



Examiners' Report January 2013

GCE Chemistry 6CH02 01



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Introduction

This paper tested a wide range of Unit 2 material and provided good opportunities for candidates to show their knowledge and understanding of the chemistry covered by the Specification.

The quality of written communication was very variable, and in some cases, responses were unable to be rewarded as the meaning was unclear. In the calculations on this paper, the routine calculations for the iodine/thiosulfate titrations were well done. One calculation tested understanding of the procedure by asking why a certain amount of potassium iodide was suitable; this was often not attempted, and when a calculation was done it was often not relevant to the question. The calculation of percentage difference in titration readings produced a lot of rounding errors. One question asked for an experimental method to compare the thermal stability of two carbonates. Some answers were excellent but many of the procedures proposed would be impossible to carry out in practice, or would not give the data necessary for the comparison. The answers suggested that many candidates have little experience of doing experiments, or attempting to solve practical problems.

Question 21 (a) (i)

This question was well answered. Most candidates were able to identify the three oxidation numbers of chlorine, but some did not specify the substance in which they were found. Other candidates gave charges instead of oxidation numbers, saying that Cl⁻ and Cl⁺ formed. This was not given credit. Only a very small minority thought that the oxidation numbers of hydrogen and oxygen were changing in the reaction.

Careful use of language was needed to gain the third mark. The definition of disproportionation was generally well known, but some answers just said that oxidation and reduction were occurring. It was necessary to say that the same species, or element, was both oxidized and reduced.

Answers saying that that "disproportionation has occurred because chlorine has gone from 0 to +1 and -1", were not allowed as this statement is just repeating the values of the oxidation numbers.

21 Chlorine is used to prevent the growth of bacteria in swimming pool water. It reacts as shown below. $Cl_2(aq) + H_2O(l) = HCl(aq) + HClO(aq)$ 4+1-2 +2 -2 +1-1 0 (a) (i) By giving appropriate oxidation numbers, explain why this is a disproportionation reaction. (3)Pispapartionation reaction occurs when an eter atom of an element is oriched reduced in the same reaction. In this case Chlorine is reduced from 0 to -1. 0 to +1 and is also from chlorine has been oxilized and & reduced, it is a disproportion since it has both accepted and lost electrons.

Results lus Examiner Comments

This candidate has given the three oxidation numbers correctly but not identified the compounds in which the oxidation number is -1 and +1.



In disproportionation the same type of atom is both oxidized and reduced, but remember that a single atom cannot be oxidized and reduced simultaneously. This answer was allowed but it would have been better to make it clear that the changes are not happening to the same atom.

21 Chlorine is used to prevent the growth of bacteria in swimming pool water. It reacts as shown below. $Cl_2(aq) + H_2O(l) \implies HCl(aq) + HClO(aq)$ (a) (i) By giving appropriate oxidation numbers, explain why this is a disproportionation reaction. (3)to start with is a because the indecute is oxidation male re's HCI and reduced Solit oxidised to diatomic then and HUO This 'n reaction disproportination J a. and Chlorie poth. reduces the oxdised J Examiner Comments This candidate has confused the oxidation number changes associated with oxidation and reduction, by saying that production of HCl is oxidation. However the oxidation numbers are correct. Kesut S **Examiner Tip**

During reduction the oxidation number is reduced i.e. gets less.

Question 21 (a) (ii)

Many candidates correctly stated that the equilibrium would move to the left, and then said this meant that more reactants would form. This simply rephrases the first statement, and was not allowed as an explanation. A comment about using up some of the added hydrochloric acid or restoring equilibrium was needed. Le Chatelier's principle was sometimes quoted, but to gain credit, it had to be applied to this equilibrium, and just saying that the equilibrium moves to counteract the change in the system did not do this.

(ii) State and explain the effect on the position of equilibrium if concentrated hydrochloric acid is added to a sample of chlorinated swimming pool water. (2)n would shift to the left as system 2, so an increase in MCI would mean river shifts to the left due to an increase in pressure **Examiner Comments** This answer gives the correct effect on the position of equilibrium. However, more is required in the explanation than saying that when HCl is added, the equilibrium moves to the left because the HCl concentration has increased. **Examiner Tip** The reverse reaction will occur when the HCl concentration is increased. This restores equilibrium by using up some of the added HCl and converting it to chlorine and water. All the added HCI will not be used up - if it was, the concentration of the substances on the left of the equation would be higher than

they were in the original equilibrium state.

(ii) State and explain the effect on the position of equilibrium if concentra hydrochloric acid is added to a sample of chlorinated swimming pool	
	(2)
The position would not change because an	increase
in moles would increase the concentration equ	
the number of moles on each side are equal	4

ResultsPlus

Examiner Comments

When equilibrium is reached again after concentrated hydrochloric acid is added, both reactants and products will be more concentrated but it is incorrect to say that the equilibrium position would not change.

The number of moles on each side of the equation is important in equilibrium reactions involving gases, as a change in pressure will cause a change in equilibrium position. However, in this reaction the species are all in solution.



You need to consider the number of moles on each side of the equation in equilibrium reactions involving gases, as a change in pressure will cause a change in equilibrium position. However, in this reaction the species are all in solution so you just have to think about the effect of a change in concentration of one of them.

Question 21 (b) (i)-(vi)

There were many incorrect half-equations given in (i), some with incorrect numbers of electrons and others showing no electrons at all. Candidates are more likely to balance the charges correctly if the charge on the electrons is shown. However it appeared as if some candidates did not know that half-equations included electrons.

The equations given in (ii) were more often correct, and were marked independently of the half-equations in (i). Overall equations should not show electrons. The common error was in the number of hydrogen ions, and if candidates had checked the total charge on each side of the equation they might have realised their mistake.

The calculation of the number of moles of iodine in (iii) was usually well done, though a few candidates stopped at the stage of calculating moles of thiosulfate ions. These candidates rarely said what their numbers referred to, making it difficult for them to gain marks as transferred errors. Very few errors in significant figures or rounding of values were seen here. If an equation was given in (ii), the calculation of the number of moles of CIO⁻ should have been based on this and the number of moles of iodine calculated in (iii).

Calculation of the concentration in (iv) sometimes caused difficulty; a few were out by a factor of ten and volumes of bleach other than 25 cm³ were used. The number of moles was sometimes multiplied by the volume instead of being divided by it. Candidates are advised to give numbers in standard form, otherwise zeros were often lost going from one part of a question to another.

In (vi), many candidates simply confirmed that 1.5 g of potassium iodide contained 9.04×10^{-3} moles, as stated in the question. They needed to show that this amount of potassium iodide was an excess, either by calculating the number of moles of iodide ions which reacted with the ClO⁻, or comparing the number of moles of ClO⁻ which could react with 9.04×10^{-3} moles of iodide with the number present in the bleach.

(b) In a similar reaction, chlorine reacts with sodium hydroxide to make household bleach.

 $Cl_{2}(aq) + 2NaOH(aq) \rightarrow NaCl(aq) + NaClO(aq) + H_{2}O(l)$

The concentration of NaClO in diluted bleach was measured by titration. A 25.0 cm³ sample of bleach was pipetted into a conical flask. Approximately 1.5 g of solid potassium iodide and 10 cm³ of hydrochloric acid with concentration 2.00 mol dm⁻³ were added. Each mole of ClO⁻, from the NaClO in the solution of bleach, produced one mole of iodine, $l_{2'}$ which was titrated with sodium thiosulfate solution.

(i) Complete the ionic half-equations below for the reaction of ClO⁻ with acidified potassium iodide by balancing them and **adding electrons** where required.

(2)

 $ClO^{-} + \dots 2 \dots H^{+} + 2e^{-} \rightarrow Cl^{-} + H_2O$ $\dots 2 \dots l^{-} \qquad \rightarrow l_2 + 2e^{-}$

(ii) Use your answer to (a)(i) to write the overall ionic equation for the reaction between CIO⁻ and I⁻ ions in acidic conditions.

(1)

 $CIO^{-} + 2H^{+} + 2I^{-} \rightarrow I_{2} + CI^{-} + H_{2}O$

(iii) The iodine in the sample required a mean (average) titre of 24.20 cm³ of 0.0500 mol dm⁻³ sodium thiosulfate solution. Thiosulfate ions react with iodine as shown below.

$$2S_2O_3^{2-} + I_2 \rightarrow S_4O_6^{2-} + 2I^{-}$$

Calculate the number of moles of iodine in the solution.

$$n(5_20_3^{2^-}) = 0.05 \times 0.02420$$

= 1.21 \times 10^{-3}
$$n(1_2) = (1.21 \times 10^{-3}) \div 2$$

= 6.05 \times 10^{-4}

(iv) What is the number of moles of CIO⁻ ions in the sample of diluted bleach?

 $n(cio^{-}) = 6.05 \times 10^{-4}$

(v) Hence calculate the concentration, in mol dm⁻³, of CIO⁻ in the diluted bleach.

 $C = \frac{6.05 \times 10^{-4}}{0.025} = 0.0242 \text{ moldm}^{-3}$

(vi) 1.5 g of potassium iodide, KI, contains 9.04×10^{-3} mol of I⁻. Use your answers to (b)(ii) and (b)(iv) to show by calculation why this amount was suitable.

mole ratio 1:2 $(6.05 \times 10^{-4} \times 2) = 1.21 \times 10^{-3}$ = 0.00121 moles (1 required)

9.04 × 10-3 = 0.00904 (added)

0.00904 moles added of I - which is more than required.

Results Plus Examiner Comments This is an example of an excellent answer. The equations are clearly set out: in the calculations it is clear what each number

clearly set out; in the calculations it is clear what each number refers to.

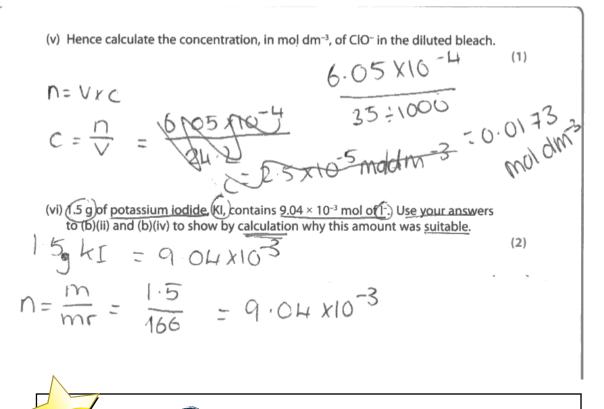
(2)

(1)

(1)

(b) In a similar reaction, chlorine reacts with sodium hydroxide to make household
beach
$$25$$

 $C(4a) + 2NaOH(a) + NaCl(a) + NaClO(a) + H_0(1)$
The concentration of NaClO in diluted bleach was measured by titration.
Ac50 cm Sample of bleach, was pipetted into a conical flask, Approximately
(15) poil solid notassium iodide and 00 mb hydrocholoris add with concentration
of bleach, produced fine mole of COO from the NaClO in the solution
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of bleach produced fine mole of cOO from the NaClO in the solution
(2)
 $CO = -2 - - - - - - -1$
(4) Use your apswer to (a)(1) to write the overall ionic equation for the reaction
betweer (CO and f) ons in acidic conditions.
 $C = 0 + 2 + 1 + 2 = - - C + 1 + 1_2 = 0 + 1_2$.
(4) The joding in the sample required a mean (average) titre of (4.20 cm) of
(0.0500 mol dm) sodium thiosulfate solution. Thiosulfate ions react with joding
as shown below.
 $250^{+} + 1_{-} + 50^{+} + 21$
Calculate the number of moles of iodine in the solution.
 $N = 2 + 2 + 2 - CM^{3}$
 $N = N = I_2 = \frac{1}{2}(1 \times 10^{-3})$
 $= 1 + 2 \times 10^{-3}$
(v) What is the number of moles of CIO ions in the sample of diluted bleach?
 $n = \frac{1}{2} + \frac{1}{2} \times 10^{-4}$.
(1)
 $= \frac{1}{2} + \frac{1}{2} - \frac{1}{2} + \frac{1}{2} - \frac{1}{2}$



ResultsPlus Examiner Comments

In this answer no electrons are shown in the half-equations in (i).

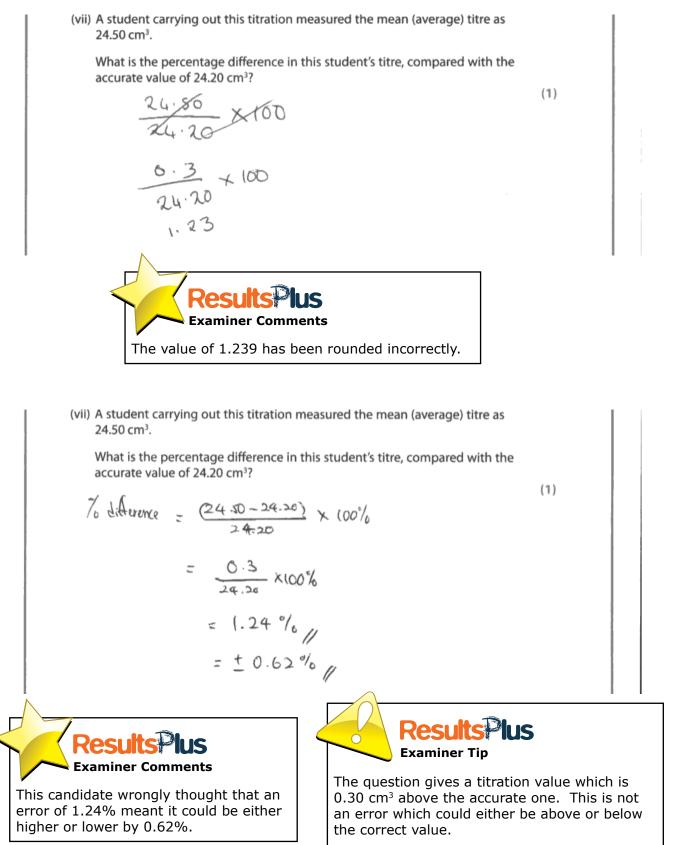
In (iii) the number of moles of iodine is labelled as such, and ideally the number of moles of thiosulfate ions would be labelled too. The volume used to calculate the concentration in (v) is incorrect, and the calculation in (vi) is not based on the reacting ratios of iodide and CIO^{-} ions.



The charge on each side of an equation should be balanced. The first equation here has a total of one positive charge on the left and one negative charge on the right. When writing half-equations, balance the numbers of atom of each element first, then balance the charge by adding electrons to the side where they are needed.

Question 21 (b) (vii)

This was a simple calculation, but errors occurred when candidates rounded the answer wrongly. The correct value was sometimes rounded to 1%, which was not allowed. Some candidates used very complicated methods for the calculation. This increased their chances of making an error.



Question 21 (b) (viii)

The question was about a titration value which was higher than the accurate one, so reasons for the difference had to be consistent with a higher titre. Suggestions such as impurities in the solution would affect both a student and a more accurate worker, so these were not allowed. Simply saying that the student had overshot the end-point was rephrasing the fact that the titre was too high, and a reason why this might have happened was required. Many answers referred to human error, but something more specific was required.

(viii) The difference between the student's mean titre and the accurate value was **not** due to the limitations in the accuracy of the measuring instruments. Suggest one possible reason for this difference. (1)subjective nature of when the titestion changel colaur accortain **Examiner Comments** The judgement of when the colour changes is subjective, and was one of the allowed answers. (viii) The difference between the student's mean titre and the accurate value was **not** due to the limitations in the accuracy of the measuring instruments. Suggest one possible reason for this difference. (1)amount prosted with the & air got l'ost **Examiner Comments** If the chlorine in the bleach had evaporated the titre would have been lower. Reaction with the air would not occur. **Examiner Tip** In a question like this you have to think why the titration reading might be too high, but factors which affect an accurate worker as well as a student could not cause the difference.

Question 21 (c)

This was answered correctly by most candidates, though a few just said that the ozone layer would be affected, rather than making clear that it would be depleted. Many added comments saying that chlorine radicals were involved, or stated the consequences of the thinning of the ozone layer.

A minority of answers said, incorrectly, that the damaging effect was global warming or acid rain.

(c) Suggest one damaging effect to the upper atmosphere which could be caused by the presence of chlorine compounds. develop which (1) free radicals/ deplete the ozone Chlorine which means nove UV, so more skin cancer Results **Examiner Comments** This was a typical correct answer. (c) Suggest **one** damaging effect to the upper atmosphere which could be caused by the presence of chlorine compounds. develop which (1) Free radicals/ depiete the ozone Chlorine Which means more UV, so more skin cancer auer **Examiner Comments** This answer is typical of candidates who are not clear about the different environmental problems caused by chemicals in the atmosphere. **Results Plus Examiner Tip** Make sure you know about the causes of acid rain, global warming and ozone depletion. Each of these problems has a different cause.

Question 22 (a) (iii)

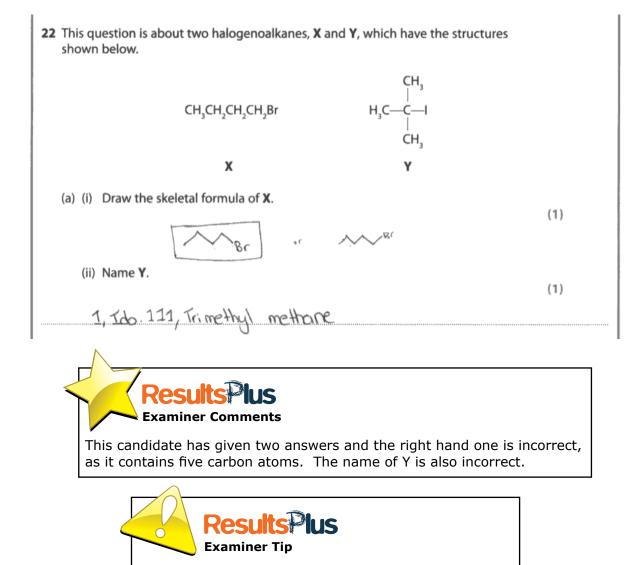
The reaction of ammonia with a halogenoalkane was not well known. Even when the products were correct many unbalanced equations were given. If the equation showed formation of the amine plus ammonium bromide, rather than just hydrogen bromide, then two moles of ammonia were needed. The product was often thought to be an amide or a nitrile, and when candidates attempted an IUPAC name for an amine they often omitted the number showing the position of the amine group. Other incorrect suggestions such as butamine appeared regularly.

(iii) Write an equation for the reaction of X with an alcoholic solution of ammonia, and name the organic product. CH3CH2 CH2 CH2Br + NH3 my CH3(H2 (42(H2NH2 (2) m)) + HBring) Bulylamine KOH ... annad Name of product. **Examiner Comments** This answer was given both marks. (iii) Write an equation for the reaction of **X** with an alcoholic solution of ammonia, and name the organic product. (2)CH3CH2CH2CH2Br + NH3 -> CH3CH2CH2CH2NH2 + NH2Br. putan-1- amore. Name of product. **Examiner Comments** This equation is not balanced. Two moles of ammonia are needed. A mark was given for this version of the name as the number was included.

Question 22 (a) (i)-(ii)

The skeletal formula for bromobutane, X, should have shown a zigzag line representing the four carbon atoms, followed by another bond to the bromine atom. Bonds were often drawn of different lengths; this was not penalised. Some very short bonds to Br were shown, but were given the mark as long as they were present. Bonds were often shown at unusual angles, and again this was accepted. A few candidates misread the question, and gave a displayed formula, and displayed or skeletal formulae for Y were sometimes given instead of the name.

22 This question is about two halogenoalkanes, X and Y, which have the structures shown below.	
CH ₃ CH ₃ CH ₂ CH ₂ CH ₂ Br H ₃ CCI CH ₃	
X Y	
(a) (i) Draw the skeletal formula of \mathbf{X} .	(1)
(ii) Name Y. 2-methyl-2-iodo propone	(1)
Results Plus Examiner Comments This shows a common mistake in the skeletal formula, as it represents a molecule with only three carbon atoms.	
After drawing the lines representing the carbon skeleton, you need to add another bond going to the other atom or atoms attached to the carbon.	

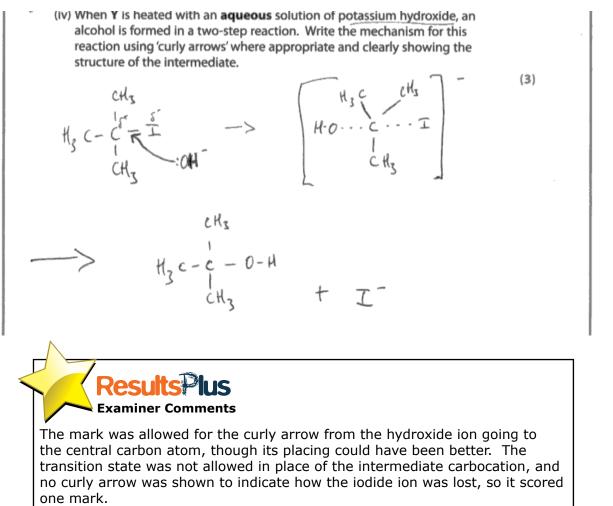


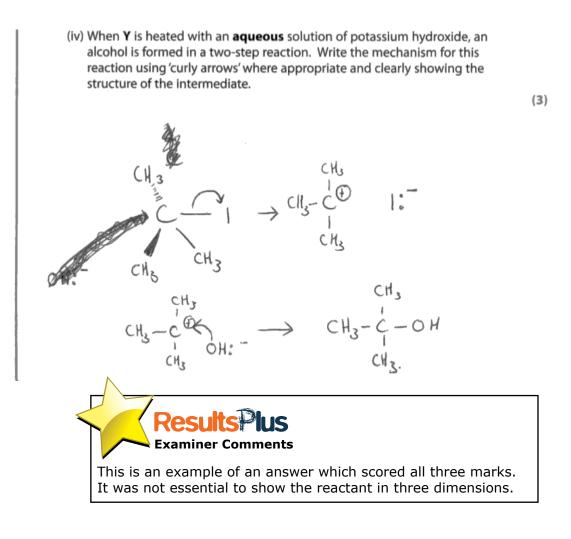
If you do not want an answer to be marked you must strike it out clearly.

Question 22 (a) (iv)

Most candidates had the correct ideas for this mechanism, though some interpreted a twostep mechanism as being the SN2 mechanism. In this case the transition state was not allowed in place of the carbocation intermediate, but two marks could be gained for showing two correct curly arrows. Some candidates were careless in the positioning of curly arrows, such as the first one coming from the C atom and not the C-I bond. This arrow was often shown going past the I atom into space. A significant minority showed the hydroxide ion attacking the central carbon concurrently with the loss of the iodide ion, but then formed the carbocation intermediate. The intermediate carbocation was often shown with a δ +, rather than a full charge, and the charge on the hydroxide ion was often missing.

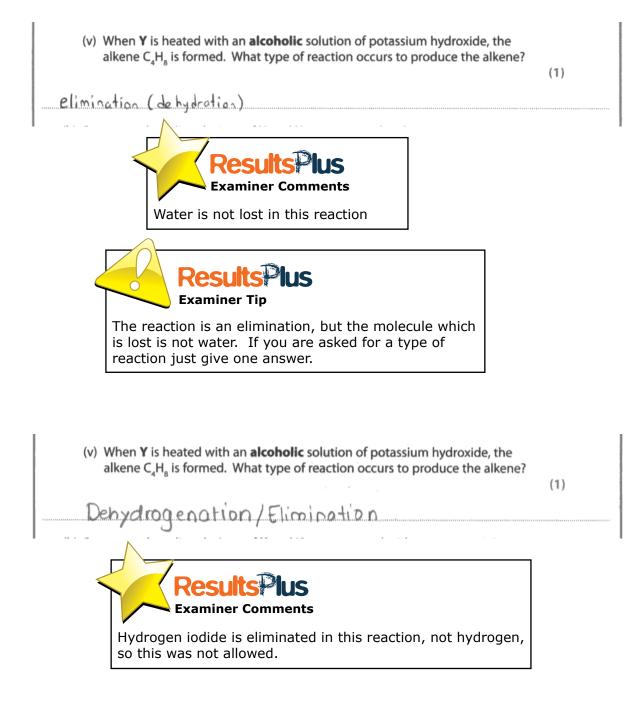
There was clearly a lack of understanding that the mechanism represented a sequence of events. Many candidates were confused about the role of potassium, and showed KOH as a covalent molecule, with a curly arrow causing formation of the hydroxide ion.





Question 22 (a) (v)

The reaction was an elimination, but many candidates expanded on this incorrectly, for example saying that it was elimination/oxidation or electrophilic elimination.



Question 22 (b)

The silver bromide formed with X would be seen as a cream, off-white or pale yellow precipitate. The description "white" was not allowed. The silver iodide formed with Y would be seen as a yellow precipitate. For full marks it had to be stated that a precipitate formed; the colours alone were not enough for both marks. Some candidates suggested several observations, such as a cream precipitate and evolution of brown fumes or fizzing. This made the overall observation incorrect and was not allowed.

(b) Separate ethanolic solutions of X and Y were dissolved silver nitrate. Describe what would differences in the rates of reaction.	
Observation with X	
silver bromide silver brom	ide (AgBr)
Observation with Y	
ายการการการการการการการการการการการการการก	(Ag I)
Results Ius Examiner Comments	
The answers given are not obse	rvations and did not score.
Results Plus Examiner Tip	
If you are asked what you would should give observations, and ne	
(b) Separate ethanolic solutions of X and Y were dissolved silver nitrate. Describe what would differences in the rates of reaction.	
Observation with X	
Pale - yellow solution	
Observation with Y yellow cloudy soluti	
Results Plus Examiner Comments	Results Plus Examiner Tip When you describe the product of a
One mark was allowed here for two correct colours, as there was a contrast between pale yellow and yellow.	reaction, give its colour and its physical state e.g. green gas or, as in this guestion, vellow precipitate.

Question 22 (c)

The majority of answers stated that the C-Br bond was stronger than the C-I bond, but understanding of the reason for the difference was very limited. As the bromine atom is smaller, the nuclei of the two elements in the bond are closer, but candidates who referred to the ionic radius of bromine did not get the mark. Other candidates thought that intermolecular forces were involved and said that, because the iodine atom had more electrons, its London forces would be greater. Some candidates thought the explanation was the reactivity of bromine compared to iodine. Many candidates answered in terms of the greater electronegativity of bromine causing a greater dipole and increasing the strength of the bond, and this was allowed.

			nary halogenoa n and the halog		ected by the stren	gth of
	C—Br bon gth differs.	d weaker or s	tronger than th	e C—I bond?	Explain why the b	oond
						(2)
Iodule	has ,	a larger	ionic ro	dins the	n bromide	This
means	the .	abtraction	between	the out	electrons	in the
condent	bond	and the	positively	changed n	unders is h	exter Calso
due to	more chu	elding). Th	is means	C- Bi	r is stron	ger.

This candidate refers to the ionic radius of bromine, and then makes a statement which describes why iodine is less electronegative than bromine. Neither comment explains the



strength of the bond with carbon.

Short bonds are stronger than long bonds because when the nuclei of one atom are close to the electrons of the other atom they have a stronger attraction. (c) The rates of hydrolysis of primary halogenoalkanes are affected by the strength of the bond between the carbon and the halogen atom.

Is the C—Br bond weaker or stronger than the C—I bond? Explain why the bond strength differs.

(2)C- to bond stronger, less repulsion of nucleus as electrons shelld repelling and have a stronger attraction to the other elements nucleus. esults **Examiner Comments** This candidate was wrong in choosing C-I as the stronger bond. There is confusion about what two things are attracted in a bond. The nuclei are not attracted, but are held together by their attraction for the shared electrons.

Question 23 (a) (i)-(ii)

The first part of the question asked for a bond angle in cyclohexanol and stated that it differed from the angle in the two dimensional diagram. Many candidates did not realise that the carbon atom involved is bonded to two hydrogen atoms not shown in the skeletal formula, and the most common answer was 120 degrees, though angles from 60 to 200 degrees were seen. As it forms four bonds, the angle would be the normal tetrahedral angle as in methane.

In the second part most candidates realised that the bond angle would change as nonbonding electrons were present on oxygen, but some thought there would only be one lone pair, and answers saying that the bond contained two lone pairs, without specifying that they were on the oxygen did not get the mark. A significant number found difficulty in expressing their answer coherently, and just mentioning maximum separation or minimum repulsion did not get a mark unless it was in a suitable context. In some cases candidates wrote about lone pairs pushing bonding pairs together despite giving an angle in (ii) which was bigger than the angle in (i).

Some candidates did not read the question carefully enough and tried to explain why the bond angle in (i) was 109.5 in terms of bond pairs & lone pairs.

23 The skeletal formula of cyclohexanol is shown below. c-10 = M nd с с-с он а ОН (a) (i) The actual bond angles differ from the angles in the two dimensional diagram shown. What is the angle of the C—C—C bond labelled a? (1)120° Angle *(ii) What is the angle of the C—O—H bond? Justify your answer, explaining why the size of the angle is different from the angle in (i). (3) 180 Angle Explanation Or Oxygen has 2 lone pairs non-bonding pairs of electrons. These repel eachothe to a point of maximum seperation. non-bonding pairs as repel more more than bonding pairs. **Results Plus Examiner Comments** This candidate has drawn the carbon skeleton of cyclohexane, but chosen an incorrect bond angle in (i) as the hydrogen **Examiner Tip** atoms have been forgotten. When you have to predict bond In the second part the bonding electrons have been forgotten, angles in skeletal formulae, and though the non-bonding electrons are said to repel each remember there are hydrogen atoms other to maximum separation, the bond angle chosen is greater which are not shown. than in (i). The answer got one mark for stating that the

oxygen atom had two lone pairs.

23 The skeletal formula of cyclohexanol is shown below. OH (a) (i) The actual bond angles differ from the angles in the two dimensional diagram shown. What is the angle of the C-C-C bond labelled a? (1)60° Angle *(ii) What is the angle of the C—O—H bond? Justify your answer, explaining why the size of the angle is different from the angle in (i). (3) 104.5 Angle 60 Explanation Concentrating only on the carbon atom which is bonded The oxygen atom created 2 bonds and has 2 lone pairs. IF we treat the lone pairs as bonds then the angle would pereis slightly more repulsion from the two lone 109.5 is taken off for each. Therefore the ang 510 Examiner Comments This candidate has also selected an incorrect bond angle in (i). In (ii) the lone pairs are said to cause more repulsion, presumably meaning more than between bonding pairs, even though the angle suggested is bigger than in (ii) and the third mark was not given here.

Question 23 (b) (i)

There were many very full answers, and only a few identified the products without describing what would be seen. Candidates are likely to have seen this reaction with ethanol and are not expected to know how it might differ with cyclohexanol, so they should have concentrated on changes due to the chemical reaction such as bubbles and a white solid forming along with the disappearance of the sodium. Comments about sodium sinking or floating or going on fire were ignored. Temperature change was not allowed as it is not something which is seen.

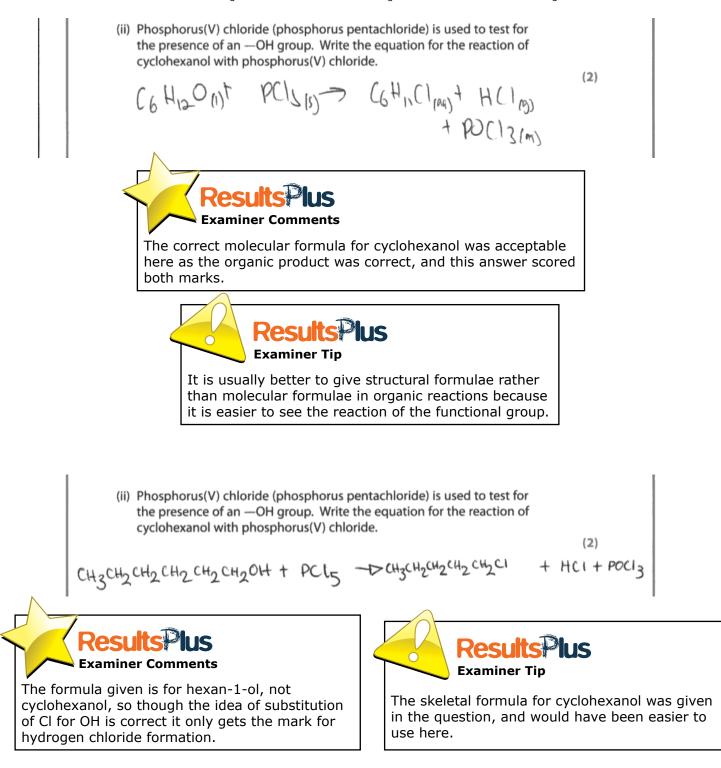
(b) (i) Suggest what you would expect to see when cyclohexanol reacts with sodium.
(2)
The No doisplaces the to buildeles of loydragers Jas
and a sodium sout / proceipiture form (white)
Results Plus Examiner Comments
This answer gained both marks for the bubbles and the white precipitate. It was not necessary to identify the chemical products.
Saying that the sodium became coated with a white solid would have been allowed.
(b) (i) Suggest what you would expect to see when cyclohexanol reacts with sodium. (2)
You would see fizzing as the sodium dissoured
and began to react with the cyclohexanal.
Results lus Examiner Comments
This also was given both marks for the fizzing and dissolving of sodium.
Candidates who said that the cyclohexanol dissolved were not given the mark.
Results Ius Examiner Tip

Remember that fizzing is another word for effervescence, so "fizzing and effervescence" would count as one observation.

Question 23 (b) (ii)

Candidates who used the skeletal formula for cyclohexanol were more likely to get this correct than those who tried to write a molecular formula. Some candidates gave the formula for phenol, C_6H_5OH , rather than $C_6H_{11}OH$. A formula written as $CH_2CH_2CH_2CH_2CH_2CH_0H$ was not accepted unless a bond was shown from the first to the sixth carbon. A fully displayed formula was accepted, but would have taken the candidates more time to write than the skeletal formula!

Most candidates got the mark in this question for hydrogen chloride formation, but there were many errors with $POCI_3$ being written as PCI_3 , or occasionally $PCIO_3$.



Question 23 (b) (iii)

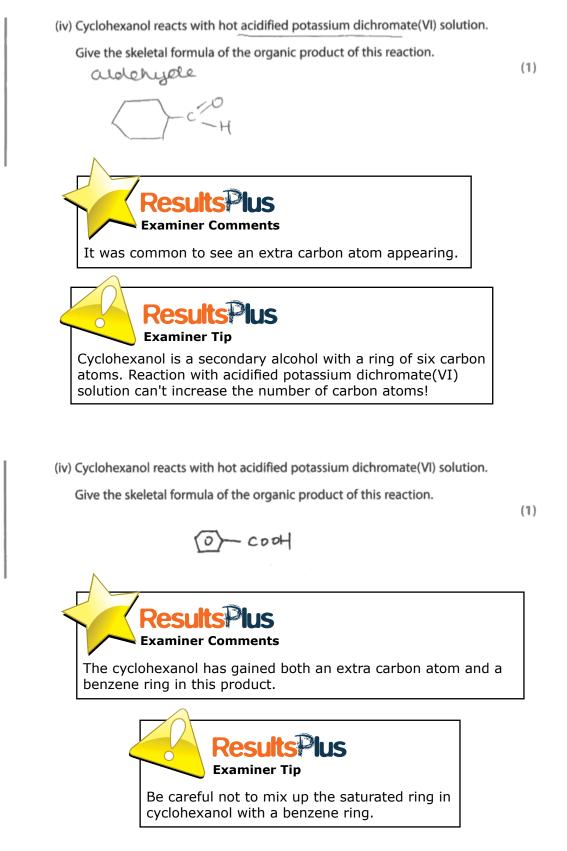
Many candidates suggested testing hydrogen chloride with litmus, but this just detects an acidic gas and was not allowed here. When the question asks for a chemical test the answer expected is one where the candidate could write an equation, if asked.

Many answers said that the hydrogen chloride would appear as steamy fumes, which is correct but not a chemical test. Some candidates suggested bubbling the gas into ammonia, rather than allowing it to mix with fumes from ammonia, in which case the dense white smoke would not be seen. Bubbling the gas into silver nitrate solution and seeing a white precipitate was allowed. However if this was in conjunction with the gas bleaching litmus paper it was not.

 (iii) Give the chemical test you could use to identify the gas produced, and the observation you would make. (1) blue Moist, lithus paper would turn red then bleach to identify the could be produced 	
Results Lus Examiner Comments The bleaching here would indicate chlorine, and there is no reference to the dense white steamy fumes being formed in a reaction with ammonia.	
Results Pus Examiner Tip If you are asked to give a chemical test, try to think of something more specific than using an indicator.	
(iii) Give the chemical test you could use to identify the gas produced, and the observation you would make. (1) React with Amnonia, dense whete funes. Results Plus Examiner Comments This is an example of one way to score the mark.)

Question 23 (b) (iv)

Cyclohexanol is a secondary alcohol, so would form a ketone on oxidation. Many answers showed formation of a carboxylic acid, with either an extra -CO₂H attached to the ring, or a 5-bonded carbon on the ring connected to an oxygen and an -OH group. Another common error was to break the ring and put a C=O bond either at the end or in the middle of the chain.



Question 23 (b) (v)

The colour change of orange to green in this redox reaction was well known.

(v) What colour change would you observe as this reaction takes place?	(1)
From Yellow Orange to Green.	
Results Plus Examiner Comments	
As potassium chromate(VI) is yellow, the initial colour of yellow-orange was not allowed here.	
(v) What colour change would you observe as this reaction takes place?	(1)
From green to ocange	******
Results Plus Examiner Comments The colour change must be given the right way round to	

Question 23 (c)

The question asked for the molecular formula of a fragment of mass 82, so the answer $C_6H_{10}^{(+)}$ should have been deduced from the formula and structure of cyclohexanol. Structural formulae were not accepted, and other formulae with mass 82 such as C_5H_6O were not allowed as they would be unlikely to form from cyclohexanol. If the charge was missing it was not penalised.

(c) The mass spectrum of cyclohexanol has a prominent peak at mass / charge ratio 82. Suggest the molecular formula of the fragment which causes this peak. (1)CH2CH2CH2CH2CH2C ar C6H10 (Total for Question 23 = 12 marks) TOTAL FOR SECTION B = 41 MARKS C₆ H₁₀. C₅ H H WI = 11 / 12 C5 . 60 $C_6 = 72$. 0=16. CH2 CH2 CH2, CH2 CH2 CH2 CH2 140 **Examiner Comments** The question asked for a molecular formula, and the answer does not say which of the two versions is the molecular one. **ResultsPlus Examiner Tip** Make sure you know the difference between a molecular and a structural formula, as a question may ask for a specific type of formula.

$[C_{6}H_{13}]^{-1}$	[C6H10	,] ⁺
		(Total for Question 23 = 12 marks
GHZZ+104		TOTAL FOR SECTION B = 41 MARK
48+9+1+16 50+12+16 = 88 u(12) = 48 19	65	OH). CH3CH2CH2CH2CH2CH2OH
Examin		ents ers are given the mark cannot be

Question 24 (a)

The link between polar bonds and absorption of infrared radiation due to changes in bond polarity was not well known, though many candidates managed to get one of the two marks.

Many candidates said that carbon dioxide is a polar molecule which is incorrect. It contains polar bonds, which is a necessary condition for absorbing infrared radiation. Also there is a change in polarity when infrared radiation is absorbed. A significant number of candidates did refer to this change in polarity, even though they had made no statement about the presence of polar bonds.

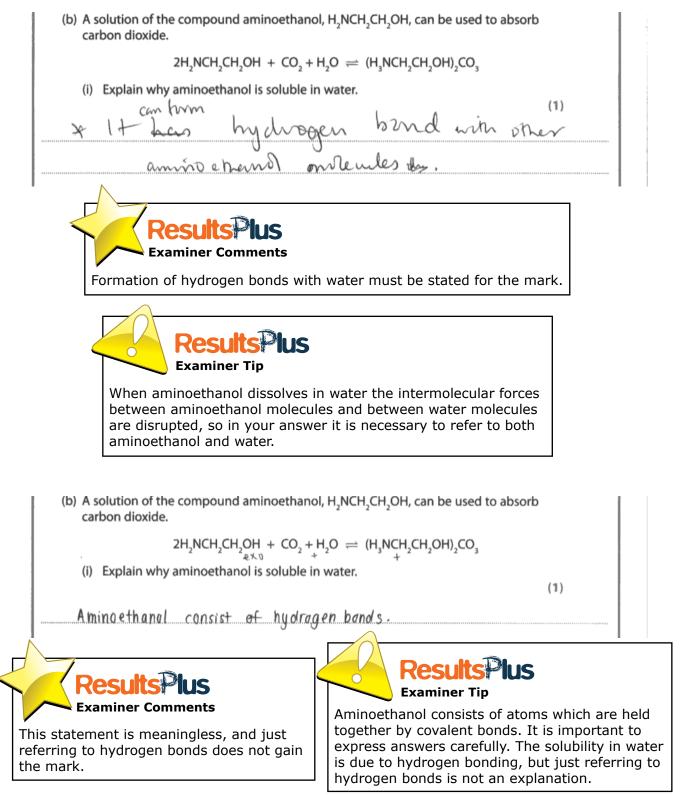
(a) Greenhouse gases can absorb infrared radiation. Explain why carbon dioxide absorbs infrared radiation but oxygen cannot. (2)Because Carbon dioxide has bolar bonds, & can & vibrate - And Vibrati with pon therefore land absorb infared **Examiner Comments** This candidate knows part of the answer, but does not state that there is a change in polarity when the molecule vibrates. (a) Greenhouse gases can absorb infrared radiation. Explain why carbon dioxide absorbs infrared radiation but oxygen cannot. (2)has bandy carbon dioxide is a polar melequice as the dipoles do not cancel vibrate when absorbing can out a change in dipole moment. But Oxyge ther bonds and has no change in dipole Dolar intraved counted absorb **Examiner Comments**

This answer was allowed both marks, for saying that carbon dioxide has polar bonds, and that there is a change of dipole moment when it vibrates. The comment that it has polar bonds as the dipoles do not cancel was not rewarded - it could refer to the different electronegativities of carbon and oxygen.

Question 24 (b) (i)

Aminoethanol is soluble in water because the energy needed to break the hydrogen bonds between water molecules is compensated by energy released when aminoethanol forms hydrogen bonds with water.

All that was required here was to say that aminoethanol can form hydrogen bonds with water. However some candidates said there was hydrogen bonding between aminoethanol molecules which, though true, does not answer the question. Others thought that being a polar molecule was enough to make it water soluble.



Question 24 (b) (ii)

For this question there were some excellent well-reasoned answers. Many candidates had clearly been taught and had learned the correct phraseology. Answers were usually in terms of the reverse reaction being endothermic, making the forward reaction exothermic. However, vague answers saying that the equilibrium moved to resist the change were also common.

(ii) The position of this equilibrium moves to the left on heating. This frees the captured carbon dioxide for storage. Use this information to decide whether the forward reaction is exothermic or endothermic. Explain your answer. (2)The formare reaction is exotherance because increasing tensproture would more the position of equilibrium to the adothermis direction and since it nover to we left reverse reaction must be and a Musimic the **Examiner Comments** This was typical of many of the answers seen and scored both marks. (ii) The position of this equilibrium moves to the left on heating. This frees the captured carbon dioxide for storage. Use this information to decide whether the forward reaction is exothermic or endothermic. Explain your answer. (2)mi, as by Le Chatchin is endether librin will more to the en and the stand of the 10.00 142 8 10 **Examiner Comments** This candidate has not applied Le Chatelier's principle correctly. **Examiner Tip** The equilibrium will move to use up the heat supplied. As the equilibrium moves to the left,

the reverse reaction is the one using up heat.

Question 24 (c) (i)

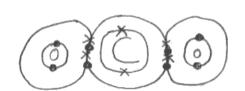
The question did not specify whether the electrons in the inner shells of carbon and oxygen had to be shown. However if the candidate chose to show them they had to be correct. The electrons in the double bonds were often shown correctly, but mistakes were often seen in the numbers of non-bonding electrons in oxygen.

	Percentage	Melting temperature / K
Methane	95.2	91.1
2-methylpropane	0.8	113.7
Butane	0.9	134.7
Other hydrocarbons	2.4	
Carbon dioxide	0.7	216.5

(c) The composition of a sample of natural gas and the melting temperatures of four of its components are shown below.

(i) Draw a dot and cross diagram for carbon dioxide.

(2)





This candidate has included the inner shell electrons correctly, but has forgotten the two lone pairs on each oxygen atom.



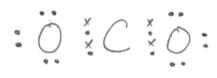
Remember to check the number of non-bonding electrons on each atom, as well as the electrons in the bonds.

(c) The composition of a sample of natural gas and the melting temperatures of four of its components are shown below.

	Percentage	Melting temperature / K			
Methane	95.2	91.1 113.7			
2-methylpropane	0.8				
Butane	0.9	134.7			
Other hydrocarbons	2.4				
Carbon dioxide	0.7	216.5			

(i) Draw a dot and cross diagram for carbon dioxide.

(2)



Results Plus Examiner Comments This mistake occurred quite often. Oxygen was shown with eight electrons in its outer shell before it formed the bonds with carbon.



The atomic number tells you the total number of electrons in the atom, not the number in the outer shell.

Question 24 (c) (ii)-(iii)

Part (i) of this question was specifically about the different strengths of London forces. This did not stop candidates writing about polar bonds, dipole-dipole interactions, the strength of double bonds and the presence of lone pairs. Many stated that oxygen has more electrons than hydrogen, without making it clear that two oxygens within CO_2 contribute more electrons to the molecule than four hydrogen atoms in methane do. Some candidates calculated the number of electrons in each molecule, but these attempts were often incorrect as they were based on mass numbers instead of atomic numbers.

When contrasting butane and methylpropane, some said that they had the same number of electrons. This is true, but does not answer the question about why their melting temperatures differ. Many wrote about molecules of butane being able to pack more closely, but they did not always associate this with the shape of the molecules or their surface area.

(ii) The London forces between molecules of carbon dioxide are stronger than the London forces between molecules of methane. Suggest a reason for this. (1)carbon disside than more = 58 (iii) Use your knowledge of intermolecular forces to suggest why butane has a higher melting temperature than 2-methylpropane. (2)corres moleuler porces of a thertin **Examiner Comments** This lost the first mark because the number of electrons given was incorrect, but it scored both marks in (ii). **Examiner Tip**

When you calculate the number of electrons in a molecule, you need to use the atomic numbers. Methane has 10 electrons and carbon dioxide has 22.

(ii) The London forces between molecules of carbon dioxide are stronger than the London forces between molecules of methane. Suggest a reason for this. (1) There are more electrons in carbon dioxide than in methane. C-C-C-C-C-C-C (iii) Use your knowledge of intermolecular forces to suggest why butane has a higher melting temperature than 2-methylpropane. (2)Straight chain molecules have high melting temperatures than branched molecules because branched molecules do not fit together as well as straight chain molecules so less London forces are present. **Examiner Comments**

This is an example of an answer which gained all three marks.

Question 24 (d) (i)

Examiners were surprised by the number of wrong answers given for the flame test on magnesium carbonate. Many candidates said that a bright white light would be seen, the observation made when burning the element. "White" was not allowed as the answer here; it had to be clear that magnesium carbonate would not colour a flame. The result for calcium carbonate was better known.

(d) When carbon dioxide dissolves, it may eventually form minerals such as magnesium carbonate and calcium carbonate.
(i) State the results of flame tests carried out on these two minerals. (2)
Magnesium carbonate white
Calcium carbonate BBcu -red
Results in the second mark only.
 (d) When carbon dioxide dissolves, it may eventually form minerals such as magnesium carbonate and calcium carbonate. こううてんしょう (i) State the results of flame tests carried out on these two minerals.
(2)
Magnesium carbonate bright white light flame Calcium carbonate brick ved plame
Calcium carbonate brick wed flame
Results Less Examiner Comments This also scored the second mark only. A range of colours was
allowed for the calcium flame, but red had to be included.

Question 24 (d) (ii)

Many answers to this question suggested a lack of practical experience or knowledge. Suggestions included soaking iron wool in calcium carbonate, heating carbonate solutions in a water bath, measuring the melting points of the carbonates, measuring enthalpy changes and many more unworkable or chemically impossible ideas. Few answers were succinct. Diagrams could be included as an aid to describing the method, but fortunately for the candidates the quality of the diagrams was not marked, and very few diagrams actually enhanced the candidates written responses.

The first mark was for suggesting a suitable method. This could involve collecting the gas evolved, passing the gas into limewater, or setting up an experiment in which change of mass could be measured. However different methods were often combined making the suggestions unworkable, for example measuring the volume of gas after passing it through lime water. Many candidates seemed to think that the first appearance of bubbles when the carbonates were heated indicated thermal decomposition, and had forgotten that expansion of the air in the tube would cause this.

The second mark had to state the measurement which would be made. "Measure the reaction" or even "measure the rate at which the gas is produced" did not score a mark. Measuring the temperature at which the carbonate decomposed was not a practical proposition, especially when no way of detecting the decomposition was suggested.

The most common suggestion for making the comparison fair was to take equal masses of the carbonate, but equal numbers of moles should have been used. Many tried to describe the need for a constant strength of Bunsen flame, but did not score by saying just "use the same bunsen" or "use the same heat". Many candidates wrongly thought that repetition of the experiments produced a fair test.

*(ii) Magnesium carbonate and calcium carbonate both undergo thermal decomposition, but they have different stability to heat. The difference in stability to heat can be compared in an experiment. Suggest how this experiment could be carried out. You should indicate how to detect when the thermal decomposition occurs the measurement you would make to compare the stability to heat how to make the comparison fair. You may include a diagram if you wish but it is not essential. (4) MgO()+ CO2(g) decomposition occurs, CO2 gos is produced by the gas syringe.

The measurement would be the fime taken for a co decided volume of CO, gas to produced. - The quicker it happens, the less thermal stability. Lepeat experiment three times withe the the same the and some quantity of carbonate each heat time. Ensure the gas syringe is completely empty ofter experiment. each **Examiner Comments** This answer scored the first two marks, but the description of how to make the comparison fair was not good enough. **Examiner Tip** When you repeat an experiment it makes your results more reliable, but it does not make the tests you are doing fair comparisons. and Heat in boiling tubes both Magnesum Carbonate + Calcium corponete. Time how long it takes for either to show pust signs of venetion (decomposition) any publies parminez -+ turning into liquid etc. To note it pair, we the same bursen purer, look per similar signs of reactions in both hubes. **Examiner Comments** Answers like this did not score. The method for detecting decomposition implies that the solid carbonates will start to bubble. The use of the same Bunsen burner is not good enough for a mark for a fair test.

Question 24 (d) (iii)

Many candidates gained a mark by knowing that calcium carbonate was more stable to heat than magnesium carbonate. After this, marks were often lost because it was not clear that the differences were due to the calcium and magnesium ions. Many answers said that magnesium atoms were smaller than calcium atoms, or just that "magnesium is smaller than calcium" or even "magnesium carbonate is smaller than calcium carbonate". Some of the answers that correctly compared ionic radii and polarising power concluded that magnesium carbonate was more stable because the ions were closer to each other. Some answers were in very general terms, saying that the ionic radius increases and polarising power of a cation decreases going down a Group in the Periodic Table. These ideas had to be clearly applied to the two carbonates in the question to earn full marks.

Very few candidates gave an answer in terms of the energy released on forming the oxide.

*(iii) State and explain which of the two carbonates is more stable to heat. (3) Magnesium any bon afe is the weaker shan and imm curbonate because it is much smaller and so has a high chape density and so weaken the bond and less heat is needed to becak the bond in in majensium curbonat



This candidate is probably thinking along the right lines. There is no clear statement that magnesium carbonate is less stable, though this is implied by saying that less heat is needed to break the bonding so the first mark was allowed.

There is no reference to ions so the second and third marks were not given. "Weakening the bonding" could refer to the bonds in the carbonate ion or to the ionic bonds between the magnesium and the carbonate.

This candidate may have had the right ideas, but they were not expressed clearly enough to get the marks.

Cardum	carbonate	is more	Stable	H	heating.	This	is be	cause	Galaro
calcium	ione, Ca	2+ have	a la	iger	MAN	radius	and	thus	a
smaller	charge	density.	This	BAAK	e make	s Ca ²¹	ions	leu	able t
polarise	the and	distort	tre	elicti	on cia	uld cli	oud o	t the	carbo
anions.	More here	t energy	is M	equire	d to	decomp	085	alciu	im cail



Paper Summary

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

- Reports by examiners almost always advise candidates to read the question carefully, and this one is no exception. Marks are regularly lost because candidates do not actually answer the question which is asked. One example of this was in 24(d)(ii) where candidates discussed the theory without suggesting how to carry out the experiment.
- Candidates often lose marks because their answers are not expressed clearly, and more practice in giving explanations is needed. Answers including the right technical terms cannot be rewarded unless their meaning is clear
- Calculations should also be set out clearly. Candidates should be encouraged to explain what they are doing in numerical work. Many scripts just gave sets of numbers with no explanation in the titration calculation. Use of standard form should also be encouraged, as errors arise in transferring numbers with many zeros at the start, such as 0.000605 in the titration.

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