

GCE MARKING SCHEME

CHEMISTRY AS/Advanced

SUMMER 2013

INTRODUCTION

The marking schemes which follow were those used by WJEC for the Summer 2013 examination in GCE CHEMISTRY. They were finalised after detailed discussion at examiners' conferences by all the examiners involved in the assessment. The conferences were held shortly after the papers were taken so that reference could be made to the full range of candidates' responses, with photocopied scripts forming the basis of discussion. The aim of the conferences was to ensure that the marking schemes were interpreted and applied in the same way by all examiners.

It is hoped that this information will be of assistance to centres but it is recognised at the same time that, without the benefit of participation in the examiners' conferences, teachers may have different views on certain matters of detail or interpretation.

WJEC regrets that it cannot enter into any discussion or correspondence about these marking schemes.

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GCE CHEMISTRY - CH1

SUMMER 2013 MARK SCHEME

SECTION A

					Tota
	(b)	2.5-6.0			[1]
Q.6	(a)	(dissociates t	o) release H⁺ io	ons	[1]
Q.5	nitrog	en / phosphoru	s (or any other	Group 5 element)	[1]
Q.4					[1]
Q.3	Provides an alternative pathway (1) with lower activation energy / more particles have energy above E_A (1)				
Q.2		ron (1) ticle (1)		(max 1 if three circled, 0 if four or more)	[2]
	numb	er of electrons	6	(all correct 2 marks, 2 correct 1 mark)	[2]
	numb	er of neutrons	8		
Q.1	numb	er of protons	6		

Total [10]

SECTION B

Q.7 (a) percentage Be by mass = 5.03% (1)

division of percentage by A_r for Be and at least one other element as shown below (1)

- AI 10.04 ÷ 27 = 0.3719 → 1.00
- Be 5.03 ÷ 9.01 = 0.5583 → 1.50
- O $53.58 \div 16 = 3.3488 \rightarrow 9.00$
- Si 31.35 ÷ 28.1 = 1.1566 → 3.10

molecular formula = $AI_2Be_3O_{18}Si_6$ or x=3 (1)

- (b) (i) Hess' Law states that where a reaction can occur by more than one route the total enthalpy **change** for each route will be the same [1]
 - (ii) $\Delta H = -393.5 (-395.4) (1) = +1.9 \text{ kJ mol}^{-1} (1)$ [2]
 - (iii) Kyran is **incorrect** as diamond is not the **standard state** of carbon [1]
 - (iv) I mass of diamond = 7.30 g [1]
 - II mass of graphite = $7.30 \div (93/100) (1) = 7.85 g (1)$ [2]

Total [10]

[3]

large jump after 8 electrons (1) [2] (ii) eighth and ninth electrons come from different shells (1) ninth electron is closer to nucleus / has less or no shielding / has greater effective nuclear charge (1) [2] (b) the compound formation has the noble gas atom being ionised (1) ionisation energy of argon is much higher than that of xenon (1) because the outer electron in argon is closer to the nucleus / has greater effective nuclear charge / shielding (1) - 2 max [2] (c) electrons move from lower energy levels to higher energy levels (1) by absorbing specific frequencies of light (1) [2] (d) 1 mol of XeO₃ released 2.5 mol gas products (1) 2.5 mol of gas occupies 24.0 x 2.5 = 60.0 dm³ (1) – follow through error (ft) if candidates calculate the volumes of the two gases separately, then (1) for one gas volume correct and (1) for total volume correct [2] Total [10]

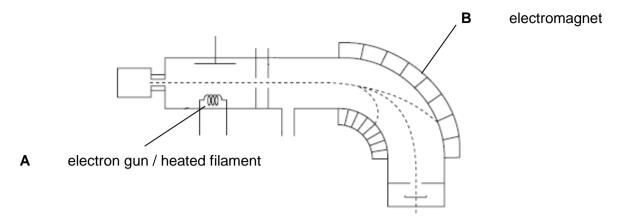
all ionisation energies showing gradual increase and one large jump (1)

Q.8

(a)

(i)

Q.9 (a) (i) both needed



[1]

(ii) electron gun bombards sample and **ionises** atoms/molecules (1)

negatively charged plates / electric field accelerates (positive ions in) sample (1)

electromagnet deflects ions according to mass and charge / m/z (1)

current in electromagnet / electromagnetic field is varied so different mass ions hit detector (1)

[4]

QWC: selection of a form and style of writing appropriate to purpose and to complexity of subject matter (1)

QWC: legibility of text, accuracy of spelling, punctuation and grammar, clarity of meaning (1)

QWC [2]

(b)
$$A_r = (78 \times 12.2) + (79 \times 26.4) + (80 \times 61.4) \div 100$$
 (1) [for method] = 79.5 (1)

(1) for 3 sig figs for sensible answer (above 79.0 and below 80.0) (1) [3]

(c)	(i)	а	81		
		Х	Br /bromine	both needed	[1]
	(ii)	75 mi	nutes = 4 half-l	ives (1)	
		2.72g	\rightarrow 1.36g \rightarrow 0.6	68g → 0.34g → 0.17g (1) – no ft	[2]
					Total [13]

Q.10 (a) *x* = 10

(i) number of moles = $250 \times 0.200 \div 1000 = 0.05 \text{ mol}(1) - \text{ft}$ (b) mass of sodium carbonate = $0.05 \times M_r(Na_2CO_3) = 0.05 \times 286.2$ = 14.31g(1)[2] (ii) any two points from: weigh by difference (1) add less water initially (1) wash out beaker / glass rod / funnel and put water into volumetric flask (1) add water up to mark in volumetric flask (1) - 2 max [2] (c) add few drops of indicator (1) do not accept 'universal indicator' take initial and final reading on burette (1) swirl the conical flask (1) add acid until the indicator changes colour (1) [4] QWC: organisation of information clearly and coherently; use of specialist vocabulary QWC [1] where appropriate. Total [10]

[1]

Q.11 (a) (i)
$$\Delta H = 9 \times (-394) + 10 \times (-286) - (-275)$$
 (1)
= -6131 kJ mol⁻¹ (1) for correct value and (1) for correct sign [3]
(ii) temperature 298K, 25°C (1) pressure 1 atm, 101 kPa (1) [2]
(b) (i) $M_r = (9 \times 12) + (20 \times 1.01) = 128.2$ (1)
number of moles = 1.56 x 10⁻³ mol (1) [2]
(ii) $\Delta H = -50 \times 4.18 \times 42 \div 1.56 \times 10^{-3}$ (1)

$$= -5626698 \text{ J mol}^{-1} = -5627 \text{ kJ mol}^{-1} (1)$$
 [2]

(iii) heat loss to environment / incomplete combustion / not standard conditions [1]

Total [10]

Q.12 (a) killing marine life / killing trees [1] either gas syringe or inverted burette attached to sealed vessel (b) (i) [1] (ii) different surface area would affect rate of reaction [1] (iii) concentration / volume / nature of acid (1) [2] temperature (1) (c) (i) increasing pressure will shift the reaction to side with fewer gas molecules (1) increasing yield of $SO_3(1)$ – reason must be given [2] (ii) L increasing temperature shifts equilibrium in endothermic direction (1) as SO_3 yield is decreased forward reaction must be exothermic (1) [2] Ш increasing temperature increases energy of particles (1) more collisions have energy above activation energy (1) successful collisions occur more frequently (1) can gain first two points from labelled Boltzmann distribution curve [3] Ш e.g. iron in production of ammonia or any valid example [1] (d) atom economy = 100% [1] (i) (ii) any two points from: lower pressure used in B (1) methanol is a renewable starting material (1) higher atom economy in B or less waste in B (1) [ignore reference to cost] 2 max [2] (iii) no effect on position of equilibrium [1] Total [17]

GCE CHEMISTRY - CH2

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Section A

Q.1	С					[1]
Q.2	В					[1]
Q.3	(a)	Calcium chlo	oride			[1]
	(b)	Magnesium	carbonate			[1]
	(c)	Sodium sulf	ate			[1]
Q.4						
		Species	CI●	NH ₃		
	Cla	ssification	Radical	Nucleophile		
	(1 for	each box)				[2]
Q.5	-		g/sterilising spr s/anti-perspira	rays/deodorant soc nts	ks/	[1]
Q.6	Potas	ssium and chl	orine (1)			

They have the largest electronegativity difference (1)	[2]

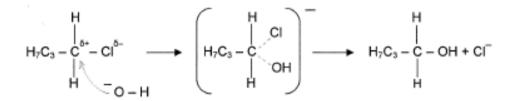
Q.7	(a)	(i)	H H H—C——C- H H	—0—н							[1]
		(ii)	Nickel / plati	num / palladiu	ım						[1]
		(iii)	Potassium / in ethanol ar	sodium hydro nd heat (1)	xide (1))					[2]
		(iv)	Elimination								[1]
	(b)	(i)	H C H	CH ₃ -C H							[1]
		(ii)		ene) unit = 42 nits = <u>1.05 × 7</u> 42		5000	(1)				[2]
	(c)	(i)	Percentage	hydrogen = 4.	6% (1)						
			C <u>22.0</u> 12	H <u>4.6</u> 1.01	Br <u>73.4</u> 79.9	(1)					
			1.83	4.55	0.92						
			2	5	1						
			Formula = C	₂H₅Br (1)							[3]
		(ii)	M _r of compo	und / number	of atoms	s of a	ny elen	nent in o	compoun	d	[1]

Total [12]

Q.8 (a) e.g. damages liver/ damages pancreas/causes cancer/causes skin disorders/ short-term effects (1)

e.g. more traffic accidents/violent behaviour/criminal behaviour (1) [2]

(b) (i) Nucleophilic substitution / hydrolysis (1)



Reactants: Polarisation curly arrow (Incorrect st	Intermediate (1) (1) (accept curly arrow to show (1) C – CI breaking instead of intermediate) arting material or product maximum 2 marks from 3 for mechanism)	[4]
(ii)	Peak at 650–800 cm ⁻¹ due to C – Cl bond will be gone (1) Peak at 2500–3500 cm ⁻¹ due to O – H bond / 1000–1300 cm ⁻¹ due to C – O bond will be present (1)	[2]
(c) (i)	ОН	[1]
(ii)	Structural / positional / chain	[1]
(iii)	Colour change from orange to green	[1]
(iv)	Concentrated sulfuric acid / aluminium oxide (1) $CH_3CH_2CH_2CH_2OH \longrightarrow CH_3CH_2CHCH_2 + H_2O$ (1)	[2]
(d) (i)	C – F bond stronger than C – CI bond (1) C – CI bond breaks (in stratosphere) forming CI● which reacts with ozone (1)	[2]
(ii)	Some CFCs still being used / CFCs take a very long time to reach the ozone layer / other substances deplete the ozone layer	[1]
	Total	[16]

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Q.9	(a)	A mixture of (many) hydrocarbons / alkanes	[1]

(b)
$$C_4H_{10} + 6\frac{1}{2}O_2 \longrightarrow 4CO_2 + 5H_2O$$
 [1]

- (c) $109\frac{1}{2}^{\circ}$ [1]
- (d) H_2O has 2 bonding and 2 lone pair of electrons (1)

 CH_4 has 4 bonding pairs only (1)

Repulsion between lone pairs and bond pairs is greater than between bond pairs and bond pairs (1) [3]

QWC The information is organised clearly and coherently, using specialist vocabulary where appropriate QWC [1]

- (e) (i) Butane is higher because it has more van der Waals' forces between molecules [1]
 - (ii) Regular array of metal ions surrounded by a 'sea' of delocalised valence electrons (1)

Strong attraction between the positive ions and the delocalised electrons (1) (Can be obtained from labelled diagrams)

Malleable because when a force is applied the layer of metal ions slide over each other forming a new shape (1)

Conduct electricity since under a potential difference the delocalised electrons flow / the delocalised electrons flow towards the positive potential

(1) **[4]**

QWC Legibility of text; accuracy of spelling, punctuation and grammar, clarity of meaning QWC [1]

Total [13]

Q.10	(a)	(i)	Chlorine – gas Iodine – solid	[1]
		(ii)	Chlorine – brown/orange solution (1) lodine – no change / no reaction (1) $Cl_2 + 2KBr \longrightarrow Br_2 + 2KCI$ (1) (Accept ionic equation)	[3]
	(b)		en loses electrons therefore oxidised / oxidation state changes from -2 to fore is oxidised (1)	0
			ine gains electrons therefore reduced / oxidation state changes from 0 to fore is reduced (1)	-1 [2]
	(c)	(i)	Boiling temperatures increase as relative molecular mass increases / number of electrons increases / down group (1)	
			HF has a higher boiling point than expected (1)	[2]
		(ii)	Group 7 hydrides contain more dipole-dipole forces as group descende	
			but HF contains hydrogen bonding between molecules (1)	(1)
			Hydrogen bonds are stronger therefore HF's boiling temperature is grea / need more energy to break (1)	ter [3]
			QWC Selection of a form and style of writing appropriate to purpose and complexity of subject matter QWC	

(iii) HCI more polar than SiH₄ therefore intermolecular forces are stronger / dipole greater in HCI / CI more electronegative than Si [1]

Total [13]

12

Q.11	(a)	(i)	2Ca + O ₂ → 2CaO	[1]
		(ii)	Ca x 0	
			(1)	
			forming Ca^{2+} and O^{2-} ions (1)	[2]
	(b)	(i)	Ca(OH) ₂	[1]
		(ii)	8 – 14	[1]
	(C)	Ca ²⁺ (aq) + $CO_3^{2-}(aq) \longrightarrow CaCO_3(s)$	[1]
	(d)	(i)	Magnesium disappears / gets smaller (1) Effervescence / bubbles (of hydrogen) (1) Heat given off (1) (Accept any 2 points)	[2]
		(ii)	Moles Mg = $\frac{0.503}{24.3}$ = 0.0207 (1) Moles HCI = 0.0414 (1) Volume HCI = $\frac{0.0414}{1.6}$ = 0.0259 dm ³ (1)	[3]
		(iii)	Volume $H_2 = 0.0207 \times 24 = 0.497 \text{ dm}^3$	[1]
		(iv)	Add aqueous silver nitrate (1) White precipitate forms (1)	[2]
	(e)	Less	reactive (1)	
		Elect	rons in beryllium more difficult to lose / ionisation energy is higher (1)	[2]

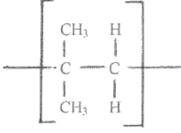
(Need reason to get first mark but accept less reactive as reactivity increases down group / outer electron has less shielding etc. for 1 mark)

Total [16]

GCE CHEMISTRY – CH4

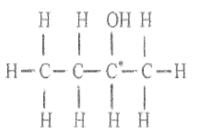
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Q.1	(a)	(i)	(2-)Methylpropan-2-ol	[1]
		(ii)	30.1 / 30	[1]
		(iii)	(Concentrated) sulfuric acid / phosphoric acid / aluminium oxide / pumice	[1]
		(iv)		



(with or without n)

(v)



(1) for structure, (1) for asterisk	[2]

[1]

	(vi)	I	acidified potassium dichromate / H^+ , $Cr_2O_7^{2-}(aq)$ [1]	
		II	ethanal has a C = O bond at 1650-1750 cm ⁻¹ (metaldehyde does not have this bond) (1)	
			metaldehyde has a C – O bond at 1000-1300 cm ⁻¹ (1) [2] (ethanal does not have this bond)	
(b)	(b) (i) Reagent 2,4-dinitrophenylhydrazine / 2,4-DNP OR iodine / NaOH or KI / NaOCI (1)			
	(Observati	ion yellow / orange / red precipitate OR yellow precipitate (1) [2]	
	(ii) I	Reagent	ethanol / sulfuric acid OR NaHCO ₃ OR Ag^+/NH_3 / Tollens' (1)	
	(Observat	ion sweet smelling liquid OR effervescence OR silver mirror (1) [2]	
	Total [13			

Q.2	(a)	React with iron(III) chloride solution Purple solution with phenol, no reaction with methyl propenoate OR React with aqueous bromine / bromine water White precipitate with phenol (and bromine decolourised), bromine decolourised with methyl propenoate			
		(1) for reagent and (1) for observation with compound	[2]	
	(b)	(i)	It absorbs all colours except yellow / absorbs the blue end of the spectrum and reflects yellow – do not accept 'emits'	[1]	
		(ii)	Tin / iron and concentrated hydrochloric acid	[1]	
	(c)	(i) Moles of 2,4-dinitrophenol = $7.36/184 = 0.040$ (1) Moles of 2,4-dinitrophenyl ethanoate = $7.91/226 = 0.035$ (1)			
			Percentage yield = $0.035 \times 100 / 0.040 = 87.5 / 88 \%$ (1)	[3]	
		(ii)	R _f value is given bydistance travelled by the 2,4-dinitrophenol distance travelled by the solvent front(1)		
			$= \frac{2.8}{5.0} = 0.56 $ (1)	[2]	
	(d)	(i)	Nickel / platinum	[1]	
		(ii)	The –OH groups are able to hydrogen bond with water (1)	[2]	

The –OH groups are able to hydrogen bond with water (1) but these are a very small part of the 'urushiol' molecule (1) [2]

Total [12]

Q.3 (a) (i) 48.5 / 49 %

(ii) Find a use for the calcium sulfate

(b) Total volume of aqueous sodium hydroxide needed = $\frac{26.40 \times 250}{25.00}$ = 264.0 cm³ (1)

from the graph this is equivalent to 0.011 mole of the acid (1)

$$\therefore M_{r} \text{ of the acid} = \frac{\text{mass}}{\text{no. of moles}} = \frac{2.31}{0.011} = 210 \quad (1)$$

$$C_{6}H_{8}O_{7} \cdot n H_{2}O = 210$$

$$\uparrow \\ 192 \quad \therefore n = 18 \quad (1)$$
since M_r of water is 18 \quad n = 1 \quad (1) \quad [5]

[1]

[1]

[1]

 (c) The two 'ends' of the double bond have different groups bonded to the carbon atoms (of the double bond) / they have different structural formulae, so cannot be stereo / geometric isomers [1]

(e)
$$C_5H_6O_5 \rightarrow CH_3COCH_3 + 2CO_2$$
 [1]

$$H - C - C - O - H$$

$$H - H$$

+ +

* *

(g) (Fractional) distillation / (preparative) gas chromatography / HPLC [1]
 (h) (i) eg An optically active isomer that will rotate the plane of polarised light / an isomer with a chiral centre [1]
 (ii) An equimolar mixture of both enantiomers (that has no apparent effect on the plane of polarised light) [1]
 Total [15]

Q.4 (a) Benzene is a compound whose molecules contain six carbon atoms bonded in a (hexagonal) ring (1) All the carbon to carbon bond lengths are equal / intermediate (1) Each carbon atom is bonded to two other carbon atoms and a hydrogen atom (1) by σ -bonds (1) All the C - Ĉ - C angles are the same / 120° (1) The remaining p electron of each carbon atom / overlap of p orbitals forms a delocalised cloud of electrons / π -system (1) above and below the plane (1) Credit can be gained from labelled diagram [*Candidates can gain a maximum of (4) for this part*]

This delocalisation increases the **stability** (1) of the molecule and this stability is maintained by benzene undergoing substitution reactions in preference to addition reactions (that would destroy the delocalised system)

The π -cloud is **electron rich** and will be attracted to electron deficient electrophiles (1) [*Candidates can gain (2) for this part*]

QWC Selection of a form and style of writing appropriate to purpose and to complexity of subject matter (1)

Legibility of text; accuracy of spelling, punctuation and grammar; clarity of meaning. (1) QWC [2]



catalyst eg AlCl₃ (anhydrous) (1)

[2]

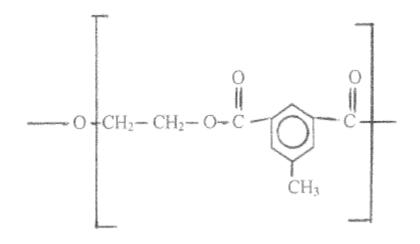
[6]

- (c) (i) (There are two environments for the protons), the 3 aromatic protons at ~6.8 δ and the 9 methyl / aliphatic protons at ~ 2.3 δ (1) These give a peak area of 3:9, ie.1:3 (1) These environments are separate / discrete (1) therefore no splitting pattern [3]
 - (ii) Dissolve in the minimum volume (1) Of hot water (1) (Filter hot) (1) Cool (1) Filter (1) Dry (1)

(up to 4 max but candidates must give the first two points in order to gain full credit)

[4]

(iii)



[1]

(iv) Reagent S is alkaline potassium manganate(VII) (1)
 Reagent T is eg hydrochloric acid (1)

[2]

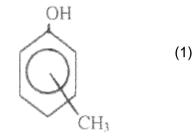
Total [20]

- **Q.5** (a) (i) The **nitrogen atom** has a **lone pair** of electrons making it an electron pair donor / proton acceptor
 - (ii) Compound **L** must contain the grouping -N-C (1)

The nitrogen atom must be bonded directly to the ring as a (primary) aromatic amine is formed on hydrolysis (1)

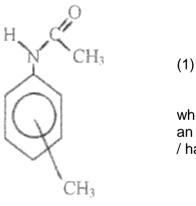
As the hydrolysis compound is a phenol (and has an OH group directly bonded to the ring) a methyl group must also be bonded directly to the ring, as the molecular formula is C_7H_8O / the compound has the structure

ΗΟ



The compound is likely to be an amide, as these are hydrolysed by bases to amines (1)

A suggested formula is

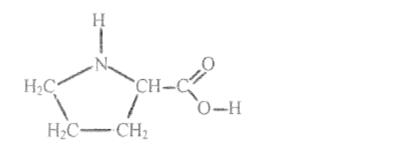


which is $C_9H_{11}NO$, an isomer of cathinone / has M_r of 149(1)

[6]

QWC Information organised clearly and coherently, using specialist vocabulary where appropriate QWC [1]

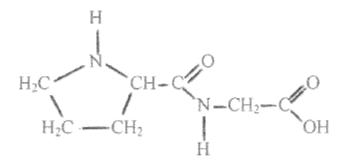
(b) (i)



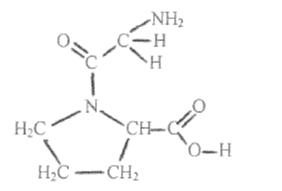
[1]

[1]

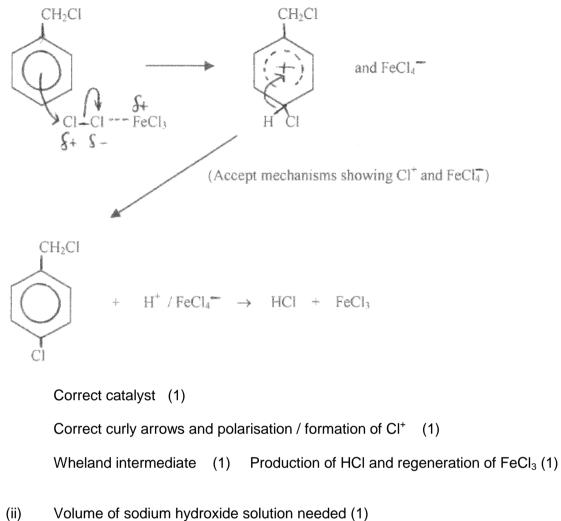
(ii)



OR



(i)



- Volume of sodium hydroxide solution needed (1)
 How long to reflux (1)
- (iii) The aromatic C Cl bond is stronger than the aliphatic C Cl bond (1) This is because a p-electron(s) of the chlorine atom in the aromatic compound becomes part of / incorporated into the delocalised π system of the ring (1) [2]
- (iv)



(1)

chlorine has two isotopes 35/37 in a 3:1 ratio (1)

[2]

[4]

[2]

Total [20]

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Q.1	(a)	(sodiu	Name of any commercially/ industrially important chlorine containing compound e (sodium) chlorate(I) as bleach/ (sodium) chlorate(V) as weedkiller/ aluminium chlo catalyst in halogenation			
		,	- do not accept (CFCs	[1]	
	(b)	(i)	$\mathcal{K}_{c} = \frac{[HI]^{2}}{[H_{2}][I_{2}]}$	must be square brackets	[1]	
		(ii)	$K_{\rm c} = \frac{0.11^2}{3.11^2} = 1.25 \times 10^7$	³ follow through error (ft)	[1]	
		(iii)	$K_{\rm c}$ has no units	ft	[1]	
		(iv)	when temperature incre	eases $K_{\rm c}$ increases (1)		
			this means equilibrium has moved to RHS / increasing temperature favours endothermic reaction (1)			
			therefore ∆H for forward (mark only awarded if n		[3]	
<i>(c)</i> (i)		(i)	+2		[1]	
		(ii)	co-ordinate/ dative (cov	alent)	[1]	
		(iii)	pink is $[Co(H_2O)_6]^{2+}$ and	blue is $[CoCl_4]^{2-}$ (1)		
			(ligand is) CI^{-} (1) (addition of HCl sends) equilibrium to RHS (1) [Co(H ₂ O) ₆] ²⁺ shown as octahedral [with attempt at 3D] (1)			
		(iv)				
			$[CoCl_4]^{2-}$ shown as tetra	ahedral/ square planar (1)	[2]	

Total [14]

Q.2	(a)	(i)	tangent drawn at t = 40 (1)		
			rate calculated 0.017 to 0.027 (ign	ore units) (1)	[2]
		(ii)	as reaction proceeds less collision	s (per unit time) occur	[1]
	(b)	(i)	1 st order shown by:		
			calculation of rates at at least 2 co	ncentrations (1)	
			statement rate α concentration (1)		
			OR		
			constant half-life (1)		
			half-life is 24 minutes (1)		[2]
		(ii)	rate = $k[N_2O_5]$ (1)		[1]
		(iii)	ii) $k = rate (from (i))/ [N_2O_5] (from graph) (1) (mark correct numbers – no need to check evaluation)$		
			units = minutes ⁻¹ (1)	ft from (ii)	[2]
		(iv) (student A more likely to be correct) reaction is 1^{st} order and $1 [N_2O_5]$ rate determining step			nvolved in [1]
	(c)	correct curve starting at 100 kPa and becoming horizontal (1)			
		horizontal at 250 kPa (1)			[2]
				То	tal [11]

- **Q.3** (a) an acid is a proton / H^+ donor
 - (b) $pH = -log[H^+] / negative log of hydrogen ion concentration$
 - (c) a low pH corresponds to a high concentration of $H^+(1)$

a strong acid is totally dissociated whilst a weak acid is partially dissociated (1)

need to consider concentration (of acid solution) as well as strength of the acid (1)

a concentrated solution of a weak acid could have a lower pH than a dilute solution of a strong acid (1) [4]

QWC Accuracy of spelling, punctuation and grammar QWC [1]

(d) (i)
$$K_{a} = \frac{[HCOO^{-}][H^{+}]}{[HCOOH]}$$
 [1]

(ii)
$$1.75 \times 10^{-4} = \frac{x^2}{0.1}$$
 (1)
x = 4.183 × 10⁻³ (1)

(ii) RCOOH
$$\rightleftharpoons$$
 RCOO⁻ + H⁺ and RCOONa \rightarrow RCOO⁻ + Na⁺ (1)

added H^+ removed by salt anion/ A^- + $H^+ \rightarrow HA$ (1)

added OH⁻ removed by acid/ OH⁻ + HA
$$\rightarrow$$
 A⁻ + H₂O (1) [3]

Total [15]

[1]

[1]

Q.4 (a) diagram with labels to show

(b)

(C)

 H_2/H^+ shown in electrode (1) platinum (in both electrodes) (1) $Fe^{2+}(aq)$ and $Fe^{3+}(aq)$ (1) high resistance voltmeter (1) salt bridge (1) gas at 1atm pressure, solutions of concentration 1 mol dm⁻³, temperature 298K (1) [any 5] [5] (i) successive ionisation energies increase gradually/ the energies of the d orbitals are similar [1] 1s²2s²2p⁶3s²3p⁶4s²3d¹⁰ / 3d¹⁰4s² (ii) [1] (iii) after 4s electrons lost 3d is full/ stable/ d electrons ionisation energy very high [1] violet solution contains V^{2+} (1) (i) SEP Zn²⁺/Zn is more negative than VO_3^{-}/VO^{2+} and VO^{2+}/V^{3+} and therefore releases electrons/ VO_3^{-}/VO^{2+} and VO^{2+}/V^{3+} are more positive than Zn^{2+}/Zn and are stronger oxidising agents (1) V^{2+} cannot be reduced (to V) since SEP is more negative than Zn^{2+}/Zn (1) [3] (ii) [1] 1.1V (ignore sign) $Zn(s) \rightarrow Zn^{2+}(aq) + 2e / Zn(s) \rightleftharpoons Zn^{2+}(aq) + 2e$ with some indication of direction (iii) [1] (iv) if Zn²⁺(aq) concentration increased equilibrium moves to LHS (1) so electrode potential becomes less negative (1) [2]

(d) (i) 2.74×10^{-3} (mol) (ii) 1.37×10^{-3} (mol)

(ii)
$$1.37 \times 10^{-3}$$
 (mol) [1]

(iii)
$$M_r KIO_3 = 214.1$$

moles KIO_3 = 0.978/ 214.1 = 4.57 \times 10 3 in 250 cm^3

$$4.57 \times 10^{-4}$$
 in 25 cm³ [1]

(iv)
$$1.37 \times 10^{-3}/4.57 \times 10^{-4} = 3$$
 (1)

equation 1 is correct since 3 moles of iodine formed (mark awarded for reason) (1) [2]

Total [20]

[1]

Q.5	(a)	(i)	atomisation of magnesium / vaporisation of magnesium	[1]	
		(ii)	increased ratio positive charge on nucleus: number of electrons	[1]	
		(iii)	is positive because the (negative) electron is repelled by negative species	[1]	
		(iv)	lattice enthalpy is –3835(kJ mol ⁻¹) numerical value (1) negative sign (1)	[2]	
	(b)	(i)	gases are more random/ have more disorder / move more freely and therefore have higher entropy	e a [1]	
		(ii)	$\Delta S = 21.8 (JK^{-1}mol^{-1})$	[1]	
		(iii)	$\Delta G = \Delta H - T \Delta S (1) $ ft from (ii)		
			ΔG must be –ve if reaction to be spontaneous/ to calculate T make ΔG = 0 (1)		
			0 = 318000 – T 21.8 T = 14587/14600 (K) (1)	[3]	
	(C)	use of	f aqueous sodium hydroxide (1)		
		white	precipitate for all possible ions (1)		
		excess aqueous sodium hydroxide – precipitate dissolves for Pb^{2+} and Al^{3+} (1)			
		use of aqueous (potassium) iodide/ hydrochloric acid/ sulfuric acid / soluble			
		chloride/ soluble sulfate (1)			
		result – yellow ppt for $Pb^{2+} + I^{-}$ and no ppt for AI^{3+} / white ppt for $Pb^{2+} + CI^{-}$ or SO_4^{2-}			
		and no ppt for Al ³⁺ [result for both needed] (1)			
		QWC Organisation of information clear and coherent (1)			
		Use of specialist vocabulary (1)			
	(d)	(i)	diagram to show central AI, 4 CI and 4 shared pairs of electrons, all CI outer electrons, dative pair identifiable	[1]	
		(ii)	chlorination of benzene (1) produces CI^{+} as electrophile (1)		
		OR gives ionic liquids (1) with low vapour pressure/ non-volatile/ do not evaporate			
		in use (1)			
			OR catalyst (1) in polymerisation of alkenes (1)	[2]	
			Total	[20]	



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