{O}_4 = (9.35 \times 10^{-3}) \times 170 = 1.59 \text{ g}$

Mass of $\text{FeC}_4\text{H}_2\text{O}_4$ in 1 tablet = $1.59/5 = 0.318 \text{ g}$

(4 marks, [-1] for each mistake)

[4]

14

12 (a) (i) Temperature: $20-75^\circ\text{C}$ [1]

Pressure: 1–25 atmospheres [1]

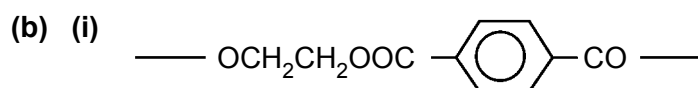
Catalyst: Ziegler/titanium(IV) chloride and triethylaluminium [1]

[3]

(ii) High density: (little branching/high crystallinity)/low flexibility
or

Low density: (high branching/low crystallinity)/high flexibility

[2]



2 marks ([-1] for each mistake)

[2]

(ii) e.g. clothing/plastic bottles

[1]

8

13 (a) Polydentate ligand: a ligand with more than one lone pair of electrons which forms more than one central bond (with a coordinate/dative metal) [2]

(b) (i) 4 [1]

(ii) Square planar [1]

(iii) — : Covalent bond [1]
 ----- : Hydrogen bond [1]
 —→ : Dative/Co-ordinate bond [1] [3]

(c) (dimethylglyoxime) replaces 6 water molecules/
 or 3 molecules → 7 molecules. [1]
 This increases the overall entropy [1] [2]

9

14 (a) methyl -2, 4, 6 - trinitrobenzene [1]

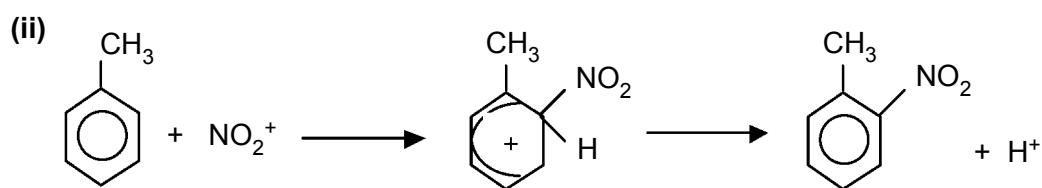
(b) $2\text{C}_7\text{H}_5\text{N}_3\text{O}_6 + 10\frac{1}{2}\text{O}_2 \rightarrow 14\text{CO}_2 + 3\text{N}_2 + 5\text{H}_2\text{O}$
 Formulae [1], balancing [1] [2]

(c) (i) Concentrated nitric acid [1] and
 concentrated sulphuric acid [1] [2]

(ii) $\text{HNO}_3 + 2\text{H}_2\text{SO}_4 \rightarrow \text{NO}_2^+ + 2\text{HSO}_4^- + \text{H}_3\text{O}^+$
 Formulae [1], balancing [1] [2]

(iii) Nitronium ion [1]

(d) (i) Electrophilic [1] substitution [1] [2]



([-1] for each mistake) [2]

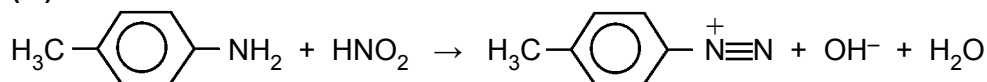
(e) (i) Tin [1] in (concentrated) hydrochloric acid [1] [2]

(ii) Addition of alkali to the salt [1]
 ([-1] for each mistake)

(f) (i) $\text{NaNO}_2 + \text{HCl} \rightarrow \text{HNO}_2 + \text{HCl}$ [1]

(ii) Below 10°C [1]

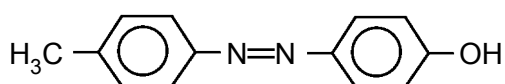
(iii)



([−1] for each mistake) [2]

(g) (i) Coupling [1]

(ii)



([−1] for each mistake) [2]

(iii) Conjugated (double bonds)/Delocalised/Energy levels close together [1]

Hence electron excited/move to higher energy level [1]

Removes a colour from light [1] [3]

25

15 (a) Transition metal atoms/ions have an incomplete d-subshell. [1]

(b) (i) $2\text{SO}_2 + \text{O}_2 \rightarrow 2\text{SO}_3$ [1]

(ii) It is in a different physical state from the reactants. [1]

(iii) Reactants adsorb onto the surface [1]

Bonds weakened in the reactants [1]

Bonds form in products and products are desorbed from the surface [1]

Orientation/closer together/lower activation energy [1]

Any 3 from 4 [3]

(c) (i)

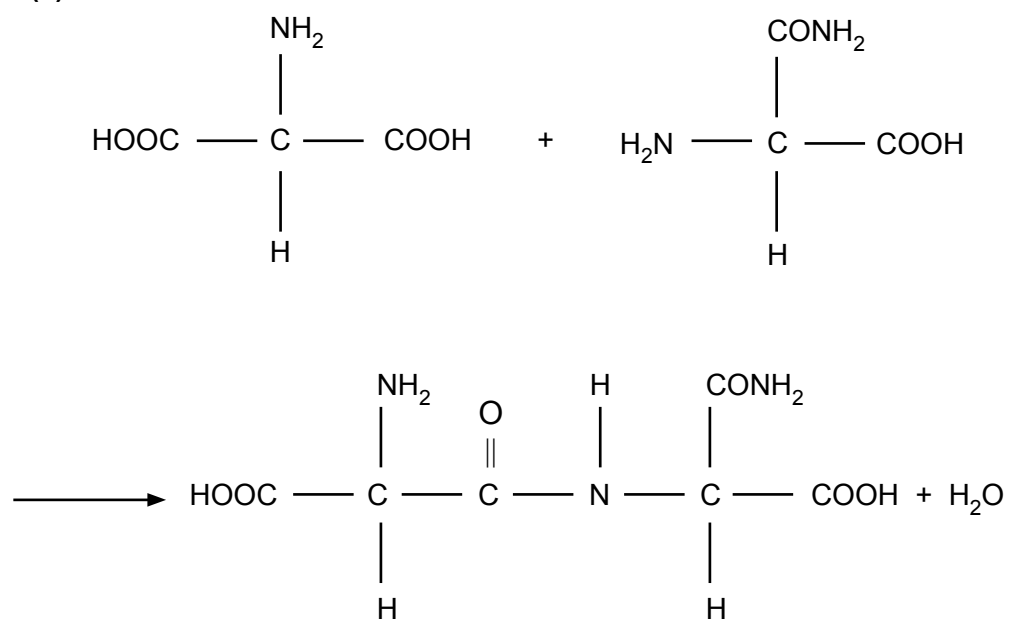
Ion	Colour
VO_2^+ (aq)	
VO^{2+} (aq)	Blue
V^{3+} (aq)	Green
V^{2+} (aq)	

([1] each) [2]

			AVAILABLE MARKS
(ii) $3\text{VO}_2^+(\text{aq}) + 6\text{H}^+(\text{aq}) + \text{Cr}(\text{s}) \rightarrow 3\text{VO}^{2+}(\text{aq}) + 3\text{H}_2\text{O}(\text{l}) + \text{Cr}^{3+}(\text{aq})$ Formulae [1], balancing [1] e.m.f. +1.74 V [1]			[3]
(d)	(i) Hydrogen peroxide		[1]
	(ii) Green [1] to yellow [1]		[2]
	(iii) Colour change from orange to yellow [1] H^+ ions will be removed [1] Equilibrium moves to the left [1]		[3]
16	(a) (i) $\text{C}_{18}\text{H}_{27}\text{O}_3\text{N}$		[1]
	(ii) 100 cm^3 contain 5 g 1000 cm^3 contain 50 g $\frac{50}{305} = 0.164 \text{ mol dm}^{-3}$		[3]
	(b) Place sample at the corner of a chromatogram [1] Run the chromatogram in a suitable solvent [1] (Dry chromatogram and) run at right angles in a different solvent [1] Development + Compare R_f values or chromatogram run with capsaicin [1] Quality of written communication [2]		[6]
17	(a) (i) Primary: sequence of amino acids [1] Secondary: alpha helix/beta pleated sheet [1] due to the formation of a hydrogen bond between the nitrogen of one peptide bond and the oxygen of another further along the chain [1] Tertiary: cross links between amino acids [1] due to H-bonds between amino acids/electrostatic attractions between polar groups/dipole–dipole interactions/Van der Waals forces between non-polar groups/disulphide bridges [1]		[5]
	(ii) The enzyme provides an active site [1]/lock and key mechanism [1]		[2]
	(iii) High temperature denatures the enzyme [1] the structure and the active site is no longer functional [1] (disrupts the secondary and tertiary structures by breaking bonds [1]) Any two		[2]

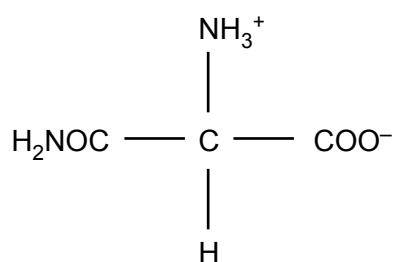
(b) (i) React with ammonia and heat the ammonium salt [1]

(ii)



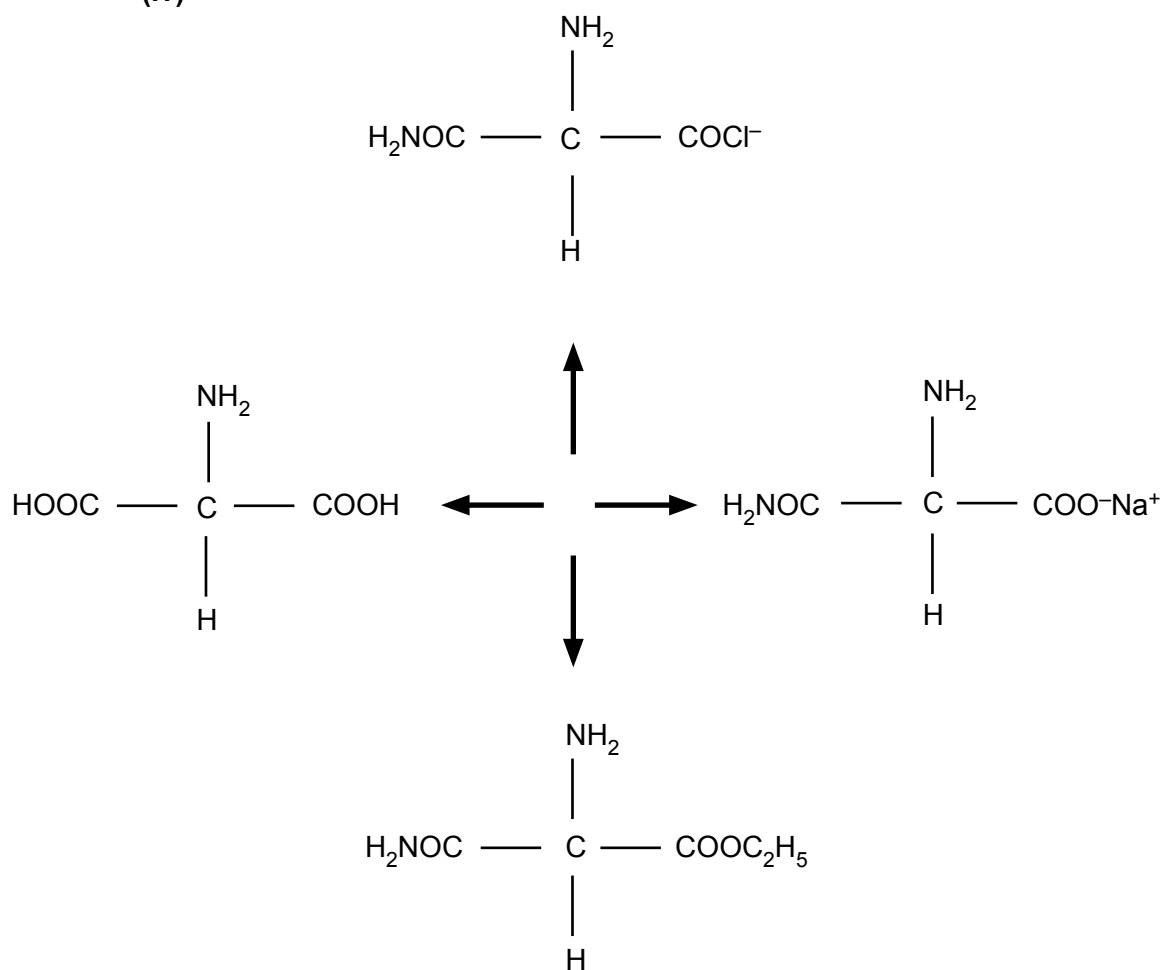
([−1] for each mistake) [2]

(iii)

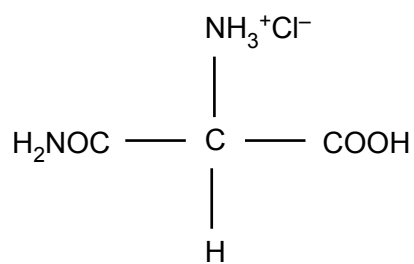


[1]

(iv)



or



([1] each)

[4]

17

Section B

100

Total

120

New
Specification



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General Certificate of Education
2010

Chemistry

Assessment Unit A2 3

Internal Assessment
Practical Examination 1

[AC231]

THURSDAY 20 MAY

MARK SCHEME

Annotation

1. Please do all marking in red ink.
2. All scripts are checked for mathematical errors. Please adopt the system of one tick (✓) equals [1] mark e.g. if you have awarded 4 marks for part of a question then 4 ticks (✓) should be on this candidate's answer.
3. As candidates have access to scripts please do not write any inappropriate comments on their scripts.

1 Titration exercise

- (a) **Rinse** out a pipette with the solution of potassium iodate (V) and **transfer** a known volume of the solution into a (conical) flask [1]
 Add a portion of sulphuric acid and a sample of potassium iodide [1]
Rinse out the burette with the solution of sodium thiosulphate and **fill** the burette [1]
 Add the solution from the burette until the solution turns a straw yellow colour [1]
 Add starch indicator [1]
 Continue to add the solution from the burette one drop at a time until the solution changes from blue/black to colourless [1]
 Repeat for reliability [1]

To a maximum of [6]

Mark denied if:

- (i) there is no mention of rinsing pipette
- (ii) there is no mention of rinsing burette

- (b) Table [1]
 Significant figures [2]
 Calculation of average titre [2]
 Titration consistency [2]
 Agreement with supervisor's titre [3] [10]

NOTES:

Table:

Table should include initial burette reading, final burette reading, and volume delivered. Units should be included for volume delivered (may be omitted in the other readings). Mark denied if no indication of units.

Significant figures:

All burette readings should be to at least one decimal place – each mistake is penalised by 1 mark.

(However, initial burette readings of 0 are penalised once only)

If used, the second decimal position should be 0 or 5 only – other values are penalised by 1 mark for each.

Average titre:

Values for accurate titrations only should be used.

The use of the rough value is [-1].

The average value can be two decimal places, e.g. 25.37

An incorrect calculation is 0. Units must be included.

Mark denied if:

- (i) only one accurate titration done or if the titre is not calculated correctly
- (ii) units not included loses one mark

Titration consistency:

This is the difference between the first and second accurate readings

Difference	Mark
------------	------

±0.1	[2]
------	-----

±0.2	[1]
------	-----

±0.3	[0]
------	-----

Titration agreement with supervisor:

This is the difference between the average titre and the supervisor's value

Difference	Marks
------------	-------

±0.1	[3]
------	-----

±0.2	[2]
------	-----

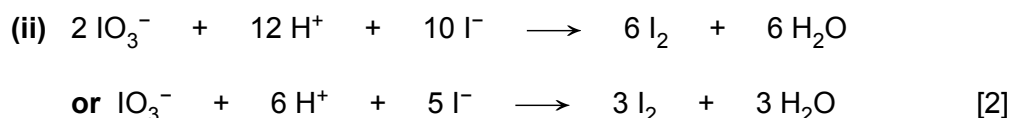
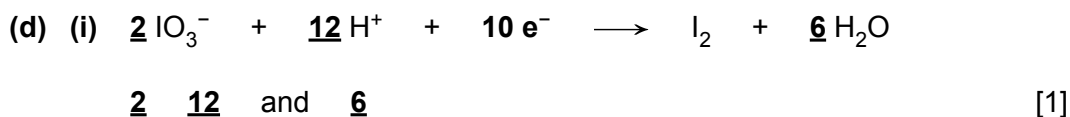
±0.3	[1]
------	-----

±0.4	[0]
------	-----

Please note that the supervisor's titre should be recorded at the bottom of page 3 in the candidate's script in RED INK.

The marks for table, significant figures etc should be recorded on the left-hand side of the candidate's table of results.

- (c) Correct calculation of moles of sodium thiosulphate [1]
 Number of moles of sodium thiosulphate divided by two [1]



- (e) Moles of iodine from (c) divided by 3 [1]
 then multiplied by 40 (1000/25) [1]
 then multiplied by RFM of KIO₃ which is 214 [1]

25

Consequential marking/carry error through (CET) to be applied in calculations e.g. incorrect ratio in (d)(ii) can be carried through into part (e).

2 Observation/deduction

**There are 28 scoring points available in question 2.
However the maximum marks for this question is 25.**

In Tests 3 and 5 candidates can score additional marks to those indicated
– see below.

If the candidate scores more than 25 then MAX 25 should be written at beginning
of question in the teacher mark column.

(a) Test 1 *Blue [1] solid*

Test 2 *Blue solution [1]*

Test 3 *White [1] precipitate [1] blue solution remains [1]
Candidates can score all three marks*

Test 4 *Blue [1] precipitate [1]*

Test 5 *Blue [1] precipitate [1]
precipitate dissolves [1]
deep blue solution [1]
Candidates can score all four marks*

Test 6 *Green [1] solution [1]
or yellow-green*

Test 2 *$[\text{Cu}(\text{H}_2\text{O})_6]^{2+}$ [1] – square bracket essential*

Test 5 *$[\text{Cu}(\text{NH}_3)_4(\text{H}_2\text{O})_2]^{2+}$ [1] – square bracket essential*

Test 6 *$[\text{CuCl}_4]^{2-}$ [1] – square bracket essential*

Hydrated [1]

Copper (II) sulphate [1] (Accept copper sulphate.)

(b) *2 layers [1]
red/red-brown [1]
solid [1]*

structure of propanal [1]

structure of propanone [1]

(c) *solid disappears/dissolves [1]
bubbles/fizzes/effervescence [1]
colourless gas/solution [1]
becomes warm [1]*

Max 3

structure of propanoic acid [1]

structure of ethyl methanoate [1]

25

3 Planning exercise

(a) $\text{Sn} + 2 \text{I}_2 \longrightarrow \text{SnI}_4$ [2]
(unbalanced [-1])

(b) Want actual yield of 6.0 g (0.00957 moles)

∴ theoretical yield is 6.67g [1] (or 0.0106 moles)

moles of iodine needed 0.0212 moles [1]

mass of iodine 5.40 [1] g [1]

Correct answer gets 4 marks. Units missing from final answer [-1] [4]

(c) (i) gloves since iodine is corrosive or DCM is toxic [1]
no naked flame/use electrical heater since DCM is flammable [1] [2]

(ii) repeated [1] boiling and condensing [1] of a reaction mixture
(without loss of material) [2]

(d) (i) colour (of iodine) [1] disappears [1] [2]

(ii) filter [1] [1]

(iii) evaporate the solvent until crystals begin to appear [1]
allow to cool and then filter [1] [2]

or

distil off the solvent [1]

using a water bath/fume cupboard / crystals in flask [1]

(iv) dissolve in minimum of the hot DCM [1]

filter while hot [1]

allow filtrate to cool [1]

filter again [1]

to a maximum of [3]

Quality of written communication:

- 2 marks The candidate expresses ideas clearly and fluently through well-linked sentences and paragraphs. Arguments are generally relevant and well-structured. There are few errors of grammar, punctuation and spelling.
- 1 mark The candidate expresses ideas clearly, if not always fluently. Arguments may sometimes stray from the point. There may be some errors of grammar, punctuation and spelling, but not such as to suggest a weakness in these areas.
- 0 marks The candidate expresses ideas satisfactorily, but without precision. Arguments may be of doubtful relevance or obscurely presented. Errors in grammar, punctuation and spelling are sufficiently intrusive to disrupt the understanding of the passage. [2]

Total**AVAILABLE
MARKS**

20

70



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2010

Chemistry

Assessment Unit A2 3

Internal Assessment
Practical Examination 2

[AC232]

FRIDAY 21 MAY

MARK SCHEME

Annotation

1. Please do all marking in red ink.
2. All scripts are checked for mathematical errors. Please adopt the system of one tick (✓) equals [1] mark e.g. if you have awarded 4 marks for part of a question then 4 ticks (✓) should be on this candidate's answer.
3. As candidates have access to scripts please do not write any inappropriate comments on their scripts.

1 Titration exercise

- (a) **Rinse** out a pipette with the solution of sodium iodate (V) and **transfer** a known volume of the solution into a (conical) flask [1]

Add a portion of sulphuric acid and a sample of potassium iodide [1]

Rinse out the burette with the solution of sodium thiosulphate and **fill** the burette [1]

Add the solution from the burette until the solution turns a straw yellow colour [1]

Add starch indicator [1]

Continue to add the solution from the burette one drop at a time until the solution changes from blue/black to colourless [1]

Repeat for reliability [1]

To a maximum of [6]

Mark denied if:

- (i) there is no mention of rinsing pipette
- (ii) there is no mention of rinsing burette

- (b) Table [1]

Significant figures [2]

Calculation of average titre [2]

Titration consistency [2]

Agreement with supervisor's titre [3]

[10]

NOTES:

Table:

Table should include initial burette reading, final burette reading, and volume delivered. Units should be included for volume delivered (may be omitted in the other readings). Mark denied if no indication of units.

Significant figures:

All burette readings should be to at least one decimal place – each mistake is penalised by 1 mark.

(However, initial burette readings of 0 are penalised once only)

If used, the second decimal position should be 0 or 5 only – other values are penalised by 1 mark for each.

Average titre:

Values for accurate titrations only should be used.

The use of the rough value is [-1].

The average value can be two decimal places, e.g. 25.37

An incorrect calculation is 0. Units must be included.

Mark denied if only one accurate titration done or if the titre is not calculated correctly.

Units not included loses one mark.

Titration consistency:

This is the difference between the first and second accurate readings

Difference	Mark
------------	------

±0.1	[2]
------	-----

±0.2	[1]
------	-----

±0.3	[0]
------	-----

Titration agreement with supervisor:

This is the difference between the average titre and the supervisor's value

Difference	Marks
------------	-------

±0.1	[3]
------	-----

±0.2	[2]
------	-----

±0.3	[1]
------	-----

±0.4	[0]
------	-----

Please note that the supervisor's titre should be recorded at the bottom of page 3 in the candidate's script in RED INK.

The marks for table, significant figures etc should be recorded on the left-hand side of the candidate's table of results.

(c) Correct calculation of moles of sodium thiosulphate [1]

Number of moles of sodium thiosulphate divided by two [1]

(d) (i) $2 \text{IO}_3^- + 12 \text{H}^+ + 10 \text{e}^- \longrightarrow \text{I}_2 + 6 \text{H}_2\text{O}$ [1]
 $2 \quad 12 \quad \text{and} \quad 6$

10e^- [1]

(ii) $2 \text{IO}_3^- + 12 \text{H}^+ + 10 \text{I}^- \longrightarrow 6 \text{I}_2 + 6 \text{H}_2\text{O}$
 or $\text{IO}_3^- + 6 \text{H}^+ + 5 \text{I}^- \longrightarrow 3 \text{I}_2 + 3 \text{H}_2\text{O}$ [2]

(e) Moles of iodine from (c) divided by 3 [1]

then multiplied by 40 (1000/25) [1]

then multiplied by RFM of NaIO_3 which is 198 [1]

25

Consequential marking/carry error through (CET) to be applied in calculations e.g. incorrect ratio in (d)(ii) can be carried through into part (e).

2 Observation/deduction

**There are 28 scoring points available in question 2.
However the maximum marks for this question is 25.**

In Tests 3 and 5 candidates can score additional marks to those indicated
– see below.

If the candidate scores more than 25 then MAX 25 should be written at beginning
of question in the teacher mark column.

(a) Test 1 *Pink [1] solid – accept red or red-brown*

Test 2 *Pink solution[1]*

Test 3 *White [1] precipitate [1] pink solution remains [1]
Candidates can score all three marks*

Test 4 *Blue [1] precipitate [1]*

Test 5 *Blue [1] precipitate [1]
precipitate dissolves [1]
yellow-brown solution [1]
Candidates can score all four marks*

Test 6 *Blue [1] solution [1]*

Test 2 *$[\text{Co}(\text{H}_2\text{O})_6]^{2+}$ [1] – square bracket essential*

Test 5 *$[\text{Co}(\text{NH}_3)_6]^{2+}$ [1] – square bracket essential*

Test 6 *$[\text{CoCl}_4]^{2-}$ [1] – square bracket essential*

Hydrated [1]

Cobalt (II) sulphate [1] (Accept cobalt sulphate.)

(b) *2 layers [1]
red/red-brown [1]
solid [1]*

*structure of propanal [1]
structure of propanone [1]*

(c) *solid disappears/dissolves [1]
bubbles/fizzes/effervescence [1]
colourless gas/solution [1]
becomes warm [1]
Max 3*

*structure of propanoic acid [1]
structure of ethyl methanoate [1]*

25

3 Planning exercise

(a) $\text{C}_3\text{H}_7\text{COOH} + \text{C}_4\text{H}_9\text{OH} \rightleftharpoons \text{C}_3\text{H}_7\text{COOC}_4\text{H}_9 + \text{H}_2\text{O}$
(equilibrium arrows missing [−1]) [2]

(b) Want actual yield of 0.181 moles/26.1 g [1]
∴ theoretical yield is 0.259 moles/37.3 g [1]
moles of butanoic acid needed 0.259 moles [1]
mass of butanoic acid 22.8 [1] g [1]
Correct answer gets 4 marks. Units missing from final answer [−1] [4]

(c) Procedure:

Add (excess) butan-1-ol, butanoic acid and concentrated sulphuric acid
to a (round-bottomed or pear-shaped) flask [1]
Reflux the mixture [1]
distil [1]
collect at 163–167°C [1]

Safety:

Either: gloves since concentrated sulphuric acid is corrosive
Or: add concentrated sulphuric acid slowly since exothermic
Or: add anti-bump granules to promote smooth boiling [1] [5]

(d) (i) shake with solution of sodium carbonate / hydrogencarbonate [1]
using a separating funnel [1]
separate the two layers/release pressure [1] [3]

(ii) (shake with) suitable named anhydrous solid [1]
filter/decant [1] [2]

(iii) distillation [1]
collect at 165°C (164°–166°) [1] [2]

Quality of written communication:

- 2 marks The candidate expresses ideas clearly and fluently through well-linked sentences and paragraphs. Arguments are generally relevant and well-structured. There are few errors of grammar, punctuation and spelling.
- 1 mark The candidate expresses ideas clearly, if not always fluently. Arguments may sometimes stray from the point. There may be some errors of grammar, punctuation and spelling, but not such as to suggest a weakness in these areas.
- 0 marks The candidate expresses ideas satisfactorily, but without precision. Arguments may be of doubtful relevance or obscurely presented. Errors in grammar, punctuation and spelling are sufficiently intrusive to disrupt the understanding of the passage. [2]

Total**AVAILABLE
MARKS**

20

70



Rewarding Learning

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General Certificate of Education
2011

Centre Number

71

Candidate Number

Chemistry

Assessment Unit A2 2

assessing

Analytical, Transition Metals, Electrochemistry
and Further Organic Chemistry

[AC222]

FRIDAY 27 MAY, AFTERNOON



AC222

TIME

2 hours.

INSTRUCTIONS TO CANDIDATES

Write your Centre Number and Candidate Number in the spaces provided at the top of this page.

Answer **all fifteen** questions.

Answer **all ten** questions in **Section A**. Record your answers by marking the appropriate letter on the answer sheet provided. Use only the spaces numbered 1 to 10. Keep in sequence when answering.

Answer **all five** questions in **Section B**. Write your answers in the spaces provided in this question paper.

INFORMATION FOR CANDIDATES

The total mark for this paper is 120.

Quality of written communication will be assessed in question **13(d)(iii)**.

In Section A all questions carry equal marks, i.e. **two** marks for each question.

In Section B the figures printed down the right-hand side of pages indicate the marks awarded to each question or part question.

A Periodic Table of Elements (including some data) is provided.

For Examiner's
use only

Question Number	Marks
--------------------	-------

Section A

1–10

Section B

11

12

13

14

15

Total
Marks

Section A

For each of the questions only **one** of the lettered responses (A–D) is correct.

Select the correct response in each case and mark its code letter by connecting the dots as illustrated on the answer sheet.

- 1 Chlorine has two isotopes, chlorine-35 and chlorine-37.
Which one of the following is the number of peaks found in the mass spectrum of chlorine gas?

A 2
B 3
C 4
D 5

- 2 The mechanism for the nitration of benzene is described as

A electrophilic addition.
B electrophilic substitution.
C nucleophilic addition.
D nucleophilic substitution.

- 3 Copper(II) ions form a coloured complex with the ligand L. The following absorbances were recorded on mixing different volumes of 0.05 M copper(II) sulfate and 0.1 M ligand L.

volume of 0.05 M $\text{CuSO}_4(\text{aq})/\text{cm}^3$	volume of 0.1 M $\text{L}(\text{aq})/\text{cm}^3$	absorbance
3.0	7.0	0.412
4.0	6.0	0.457
5.0	5.0	0.406
6.0	4.0	0.335
7.0	3.0	0.251

Which one of the following is the cation to ligand ratio in this complex?

A 1:2
B 1:3
C 2:3
D 3:2

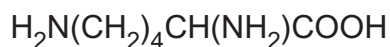
Examiner Only

Marks Remark

Section B

Answer **all five** questions in the spaces provided.

- 11** Lysine (2,6-diaminohexanoic acid) has the formula



The molecule is optically active and may undergo polymerisation.

- (a) (i) Explain the term **optically active**.

 [2]

- (ii) Draw the 3D structure of lysine labelling the asymmetric carbon with an asterisk (*).

[2]

- (b)** Explain why lysine has a relatively high melting point.

[2]

- (c) Write the formula of the organic ion present when lysine is dissolved in an alkaline solution.

 [1]

Examiner Only	
Marks	Remark

(ii) Name the reagents required for step A.

[1]

(iii) State and explain the condition necessary for the first step.

[2]

(iv) Suggest a structure of the dye Prontosil which is formed by a coupling reaction.

[2]

(v) Explain why Prontosil is coloured.

[3]

(ii) Hexaaquanickel(II) ions react with en in solution. Write the equation for this reaction in which all the water ligands are replaced.

[2]

(iii) Explain why this ligand replacement takes place.

[2]

(d) (i) Explain the role of iron(II) in haemoglobin.

[2]

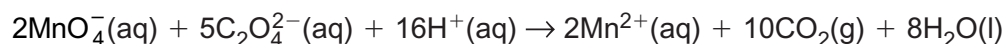
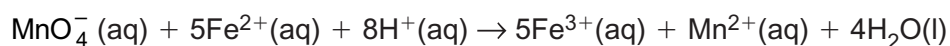
(ii) Explain the effect of the inhalation of carbon monoxide on haemoglobin.

 [1]

(e) Aqueous iron(III) ions form a stable complex with the bidentate ligand ethanedioate, $\text{C}_2\text{O}_4^{2-}$.
The iron(III) ions combine with three ethanedioate ions. Deduce the formula of the complex formed.

[1]

(f) Potassium manganate(VII) oxidises iron(II) ions and ethanedioate ions according to the equations:



(i) Describe, with observations, how you could confirm the presence of aqueous iron(III) ions, following the oxidation of iron(II) ions, without interference from manganese(II) ions.

[3]

(ii) 25.0 cm^3 of an acidified iron(II) ethanedioate solution required 32.2 cm^3 of 0.025 mol dm^{-3} of potassium manganate(VII) solution for complete reaction. Calculate the concentration, in mol dm^{-3} , of the iron(II) ethanedioate solution.

[4]

THIS IS THE END OF THE QUESTION PAPER

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2011

Chemistry

Assessment Unit A2 2

assessing

Analytical, Transition Metals, Electrochemistry
and Further Organic Chemistry

[AC222]

FRIDAY 27 MAY, AFTERNOON

MARK SCHEME

Section A

- 1 D
- 2 B
- 3 B
- 4 C
- 5 D
- 6 B
- 7 B
- 8 B
- 9 D
- 10 C

[2] for each correct answer

[20]

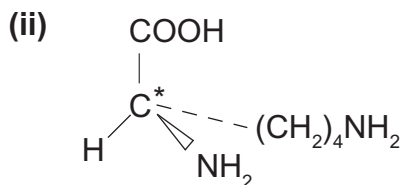
Section A

AVAILABLE MARKS	
	20
	20

Section B

AVAILABLE
MARKS

- 11 (a) (i) ability to rotate plane [1]
of plane polarised light [1] [2]



correct structure (3D) [1]
asymmetric carbon labelled correctly [1] [2]

- (b) zwitterion/ionic [1]
strong attractions between oppositely charged ions/molecules [1] [2]

- (c) $\text{H}_2\text{N}(\text{CH}_2)_4\text{CH}(\text{NH}_2)\text{COO}^-$ [1]

- (d) $\text{H}_2\text{N}(\text{CH}_2)_4\text{CH}(\text{NH}_2)\text{CONH}(\text{CH}_2)_4\text{CH}(\text{NH}_2)\text{COOH}/$
 $\text{H}_2\text{N}(\text{CH}_2)_4\text{CH}(\text{NH}_2)\text{CONHCH}(\text{CH}_2)_4\text{NH}_2$

|

COOH

 [2]

- (e) (i) distance travelled by solute
divided by distance travelled by solvent (front) [1]

- (ii) more soluble in water/stationary phase [1]
than mobile phase/solvent [1] [2]

(f)

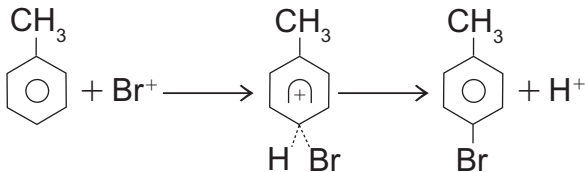
	N	C	H	O
	10.5	36.1	5.3	48.1
÷ RAM	0.75	3.01	5.3	3.01
÷ 0.75	1	4	7	4

empirical formula = $\text{C}_4\text{H}_7\text{O}_4\text{N}$ (each error [−1], cet) [3]

15

12 (a)	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^5 4s^1$ [1] half-filled d subshell stable [1]	[2]
(b)	moles Al = $25/27 = 0.926$ moles $Cr_2O_3 = 100/152 = 0.658$ 2 moles Al = 1 mole Cr_2O_3 therefore Cr_2O_3 in excess 1 mole Al = 1 mole Cr 0.926 mole Al = 0.926 mole Cr mass Cr = $0.926 \times 52 = 48.15$ g % yield = $42.5 \times 100/48.15 = 88.3\%$ Award [3] for correct answer. Each error [−1], carry error through	[3]
(c)	$K_2Cr_2O_7 + H_2SO_4 \rightarrow K_2SO_4 + 2CrO_3 + H_2O$ unbalanced [−1]	[2]
(d) (i)	$Cr^{2+} + Ag^+ \rightarrow Cr^{3+} + Ag$	[1]
	(ii) $0.8 - (-0.41) = (+)1.21$ (V)	[1]
(e) (i)	ethanol/alcohol	[1]
	(ii) purple/violet [1] $K_2SO_4 \cdot Cr_2(SO_4)_3 \cdot 24H_2O$ [1]	[2]
(f) (i)	$C_6SN_2H_8O_2$	[1]
	(ii) sodium nitrite and hydrochloric acid	[1]
	(iii) $\leq 10^\circ C$ [1] diazonium ion decomposes above this [1]	[2]
(iv)	H_2NSO_2  $N=N$  NH_2	[2]
	structure [2], each error [−1]	[2]
(v)	highly conjugated/delocalised [1] electronic energy levels close together [1] (electron transitions) absorb visible light [1]	[3]

13 (a)	CH_3COO^+	[1]
(b) (i)	$\left\{ \begin{array}{l} \text{ratio } 2:3:3 \text{ or } 3:2:3 \text{ [1]} \\ \text{CH}_3:\text{CH}_2:\text{CH}_3/\text{no. of Hs in each environment [1]} \end{array} \right.$	[2]
(ii)	high value quartet near electronegative O lowest CH_3 from CH_2CH_3 intermediate singlet proximity to COO all three [2] errors [−1]	[2]
(iii)	(quartet) due to split by three chemically equivalent H ($n + 1$)	[1]
(iv)	no H on adjacent atom (to split signal)	[1]
(c) (i)	$\text{CH}_3\text{COOH} + \text{NH}_3 \rightarrow \text{CH}_3\text{COONH}_4$	[1]
(ii)	$\text{CH}_3\text{COONH}_4 \rightarrow \text{CH}_3\text{CONH}_2 + \text{H}_2\text{O}$	[1]
(iii)	acid: CH_3COOH [1] alkaline: CH_3COONa [1]	[2]
(d) (i)	A: P_4O_{10} [1] B: LiAlH_4 [1]	[2]
(ii)	$\text{RNH}_2 + \text{CH}_3\text{COCl} \rightarrow \text{CH}_3\text{CONHR} + \text{HCl}$	[1]
(iii)	determine melting point/description of apparatus heat slowly record temperature when melting starts and stops/range compare to tables Any four	[4]
	Quality of written communication	[2]
(e) (i)	two coordinate (dative) bonds formed by lone pairs	[2]
(ii)	$[\text{Ni}(\text{H}_2\text{O})_6]^{2+} + 3\text{en} \rightarrow [\text{Ni}(\text{en})_3]^{2+} + 6\text{H}_2\text{O}$ unbalanced [1]	[2]
(iii)	increase in entropy [1] 4 particles to 7 particles [1]	[2]

- 14 (a)** Fe^{2+} : $1s^2 2s^2 2p^6 3s^2 3p^6 3d^6$ [1]
 Fe^{3+} : $1s^2 2s^2 2p^6 3s^2 3p^6 3d^5$ [1] [2]
- (b)** molecules adsorbed/attached to surface [1]
 weakens bonds/alignment/bonds formed with surface [1]
 lowers activation energy [1] [3]
- (c) (i)**  [3]
- Each error [− 1] [3]
- (ii)** 2-bromo-1,4-dimethylbenzene [2]
 Each error [− 1]
- (d) (i)** oxygen forms bond with Fe^{2+} [1]
 carried around the body [1] [2]
- (ii)** CO combines irreversibly [1]
- (e)** $[\text{Fe}(\text{C}_2\text{O}_4)_3]^{3-}$ [1]
- (f) (i)** thiocyanate ions/hydroxide ions/ $\text{NH}_3(\text{aq})$ [1]
 blood red/rust/brown [1]
 solution/ppt [1] [3]
- (ii)** 1 mole $\text{MnO}_4^- = 5$ moles Fe^{2+}
 2 moles $\text{MnO}_4^- = 5$ moles $\text{C}_2\text{O}_4^{2-}$
 $\Rightarrow 3$ moles $\text{MnO}_4^- = 5$ moles FeC_2O_4
 Number of moles iron(II) ethanedioate = $\frac{32.2 \times 0.025}{1000}$
 $= 8.05 \times 10^{-4}$
 Number of moles $\text{MnO}_4^- = 8.05 \times 10^{-4} \times \frac{5}{3}$
 $= 1.34 \times 10^{-3}$ in 25.0 cm^3
 Concentration $\text{MnO}_4^- = 1.34 \times 10^{-3} \times 40 = 0.0536$
 $= 0.054 \text{ mol dm}^{-3}$
 Award [4] directly for correct answer
 Each error [− 1], carry error through [4]

AVAILABLE
MARKS

21

- 15 (a) (i)** Ziegler 50–75° 1–10 atms./low TiCl_4 /aluminium (Ziegler catalyst) [1] [1] [1] [3]
- Philips 150–180° 30–40 atms. Cr_2O_3 [1] [1] [1]
- (ii)** rigid and high softening temperature [1]
 little branching [1]
 pack close together/crystalline/strong intermolecular forces [1] [3]
- (b)**
- $$\begin{array}{ccccccc}
 \text{H} & \text{CH}_3 & & \text{H} & \text{CH}_3 \\
 | & | & & | & | \\
 -\text{C}- & \text{C}- & & \text{C}- & \text{C}- \\
 | & | & & | & | \\
 \text{H} & \text{COOCH}_3 & & \text{H} & \text{COOCH}_3
 \end{array}$$
- Structure [2]
- (c) (i)** (polymerisation involves) loss of water/small molecule [1]
- (ii)**
- $$\left[\begin{array}{c} \text{H} \\ | \\ -\text{N}- \end{array} (\text{CH}_2)_6 \begin{array}{c} \text{H} \quad \text{O} \\ | \quad || \\ -\text{N}- \text{C}- \end{array} (\text{CH}_2)_4 \begin{array}{c} \text{O} \\ || \\ -\text{C}- \end{array} \right]$$
- structure [2]
 peptide link [1] [3]
- (d)** polyesters are biodegradable [1]
 as they can be hydrolysed [1] [2]
- (e) (i)** sequence of amino acids [1]
- (ii)** lock and key/active site [1]
 lowers activation energy/lower energy pathway [1] [2]

AVAILABLE
MARKS

17

Section B

100

Total

120



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Centre Number

71

Candidate Number

Chemistry

Assessment Unit A2 2

assessing

Analytical, Transition Metals, Electrochemistry
and Further Organic Chemistry

[AC222]



WEDNESDAY 23 MAY, AFTERNOON

TIME

2 hours.

INSTRUCTIONS TO CANDIDATES

Write your Centre Number and Candidate Number in the spaces provided at the top of this page.

Answer **all seventeen** questions.

Answer **all ten** questions in **Section A**. Record your answers by marking the appropriate letter on the answer sheet provided. Use only the spaces numbered 1 to 10. Keep in sequence when answering.

Answer **all seven** questions in **Section B**. Write your answers in the spaces provided in this question paper.

INFORMATION FOR CANDIDATES

The total mark for this paper is 120.

Quality of written communication will be assessed in question **16(c)(i)**.

In Section A all questions carry equal marks, i.e. **two** marks for each question.

In Section B the figures printed down the right-hand side of pages indicate the marks awarded to each question or part question.

A Periodic Table of Elements (including some data) is provided.

For Examiner's
use only

Question Number	Marks
Section A	
1–10	
Section B	
11	
12	
13	
14	
15	
16	
17	

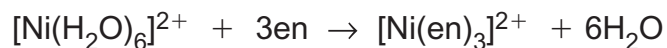
Total
Marks

Section A

For each of the following questions only **one** of the lettered responses (A-D) is correct.

Select the correct response in each case and mark its code letter by connecting the dots as illustrated on the answer sheet.

- 1 Which one of the following applies to the ligand substitution reaction shown?

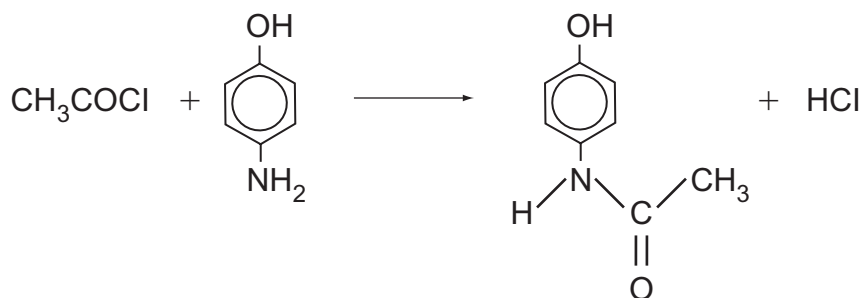


	Change in Coordination Number	$\Delta S^\ominus$
A	6 to 3	negative
B	6 to 3	positive
C	none	negative
D	none	positive

- 2 Which one of the following lists the compounds in order of increasing base strength?

- A ethanamide, methylamine, phenylamine
- B ethanamide, phenylamine, methylamine
- C methylamine, ethanamide, phenylamine
- D phenylamine, ethanamide, methylamine

- 3 The reaction of 4-hydroxyphenylamine to produce paracetamol is shown below.



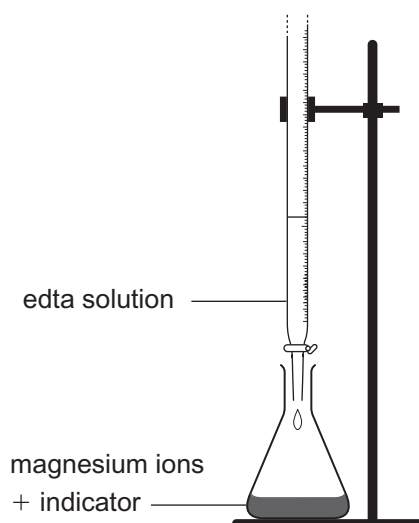
If the reaction has an 80% yield, 10.9 g of 4-hydroxyphenylamine produces

- A 12.1 g of paracetamol.
- B 13.6 g of paracetamol.
- C 15.1 g of paracetamol.
- D 18.9 g of paracetamol.

- 4 Which one of the following methods may be used to separate a mixture of amino acids obtained from protein hydrolysis?
- A distillation
B recrystallisation
C solvent extraction
D thin-layer chromatography
- 5 Which one of the following is a correct statement about the stereochemistry of the complex $[\text{Pt}(\text{NH}_3)_2\text{Cl}_2]$?
- A It is square planar and has cis/trans isomers.
B It is square planar and has two optical isomers.
C It is tetrahedral and has cis/trans isomers.
D It is tetrahedral and has two optical isomers.
- 6 25.0 cm^3 of potassium iodate(V) solution were added to excess potassium iodide solution dissolved in sulfuric acid. The iodine liberated required 30.0 cm^3 of 0.05 mol dm^{-3} $\text{Na}_2\text{S}_2\text{O}_3$ solution. Which one of the following is the concentration of the potassium iodate(V) solution?
- A 0.01 mol dm^{-3}
B 0.02 mol dm^{-3}
C 0.04 mol dm^{-3}
D 0.05 mol dm^{-3}
- 7 Which one of the following gives the ground state electronic configuration for the copper atom and the copper(II) ion?

	copper atom	copper(II) ion
A	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^9 4s^2$	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^9$
B	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^9 4s^2$	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^7 4s^2$
C	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^{10} 4s^1$	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^8 4s^1$
D	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^{10} 4s^1$	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^9$

- 8 The diagram below shows the titration of a solution of magnesium ions with edta using Eriochrome Black T as indicator.



What is the colour change at the end point?

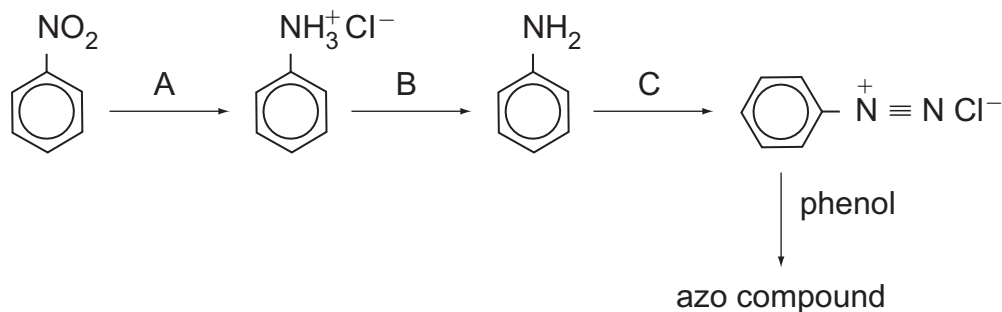
- A blue to red
 - B green to blue
 - C red to green
 - D red to blue
- 9 Which one of the following statements about glycine is **not** correct?
- A It has a relatively high melting point.
 - B It contains 32% carbon by mass.
 - C It exists as optical isomers.
 - D It is soluble in water.
- 10 Which one of the following statements about propanamide is **not** correct?
- A It produces an $M+1$ peak at 73 in its mass spectrum.
 - B It can be dehydrated to form propanenitrile.
 - C It has the molecular formula C_3H_7NO .
 - D It is a weaker base than ammonia.

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(Questions continue overleaf)

Section B

Answer **all seven** questions in the spaces provided.

- 11** The amino group is found in amines. Phenylamine is used in the synthesis of azo compounds. Consider the following sequence of steps:



- (a) (i)** Give the names of the reagents used in the following steps.

A _____ [2]

B _____ [1]

C _____ [2]

- (ii) Give the conditions used in step C and name the product.

[2]

- (iii) Draw the structure of the azo compound. Describe its appearance and name the compound.

[3]

Examiner Only	
Marks	Remark

12 Standard electrode potentials can be used to predict the feasibility of reactions.

	$E^\ominus/V$
$\text{Na}^+(\text{aq}) + \text{e}^- \rightleftharpoons \text{Na}(\text{s})$	-2.71
$\text{Mg}^{2+}(\text{aq}) + 2\text{e}^- \rightleftharpoons \text{Mg}(\text{s})$	-2.37
$\text{Al}^{3+}(\text{aq}) + 3\text{e}^- \rightleftharpoons \text{Al}(\text{s})$	-1.66
$\text{Zn}^{2+}(\text{aq}) + 2\text{e}^- \rightleftharpoons \text{Zn}(\text{s})$	-0.76
$\text{Cr}^{3+}(\text{aq}) + \text{e}^- \rightleftharpoons \text{Cr}^{2+}(\text{aq})$	-0.41
$2\text{H}^+(\text{aq}) + 2\text{e}^- \rightleftharpoons \text{H}_2(\text{g})$	0.00
$\text{Fe}^{3+}(\text{aq}) + \text{e}^- \rightleftharpoons \text{Fe}^{2+}(\text{aq})$	+0.77

- (a) Define the term **standard electrode potential**.

[3]

- (b)** From the table, select the species which is the most powerful reducing agent.

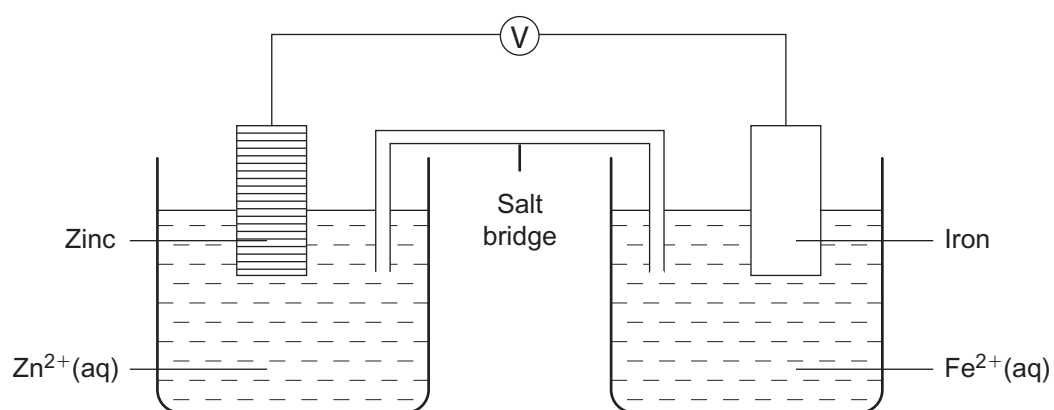
[1]

- (c)** Write the equation for the reaction of aluminium with aqueous zinc ions and calculate the e.m.f.

[3]

Examiner Only	
Marks	Remark

- (d) Under standard conditions, the e.m.f. of the cell shown below is +0.32 V.



Calculate the standard electrode potential for the iron half-cell.

[1]

(b) Mass spectrometry is another important analytical technique.

2-chloropropanoic acid produces molecular ion peaks at 108 and 110. It also produces a significant fragment peak at 91.

(i) Suggest why there are **two** molecular ion peaks.

_____ [2]

(ii) Identify the fragment ion.

_____ [2]

(iii) Complete the table giving the integration values and the splitting of each peak in the nmr spectrum of 2-chloropropanoic acid:

	Peak 1	Peak 2	Peak 3
Integration	3		
Splitting			singlet

[4]

Examiner Only

Marks Remark

(b) The $[\text{Fe}(\text{H}_2\text{O})_6]^{3+}$ ion behaves as a Brønsted acid by the loss of one hydrogen ion.

(i) Write an equation to show $[\text{Fe}(\text{H}_2\text{O})_6]^{3+}$ behaving as a Brønsted acid.

[2]

(ii) Write the expression for the acid dissociation constant of the $[\text{Fe}(\text{H}_2\text{O})_6]^{3+}$ ion.

[1]

(iii) What is observed when sodium hydroxide solution is added to a solution containing $[\text{Fe}(\text{H}_2\text{O})_6]^{3+}$ ions?

[2]

(iv) Describe a different chemical test, including observations, which can be used to detect the presence of low concentrations of $[\text{Fe}(\text{H}_2\text{O})_6]^{3+}$ ions. Give the formula of any new complex formed.

[4]

(c) With reference to the iron(II) ions in haemoglobin, explain why breathing carbon monoxide can result in death.

[2]

Examiner Only	
Marks	Remark

(iv) Name the organic product of this reaction.

_____ [1]

(v) Describe the appearance of this organic product.

_____ [2]

Examiner Only	
Marks	Remark

16 Transition metals form coloured complex ions and exist in a range of oxidation states.

(a) In terms of electron structures, explain why zinc is not a transition metal.

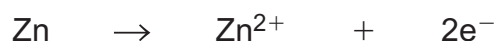
_____ [2]

(b) The VO_2^+ ion can be reduced to V^{2+} using zinc in the presence of an acid. Zinc is oxidised to form Zn^{2+} ions.

(i) Write a half-equation for the reduction of VO_2^+ to V^{2+} .

_____ [2]

(ii) Combine the above half-equation with the following oxidation half-equation:



to give the ionic equation for the reaction.

_____ [2]

(iii) When this reduction is carried out in the laboratory a series of colour changes are observed. Complete the following table:

Ion	Colour
VO_2^+	
	Blue
	Green
V^{2+}	

[4]

Examiner Only

Marks Remark

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Chemistry

Assessment Unit A2 2

assessing

Analytical, Transition Metals, Electrochemistry
and Further Organic Chemistry

[AC222]

WEDNESDAY 23 MAY, AFTERNOON

**MARK
SCHEME**

Section A

- 1** D
- 2** B
- 3** A
- 4** D
- 5** A
- 6** A
- 7** D
- 8** D
- 9** C
- 10** A

[2] for each correct answer

[20]

Section A

**AVAILABLE
MARKS**

20

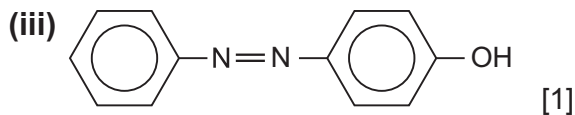
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Section B

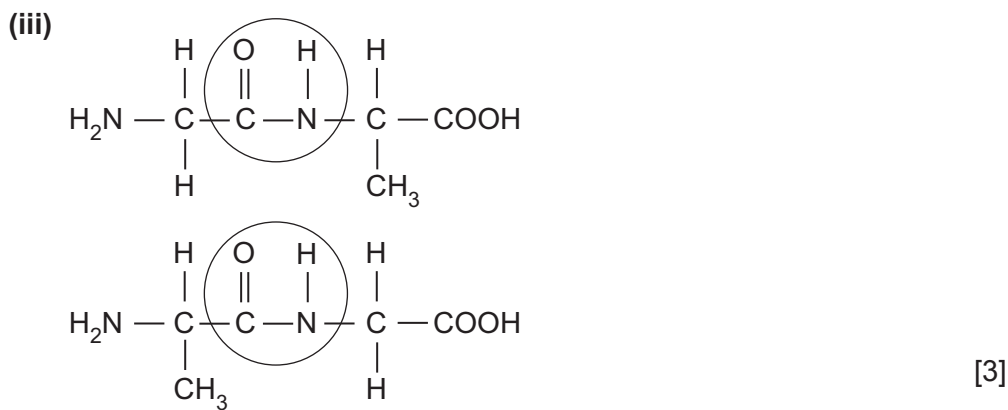
AVAILABLE
MARKS

- 11 (a) (i) A tin/concentrated hydrochloric acid [2]
 B sodium hydroxide [1]
 C sodium nitrite/hydrochloric acid [2] [5]

- (ii) 0–10°C [1]
 benzenediazonium chloride [1] [2]



orange solid [1]
 4-hydroxyazobenzene/4-hydroxyphenylazobenzene [1] [3]



16

- 12 (a)** The potential difference [1] measured when a half-cell is connected to a hydrogen electrode [1] under standard conditions [1]

[3]

- (b)** Na(s)

[1]

- (c)** $2\text{Al} + 3\text{Zn}^{2+} \rightarrow 2\text{Al}^{3+} + 3\text{Zn}$

[2]

$$-0.76 - (-1.66)$$

$$+ 1.66 - 0.76 = +0.90 \text{ V}$$

[1]

[3]

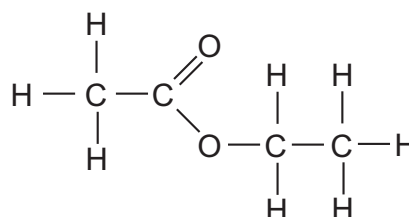
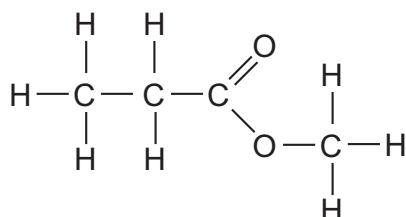
- (d)** $+0.32 = x - (-0.76)$

$$-0.44 \text{ V}$$

[1]

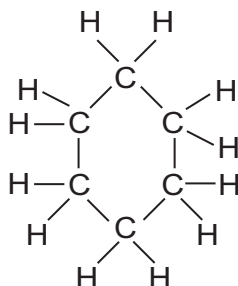
8

- 13 (a) (i)**

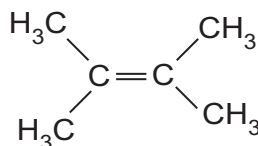


[2]

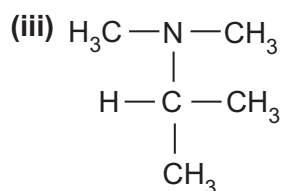
- (ii)**



or



[1]



[1]

- (b) (i)** 2 isotopes [1] of chlorine [1]
or ^{35}Cl and ^{37}Cl [1]

[2]

- (ii)** $\text{C}_3\text{H}_4^{35}\text{ClO}^+$

[2]

- (iii)**

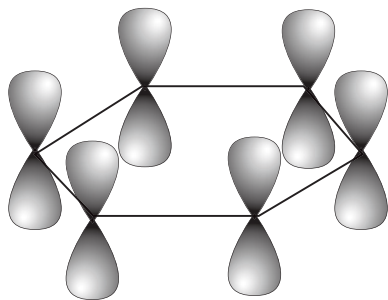
	Peak 1	Peak 2	Peak 3
		1 [1]	1 [1]
	Doublet [1]	Quartet [1]	

[4]

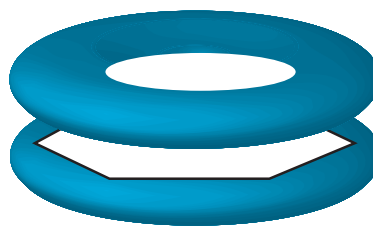
12

14	(a)	(i) $\text{MnO}_4^- + 8\text{H}^+ + 5\text{Fe}^{2+} \rightarrow \text{Mn}^{2+} + 4\text{H}_2\text{O} + 5\text{Fe}^{3+}$	[2]	AVAILABLE MARKS
		(ii) colourless to pink	[2]	
		(iii) moles of manganate(VII) = 0.00048 moles of iron(II) in 25 cm ³ = 0.0024 moles of iron(II) in 250 cm ³ = 0.024 mass of $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ = 0.024 × 278 = 6.672 g % hydrated iron(II) sulfate = 83.4% Each error [−1]	[5]	
	(b)	(i) $[\text{Fe}(\text{H}_2\text{O})_6]^{3+} \rightleftharpoons [\text{Fe}(\text{OH})(\text{H}_2\text{O})_5]^{2+} + \text{H}^+$	[2]	
		(ii) $K_a = \frac{[\text{Fe}(\text{OH})(\text{H}_2\text{O})_5^{2+}][\text{H}^+]}{[\text{Fe}(\text{H}_2\text{O})_6^{3+}]}$	[1]	
		(iii) a rust/brown [1] precipitate [1]	[2]	
		(iv) (potassium) thiocyanate solution [1] blood red [1] solution [1] $[\text{Fe}(\text{SCN})(\text{H}_2\text{O})_5]^{2+}$ [1]	[4]	
	(c)	CO forms a very strong bond with the iron(II) ion in haemoglobin [1] this prevents the transportation of oxygen [1]	[2]	
				20

15 (a)

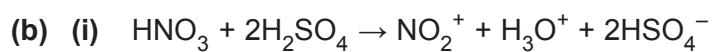


before



after

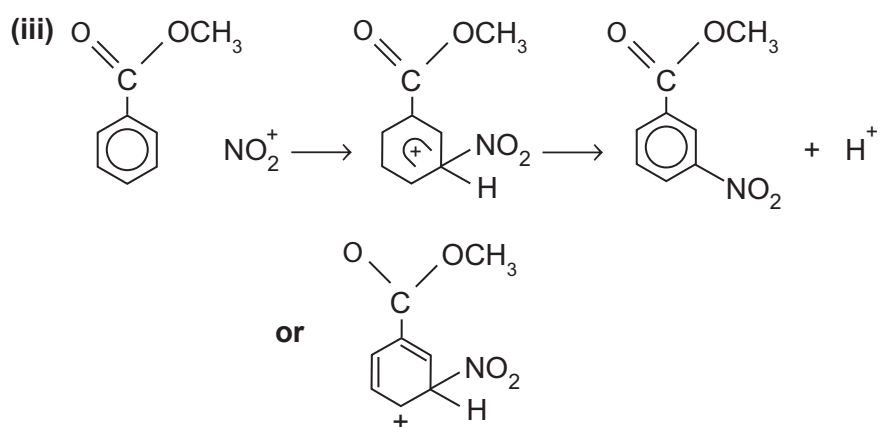
[2]



[2]

(ii) nitronium ion

[1]



[3]

electrophilic substitution

[1]

[4]

(iv) methyl 3-nitrobenzoate

[1]

(v) cream [1] solid [1]

[2]

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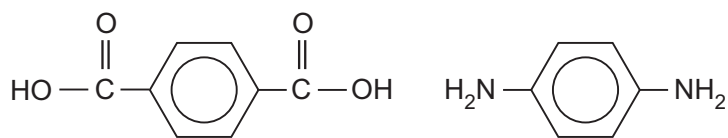
12

- 16 (a)** Does not have an incomplete d-subshell in its (stable) ion Zn^{2+} is $3d^{10}$ [1]
[1] [2]
- (b) (i)** $\text{VO}_2^+ + 4\text{H}^+ + 3\text{e}^- \rightarrow \text{V}^{2+} + 2\text{H}_2\text{O}$ [2]
- (ii)** $2\text{VO}_2^+ + 8\text{H}^+ + 3\text{Zn} \rightarrow 2\text{V}^{2+} + 4\text{H}_2\text{O} + 3\text{Zn}^{2+}$ [2]
- (iii)**
- | Ion | Colour |
|--|-------------------|
| VO_2^+ | Yellow [1] |
| VO^{2+} [1] | Blue |
| V^{3+} [1] | Green |
| V^{2+} | Violet [1] |
- [4]
- (c) (i)** dissolve in water and add (excess) potassium hydroxide [1]
a green-blue precipitate will form and then dissolve to give a deep green solution [1]
add excess hydrogen peroxide and heat [1]
the (green) solution turns yellow [1]
boil the solution (and reduce the volume) [1]
add named acid [1]
orange solution/orange crystals [1]
maximum [6] [6]
- Quality of written communication [2]
- (ii)** moles of chromium(III) chloride hexahydrate = 0.05 [1]
theoretical yield of potassium dichromate = 0.025 mole [1]
actual yield = 0.00997 mole [1]
% yield = 40% [1] [4]

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MARKS

22

- 17 (a)
$$\begin{array}{ccccccccccc} & \text{O} & & \text{O} & \text{H} & & \text{H} & \text{O} & & \text{O} & \text{H} & & \text{H} \\ & || & & || & | & & | & || & & || & | & & | \\ - & \text{C} & - (\text{CH}_2)_4 & - & \text{C} & - & \text{N} & - (\text{CH}_2)_6 & - & \text{N} & - & \text{C} & - (\text{CH}_2)_4 & - & \text{C} & - & \text{N} & - (\text{CH}_2)_6 & - & \text{N} & - \end{array}$$
- Each error [-1] [3]
- (b) (i) $\text{HOCH}_2\text{CH}_2\text{OH}$ [1]
- (ii) ethane-1,2-diol [1]
- (c) (i) 2 [1]
- (ii)



[2]

- (d) 3-methylpent-2-ene [2]

Section B

Total

**AVAILABLE
MARKS**

10

100

120



Rewarding Learning

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General Certificate of Education
2014

Centre Number

71

Candidate Number

Chemistry

Assessment Unit A2 2

assessing

Analytical, Transition Metals, Electrochemistry
and Further Organic Chemistry

[AC222]

TUESDAY 3 JUNE, AFTERNOON



TIME

2 hours.

INSTRUCTIONS TO CANDIDATES

Write your Centre Number and Candidate Number in the spaces provided at the top of this page.

Answer **all fifteen** questions.

Answer **all ten** questions in **Section A**. Record your answers by marking the appropriate letter on the answer sheet provided. Use only the spaces numbered 1 to 10. Keep in sequence when answering.

Answer **all five** questions in **Section B**. Write your answers in the spaces provided in this question paper.

INFORMATION FOR CANDIDATES

The total mark for this paper is 120.

Quality of written communication will be assessed in question **13(d)(i)**.

In Section A all questions carry equal marks, i.e. **two** marks for each question.

In Section B the figures printed down the right-hand side of pages indicate the marks awarded to each question or part question.

A Periodic Table of the Elements, containing some data, is included in this question paper.

**For Examiner's
use only**

Question Number	Marks
Section A	
1–10	
Section B	
11	
12	
13	
14	
15	

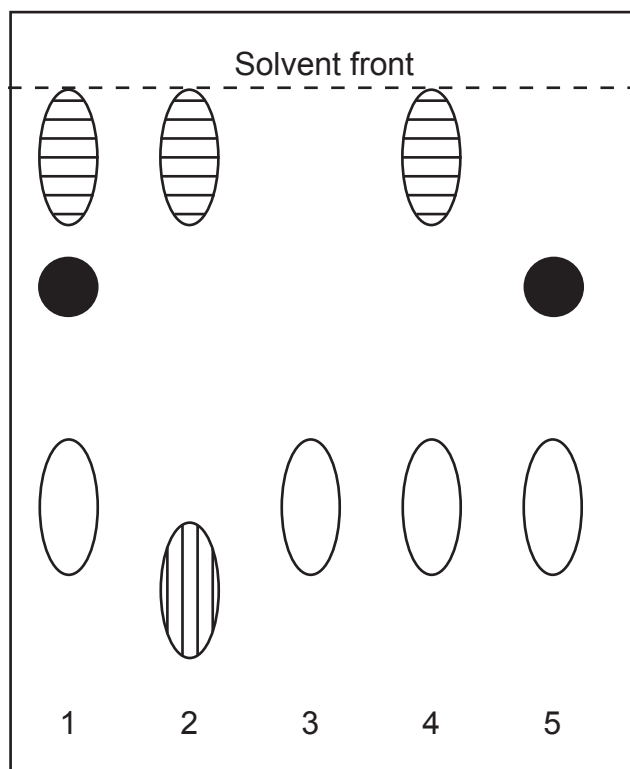
**Total
Marks**

Section A

For each of the following questions only **one** of the lettered responses (A–D) is correct.

Select the correct response in each case and mark its code letter by connecting the dots as illustrated on the answer sheet.

- 1 Paper chromatography was used to identify the amino acids in a polypeptide. The hydrolysed polypeptide is at point 1. The starting point for each amino acid or dipeptide is shown at points 2–5 in the chromatogram below.



Which one of the following mixtures makes up the polypeptide?

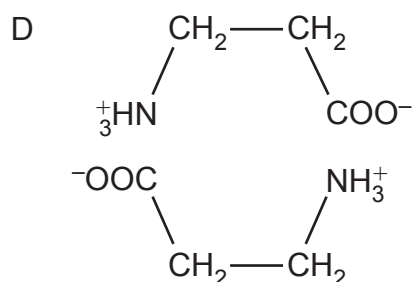
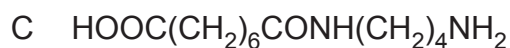
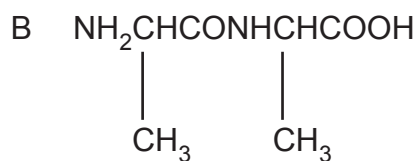
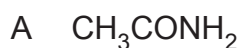
- A 2 + 5
- B 3 + 4
- C 3 + 5
- D 4 + 5

- 2 Oxygen has three isotopes ^{16}O , ^{17}O and ^{18}O . A sample of oxygen was analysed in a mass spectrometer and three groups of peaks were obtained:

group P corresponding to the ion O_2^+
 group Q corresponding to the ion O^{2+}
 group R corresponding to the ion O^+

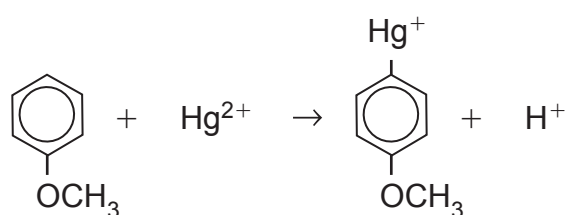
Which one of the following is the order on the mass/charge axis, from left to right of the groups?

- A P Q R
 B P R Q
 C Q P R
 D Q R P
- 3 Which one of the following compounds is a peptide?



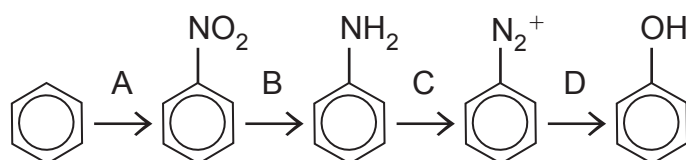
- 4 When 2.6 g of a metal X are added to copper(II) sulfate solution 4.8 g of copper are obtained. The relative atomic mass of X is 52. Which one of the following cations of X is produced?
- A X^+
- B X^{2+}
- C X^{3+}
- D X^{4+}

- 5 The reaction below shows the mercury(II) ion reacting with anisole.

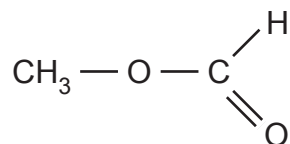


Which one of the following is the role of the mercury ion?

- A Catalyst
- B Electrophile
- C Nucleophile
- D Oxidising agent
- 6 Which one of the steps in the following synthesis requires the reagent $\text{NaNO}_2(\text{s})/\text{HCl}(\text{aq})$?

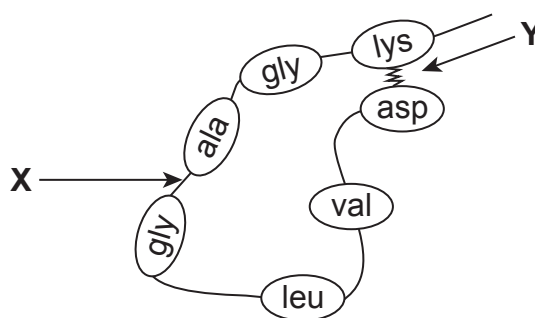


- 7 The structure below represents an organic compound.



This compound is an example of

- A an acid.
 - B an aldehyde.
 - C an ester
 - D a ketone
- 8 The following diagram shows a section of a protein chain. The three-letter words represent the names of amino acids.



Different types of bonds are formed at X and Y.

Which one of the following represents the bonds X and Y?

- | X | Y |
|-----------------|---------------|
| A hydrogen bond | hydrogen bond |
| B hydrogen bond | ionic bond |
| C peptide bond | hydrogen bond |
| D peptide bond | ionic bond |

- 9** A sample of DDT, $C_{14}H_9Cl_5$ was found to contain 0.120 g of carbon. Which one of the following was the mass of chlorine in the sample?
- A 0.127 g
 - B 0.335 g
 - C 0.994 g
 - D 1.01 g
- 10** Which one of the following industrial processes is catalysed by a metal compound?
- A hydrogenation of alkenes
 - B manufacture of sulfuric acid
 - C oxidation of ammonia
 - D reaction of nitrogen with hydrogen

Section B

Answer **all five** questions in the spaces provided.

- 11 Complete the table below to give the colours of the metal ions.

metal ion	colour of aqueous complex
$\text{Cu}^{2+}(\text{aq})$	
$\text{Cr}^{3+}(\text{aq})$	
$\text{Co}^{2+}(\text{aq})$	
$\text{Fe}^{3+}(\text{aq})$	

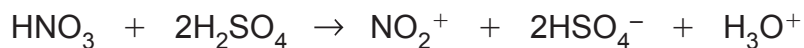
[3]

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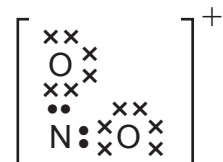
Marks

Remark

- 12** Benzene and its derivatives are nitrated by the nitronium ion which is formed when nitric acid reacts with sulfuric acid.



The nitronium ion has the following electron structure:



- (a) (i)** How does this structure explain the positive charge on the nitronium ion?

_____ [1]

- (ii)** Explain whether the nitronium ion is an electrophile or a nucleophile.

_____ [2]

- (iii)** Draw a flow scheme to show the mechanism for the reaction of benzene with the nitronium ion to form nitrobenzene.

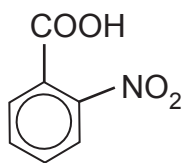
[3]

- (iv)** Explain whether this mechanism is addition or substitution.

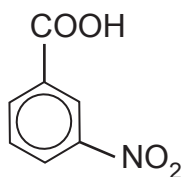
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Marks	Remark

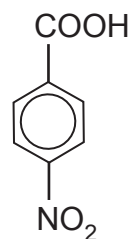
- (b) Benzoic acid is nitrated to produce three products which are shown below together with their melting points and their former names.



ortho-nitrobenzoic acid
145 °C



meta-nitrobenzoic acid
174 °C



para-nitrobenzoic acid
186 °C

- (i) Suggest the systematic names for *ortho*, *meta* and *para* nitrobenzoic acids.

ortho nitrobenzoic acid _____

meta nitrobenzoic acid _____

para nitrobenzoic acid _____ [2]

- (ii) Suggest, using hydrogen bonding, why the melting points of these nitrobenzoic acids increase as the nitro group is further away from the carboxylic acid group.

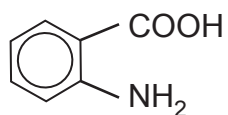
_____ [2]

- (iii) Each of these nitrobenzoic acids can be reduced to amino compounds. Name the reagent used and explain how the amines can be liberated from the reaction mixture.

_____ [2]

Examiner Only	
Marks	Remark

(c) The best known aminobenzoic acid is anthranilic acid.



- (i) When distilled it is decarboxylated to form phenylamine. Suggest what is meant by the term **decarboxylated** and write an equation for the reaction.

[3]

- (ii) Anthranilic acid could be described as an amino acid but it is not part of any protein. Suggest an explanation for this contradictory statement.

[2]

- (iii) Anthranilic acid is diazotised to form a diazonium ion. State the condition necessary for this reaction.

[1]

- (iv) Suggest the structure of the product when the diazotised anthranilic acid is coupled with phenol.

[2]

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Marks	Remark

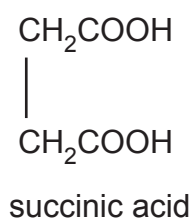
- (v) Explain why it is not possible to couple diazotised butylamine with phenol.

_____ [2]

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Marks	Remark

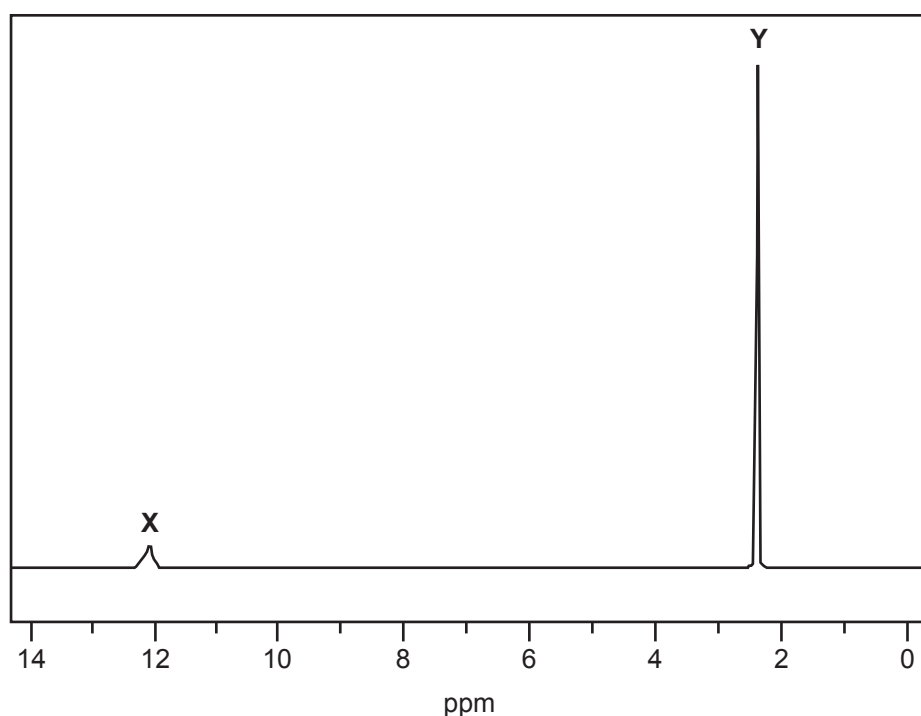
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- 14 Succinic acid is a dicarboxylic acid and was named after the Latin word for amber from which it was first obtained by distillation (Lat: succinum).



It is a crystalline solid with a melting point of 185 °C and is moderately soluble in water and ethanol.

- (a) The high resolution nmr spectrum of succinic acid is shown below.



- (i) Explain why the CH₂ groups do not interact with each other.

_____ [1]

- (ii) Identify which hydrogen atoms give rise to the peaks at X and Y.

X _____ [1]

Y _____ [1]

Examiner Only	
Marks	Remark

(c) When heated with excess ethane-1,2-diol succinic acid forms polymers known as alkyd resins.

(i) Draw the structure of the polymer showing the repeating unit.

[3]

(ii) What is the name given to this type of polymerisation?

[1]

(iii) Suggest why the polymer is acidic.

[1]

(iv) Explain why this polymer is biodegradable.

[2]

(d) Succinic acid reacts with excess ammonia to form the diammonium salt which, when heated, is dehydrated to form the diamide.

(i) Draw the flow scheme for the reactions above.

[2]

(ii) Name the reagent used to convert an amide into a nitrile.

[1]

(iii) Draw the structure of the product formed when the dinitrile is reacted with excess lithal.

[2]

(e) Succinic acid can be analysed by converting it to the diethyl ester and submitting the ester to GLC analysis.

(i) Why is it better to use the ester rather than the acid in GLC analysis?

[2]

(ii) Explain the results expected if the sample of the ester was 90% pure.

[2]

(d) (i) Describe the colour change in the vanadyl half-cell.

[2]

(ii) Describe the colour change in the permanganate half-cell.

[2]

(e) Calculate the reading that will be observed on the voltmeter.

[2]

(f) Explain the purpose of the salt bridge and how it works.

[2]

(g) The electrode potentials of the half-cells are measured using a hydrogen electrode. Describe a hydrogen electrode and state the conditions under which it operates.

[4]

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