



Examiners' Report June 2015

IAL Chemistry WCH06 01



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Introduction

There were some very well prepared candidates. Centres can be commended where the practical work is carried out with care and attention to reasons for procedures.

Some candidates seem not to have done many of the practical exercises in the specification.

The full range of marks were seen on this paper. There was no evidence that candidates had insufficient time to complete the paper.

Candidates were good at the calculation aspects, but less good when asked for descriptions or reasons.

Question 1 (a) (i)

This test for the sulfate ion was known by three quarters of the candidates.

- 1 A white solid, **A**, contains one cation and one anion. When water is added slowly, the solid turns blue and then dissolves to form a blue solution, **B**.
 - (a) When aqueous barium chloride is added to an acidified portion of solution **B**, a white precipitate forms.

(i) Give the **formula** of the anion in **B**.

Cu²⁺



- 1 A white solid, **A**, contains one cation and one anion. When water is added slowly, the solid turns blue and then dissolves to form a blue solution, **B**.
 - (a) When aqueous barium chloride is added to an acidified portion of solution **B**, a white precipitate forms.
- (i) Give the **formula** of the anion in **B**. (1)





Question 1 (a) (ii)

It was disappointing that nearly one quarter of the candidates answered with sulfuric acid.

(ii) Name a suitable acid for acidifying solution **B** in this test.





Question 1 (b) (i)

This was where many candidates began to make errors in this section with much confusion between ions and substances, and appropriate ligands.

- (b) When aqueous ammonia is added to another portion of solution **B**, a blue precipitate forms. When more aqueous ammonia is added, this precipitate dissolves to form a <u>deep</u> blue solution, **C**.
- - (b) When aqueous ammonia is added to another portion of solution **B**, a blue precipitate forms. When more aqueous ammonia is added, this precipitate dissolves to form a deep blue solution, **C**.
 - (i) Identify, by name or formula, the blue precipitate.

(1)

17.56







this does but the substance cannot have a charge.

Question 1 (b) (ii)

The ongoing confusion caused by ammonia solution continued. Quite a few candidates gave the correct numbers of ligands but no charge or remarkably, negative charges. It is important to remember that an aqueous solution of ammonia is an equilibrium.NH₃(aq) + $H_2O(I) \square NH_4^+(aq) + OH^-(aq)$

At lower concentrations of ammonia solution in a solution containing copper(II) ions the concentration of hydroxide ions is sufficient to form the precipitate, copper(II) hydroxide, which drives the above equilibrium to the right.

As the relative amount of water decreases, because the ammonia solution is more concentrated, this equilibrium is reversed, increasing the concentration of ammonia. This then is removed by reaction with copper ions.

(ii) Give the **formula** of the ion responsible for the deep blue colour of solution **C**.

(1)

(1)





(ii) Give the **formula** of the ion responsible for the deep blue colour of solution **C**.

 $[(4(NH_3)_6]^{2+}]$





attached to four ligands in complexes. The extra two waters are further from the copper ion and are best ignored.

Question 1 (c)

There were many answers with incorrect ligands or incorrect charges.

A selection of these are given.

(c) Give the **formula** of the complex ion which gives the blue colour to solution **B**. Include the ligands in your answer.



(c) Give the **formula** of the complex ion which gives the blue colour to solution **B**. Include the ligands in your answer.



Question 1 (d)

Incorrect formulae for copper(II) sulfate were quite common.

Copper(II) chloride and the tetrachlorocopper(II) ion were also popular.

(d) Give the **formula** of the white solid **A**.



(d) Give the **formula** of the white solid **A**.



Question 1 (e)

This either simply wasn't known or the question wasn't understood as there were many very poor answers. It was a good discriminator as only the really good candidates scored both marks.

It was important to realise the role of water ligands in altering the energy levels of d orbitals to differing extents so that 3d to 3d transitions absorb energy from visible light.

Question 2 (a)

Though many know it was the sodium ion, it was a shame how many candidates did not read 'the *formula* of the *ion'*.

- 2 A salt, P, contains carbon, hydrogen, oxygen and one metallic element.
 - (a) When a flame test is carried out on **P**, a yellow flame results.

Give the **formula** of the metal ion in the salt.



- 2 A salt, P, contains carbon, hydrogen, oxygen and one metallic element.
 - (a) When a flame test is carried out on **P**, a yellow flame results.

Give the **formula** of the metal ion in the salt.

no. Na



Question 2 (b)

(i) It was clear that many candidates had never calibrated a pH meter, though this is essential.

It is normal to use two buffer solutions of known pH, one alkaline and one acidic.

(ii) To show the pH of a solution universal indicator solution or universal indicator paper must be used.

This was rarely given – acid base indicators suitable for titrations were the common responses.

(iii) A pH meter measures precisely to one decimal place. Universal indicator changes colour over a range of pH values, as do other acid base indicators.

(b) P dissolves in water to form a weakly alkaline solution, Q, with pH 8.1.

The pH of **Q** can be measured by using a calibrated pH meter or an indicator.

(i) Describe how to calibrate a pH meter.

(ii) Name a suitable indicator you could use and state the colour you would expect to observe.

universal indicator.

Get the incubator

- (iii) Which of the two methods will give the more accurate value for the pH of **Q**? Justify your answer.
- (1)

(2)

(2)

pH meter is more accurate.



(ii) Read the question and underline key words.

(b) **P** dissolves in water to form a weakly alkaline solution, **Q**, with pH 8.1.

The pH of **Q** can be measured by using a calibrated pH meter or an indicator.

(i) Describe how to calibrate a pH meter.

(2) By putfing the pH meter into a strong acid and PY pH 1 and then putting it into a label as strong alkali. and and labelling as pH14 (ii) Name a suitable indicator you could use and state the colour you would expect to observe. (2) Methyl orange Orange to yellow (iii) Which of the two methods will give the more accurate value for the pH of Q? Justify your answer. (1)PH meter. Because you can identify the oxact from the pH meter RA. PH **ResultsPlus Examiner Tip** Examiner Comments (i) Though this is not best practice, it was This is a good example of a candidate accepted on this occasion. who has the good sense to apply what they know to the question, even though (ii) This gets a 'rescue' mark for an they are on unfamiliar ground. indicator with the correct alkaline colour. (iii) This is fine as per the mark scheme.

Question 2 (c) (i)

It was surprising that so few candidates seemed to be familiar with this common test for an ester.

The purpose of the sodium carbonate solution is to neutralise the carboxylic acid which often has a stronger (and often unpleasant) smell than the ester.

Many answers referred to the solid dissolving or disappearing, confusing the test with adding sodium to an alcohol.

This was probably why so many candidates gave a white solid forming.



(c) Some of the solution \mathbf{Q} is acidified with concentrated hydrochloric acid.

An organic compound, **R**, forms in the solution.

Methanol is added and the mixture warmed, forming a new organic compound **S**.

This mixture is added to sodium carbonate solution in an evaporating basin. A fruity smell is detected.

(i) Describe and explain what you would see as the mixture is added to the sodium carbonate solution.

```
white misty fumes of them
     efferveseen. & gas is evolved given off.
                                                                  US
       Examiner Comments
                                                     Examiner Tip
The candidate just manages to save
                                            It is best to fully cross out parts of
their marks.
                                             answers which you do not wish to have
                                             marked.
```

(2)

Question 2 (c) (ii)

Three quarters of candidates did recognise the test for an ester.

(i) What type of compound is \$? (i) What type of compound is \$? (i) What type of compound is \$? Αι Cebo





Question 2 (d)

This question was another good discriminator – often if they could answer they gave R as an acyl chloride.

For P, too many drew a covalent bond between O and Na, some even gave the charges and still drew in the bond.

(d) The high resolution proton nuclear magnetic resonance (hmr) spectrum of **S** has only two peaks which are both singlets and have the same area.

Deduce the structural formulae of S, R, and P. (3) S - 01 R

P





This is an instructive example. The candidate has found an ester which would give two singlet peaks in the NMR spectrum, but has failed to realise the significance of the same areas.

The second and third structures are acceptable transferred errors.



Always consider all the information in the stem of a question.

(d) The high resolution proton nuclear magnetic resonance (nmr) spectrum of **S** has only two peaks which are both singlets and have the same area, Deduce the structural formulae of S, R, and P. (3)S CH3-CH3 COO CH3 R CH3 COCL Ρ + CH3 CO Na Natoz + Chacoce Warten CHZ CH3 COOCH, oculte#lus **Results**Plus **Examiner Comments Examiner Tip** Always check each carbon has four This was a common error in **R**. It is difficult to make an acyl chloride from bonds.

Then a subscript 2 in lost from the

an ester.

formula for **P**.

Question 3 (a)

It is rarely correct to add water to concentrated acid because of the risk of a violent reaction.

3 Cupronickel is an alloy of copper and nickel. It is used to make 'silver' coins.

A coin is analysed by the following method.

- **Step 1** It is weighed on a balance which reads to two decimal places and found to have mass 4.00 g.
- **Step 2** Water is added to the coin in a beaker. Concentrated nitric and sulfuric acids are added and the coin dissolves.
- **Step 3** When the coin is completely dissolved, the solution is neutralized.
- **Step 4** The neutral solution is transferred, with the washings, to a 100 cm³ volumetric flask, made up to the mark with water and mixed thoroughly.
- **Step 5** 10 cm³ samples of the solution are taken and an excess of potassium iodide is added, producing iodine.
- **Step 6** The iodine is titrated with 0.200 mol dm⁻³ sodium thiosulfate solution.
- (a) Why, in Step 2, is water added before, rather than after, the acids?

(1)

To cool dans the soir and solution, Reduce the temperature as

addition of concentrated actal is highly valetile and vigorous reactions can occur. As it is an exothermic reaction reter is added

Results Plus Examiner Comments

Although the first sentence is incorrect, the candidate recognises the consequences of adding water to acid.



When doing experiments ask why each step is done and consider the order in which chemicals are added.

3 Cupronickel is an alloy of copper and nickel. It is used to make 'silver' coins.

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- **Step 2** Water is added to the coin in a beaker. Concentrated nitric and sulfuric acids are added and the coin dissolves.
- Step 3 When the coin is completely dissolved, the solution is neutralized.
- **Step 4** The neutral solution is transferred, with the washings, to a 100 cm³ volumetric flask, made up to the mark with water and mixed thoroughly.
- **Step 5** 10 cm³ samples of the solution are taken and an excess of potassium iodide is added, producing iodine.
- **Step 6** The iodine is titrated with 0.200 mol dm⁻³ sodium thiosulfate solution.
- (a) Why, in Step 2, is water added before, rather than after, the acids?

To provide a	medium	for the	neuHalization
reaction to take	place.	****	



added.



Question 3 (b)

It was common to see the colour of iodine in starch solution given here.

(b) What is the colour of an aqueous iodine solution?

blue black Solution.





(1)

(1)

(b) What is the colour of an aqueous iodine solution?

We Orange





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Question 3 (c) (i)

This was the easiest question on the paper.

(c) (i) To make the end point of the titration more obvious, an indicator is added just before the colour of the iodine disappears.

Name this indicator.

Nlethyl opange.





(1)

(1)

(c) (i) To make the end point of the titration more obvious, an indicator is added just before the colour of the iodine disappears.

Name this indicator.

Phinophthalein.





Question 3 (c) (ii)

It was pleasing to see how many candidates were aware that if added to early a stable iodine-starch complex forms.

(ii) Suggest why the indicator is not added to the iodine solution earlier in the titration.

(1)

(1)

Starch forms an insoluble blueblack complex

in high concentrations of iodine.





(ii) Suggest why the indicator is not added to the iodine solution earlier in the titration.

```
the fit titre reacts with indine.
```





Question 3 (c) (iii)

The common error was to fail to give the starting colour.

(iii) Give the colour change at the end point when the indicator is used in this titration.







(iii) Give the colour change at the end point when the indicator is used in this titration.

(1)

(1)



to wream

Orange

Question 3 (d) (i-iv)

(i) This was rarely incorrect.

(ii) This was another good discriminator as the weaker candidates gave 2, 3 and 4.

Several candidates weren't precise enough with their explanation for their correct choice e.g. closer or smallest distance, rather than not concordant.

(iii) Again rarely incorrect with transferred error allowed from (ii).

(iv) Here, the most common error was to forget the factor of 10.

Candidates either knew what to do or didn't know where to start!

They need to be encouraged to write chemical statements to explain their maths in order to gain credit for their working, as this was essential to gain credit for an incorrect final answer.

(d) The results for the titrations are shown below.

Titration number	1	2	3	4
Burette reading (final) / cm³	24.10	47.90	23.55	47.00
Burette reading (initial) / cm³	0.00	24.10	0.00	23.55
Titre / cm ³	24.10	23.8	29.55	23:45

(i) Complete the table.

(1)

(ii) Which titres should be used to calculate the mean? Explain your choice.

(1)

3 and 4						
* be t because	They	titoes	Cland	2)	has	
greated	Value	han	± 0.2			

(iii) Calculate the mean titre.

(iv) Calculate the percentage by mass of copper in the coin.

Use the equations below.

$$2Cu^{3}(aq) + 4i^{2}(aq) \rightarrow 2Cu(s) + i^{2}(aq)$$

$$25,0i^{2}(aq) + i^{2}(aq) \rightarrow 5,0i^{2}(aq) + 2i^{2}(aq)$$

$$moles \quad gf \quad S_{2} \quad 0_{2}^{n} \qquad (5)$$

$$0 \cdot 200 \quad -7 \quad loo \ ocm^{3}$$

$$\pi \quad -7 \quad 25 \cdot 5 \ cm^{7}$$

$$\pi = 4 \cdot 7 \times (0^{-7} \ moles$$

$$moles \quad gf \quad I_{n}$$

$$4 = 4 \cdot 7 \times (0^{-7} \ moles)$$

$$moles \quad gf \quad Cu^{3}$$

$$\pi = 2 \cdot 35 \times 16^{3} \ moles$$

$$moles \quad gf \quad Cu^{3}$$

$$1 \quad 2 \quad 7 \quad 7 \quad 7 \quad 75 \times 16^{3} \ moles$$

$$moles \quad gf \quad Cu^{3}$$

$$\pi = 2 \cdot 35 \times 16^{3} \ moles$$

$$moles \quad gf \quad Cu^{3}$$

$$\pi = 4 \cdot 7 \times 10^{-7} \ moles$$

$$mass \quad d \quad Cu^{9}$$

$$The is infinet ull the final part when the factor of ten is nonitted.$$

$$The final part when the factor in repeated threat or in repeated threat or in repeated threat or in represent the factor in repre$$

-

Question 3 (d) (v)

The factor of 2 was often omitted in the burette calculation.

(v) The uncertainty in each burette reading is ± 0.05 cm³ and the uncertainty in each reading of the balance is ± 0.005 g.

Calculate the percentage uncertainty in the third titre value and in the mass measurement. Use your results to decide whether using a balance that weighs to three decimal places would significantly improve the accuracy of the result.

burette balance (2)
% U=
$$\frac{0.05}{23.55} \times 100$$
 $\frac{0.005}{4.00} \times 100$
= 0.21% 0.13%
 \therefore uncertainity in 4.004 less than \approx uncertainity 4.00
therefore balance that reads $3d.p$ will improve the
accoracy
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ResultsPlus

Examiner Comments The balance uncertainty is fine. The burette calculation has omitted the factor of 2. A titration involves two readings, one before and the other after the titration.

Question 4 (a)

It was disappointing to see how many candidates did not appreciate that bromine is a diatomic molecule.

This was the most common method of losing a mark.

Bromobenzene can be prepared from benzene by the following steps. Reflux 20.0 cm³ of benzene with 6.0 cm³ of bromine and about 10g of Step 1 iron filings, by heating on a water bath at 50°C. After the reaction has finished, remove the water bath and heat to boiling Step 2 until no bromine vapour can be seen. Cool the mixture and add 25 cm³ of ethoxyethane (diethyl ether) to extract Step 3 the bromobenzene. Wash the ethoxyethane layer with aqueous sodium hydroxide. Separate Step 4 the ethoxyethane layer. Wash the ethoxyethane layer with water and repeat the separation. Step 5 Dry the ethoxyethane layer with a suitable drying agent. Step 6 Step 7 Decant the dried solution. Distil the separated solution, collecting the fraction boiling around the Step 8 boiling temperature of bromobenzene, 156°C. (a) Calculate the number of moles of bromine, Br₂, used in the experiment. [Density of bromine 3.1 g cm⁻³] de (2)Density= m/ n= CV 1. 1. 1. $\mathcal{M} = \frac{1}{2} \left(\frac{1}{2} - \frac{1}{2} \right)^2 \left$ 3.1 = <u>19</u>. = 6 0 × <u>25 17</u> 1000 /≫¢3×1 α the mater = 0.155 moles. duar. volum @ = 25.77 cm3





Practise density calculations connecting moles and volumes.

- Bromobenzene can be prepared from benzene by the following steps. 4
 - Reflux 20.0 cm³ of benzene with 6.0 cm³ of bromine and about 10g of Step 1 iron filings, by heating on a water bath at 50 °C.
 - After the reaction has finished, remove the water bath and heat to boiling Step 2 until no bromine vapour can be seen.
 - Cool the mixture and add 25 cm³ of ethoxyethane (diethyl ether) to extract Step 3 the bromobenzene.
 - Wash the ethoxyethane layer with aqueous sodium hydroxide. Separate Step 4 the ethoxyethane layer.
 - Step 5 Wash the ethoxyethane layer with water and repeat the separation.
 - Dry the ethoxyethane layer with a suitable drying agent. Step 6
 - Decant the dried solution. Step 7
 - Step 8 Distil the separated solution, collecting the fraction boiling around the boiling temperature of bromobenzene, 156°C.
 - (a) Calculate the number of moles of bromine, Br₂, used in the experiment.

[Density of bromine 3.1 g cm⁻³]



(2)

Question 4 (b)

Many knew the first step but balancing of this equation was poor.

In the second step – far too many didn't know it or produced Br-.

(b) Bromine reacts with iron to form iron(III) bromide, which reacts with bromine to produce the attacking electrophile in **Step 1**.

Write the chemical equations for these reactions.

(2)

that they had not answered the question

$$BI_{0} + FeBr_{3} \rightarrow FeBr_{4} + Br^{+}$$

$$Br_{1} + O \rightarrow O Br^{+} + H^{+}$$

$$Br_{1} + O \rightarrow O Br^{+} + H^{+}$$

$$Br_{1} + O Br^{+} + FeBr_{3}$$

$$Br_{1} + FeBr_{3} + FeBr_{3}$$

$$Br_{1} + FeBr_{3} + FeBr_{3}$$

$$Br_{2} + FeBr_{3} + FeBr_{3}$$

$$Br_{2} + FeBr_{3} + FeBr_{3}$$

$$Br_{3} + FeBr_{3}$$

asked.

As a result they have given the correct

equation correctly.

second equation and even given the next

(b) Bromine reacts with iron to form iron(III) bromide, which reacts with bromine to produce the attacking electrophile in **Step 1**.

Write the chemical equations for these reactions.

(2)

$$Fr(Br_3)$$

 $Fr(Br_3)$
 $3Br_2 + 2Fe \rightarrow 2FeBr_3$
 $Fr Br_3 + Br_2 \rightarrow FeBr_2 + 3Br$
Results Puss
Examiner Comments
The first equation is correct.
(2)
 $Fr(Br_3)$
 $Fr(Br_3)$
 $Fr(Br_3) + Br_2 \rightarrow FeBr_2 + 3Br$
 $Fr(Br_3) + Br_2 \rightarrow FeBr_2 + 3Br$

Question 4 (c)

(c) Why is the ethoxyethane layer washed with sodium hydroxide solution in Step 4?

(1)

```
To remove the impunities present.
```





(c) Why is the ethoxyethane layer washed with sodium hydroxide solution in Step 4?

(1)Sodium hydroxicle is added to remove HBr which disgolved in the ethoxyethane layer.



The fully correct answer, even mentioning where the HBr is dissolved.



Question 4 (d)

Some very poor drawings were seen here. The narrow neck and/or the tap was missed.

It was rare to see the layers the wrong way round which suggested the density information had been ignored.

(d) Draw a diagram of the apparatus used to separate the ethoxyethane layer from the aqueous layer in **Step 5**. Clearly label the ethoxyethane layer.

[Densities: water 1.0 g cm⁻³, ethoxyethane 0.7 g cm⁻³]





Though the layers are correct, this is quite clearly a burette.

Results Plus Examiner Tip

Be familiar with common apparatus like a separating funnel.

(2)

(d) Draw a diagram of the apparatus used to separate the ethoxyethane layer from the aqueous layer in **Step 5**. Clearly label the ethoxyethane layer.

[Densities: water 1.0 g cm⁻³, ethoxyethane 0.7 g cm⁻³]

(2)



Question 4 (e)

Omissions of one or other acid were common, as were omission of the word 'concentrated'. So only about half the candidates gained this mark.

(e) The bromobenzene formed in this reaction can be nitrated to make 2,4-dinitrobromobenzene.

Identify, by name or formula, the chemicals needed for this reaction.

		(-)
ar concentrated	hydrochloric actid and	
concentrated	Sodium hydrovide hydroc Si	ulfuric acid
Results Plus Examiner Comments This mixture tends to spit or e	xplode!	Ius nitration'.

(e) The bromobenzene formed in this reaction can be nitrated to make 2,4-dinitrobromobenzene.

Identify, by name or formula, the chemicals needed for this reaction.

(1)

(1)

concentrated HNO,



Nitric acid alone will nitrate a phenol but not benzene.



Question 4 (f)

This was a good question for the better candidates. Weaker candidates found the mean of the three values, instead of multiplying them together appropriately.

1102

(f) 2,4-dinitrobromobenzene reacts with hydrazine hydrate to make 2,4-dinitrophenylhydrazine crystals.

The percentage yields for the reactions are:

75% for the formation of bromobenzene from benzene

70% for the formation of 2,4-dinitrobromobenzene from bromobenzene

70% for the formation of 2,4-dinitrophenylhydrazine from 2,4-dinitrobromobenzene

Calculate the overall percentage yield of 2,4-dinitrophenylhydrazine from benzene, for this series of reactions.



(f) 2,4-dinitrobromobenzene reacts with hydrazine hydrate to make 2,4-dinitrophenylhydrazine crystals.

The percentage yields for the reactions are:

75% for the formation of bromobenzene from benzene

70% for the formation of 2,4-dinitrobromobenzene from bromobenzene

70% for the formation of 2,4-dinitrophenylhydrazine from 2,4-dinitrobromobenzene 38.95

Calculate the overall percentage yield of 2,4-dinitrophenylhydrazine from benzene, for this series of reactions.





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75% for the formation of bromobenzene from benzene

70% for the formation of 2,4-dinitrobromobenzene from bromobenzene

70% for the formation of 2,4-dinitrophenylhydrazine from 2,4-dinitrobromobenzene

Calculate the overall percentage yield of 2,4-dinitrophenylhydrazine from benzene, for this series of reactions.

70 +25

= 95%



Question 4 (g) (i)

Very few candidates seemed familiar with this, though it is a standard procedure to melt one end of the tube in a hot Bunsen flame.

(g) The purity of the 2,4-dinitrophenylhydrazine crystals can be checked by carrying out a melting temperature determination using the Thiele tube apparatus shown below.



(i) The capillary tube must be sealed at one end. Describe how this is done.



(g) The purity of the 2,4-dinitrophenylhydrazine crystals can be checked by carrying out a melting temperature determination using the Thiele tube apparatus shown below.



(i) The capillary tube must be sealed at one end. Describe how this is done.

The end that is to be sealed is heated above a (intrablue part of the flame) (intrablue part of the flame) bursen burner (to melt it together The end **Examiner Comments**

(1)

An excellent answer.

Question 4 (g) (ii)

Again, very few candidates seem to have done this. Often the use of wires or rods were suggested.

Many heated the tube!

- (ii) When crystals are placed in the capillary tube they often stick in the top. Describe how to ensure the crystals reach the bottom of the capillary tube.
- (1)

Shake the capillary tube vigorously to ensure the crystals reach bottom. A 1 **lesuits Examiner Comments** Though this has some merit, the crystals would most likely spill out.

(ii) When crystals are placed in the capillary tube they often stick in the top. Describe how to ensure the crystals reach the bottom of the capillary tube.

(1)Shake the capillary tube and lightly tap the abide of the type with a stirrer to make themfall





Question 4 (g) (iii)

Weaker candidates managed to get the first mark for high boiling temperature. Only more able candidates were able to get the second mark. The better candidates saying that the solvent must not decompose at high temperature.

(iii) Dibutyl phthalate is often used as the liquid in the Thiele tube.

Suggest **two** properties of dibutyl phthalate that make it a suitable liquid for this purpose.

(2)

(2)

higher boiling point. high specific neat capac	-ાન્યુ.
Results Plus Examiner Comments Though arguably both of these reasons are correct, the second is a physical property and this is a chemical examination.	Results Plus Examiner Tip Always give chemical reasons first, in this type of question.

(iii) Dibutyl phthalate is often used as the liquid in the Thiele tube.

Suggest **two** properties of dibutyl phthalate that make it a suitable liquid for this purpose.

It must be having a boiling	ng temperature higher than
that of the sample.	
It stocked must be a clea	r liquid, so that the any
changes to the sample won't be	e obscured by the liquid.
Results Plus Examiner Comments	Results lus
The high boiling temperature is fine. The clarity of the liquid was deemed just sufficient - especially with the reason given.	Always answer questions as fully as possible.

Question 4 (g) (iv)

Answers here showed the lack of practical experience of many candidates. The word 'range' in the question was often ignored.

Values were often ridiculous.

(iv) The melting temperature for crystals of Suggest the temperature range over w melt before and after purification by red	2,4-dinitrophenylhydrazine is 201°C. hich you would expect the crystals to crystallization. (2)
Before recrystallization 1582-2	2500 - 170°C - 220°C
After recrystallization 199°C - 2	03°C
Results Plus Examiner Comments The first range is far too large. The second is just acceptable - though it is unlikely that crystals will melt above their melting temperature.	Results Lines Examiner Tip Remember what you found when finding a melting temperature in the laboratory.

Suggest the temperature **range** over which you would expect the crystals to melt before and after purification by recrystallization.

(2)

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After recrystallization 203°c

Before recrystallization 180°C





Twice before answering and then afterwards to check you have answered it.

Paper Summary

To improve on this paper candidates should:

- Treat practical work seriously
- Constantly be considering the reasons for all the procedures that they use
- Record and remember observations that they make during experiments
- Practise drawing fully labelled diagrams of apparatus.

Grade Boundaries

Grade boundaries for this, and all other papers, can be found on the website on this link: http://www.edexcel.com/iwantto/Pages/grade-boundaries.aspx





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