



Examiners' Report January 2013

GCE Chemistry 6CH08 01



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January 2013

Publications Code UA034343

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Introduction

To score high marks on this paper candidates need to be familiar with standard practical techniques. Questions focus on observations, measurement and the logical deduction of valid conclusions from data. In general candidates scored very well on questions involving routine techniques, observations and calculations, but found it more difficult to apply their knowledge to unfamiliar situations and to explain the rationale behind some standard techniques.

Question 1

This question was generally well answered. Candidates found parts (b) and (c) most difficult. In part (b) few could identify the correct reaction type and in part (c) few realised that the sodium hydroxide was acting as a base.

A good answer with just one error.

(b)	Measure the pH of a dilute aqueous solution of A using a pH meter.	The pH is 6.0.	The type of reaction that has occurred when A dissolved in water is $actd - b_{500} \epsilon$.	(1)
(c)	Add a few drops of dilute sodium hydroxide solution to a solution of A .	A green precipitate forms.	The sodium hydroxide is acting as <u>a ligand</u> The formula of the green precipitate is	
			[fe(HJO)4(он)]]	(2)



Part (b): Transition metal salts form acid solutions because of deprotonation reactions. These are a kind of acid-base reaction so the mark is gained.

Part (c): When hydroxide precipitates are formed the sodium hydroxide is acting as a base not as a ligand.



Make sure you understand the difference between deprotonation and ligand-exchange reactions.

Again a good answer but the reactions in part (b) and (c) were not understood.

		L		1
(a)	Observe the appearance of A .	Pale green solid.	The cation By may be Fe ^{2t} or N ²⁺	(1)
(b)	Measure the pH of a dilute aqueous solution of A using a pH meter.	The pH is 6.0.	The type of reaction that has occurred when A dissolved in water is in the transformed in the state of the	(1)
(c)	Add a few drops of dilute sodium hydroxide solution to a solution of A .	A green precipitate forms.	The sodium hydroxide is acting as Pactor and The formula of the green precipitate is	
			Fe (OH)2 OF Ni(OH)2	(2)



e) Add excess so hydroxide sol a sample of th precipitate fo	dium The green ution to precipitate does not rmed in (c). dissolve.	It is not The cation i	(+34 5 Fe ²⁺	(1)
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lesuite US **Examiner Comments**

In part (a), it is perfectly acceptable to suggest that the pale green salt might contain Ni²⁺.

In both parts (b) and (c) the reaction was wrongly identified as ligand exchange.

Part (e) is particularly well answered - the lack of amphoteric behaviour does indeed rule out Cr³⁺.



Make sure you understand the difference between deprotonation and ligand exchange reactions.

Question 2 (a)

This was generally very well answered.

One mark lost through an incorrect name.



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In this example the candidate fails to appreciate that the test is for the carbonyl group.

2 Two organic compounds, X and Y, are colourless liquids. Both compounds contain four carbon atoms and one functional group. (a) A series of tests was carried out on compound X. (i) When a few drops of 2,4-dinitrophenylhydrazine solution were added to X, an orange precipitate was formed. What deduction can be made from the result of this test alone? (1)Ketone present (ii) When X was warmed with Fehling's solution, a red precipitate was formed. What further deduction can be made from the result of this test? (1) Aldehyde present **Examiner Comments** In part (i) the positive test shows that the compound could be a ketone or an aldehyde. Resu **Examiner Tip**

Learn the organic functional group tests.

Question 2 (b)

This was well answered, though some candidates drew a ketone structure and a few wrote the same structure twice with different bond angles!

One correct and one wrong structure.



One mark lost because the formula is ambiguous.



Question 2 (c) (iii)

Most candidates appreciated that the molecule had a chiral centre. This could be expressed in a variety of ways.

It was not enough to simply say that it was optically active, since this merely repeats the information given in the question.

Note that it cannot be a racemic mixture as this would not be optically active.



If you include several points in a one mark question, they all have to be correct. It is better to limit yourself to the central point you wish to communicate to the examiner.

Question 2 (c) (i)-(ii)

Well answered, though a few candidates threw away the second mark by simply saying that Y contains OH, which is true of alcohols and carboxylic acids.

An excellent answer.

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1.418

Incorrect terminology loses a mark.



"Hydroxide" is the name of the OH^{-} ion, "hydroxyl" is the correct

Examiner Tip

Learn the correct terminology.

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name for a covalent bonded -OH group.

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Question 2 (c) (iv)

Well answered, though a few candidates gave the primary or the tertiary alcohol - neither of which is chiral.



not chiral.

Read the question.

Question 2 (c) (v)

The correct colour and physical state of the product was required.

Examiner Comments

The precipitate is yellow, not white.



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Plus

Result

Examiner Tip

Learn the results of standard tests.

Question 3 (a) (i)-(ii)

The demanding calculations in this question were generally very well done. However, a surprising number of candidates failed to round the answer in part (ii) to two significant figures. Others truncated intermediate answers to only one significant figure, causing their final answers to be very inaccurate.



(ii) A student obtained 2.97 g of 2-ethanoylaminobenzoic acid from 4.00 g of 2-aminobenzoic acid. Calculate the percentage yield obtained by this student. Give your answer to two significant figures. relative molecular mass of 2-ethanoly laminobenzoid acid = 102 + 12 + 16 + 12 + 1 = 143. theoretical yield = $143 \times 0.02919708 = 4.1752g$:. % yield = $\frac{2.97}{4.1752} \times 100\% = 71.1\%$ **Examiner Comments** Part (a) is fully correct but in part (b) the M_r of the product is wrongly calculated and then the final answer is not given to two significant figures - so two marks are lost. **NIS** Kesu **Examiner Tip** Be careful with significant figures. Note that if the only error had been the M_r of the product, two marks could still be gained even though the final answer

two marks could still be gained even though the final answe would have been wrong. This is why it is important to show your workings.

Question 3 (b) (i)

Most candidates appreciated that the ethanoic anhydride was used in excess to ensure that all the 2-aminobenzoic acid reacts.

Some lost the mark by stating that **all** the reactants are used up - which is not true of the excess ethanoic anhydride.



The answer simply repeats the information in the question, so does not score.

Rephrasing the question will not score marks - think about the reasons behind the information given.

Question 3 (b) (ii)

Many answers were too vague to gain any credit.

It was not enough to say that bubbles are formed - any boiling liquid will produce bubbles. Without anti-bumping granules the bubbles will be **large** and may cause liquid to be lost when it splashes out of the top of the condenser.

Some candidates incorrectly stated that the bubbles were made of air rather than the vapour of the reactants.

(ii) Anti-bumping granules are added in step 2. What would be observed if 'bumping' occurred? (1)large bubbles of air are produced furthich cause vioprous shaking of the pear shaped flask Reculte **Examiner Comments Examiner Tip** A promising answer - "large bubbles" would indeed Think about the processes going on in a be formed, but they are bubbles of the vapour of practical technique. the reactants, not of air - so the mark is lost. (ii) Anti-bumping granules are added in step 2. What would be observed if 'bumping' occurred? (1)There would be uneven heating , and **N**IS **Examiner Comments Examiner Tip** "Uneven heating" is not quite enough to score. The point is that it is the boiling that is uneven Avoid vague statements - try to be specific. without the anti-bumping granules.

Question 3 (b) (iii)

Most candidates appreciated the need for gloves but some thought that a face mask would be sufficient to prevent damage to the respiratory system. A simple mask would not absorb corrosive chemicals so this was not allowed. A fume cupboard is required.

(iii) Ethanoic anhydride is corrosive to both the skin and the respiratory system. Suggest two precautions to minimise the risks when using ethanoic anhydride, other than wearing eye protection and a lab coat. (2)a mai · Use gloves **Examiner Comments** A face mask alone would not give adequate protection, so a mark is lost. (iii) Ethanoic anhydride is corrosive to both the skin and the respiratory system. Suggest two precautions to minimise the risks when using ethanoic anhydride, other than wearing eye protection and a lab coat. (2)Wear a loves use and tumes Noboard **Examiner Comments** Despite the reference to "fumes cupboard", this scores the two marks.

Question 3 (b) (iv)

Many candidates were fully conversant with the recrystallization process and scored high marks.

Others had the correct processes in the wrong order - which simply would not work - so lost marks.

(iv) Outline how you would carry out the recrystallization in step 6. (4)Use hot and small amount of solvent. Solid is placed in the solvent . cond solvent is fittered using pre-heated Buchner filter. Then it is left to cold. The solid is placed in cold water. Remove solid by filter funnel and filter paper. Solid is Small amount of cold water is poured on surface of the solid. esults Plus **ResultsPlus** Examiner Comments Examiner Tip Almost all correct, but the final product has A pure product must be dry. to be dried. (iv) Outline how you would carry out the recrystallization in step 6. (4)Dissolve the solid with minimum amount of hot ethanoic acid to get saturated solution. Filter the hot solution to remove insoluble impurities. Allow the solution to cool to get crystals Wash the crystals to remove soluble impurities. Dry the crystals with filter paper **Results Plus** Examiner Comments **Examiner Tip** Almost all correct but the filtration to separate Imagine you are carrying out the process in the crystals before they are washed and dried the lab. - then you won't leave steps out. has been omitted.

Question 3 (b) (v)

Many answers simply referred to "transfer errors" without giving details. This was not enough to score; one specific source of product loss was required for the mark. Disappointingly few candidates cited the likeliest source of error - product remaining dissolved in the saturated solution after crystallization.

2 ethanoylaminobenzoic a	(1) reid may remain in
solution.	
The best answer - this is the major source of product loss	A short clear answer will get the ma
	vill slightly reduce the yield of
 (v) Suggest a reason why the recrystallization v 2-ethanoylaminobenzoic acid. 	

Question 3 (b) (vi)

Quite well done but many candidates suggested that a water bath be used to produce the high temperature required to melt the product. It is very unlikely that such a large molecule, capable of inter-molecular hydrogen bonding, would have such a low melting point - so a mark was lost. An oil or sand bath is required or an electrically heated metal block.



Question 3 (b) (vii)

melting point so the second mark is lost.

Generally well answered, however, some candidates confused 'a narrow melting point range (sharp melting point)' with a small difference between the literature and the measured value.

Others said that the measured value should be compared with a "theoretical" value but did not say how the "theoretical" value could be obtained. Reference to a data source was essential for the mark.

	(vii) State two ways you would use the results from purity of the product.	(vi)	to check the identity and	
	It should The melting point should ma	iton	to the one in the	
	data booklet. enerk for compound with	siw	ilar melting point.	
	1	-		
	Results Plus Examiner Comments		Results Plus Examiner Tip	
One book	mark is gained for the comparison with the data value but there is no reference to a "sharp"		A two mark question usually requires two separate points in the answer.	

(vii) State two ways you would use the results from (vi) to check the identity and purity of the product. (2) check the melting and builing Point of Marthethe Place obtained Product with the Pure sample of the Product. IF both has significantly very less difference. ်င္လာင္ရွိ (Total for Question 3 = 19 marks) the product 13 then **lesults Examiner Comments** The candidate appreciates that the melting temperature must be compared with

that of the pure product, but does not explain how this is to be found, so loses the mark. Reference to a sharp melting point is also missing.



Question 4 (a)

Most candidates appreciated the purpose of a trial titration however, a few wrongly thought that it was to test the effectiveness of the apparatus and the method.



(1) To act as a range finder so that the rest of a the titrations can be <u>Carried out with greater accuracy</u>. To find an approximate of how much volume of solution is required to reach end point.

(a) Explain why a trial titration (titration 1) is carried out.



When doing practical work, make sure you understand the reason for the operations you carry out.

Question 4 (b) (i)

Generally very well answered.

A few candidates expressed the 19.90 cm³ as 19.9 cm³ and lost a mark.

A few used only two of the three concordant results and so lost a mark.

Titration number 3 1 (trial) 2 4 Burette reading 21.45 41.35 21.95 2.15 (final) / cm³ Burette reading 1.95 1.20 21.45 21.95 (initial) / cm³ Volume of 17.875 20.00 19.80 19.9 Fe²⁺(ag) used / cm³ Titre used to calculate mean (✓) (a) Explain why a trial titration (titration 1) is carried out. (1)accurate. To ensure that the readin (b) (i) Complete the table and indicate with a tick (\checkmark) those titres most suitable for calculating a mean titre. Use the titres you have chosen to calculate the mean titre. (4) = 19.9+20.0+19.8 = 19.9 Mean titre = 19.9 ... cm³ **Examiner Comments** A potentially good answer spoiled by the failure to express the second titre to two decimal places in run 2 and a silly slip in calculating the final burette reading in run 4. **Examiner Tip** Always give burette readings and titres to two decimal places and check your arithmetic.

Titration number	1 (trial)	2	3	4
Burette reading (final) / cm ³	21.45	41.35	21.95	41.75
Burette reading (initial) / cm ³	1.20	21.45	1.95	21.95
Volume of Fe ²⁺ (aq) used / cm ³	20.25	19.9	20.00	19.80
Titre used to calculate mean (✓)	×	\checkmark	\bigvee	X
a) Explain why a trial	titration (titration he <u>burett</u> e	1) is carried out. 3 not a	onoppletely .	(1) empty
b) (i) Complete the t calculating a m	able and indicate	e with a tick (✓) th	ose titres most su	iitable for
Use the titres y	ou have chosen t	o calculate the m	ean titre.	(4)
	wean titr	- 199-	20	(4)
		2		
		= 19.95	cm ³	
			Mean titre =	19.95
	Results xaminer Comn	US nents		
Again, a n decimal pl	nark is lost for lace and anoth ed as concorda	expressing the per is lost beca nt.	e titre in run 2 use the titre in	to only one run 4 is
not marke				
	Res	suitsPlus ner Tip		

Question 4 (b) (ii)

This question was very well answered. Most candidates were clearly familiar with this type of calculation from titration results.

(ii) Use the equation below, and your mean titre, to calculate the concentration of the sodium dichromate(VI) solution, in mol dm⁻³. (3) $Cr_{2}O_{2}^{2-}(aq) + 6Fe^{2+}(aq) + 14H^{+}(aq) \rightarrow 2Cr^{3+}(aq) + 6Fe^{3+}(aq) + 7H_{2}O(I)$ orange green The no. of mole of Fe²⁺ (ng)=(19.9)(0.05)=9.95×10-4mol The mole ratio of Croog2 -: Fezt The no. of mole of $(r_1 O_7)^2 = \frac{9.95 \times 10^{-4}}{6} = 1.65 \# \times 10^{-4} \text{ mol}$ The concentration of the sodium dictromate (VI) solution = 1.658 ×/6-4 - 20 = 8.29 × 10-3 moldm-3 **Results**Plus **Examiner Comments** An excellent answer, the calculation is correct and set out in logical steps. **Results Plus Examiner Tip** Try to set out your calculations like this - showing

what is calculated in each step.

 (ii) Use the equation below, and your mean titre, to calculate the concentration of the sodium dichromate(VI) solution, in mol dm⁻³.

$$Cr_2O_7^{2-}(aq) + 6Fe^{2+}(aq) + 14H^+(aq) \rightarrow 2Cr^{3+}(aq) + 6Fe^{3+}(aq) + 7H_2O(I)$$



(3)

Results Plus Examiner Comments

The calculation is essentially correct but the result of the first step has been rounded to only one significant figure which makes the following results very inaccurate. One mark is lost for this inaccuracy.



In all intermediate steps in a calculation, work to one more significant figure than you will use for the final answer.

Question 4 (c)

Many candidates failed to appreciate that the error in the burette reading must be doubled since each titration requires two readings to be taken.

A few students used the wrong titre value and lost a mark.



(c) Assuming the accuracy of the burette is ±0.05 cm³ each time the burette is read, calculate the % error of the titre in **titration 3**.



(1)

Question 4 (d)

Many answers were too vague to score. The point is that the indicator makes the colour change at the end-point sharper or clearer to see.



A good answer which makes the function of the indicator clear.



When answering this type of question, imagine yourself carrying out the titration and think what the problems might be.

Question 4 (e)

Many candidates appreciated that the titre would be too high, but then failed to explain that this was a result of the air bubble in the burette at the beginning of the titration.

(e) A student carrying out one titration left an air bubble in the tip of the burette before taking the initial reading. This bubble was no longer present when the student took the final reading. State and explain what effect, if any, this would have on the titre value. What effect would the use of this titre have on the calculated concentration of sodium dichromate(VI)? (3)Wrangly. This will everymally cause 1 +400 -This will allo of sodium dichonale (VI) Cal Culated in accurate. Concombation increates enor Cante SUITS Examiner Comments This answer is too vague to score any marks. There is no indication of why the titre would be inaccurate and no discussion of whether the titre would be increased or decreased as a result. IS **Examiner Tip** Don't be vague - try to think through the exact effect

of an error like this.

(e) A student carrying out one titration left an air bubble in the tip of the burette before taking the initial reading. This bubble was no longer present when the student took the final reading. State and explain what effect, if any, this would have on the titre value. What effect would the use of this titre have on the calculated concentration of sodium dichromate(VI)? (3)than the fitre value would been compared actual He bu air bubble -catentated without value the air bubble would left 54 empty Space since as the volume used in the hitration. The measured be. concentration of Sodium Dichromate (III) wat would calcul (Total for Question 4 = 13 marks) Han actual value. the arger



An excellent answer. The candidate fully understands the effect of the bubble.

Res **Examiner Tip**

Try to give answers like this with a clear logical explanation of effect of the suggested error.

Paper Summary

Based on their performance on the paper, candidates are offered the following advice:

- Make sure you can explain the reasons for the steps in standard techniques such as titrations and recrystallization.
- Explain less familiar techniques.
- Make sure you understand the importance of using an appropriate number of significant figures in the various steps of calculations to ensure the final answer is accurate.
- Learn the tests that distinguish between functional groups and their position in molecules. e.g. primary, secondary and tertiary alcohols.

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