

GCE MARKING SCHEME

CHEMISTRY AS/Advanced

SUMMER 2012

INTRODUCTION

The marking schemes which follow were those used by WJEC for the Summer 2012 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.

CH1 Section A

1.		>	\bigcirc							
										[1]
2.	1/12 th	mass	of one	atom c	of carbo	on-12.				[1]
3.	С									[1]
4.	(a)		C <u>12.1</u> 12		O <u>16.2</u> 16		CI <u>71.7</u> 35.5		(1)	
			1.01		1.01		2.02			
			1		1		2			
			Form	ula = C	OCI ₂				(1)	[2]
	(b)	M _r / m comp	nolecul ound	ar mas	s / num	nber of	atoms o	f any	elemen	t in [1]
5.	(a)	С	в	D	Е	Α				[2]
		(1 ma	irk if on	ie mista	ake e.g	j. A in v	vrong pla	ace)		
	(b)	z							(1)	
		Si is i be aft large Group	n Grou er the jump b o 4	p 4 the fourth i efore t	erefore onisation he four	large ju on, not th ionis	Imp in ic before i ation er	onisat t / W , nergy	on ener X and V so canr (1)	rgy would Y have a not be in
										[2]
										Total [10]

Section B

6.	(a)	(i)	12				[1]
		(ii)	14				[1]
		(iii)	Percentage / each isotope	abundance / rat	io / proportion	of	[1]
	(b)	(i)	0.125 g				[1]
		(ii)	e.g. Cobalt-6 radio carbon thyroid gland	0 (1) in radiother dating (1) / lodir ls (1)	rapy (1) / Carb ne-131 (1) as a	on-14 (1) a tracer in	in [2]
	(c)	(i)	Atoms are hi an electron g	t by an electron b jun (and lose ele	oeam / electroi ctrons)	ns fired fro	om [1]
		(ii)	To be able to they can be o - no ci	accelerate the indeflected by a material deflected by a material for 'so that a	ons (to high sp agnetic field atoms can be c	eed) / so	that ' [1]
		(iii)	They are def m/z ratio	lected by a magr	netic field / acc	ording to	the [1]
	(d)	1s	2s	2р	3s	Зр	
		↓↑	↓↑	$\downarrow \uparrow \ \downarrow \uparrow \ \downarrow \uparrow$			
							[1]
	(e)	(i)	Mg ₃ N ₂ + 6⊢	l₂0 →	3 Mg(OH) ₂ +	2 NH ₃	[1]
		(ii)	moles Mg(Ol	H) ₂ = 1.75/58.32	= 0.0300 (1)		
			moles Mg ₃ N ₂	₂ = 0.0100 (1)			
			mass Mg_3N_2	= 0.01 x 100.9 =	: 1.01 g (1)		[3]

- must be 3 significant figures to gain third mark

Total [14]

7.	(a)	Plott	ing	(2)				
		Best	fit line	(1)	[3]			
	(b)	(i)	С	(1)				
			Curve steeper	(1)	[2]			
		(ii)	Concentration of acid is greatest		[1]			
	(c)	44 c	m^{3} (±1 cm ³)		[1]			
	(d)	Mole	es Mg = 0.101/24.3 = 0.00416	(1)				
		Mole	es HCI = 2 x 0.02 = 0.04	(1)	[2]			
	(e)	(i)	Mg is not the limiting factor /					
			Mg now in excess / HCI not in exces	S	[1]			
		(ii)	Moles acid = 0.5 x 0.04 = 0.02	(1)				
			Volume $H_2 = 0.01 \times 24 = 0.24 \text{ dm}^3$					
			- correct unit needed	(1)	[2]			
	(f)	Low	er the temperature of the acid	(1)				
		Rea	ctants collide with less energy	(1)				
		Few	Fewer molecules that have the required activation energy (1)[3]					
	or	Use of su	Use pieces of magnesium (1) less surface area (1) less chance of successful collisions (1)					
	QWC Selection of a form and style of writing appropriate to purpose and to complexity of subject matter. [1]							

Total [16]

8.	(a)	Oil is	non-renewable / will run out (1)					
		Contribution of CO_2 to global warming (1)						
		Oil ha	s other important uses (1)	[2]				
		(Maxii	mum 2 marks)					
	(b)	(i)	Power stations / fossil fuels used to generate the electricity needed to make $H_2^{}(1)$					
			Resulting in CO ₂ formation (global warming) / acid rai	n (1)				
			Manufacture of car produces pollution (1)	[2]				
			(Maximum 2 marks)					
			QWC Legibility of text; accuracy of spelling, punctuat and grammar, clarity of meaning	ion [1]				
		(ii)	Disagree, no fuel is 100% safe /					
			petrol can burn explosively (Accept agree if valid reason given e.g. in terms of lives					
			being lost)	[1]				
	(c)	(i)	Hydrogen since frequency is inversely proportional to wavelength / smaller wavelength	[1]				
		(ii)	Hydrogen since energy is proportional to frequency / greater frequency / E = hf	[1]				
	(d)	In Ne nuclea	greater shielding of <i>outer</i> electron (1) outweighs larger ar charge (1) / He has greater effective nuclear charge	(1) /				
			- max 1 if no reference to <i>outer</i> electron	[2]				
		(Maximum 2 marks)						
	(e)	(i)	²¹⁸ Po	[1]				
		(ii)	Since radon is a gas / inhaled, α particles will be given in the lungs (which may cause cancer)	n off [1]				

Total [12]

9. (a) Low temperature (1)As temperature is decreased equilibrium moves in exothermic direction. (1)High pressure (1)As pressure is increased equilibrium moves towards side with smaller number of gas moles (1)[4] QWCThe information is organised clearly and coherently, using specialist vocabulary where appropriate [1] Δ Hreaction = Δ H_f products – Δ H_f reactants (1)(b) $-46 = \Delta H_{f}$ ethanol – (52.3 – 242) ΔH_f ethanol = -46 - 189.7 (1) ΔH_{f} ethanol = -235.7 kJ mol⁻¹ (1) [3] Bonds broken = 1648 + 612 + 926 = 3186 kJ mol⁻¹ (C) (1)Bonds formed = $2060 + 348 + 360 + 463 = 3231 \text{ kJ mol}^{-1}$ (1) Δ H reaction = 3186 – 3231 = -45 kJ mol⁻¹ (1)[3] Average bond enthalpies used (not actual ones) (d) (i) [1] (ii) Yes, since answers are close to each other [1] Catalyst is in different (physical) state to reactants [1] (e) (f) (i) exothermic reaction [1] (ii) catalysed reaction [1] Energy

Extent of reaction

Total [16]

(a)	Weig disso	hing bottle would not have been washed / diffic lve solid in volumetric flask / final volume would	cult to d not				
	nece	ssarily be 250 cm ³		[1]			
(b)	Pipet	te		[1]			
(c)	To sh when	To show the end point / when to stop adding acid / when it's neutralised					
(d)	So th	So that a certain volume of acid can be added quickly before					
	titratio	ons / to give a rough idea of the end point	urale	[1]			
(e)	To ob	otain a more reliable value		[1]			
(f)	(i)	Moles = 0.730/36.5 = 0.0200	(1)				
		Concentration = $0.02/0.1 = 0.200 \text{ mol dm}^{-3}$	(1)	[2]			
	(ii)	Moles = 0.2 x 0.0238 = 0.00476		[1]			
	(iii)	0.00476		[1]			
	(iv)	0.00476 x 10 = 0.0476		[1]			
	(v)	$M_r = 1.14/0.0476 = 23.95$		[1]			
	(vi)	Lithium		[1]			
		 mark consequentially throughout (f) 					
			Tota	al [12]			

Section B Total [70]

10.

CH2

SECTION A

Q.1	(a)	C ₁₉ H ₄₀	[1]
	(b)	$C_{19}H_{40} \ \to \ C_8H_{18} \ + \ C_{11}H_{22} \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ $	[1]
Q.2	2-chlo	robutane	[1]
Q.3	H ₃ C H		[1]
Q.4	any ni	umber in range 1 to 6	[1]
Q.5	(a)	maximum mass = 44-45 (g)	[1]
	(b)	(less solute would form as a solid) because more will remain in the solution	[1]
Q.6	(a)	iodine force is Van der Waals/ induced dipole-induced dipole (1)	
		diamond force is covalent bond/ description of attractive forces in a covalent bond (1)	[2]
	(b)	diamond would have a higher sublimation temperature because it has stronger forces/ forces are harder to break	as [1]

Section A Total [10]

SECTION B

Q.7 (a) (i) one σ bond/ description of σ bond/ diagram to show overlap of s orbitals (1)

one π bond/ description of π bond/ diagram to show sideways overlap of p orbitals (1)

[2]

 (ii) joining of many/lots of (small) units or many alkenes / molecules to make a large/long unit/ molecule [1]



[1]

	(iv)	C ₄ H ₅ Cl	[1]
(b)	(i)	BF3 is planar triangular/ trigonal planar (1)	
		NH ₃ is pyramidal/ trigonal pyramid (1)	[2]
	(ii)	BF_3 has 3 bond pairs (1)	
		$\rm NH_3$ has 3 bond pairs and 1 lone pair (1)	[2]
		QWC the information is organised clearly and coherently, usin specialist vocabulary where appropriate	ng [1]
(c)	(i)	co-ordinate/ dative covalent/ dative - no credit for 'covalent'	[1]
	(ii)	109½° (accept any in range 109°-110°)	[1]
	(iii)	4 bond pairs/ bonds (around B) - no credit for 'tetrahedral'	[1]

Total [13]

Q.8 (a) (i) % H = 14.3 (1) C : H = $\frac{85.7}{12.0}$: $\frac{14.3}{1.01}$ = 7.14 : 14.16 (1) empirical formula = CH₂ (1) [3] (ii) M_r = 42/ largest fragment has mass 42 (1)

$$(CH_2 = 14)$$
 therefore molecular formula = $C_3H_6(1)$ [2]

(iii)
$$CH_3$$
 is present [1]







[3]

Q.9 (a) apparatus in which reaction can occur, e.g. flask/ test tube, and delivery/ rubber tube (1)

apparatus in which to measure volume of gas, e.g. over water with measuring cylinder/ gas syringe (1) [2]

- (b) (i) fewer **moles** of barium used / barium has a higher A_r [1]
 - (ii) reaction faster/ more vigorous/ less cloudy solution formed with barium (1)

because ionisation energy of barium is less/ electrons lost more easily from barium/ barium is lower in the group/ barium hydroxide is more soluble (1) [2]

(c) flame test (1) brick red for calcium **and** (apple) green for barium (1)

OR

add sulfuric acid/ sodium sulfate solution/ potassium sulfate solution (1)

white precipitate with Ba^{2+} , less precipitate/ no precipitate with Ca^{2+} (1) [2]

(d) electrons correct – oxide ion clearly shows that 2 electrons originated from calcium atom (1)

charges correct (1)

(e) (i) add sulfuric acid/ sodium sulfate solution/ potassium sulfate solution (1)

filter (1)

(ii) moles
$$Ba = 2/137(1)$$

mass BaSO₄ =
$$\frac{2 \times 233.1}{137}$$
 = 3.4 (g) (1) [2]

Total [14]

[2]

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Q.10 (a) both contain metallic bonds/ positive ions and delocalised electrons labelled on diagram (1)

those in magnesium are stronger/ harder to break/ need more energy to break (1)

because **2** electrons are involved in delocalisation/ attraction to the positive ions (1) [3]

(b) reaction is hydrolysis of halogenoalkane/ nucleophilic substitution of halogenoalkane (1)

 $C_4H_9 X + OH^- \rightarrow C_4H_9 OH + X^-$ X can be Cl or Br (1)

(white precipitate is) silver chloride and (cream precipitate is) silver bromide (1)

 $Ag^{+}(aq) + X^{-}(aq) \rightarrow AgX(s) \text{ or } AgNO_{3} + X^{-} \rightarrow AgX + NO_{3}^{-}$ (1)

- state symbols ignored [4]

QWC selection of form and style of writing appropriate to purpose and to complexity of subject matter [1]

(c) caesium ions are bigger than sodium ions – accept 'atoms' (1)

co-ordination number 6 : 6 for sodium and 8 : 8 for caesium (1)

both cubic (1)

[3]

(1)

(d) reaction is electrophilic addition (1)

two possible products are 1-bromopropane and 2-bromopropane (1)

more 2-bromopropane formed (1)

because of greater stability of intermediate positive ion/ 2° carbocation

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

Total [16]

Q.11 (a) diagram completed with at least 1 water molecule and indication of interaction between O on one molecule and H on the other (1)

interaction between δ^+ on H and lone pair on O (1) interaction labelled hydrogen bond (1) [3]

(c) (i) [1]
$$H - C - C - H$$

(ii) peak at 1650-1750 (1)

Total [9]

Q.12	(a)	incom electro in p si	plete p sub-shell/ outer electron configuration s ² p ⁵ / outer ons in p subshell/ outer electrons in p orbitals/ valence electror ubshell/ valence electrons in p orbital	าร [1]
	(b)	(i)	gaining one electron completes shell/ gives p ⁶ / takes an elect from another species/gains an electron - do not accept 'attracts an electron'	ron [1]
		(ii)	fluorine because it is the smallest/ has the greatest electron affinity/ has the least shielding/ has the greatest effective nuclear charge/ oxidising power decreases as the group is descended	
				[1]
	(c)	oxidat	tion state is (+)5/ V - do not accept '5+'	[1]
	(d)	(i)	$Cl_2 \rightarrow 2Cl^{\bullet}$ - ignore hf	[1]
		(ii)	$CH_4 + Cl^{\bullet} \rightarrow HCl + {}^{\bullet}CH_3(1)$	
			${}^{\bullet}CH_{3} + CI_{2} \rightarrow CH_{3}CI + CI^{\bullet} (1)$	[2]
	(e)	produ	cts: ${}^{\bullet}CFH_2$ and Cl ${}^{\bullet}(1)$	

C-CI bond is the weakest/ most easily broken (1) [2]

Total [9]

Section B Total [70]

(a) Any valid ester structure with formula $C_{10}H_{12}O_2$

Examples:



(b) (i) Compound X

[1]

[1]



(iii) Rotate the plane of polarised light in opposite directions

[1]

,	•
1	c١
١	ς,

Reagent(s)	Observation if the test is positive	Compound(s) that would give a positive result
I ₂ / NaOH (aq)	Yellow solid	x
Na ₂ CO ₃ (aq)	Bubbles of colourless gas / effervescence	w
FeCl₃ (aq)	Dark purple/blue/green - do not accept 'precipitate'	X, Z

(1 mark for each box) [6]

- (d) (i) Heat / Alkaline / Potassium manganate(VII) / then acidify
 (1 mark for Potassium manganate + 1 other point; 2 marks for all)
 - (ii) I. Addition polymer One large molecule formed only / Condensation polymer one large molecule with small molecules (e.g. water) lost.
 (1)

Addition polymer – one starting material / Condensation – two starting materialsOR Addition polymer – one functional group in each molecule/ Condensation polymer –
two functional groups in each molecule(1)[2]



- (e) (i) NaBH₄ / LiAlH₄ or name(1) Reduction (1) [2]
 ignore conditions unless LiACH₄ do not accept 'redox' in water
 - (ii)



Accept structures with only one –OH group reacted.

[1] [1]

(iii)



[19 marks]

(a) (i) Alanine forms a zwitterion (1)
 Forces between alanine molecules are ionic bonding (1)
 Ionic bonding much stronger than hydrogen bonding / van der Waals (1)

Max 2 marks [2]

[2]

[1]

[1]

(ii) 1 mark for each correct structure



(iii) 1 mark for correct identification of peptide link

- (b) Enzymes / Structural proteins / Hormones or specific example
- (c) 1 mark for arrows in first stage; 1 mark for correct intermediate; 1 mark for arrow giving gain of proton in second stage (from HCN or from H^+).



[10 marks]

(a) (i)



Phthalide

(ii)

[1]



(b) Distillation / Chromatography

[1]

(c) Hydrogenation of 3-butyl phthalide removes a benzene ring (1)
 Benzene ring is more stable than alkene/ Reference to delocalisation energy (1) [2]

(d)	62.1%	[1]
(e)	(i) Greater variety of different phthalides that can be produced	[1]
	(ii) Higher atom economy / less waste / carbon monoxide is toxic	[1]
	 do not accept references to yield 	

(f) Silver nitrate and ammonia / Tollen's reagent (1); Q = Silver mirror (1); R = No reaction (1)
OR 2,4,-DNP (1); Orange precipitate with Q (1); No reaction with R (1)
OR Fehling's solution (1); Orange solid with Q (1); No reaction with R (1) [3]

[11 marks]

[1]

(a)	(i)	Nucleophilic substitution / Hydrolysis	[1]
	(ii)	Dissolved in alcohol (1) Propene or unambiguous structure (1)	[2]
	(iii)	Potassium manganate(VII) / Potassium dichromate(VI) - must be name Oxidation (1)	(1) [2]
	(iv)	(Add Potassium dichromate(VI)) and distil off the propanal from the reaction mixture	[1]
(b)	(i) Step Step 2:	1: Potassium cyanide in ethanol / Heat (1) Heat with aqueous hydrochloric acid (or other acid) (1)	



- (ii) Two points from different bullet points 1 mark each.
- Atom economy / Amount of waste / Whether waste material was recyclable / Whether waste was toxic.
- Amount of energy required / temperature required / pressure required / conditions used
- Rate of production / time
- Availability of catalyst
- Cost of reactants / Availability of reactants / toxicity of reactants.
- Two step processes usually have lower yields than one step processes / percentage yield [2]
- Purification method / separation

(c) (i) Butanoic acid is $C_4H_8O_2$ so $M_r = 88$ (1) Percentage carbon = 48/88 x 100 = 54.5%; percentage hydrogen = 8/88 = 9.1%; Percentage oxygen = 32/88 = 36.4% (At least two of these for 1) OR empirical formula for butanoic acid = C_2H_4O (1) and calculate empirical formula from percentage masses = C_2H_4O (1) [2]

- (ii) Structure 1 mark + 4 marks for explanations.
 - Product is ethyl ethanoate. (1)
 - Two points from the following required for each mark– MAX 4 marks
 - Sweet-smelling = ester
 - \circ Peak at 1.0ppm implies CH₃
 - Peak area $3 = CH_3$
 - Peak area 2 = CH_2
 - Triplet shows CH_3 is next to a CH_2 group.
 - \circ Singlet shows CH₃ no hydrogen atoms bonded to adjacent carbon.
 - \circ Peak at 2.1 ppm suggests this is next to C=O.
 - $\circ \quad \text{Quartet shows CH}_2 \text{ is adjacent to a CH}_3 \text{ group.}$
 - Peak at 4.0 ppm shows it is -O-CH₂-
 - \circ IR Peak at 1752 cm⁻¹ = C=O
 - IR Peak at 2981 cm⁻¹ = C-H or O-H
 - Cannot be –OH as we know there is no –OH in NMR spectrum

[5]

- *QWC: selection of a form and style of writing appropriate to purpose and to complexity of subject matter. (1)*
- QWC: organisation of information clearly and coherently; use of specialist vocabulary where appropriate. (1) [2]

[20 marks]

(a)	 (i) (Concentrated) nitric acid / (concentrated) sulfuric acid / Temperature of 40-80°C (Any 2 = 1 mark; All 3 = 2 marks) Electrophilic substitution (1) [3] 		
	 (ii) I. Peak area is proportional to amount of substance (1) Percentage = (30 / 38) x 100 = 79% (1) (Can obtain both marks from correct percentage) 	[2]	
	II. $45 = COOH^+$, $46 = NO_2^+$, $122 = C_6H_4NO_2^+$ and $167 = C_7H_5NO_4^+$. (Any 2 = 1 mark; All 4 = 2 marks)	[2]	
	 (iii) I. Lower melting point / melts over a range II. 1 mark for each point. Dissolve in the minimum volume Of hot water Filter hot Allow to cool Filter Dry residue under suction / in oven below 142°C 	[1]	
	QWC: legibility of text, accuracy of spelling, punctuation and grammar, clarity of meaning	7.[1]	
(b)	(i) Tin and concentrated hydrochloric acid (ii) Below 10°C (1)	[1]	





[2]

(iii) N=N double bond is chromophore (1) Compound absorbs blue /green / complementary colours to red / all colours but red (1) Remaining frequencies are transmitted, giving the red colour seen. (1) Any 2 out of 3 [2]

(c) Nitrogen has a lone pair (1) which can accept a proton (1) [2]

[20 marks]

CH5

SECTION A

1. (a) 1 dm^3 at 20°C contains 52.9 g and at 0°C it contains 17.5 g (1) \therefore amount crystallised = 52.9 - 17.5 = 35.4 g (1)	[2]
(b) (i) 2 mol of $K_2S_2O_8$ give 1 mol of O_2 2 mol of $K_2S_2O_8$ give 29.0 dm ³ of O_2 (1) \therefore 0.1 mol of $K_2S_2O_8$ gives 29.0/20 = 1.45 dm ³ of oxygen (1)	[2]
 (ii) Measure the volume of oxygen produced at specified time intervals / Measure the pH of the solution at specified time intervals 	[1]
(c) (i) An (inert) electrode that is used to carry the charge / current / electron flow	[1]
 (ii) A comment on the relative values (e.g. the persulfate system is the more positive of the two systems) (1) The more positive 'reagent' / persulfate ions acts as the oxidising agent, accepting electrons via the external circuit (1) must have the first mark to get second 	g [2]
(d) (i) The experiments show that both the concentrations of iodide and persulfate have doubled (1) therefore the initial rate should increase four times $4 \times 8.64 \times 10^{-6} = 3.46 \times 10^{-5}$ (1)	[2]
(ii) Rate = $k [S_2 O_8^{2-7}] [I^-]$ (1) $\therefore k = \frac{8.64 \times 10^{-6}}{0.0400 \times 0.0100}$	
$= 0.0216 (1) dm^3 mol^{-1} s^{-1} (1)$	[3]
(iii) In the rate equation one $S_2O_8^{2-}$ ion reacts with one Γ ion. The rate-determining step therefore has to have 1 mole of each reacting, as (only) seen in step 1	[1]

Total [14]

2.	(a)	- 705 (kJ mol ⁻¹) (1) for correct sign (1) for correct number	[2]
	(b) (i)	hydration	[1]
	(ii)	e.g. add a small 'amount' of an alkali / sodium hydroxide / NaOH / OH ⁻ ions (1) this would remove / react with hydrogen ions giving water, shifting the position of equilibrium to the left (removing iodine) (1) add P6 ²⁺ / Ag ⁺ ect.	ı [2]
	(c) (i)	Any TWO from white / misty fumes (of HI) yellow solid / solution (of sulfur) brown / black solid / purple vapour (of iodine) bubbles / effervescence / fizzing	
		One mark for each correct response	[2]
	(ii)	The values show that chlorine is the best oxidising agent, as it has the most positive E^{θ} value and therefore iodide is the better reducing agent (1) and is 'strong' enough to reduce the sulfuric acid. / OWTTE (1)	[2]
	(d) (i)	$2 \text{ NaOH} + \text{Cl}_2 \rightarrow \text{NaOCl} + \text{NaCl} + \text{H}_2\text{O}$	[1]
	(ii)	e.g. bleach, kills bacteria	[1]

3. (a) (i)

Number of moles of EDTA = $\frac{19.20 \times 0.010}{1000}$ = $1.92 \times 10^{-4} / 0.000192$ [1]

- error carried forward throughout (a)

(ii)
$$1.92 \times 10^{-4} / 0.000192$$
 [1]

(iii) Concentration =
$$\frac{1.92 \times 10^{-4} \times 1000}{50}$$
 = $3.84 \times 10^{-3} / 0.00384 \text{ mol dm}^{-3}$ (1)

Concentration = $3.84 \times 10^{-3} \times 63.5 = 0.244 \text{ g dm}^{-3}$ (1) [2]

(iv) % Cu =
$$\frac{0.244 \times 100}{11.56}$$
 = 2.11 [1]

- (b) Transition elements have either a partly filled 3d sub-shell or form ions that have a partly filled 3d sub-shell (1) However copper forms Cu^{2+} ions that are '3d⁹' / partly filled 3d sub-shell (1) whereas Zn^{2+} ions are '3d¹⁰' / full 3d sub-shell (1) - any 2 from 3 [2]
 - *QWC Organisation of information clearly and coherently; use of specialist vocabulary where appropriate.* [1]

(c)

Complex ion	Shape	Colour
$[CuCl_4]^{2-}$	tetrahedral	yellow / lime green
$[Cu(NH_3)_4(H_2O)_2]^{2+}$	octahedral	deep blue

Any two correct (1) all correct (2)

- (d) The more negative the ΔH_f value the more stable the oxide (1) PbO is relatively the more stable / CuO is relatively the less stable (1) - must have the first mark to get second [2]
- (e) (i) Any TWO from variable oxidation states partially filled 3d energy levels ability to adsorb 'molecules' ability to form complexes with reacting molecules / temporary / co-ordinate bonds
 One mark for each correct response [2]
 (ii) e.g. to allow lower pressures / temperatures use recyclable catalysts - needs qualifying [1]

[2]

SECTION B

4.	(a)	$CO \rightarrow C +2 \qquad CO_2 \rightarrow +4 \qquad (1)$	
		Increase of (positive) oxidation number = oxidation / reducing agents themselves are always oxidised are always oxidised (1)	
		OR $I_2O_5 \rightarrow I +5$ $I_2 \rightarrow I_2 0$ (1)	
		Oxidation number of iodine reduced, reduction occurring, CO reducing agent (1)	[2]
	(b)	+2 state becomes mores stable down the group and +4 becomes less stable.	[1]
	(c)	 (i) Add (a little) sodium hydroxide solution (1) to each solution. A white precipitate (1) of aluminium / lead(II) hydroxide (1) is seen. When more sodium hydroxide solution is added these precipitates (dissolve giving a colourless solution). (1) 	[4]
		<i>QWC</i> Legibility of text: accuracy of spelling, punctuation and grammar; clarity of meaning.	[1]
		(ii) Yellow precipitate (1) $Pb^{2+} + 2\Gamma \rightarrow PbI_2$	[2]

(d) (i) The bonding of aluminium in the monomer has not completed the octet / suitable diagram / 6 electrons in its outer shell (1)
 When the dimer is formed this octet of bonded electrons is formed (1)

$$CI \longrightarrow AI \xleftarrow{CI} (1)$$

$$CI \longrightarrow AI \longrightarrow CI \qquad (1)$$

$$CI \longrightarrow AI \longrightarrow CI \qquad (3)$$

(iii) I The number of (gaseous) species is increasing, leading to less order [1]

II For the reaction to be just spontaneous $\Delta G = 0$ (1)

substituting $0 = 60\,000 - 88\,\mathrm{T}$

$$T = 60\,000\,/\,88 = 682\,K\,/\,409^{\circ}C \qquad (1) \qquad [2]$$

$K_{c} = [[\underline{Al(H_{2}O)_{5}(OH)}]^{2+}(aq)][[\underline{H^{+}}](aq)]] \\ [[Al(H_{2}O)_{6}]^{3+}(aq)]$

$$\therefore 1.26 \times 10^{-5} = [H^{+}]^{2} / 0.10$$

$$\therefore [H^{+}]^{2} = 1.26 \times 10^{-6} [1]$$

$$\therefore [H^{+}] = \sqrt{1.26 \times 10^{-6}} = 1.12 \times 10^{-3} / 0.00112 (1)$$

$$- \text{ error carried forward}$$

$$pH = -\log_{10}[H^{+}] = -\log_{10} 1.12 \times 10^{-3} = 2.95 (1) [3]$$

Total [20]

(e)

5.	(a) (i	(1) $K_p = \frac{pSO_3(g) \times pNO(g)}{pSO_2(g) \times pNO_2(g)}$ (1) there are no units (1)	[2]
	(ii) The line for SO_3 / NO at equilibrium should be above the SO_2 / NO_2 line (1) as K_p has a value of 2.5, the partial pressures of SO_3 and NO at equilibrium will be greater than the partial pressures of SO_2 and NO_2 . (1)	
		- accept answer in terms alternative calculated K	s of , value
		The line for equilibrium should start at 9 hours. (1) as at equilibrium the concentrations is unchanged as time progresses. (1)	p • ••••••
		There may be other acceptable forms of explanation to be discussed at the conference	[4] e
	(iii)	If the temperature rises then the position of equilibrium will move to the left,	
		(reducing the quantities of SO ₃ and NO). (1) This will make the value of K_p smaller. (1)	[2]
	(b) (i) Nitric acid is a strong acid and its pH is $low / <2 / 1.0$ (1) As aqueous ammonia is added the pH slowly rises (1) until a pH of ~3 is reached, when it rises rapidly (1)	
		At a pH of 8-9, it tails off slowly as ammonia is a weak base. (1) Accept any 3 from 4	[3]
		Selection of a form and style of writing which is appropriate to purpose and to complexity of subject matter	[1]
	(ii)	The equivalence point is reached when 20.0 cm^3 of ammonia solution has been added as this is at the mid point of the more vertical section. (1)	1
		Since both reagents have the same concentration and the volumes used are both 20 cm the same, the number of moles of each are the same (1)	n^3 /
		OR the number of moles of both nitric acid and aqueous ammonia are calculated (0.0 and shown to be the same (1)	0020)
		\therefore Mole ratio must be 1 : 1 (1)	[2]
	(iii)	 I Ammonium nitrate is the salt of a strong acid and weak base / there is a buffering in operation. II ~ 5.5 	effect [1] [1]
	(iv)	Blue, as bromophenol blue is blue at a pH of 4.7 and above	[1]
	(c)	Number of moles of ammonium nitrate $= \frac{40}{80} = 0.50$ (1)	
		- error carried forward Concentration of ammonium nitrate solution $= \frac{0.5 \times 1000}{200} = 2.5 \text{ mol dm}^{-3}$	(1)
		$\therefore \text{ Temperature drop} = 2.5 \times 6.2 = 15.5^{\circ}\text{C} (1)$	[3]
		Total	[20]
GC	E Chemis	try MS – Summer 2012	



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