



Examiners' Report January 2012

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. There was no evidence that candidates were short of time. With a mean score of 14.2/20, Section A was clearly accessible to almost all candidates. In Section B and C candidates generally set out their work clearly and made sensible use of the space provided, although, in some cases, there was wasteful repetition both of the statements in the question and of the candidate's responses. Candidates showed a good understanding of the core concepts underlying chemical calculations and there was an improvement in the use of significant figures. While many candidates used scientific vocabulary with skill and accuracy, there remain a significant number whose use of basic chemical terms appeared to lack an appreciation of their precise meaning; for example terms such as atom, ion and molecule can be taken by some candidates as interchangeable. The quality of the diagrams in 22(f)(ii) was typically rather poor. Candidates were most comfortable when dealing with questions that mainly required repetition of learned material and questions requiring them to demonstrate their understanding, notably the organic mechanism question, proved much more challenging. The practical dimension of this paper still seemed to surprise many candidates.

Question 18 (a)

There were many excellent answers to this question demonstrating a clear understanding of the nature of the intermolecular forces involved and the factors which affect their magnitude. Problems arose where candidates failed to distinguish clearly between intramolecular bonds and intermolecular forces and some clearly believed that boiling molecular substances involves the breaking of covalent bonds. Another area of difficulty was in identifying the relevant particles with many references to bromide and iodide ions and to the electronegativity of ions and of molecules.

Answer A	LL the questions. W	Vrite your answers in the spa	ces provided.
8 The boiling temperat	ures of some hydrides	s are given below.	
	Compound	Boiling temperature / K	
	HF	293	
	HC1	188	
	HBr	206	
	HI	238	
	ЦО	200	
*(a) Explain, by comm	H_2O baring the forces invol	373 ved, why HI has a higher boili	ng temperature
than HBr. • HI hs	paring the forces invol	ved, why HI has a higher boili w number of electr	(3) ons because
than HBr. • HI he Iwine has	paring the forces invol us a great o a greater	ved, why HI has a higher boili w number of electr Ar than . Browine	(3) uns because
than HBr. • HI he Iwine has • & Gream	paring the forces invol us a great o a greater Van der wo	ved, why HI has a higher boili w number of dectr Ar than Bronnine aals forces betwee	(3) uns because
than HBr. • HI he Iodine has • & Gream	paring the forces invol us a great o a greater Van der wo	ved, why HI has a higher boili w number of electr Ar than . Browine	(3) uns because

ResultsPlus

< Examiner Comments

This candidate correctly identifies the main intermolecular force involved and the factor which determines the magnitude of the force in these examples. Strictly speaking, van der Waals forces is a general term that encompasses London forces *and* permanent dipole interactions but its use was allowed for London forces in the mark scheme. Note that the candidate only makes two points so scores two marks.



Make sure that the number of points in your answer corresponds to the total marks available.

SECTION B

Answer ALL the questions. Write your answers in the spaces provided.

18 The boiling temperatures of some hydrides are given below.

Compound	Boiling temperature / K
HF	293
HCl	188
HBr	206
HI	238
H ₂ O	373

*(a) Explain, by comparing the forces involved, why HI has a higher boiling temperature than HBr.

(3)HI has a higher briling Harrow Mada temperature than HBr because it has more electrons than Bronine and Meregone has a greater number of dispersion Forces Strengthening the H-I band. - The more electrons it has, the stronger the dispersion forces and therefore



This candidate gets off to a good start with the first three lines of the answer but then spoils the answer by confusing the intermolecular forces and the covalent bond.



Great care is needed with this type of question where incorrect statements can negate correct answers.

SECTION B	
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Answer ALL the questions. Write your answers in the spaces provided.

18 The boiling temperatures of some hydrides are given below.

Compound	Boiling temperature / K
HF	293
HC1	188
HBr	206
HI	238
H ₂ O	373

*(a) Explain, by comparing the forces involved, why HI has a higher boiling temperature than HBr.

The vander waal's and lor London forces present	t in the HI
are much tigher than those in HBr." nodeallar the a the increase in the atomic size of I	so, with an
increasing size, it becomes earlier to distort	
this, therefore results in a stronger V	

(3)



Examiner Comments

This answer also starts well but the reference to the 'molecular size of the anion' reveals serious confusion about the nature of the particles present in these molecules.



Understanding the distinction between the basic particles that make up chemical compounds (atoms, ions and molecules) is fundamental to answering this type of question.

Question 18 (b)

Generally this question was more accessible to candidates than 18(a) as most were aware that the main intermolecular force in HF is hydrogen bonding. It was generally appreciated that hydrogen bonding is a particularly strong intermolecular force but few candidates showed any awareness that in HCl the London forces are much larger than the permanent dipole-dipole interactions. As in 18(a) difficulties arose due to a lack of clarity about particles and where candidates failed to distinguish bonds and intermolecular forces. This led to discussions about the electronegativity of molecules and ions. There were many answers that included detailed explanations about the formation of hydrogen bonds which were not required by the question.

*(b) Explain, by comparing the types of forces involved, why HF has a higher boiling because temperature than HCl. hydrogen bonding due to nely gative 1 bondo dipole There rds ba



This candidate makes four good points but note the incorrect statement that HCl has *only* dipole-dipole bonds.



The use of the phrase 'dipole-dipole bonds' is technically correct but it is generally a good idea to refer to intermolecular interactions as *forces* to make clear the distinction between these and ionic or covalent bonds. The exception to this is, of course, the hydrogen *bond*.

*(b) Explain, by comparing the types of forces involved, why HF has a higher boiling temperature than HCl. (3) to ion is the most Flouride electronegative element. electronegativity enables it to bond tightly to the enable HF polar bonds bonds Formpol ion. bonds which hydregen. en hance Strano Corm Examiner Comments There are some typical confusions in this response, notably the reference to the fluoride *ion* being the most electronegative element and the suggestion that HF is ionically bonded. **Examiner Tip** Only atoms have electronegativity. *(b) Explain, by comparing the types of forces involved, why HF has a higher boiling temperature than HCl. (3)As HF has hydrogen bonding which is much stronger then then the London forces in H(1, hence HF has higher Boiling tenpreture **Examiner Comments** There are only two statements in this answer that can gain credit, the final phrase being just a re-statement of part of the question. Thus, although both are fully correct, the maximum score is 2. Result **Examiner Tip** Do match your answer to the marks available.

*(b) Explain, by comparing the types of forces involved, why HF has a higher boiling temperature than HCl. (3)are hydrogen This is because there bonding present in the HE molecule. to i has a Whereas the HCL molecule weak has wres (weak intermolecular forces) present in its. The london in HF makes it the HF molecule have very strong bonds bonding these are very hard to break. This is why HF has a and weak HCL bonds are easier to break which builing boiling point compared to HF. It has a



The use of the word 'bonds' in the latter half of the question introduces an ambiguity about the forces under discussion that weakens the answer.



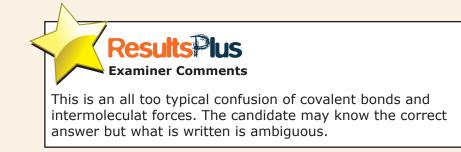
Do read through your answers to check that the meaning of what you have written is clear and is what you want to convey.

Question 18 (c)

There were many good answers to this question although too many candidates suggested that the individual hydrogen bond in water was stronger than that in hydrogen fluoride. Other candidates attempted to hedge their bets by referring to the stronger London forces in water. A surprising number of candidates explained that the different numbers of hydrogen bonds in the two molecules was due to the fluorine atom having fewer lone pairs than the oxygen atom. As elsewhere in question 18 there was confusion about the particles present.

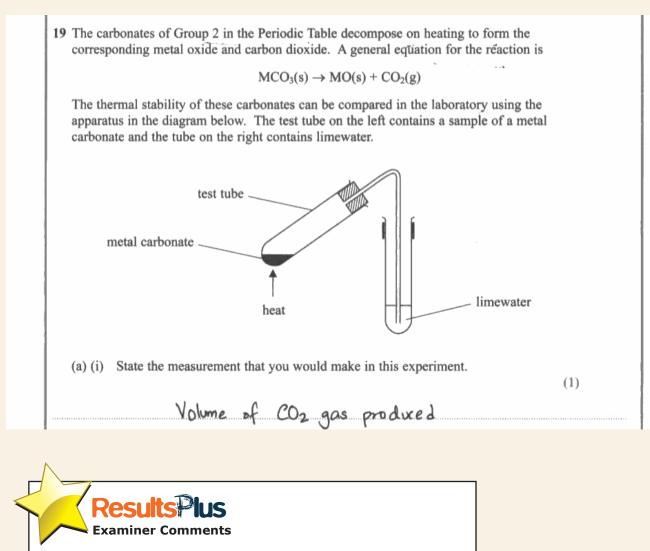
(c) Suggest why H₂O has a higher boiling temperature than HF. (1)H2O can form two hydrogen bonds within itself due to the presence of electronegative oxygen atoms, but HF can form . so mor hydrogen bond is required to break the greater extent of hydrogen bonding in 160 that that of HF. (Total for Question 18 = 7 marks) Examiner Comments ResultsPlus It is hard for an examiner to consider **Examiner Tip** the reference to *two* oxygen atoms in water as just a slip. Do read through your answers! (c) Suggest why H₂O has a higher boiling temperature than HF. (1)H20 has two bonds of hydrogen with ar highly electronegative element (oxygen) whereas HF only has one Therefore more energy is needed to break the bonds in the hence forming a higher

bailing temperature need.



Question 19 (a) (i)

The first half of question 19 was intended to test the practical understanding of candidates by considering an experiment that could easily be carried out in a school laboratory. However, many candidates envisaged experimental methods that were far too complex, such as monitoring the mass of the carbonate or the degree of cloudiness of the lime water.

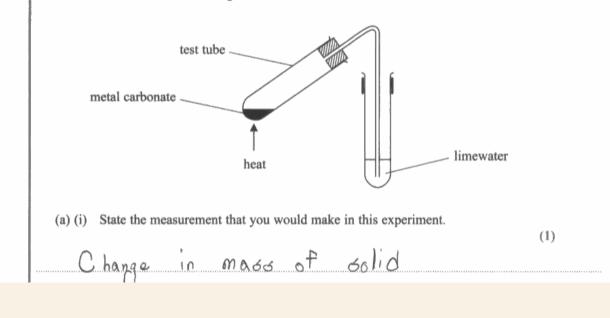


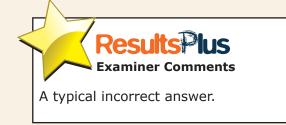
The answer bears little relation to the apparatus in the diagram.

19 The carbonates of Group 2 in the Periodic Table decompose on heating to form the corresponding metal oxide and carbon dioxide. A general equation for the reaction is

 $MCO_3(s) \rightarrow MO(s) + CO_2(g)$

The thermal stability of these carbonates can be compared in the laboratory using the apparatus in the diagram below. The test tube on the left contains a sample of a metal carbonate and the tube on the right contains limewater.





Question 19 (a) (ii)

The idea of controlling key variables to ensure a 'fair test' should be well established with candidates but many slipped into reliance on generalities such as repeating experiments and using the same apparatus. It was of some concern that quite a few candidates envisaged measuring out the solid carbonate by volume or even referred to constant *concentration* of the solid. The best answers were clearly based on careful consideration of the experiment described in the question.

(ii) Suggest three ways to make sure that, when thermal stabilities of the different carbonates	
1 Amount of metal carbonate	
2 Buse some temperature for each	experiment.
3 Use identical test type for each	metal corbonate
Results Line Examiner Comments The first point is careless; the candidate probably intends to say 'fixed amount' but has not done so. The second point shows some understanding of what is required but ignores the practical dimension of this question; the candidate needs to consider how temperature would be kept constant.	Results Pus Examiner Tip Bear in mind that your understanding of practical chemistry will be tested on the theory papers. Make sure that your meaning is clear in your answers.
 (ii) Suggest three ways to make sure that, when thermal stabilities of the different carbonate 1 Repeat the experiment 	
2 Clse some amount of experiment 3 Clse the same he	
Results Plus Examiner Comments Once again a candidate shows an appreciation of heat the carbonates in a consistent way but fails how this will be achieved.	the need to

Question 19 (b) (ii)

Most candidates appreciated that this was a question about polarising power and polarisation with many excellent answers. The mark most likely to be lost was that for identifying the species (or bond) being polarised; sometimes this was omitted although a number of candidates thought the cation was polarised. As elsewhere in this paper candidates lost marks through slipshod use of terminology, for example referring to ions or even atoms rather than cations. Weaker candidates gave often extensive answers discussing shielding and bond strength without managing to touch on relevant or correct chemistry.

(b) (i) State the trend in the thermal stability of the metal carbonates as the group is descended.
(1)
Increases
*(ii) Explain this trend in stability.
(3)
as the Cation increases in size there
is less distortion on the anion
as there is more shielding. The loss
distortion there is on the anion
the greater the sta more stable it
15 So as you go clown the group
the cotton here is less dissortion so it gets mill
(Total for Question 19 = 8 marks)
ResultsPlus
Examiner Comments

This answer covers the essential points although it does not make it absolutely clear that the cation is responsible for polarising the anion.

(b) (i) State the trend in the thermal stability of the metal carbonates as the group is descended. (1)The thormal stability decreases *(ii) Explain this trend in stability. (3)· Thermal stability of a metal depends on its polarising Power As you go down the group the ionic radius increases and so the Grons of group two become less electron opatine and so are less polar. • The lations ability to polacise the electron cloub of the Carbonate Ion decreases down the group and and hence thermal Stability noreases down the group. (Total for Question 19 = 8 marks)

Results Plus Examiner Comments

There are the elements of a good answer here, particularly the final bullet point, but note the errors and imprecision that mar the first two points. In the first point the candidate refers to the thermal stability of the metal and 'its' polarising power. In the second point the candidate writes 'ion' rather than 'cation' and then refers to the electronegativity of the cations.

(b) (i) State the trend in the thermal stability of the metal carbonates as the group is descended.
(1)
Ans. bi) As the group is descended, thermal stability of the metal catornated metalled decreates. *(ii) Explain this trend in stability.
Aus. ii). Berylium carbonate decomposes much more filletter then collecter
magnesium or calcuir cabonate do. As we have hept the temperature a
constant in this experiment, we measure the time taken for nuture each carbonate to decompose. Thurlor, as the group is descended, decreated the time taken intervent, which means that the thermal stability also decreated. This is because the fire of the ionic radii increases down
the group, making the metals mare reactive and so they easily bond
with the carbonetic ion. This means when heat is provided to this mollade,
with the calconete ion. This means when heat it provided to this molecule, because of the weak bonds, they decompose much more quickly. (Total for Question 19 = 8 marks)
Therefore berylium and magnelium carbonete are more stable than Barium carbonate.

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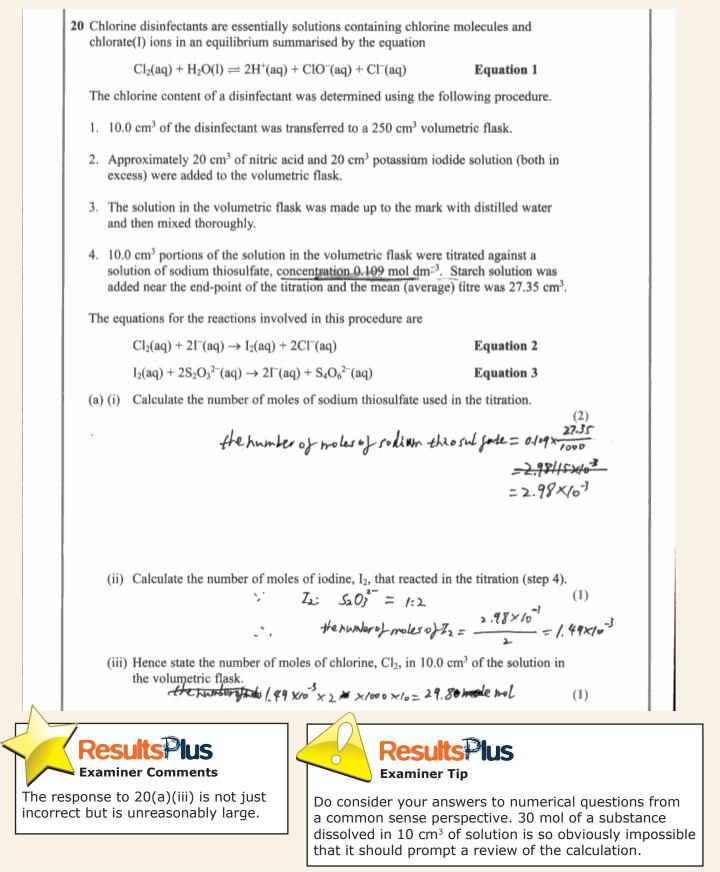
This is a weak answer but it is worth noting the inappropriate use of specific group II elements in a question that clearly refers to general trends and the extensive references to the experimental results which gain no credit.

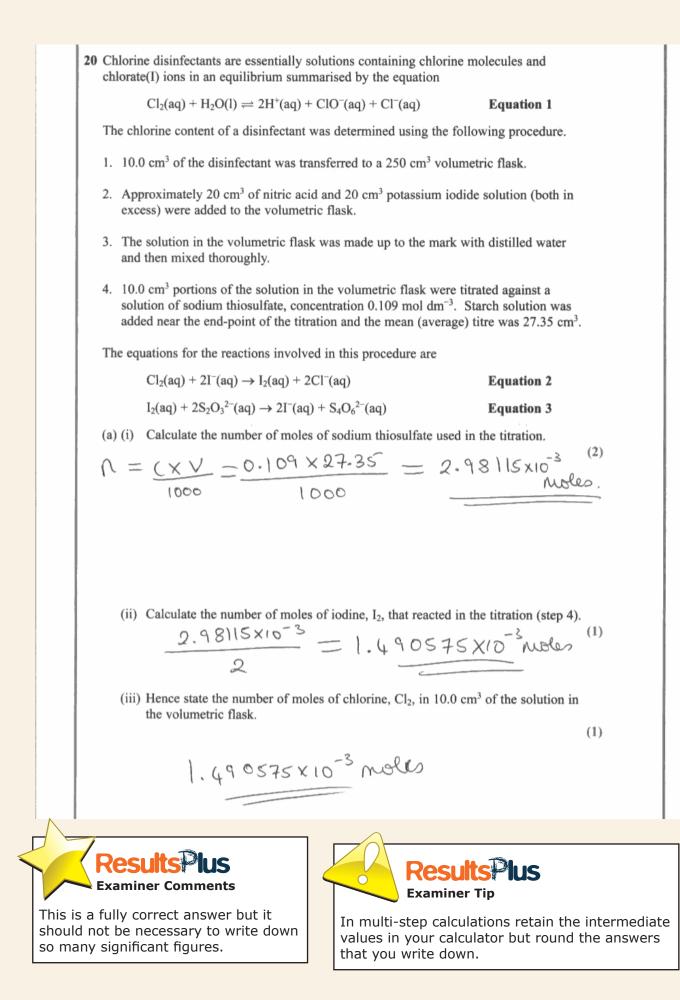
(b) (i) State the trend in the thermal stability of the metal carbonates as the group is descended. (1)In Creases *(ii) Explain this trend in stability. (3)down the group cation size increase the charge is same which is +2, so asing the the charge density decrease the polarising pou de cheasing distor tiph Cloud fort ess electron



Question 20 (a) (i-ii)

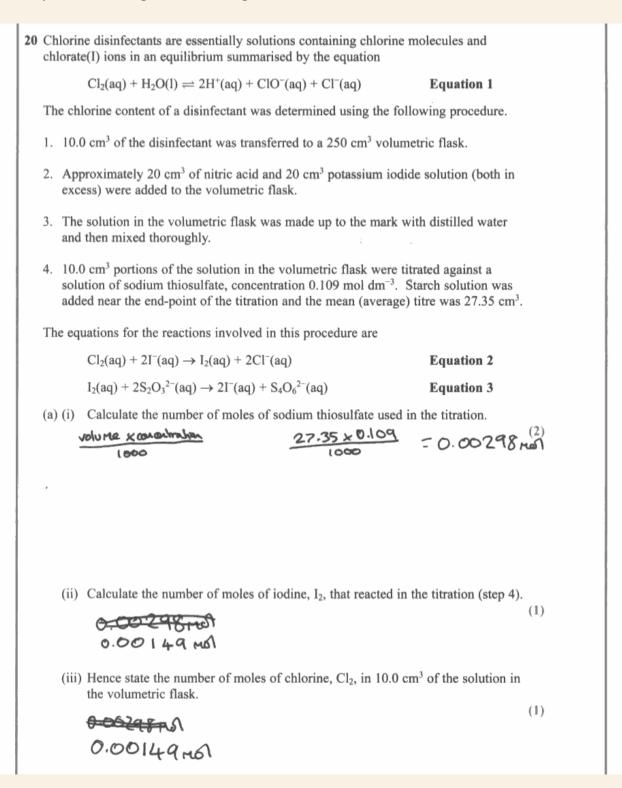
Most candidates were able to complete this calculation correctly although some failed to use the correct volume in 20(a)(i) and others did not use the appropriate stoichiometric ratio in 20(a)(i).





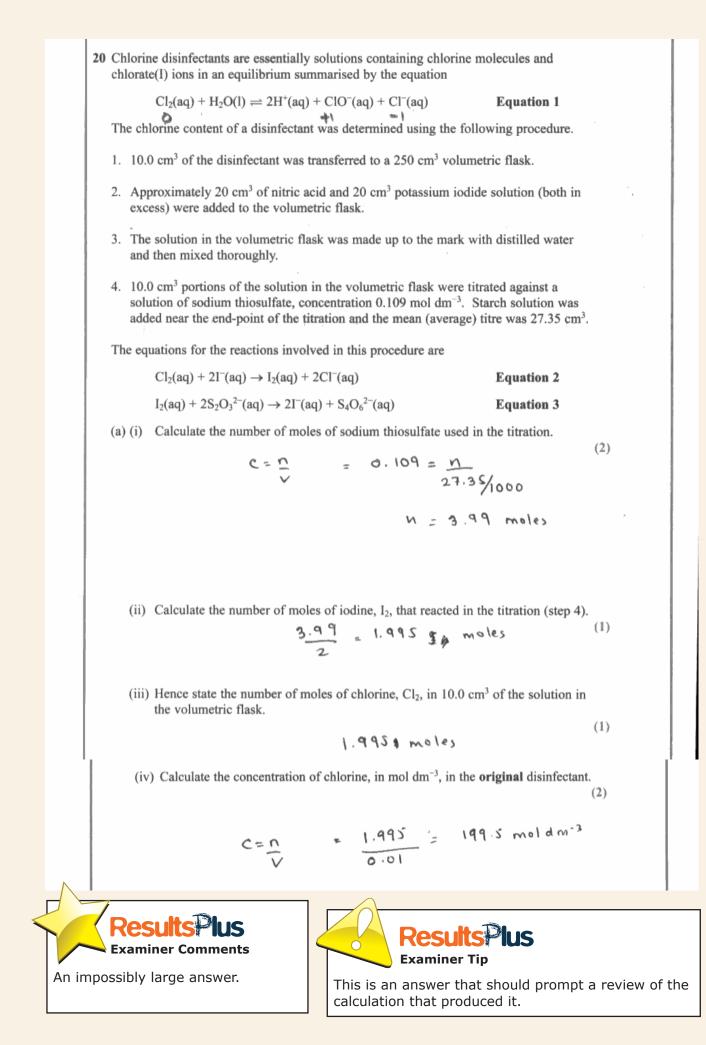
Question 20 (a) (iv)

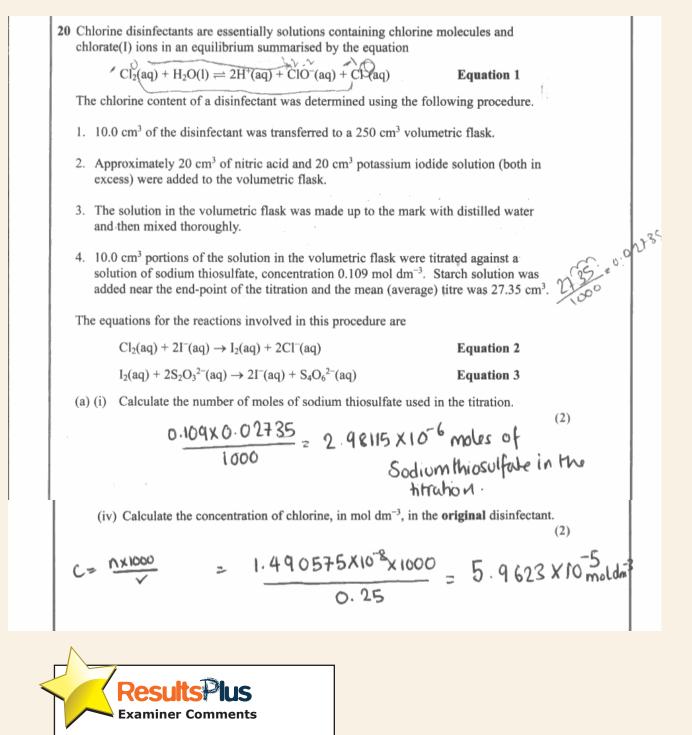
This calculation caused considerable difficulties. Most candidates realised that the answer to 20(a)(iii) needed to be multiplied by 100 but few appreciated that a further factor of 25 was needed to produce the final answer. Some candidates gave, without comment, answers that were clearly of the wrong order of magnitude.



(iv) Calculate the concentration of chlorine, in mol dm^{-3} , in the **original** disinfectant.

(2) 0190/2 RNOO DOLLANDIDM-3 valone accelertration = moles molerx 1000 1000 $\frac{0.00149 \times 1000}{L0} = 0.149 \, \text{moldm}^{-3}$ Results US **Examiner Comments** This is a typical answer gaining one mark out of two.





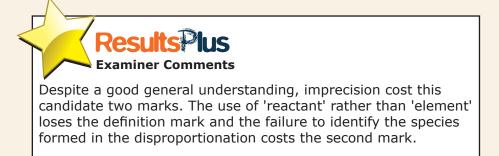
This answer is impossibly small.

Question 20 (b)

Most candidates showed a good general understanding of the term disproportionation but a lack of precision in the answers meant that relatively few were able to score full marks. In the general definition candidates frequently failed to appreciate that disproportionation involves changes in the oxidation state of a single *element* while, in the explanation of the specific example, all too frequently neither the oxidation and reduction processes nor the species involved were clearly identified.

(b) Equation 1 is an example of a disproportionation reaction. Define the term 'disproportionation' and explain, by considering the relevant oxidation numbers, why this reaction is a disproportionation. (3)Disproportionation reaction is when a species is both reduced and oxidized in the same reaction. and its reduced to - I in and also oxidused to +1 in CLOT. Examiner Comm The word species is not sufficient to gain the first mark but the subsequent explanation, despite its brevity, scores the two explanation marks. (b) Equation 1 is an example of a disproportionation reaction. Define the term 'disproportionation' and explain, by considering the relevant oxidation numbers, why this reaction is a disproportionation. (3)In a same the deaction taking place in a chemically closed system a simultanious occuring of both oxidation and reduction of the same species is known as "disproportionation". Cl. reduces to give cl. and cl oxidize to give clt **Examiner Comments** The use of Cl⁺, either as a shorthand for 'chlorine in oxidation state +1' or in place of ClO⁻, is very dubious. ResultsPlus Examiner Tip It is best to use the full formula of the species and to state the oxidation number, remembering to include the sign.

'dispropo		xplain, by c	portionation reaction reaction of the rele			(3)
Di	spropor	tiona-	tion rec	ection	occur	s
when	the sa	me n	eactant	gets	axeid is	ed
			neduced	0		
	chlopine	gets	oxidised	from	0 +0	- 그 .



(b) Equation 1 is an example of a disproportionation reaction. Define the term 'disproportionation' and explain, by considering the relevant oxidation numbers, why this reaction is a disproportionation.
(3)
(3)
Disproportionation means one element is both oxidised and reduced. Oxidation number of chlorrine in Cl_is O. To Cl_ oxidation number is -1. So chlorrine is reduced. To Clo_, oxidation number of chlorrine is +1. So a berre chlorrine is oxidised.

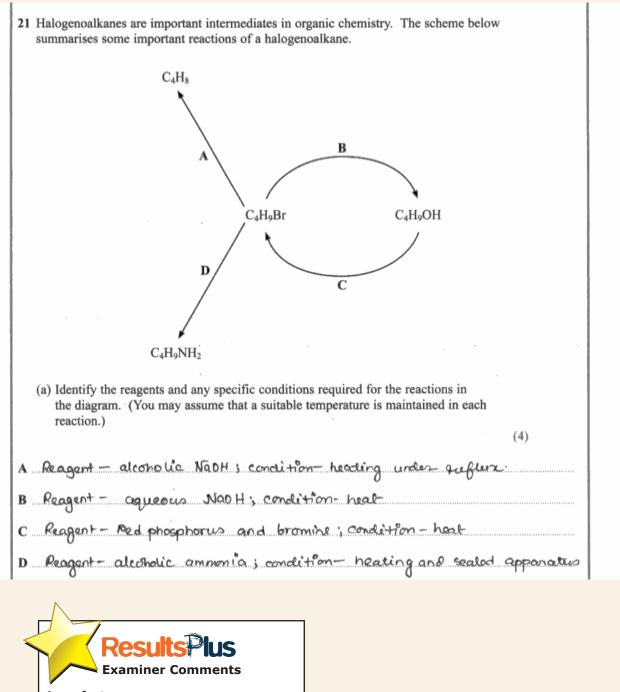


Question 20 (c)

It is important in an iodine-thiosulfate titration to add the starch solution very close to the end-point, so the *pale* in pale yellow is important.

Question 21 (a)

This question tested knowledge of the reagents and essential conditions for some basic organic reactions; these were generally well-known but a lack of precision often resulted in the loss of marks.



Question 21 (b) (ii)

This question proved very challenging. While most candidates were able to draw the tertiary carbocation, many assumed that the formation of the primary alcohol proceeded via a primary carbocation and those who were aware of the the $S_N 1$ mechanism were often unable to draw the transition state correctly. Some candidates simply described the $S_N 1$ and $S_N 2$ mechanisms without attempting to answer the actual question.

*(ii) Reaction B can proceed via two possible reaction mechanisms, depending on the structure of the original compound. For each of the two isomers of C₄H₉Br shown below, draw the structure of the intermediate or transition state which is formed during the reaction. Explain in each case why the specified structure is more favourable. (4)НННН C3H7-C5 H-C-C-C-Br НННН 5.2 Intermediate or transition state Explanation because here isn't enough alkyl groups because or it being a primary haloalkanp So it isn't at strong enough to make a Stuble Carbo Cation CH_3 CH3-C-Br CH₃ Intermediate or transition state Explanation because 15 a terhing halo alkane here is 3 alkyl graves attached to me carbon meaning it is strong enough to form a stuble corbocation **Examiner Comments** The $S_{\mathbb{N}}1$ transition state is shown with five bonds to the central carbon rather than with the attacking OH⁻ and the leaving group Br⁻ both partially bonded to the central carbon atom. The candidate appreciates that the tertiary carbocation is more stable but the explanation of this is too confused to gain a further mark.

*(ii) Reaction B can proceed via two possible reaction mechanisms, depending on the structure of the original compound. For each of the two isomers of C₄H₀Br shown below, draw the structure of the intermediate or transition state which is formed during the reaction. Explain in each case why the specified structure is more favourable. (4)0H---C-H НННН $\begin{array}{c|c} | & | & | & | \\ \hline \\ -C - C - C - C - C - Br \\ | & | & | \\ H & H & H \end{array}$ SN2 (2 species at beginning) Intermediate di lansing Explanation due 15 being a primary halogenoalleane no shielding in reaction can provi in 1 step is a transition $CH_{3} - CH_{3} - CH_{3}$ Br-CHA Intermediate or transition state Explanation SN, (1 species at beginning) due to being a tertiary halogencialleane the methyl groups create a lot op shielding in 2 steps required where a carbo cation is **J**US **Examiner Comments Examiner Tip** The transition state here has two errors: the hydrogen in the OH group is shown When writing equations involving ions, bonded to the central carbon and the keep a check on the overall charge on the transition state has no charge, the latter system, which must be constant. error was not penalised. The candidate has the idea that the methyl groups stabilise the tertiary carbocation but no real idea

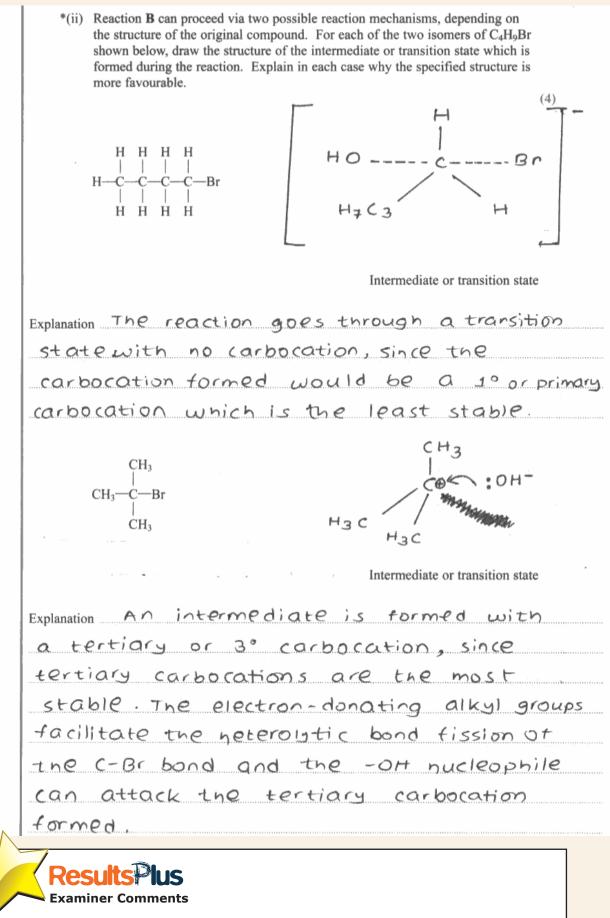
how this is effected.

*(ii) Reaction **B** can proceed via two possible reaction mechanisms, depending on the structure of the original compound. For each of the two isomers of C₄H₉Br shown below, draw the structure of the intermediate or transition state which is formed during the reaction. Explain in each case why the specified structure is more favourable. (4) $\begin{array}{c} H & H & H & H \\ H & -C & -C & -C & -C & -Br \\ H & H & H & H \end{array}$

Intermediate or transition state Explanation there there is prenty of space for the OH to attack and it does so because the carbon is not surrounded by methyl groups. This process can produce 1 bromo butome, 2 bromo butome. CH2 CH3 - est Brs OH-CH3 Intermediate or transition state Explanation Here the carbon is surrounded by 3 methy! groups so there is no space to attack the carbon so the Br has to go first and then the OHT can attack the carbon. can attack the carbon. This process can only produce one type of promo butane.

Results Plus Examiner Comments

This candidate shows a reasonable understanding of the steric factors that influence the mechanism, the transition state and intermediate have not been drawn. This type of response showing the electron movements leading to the transition state and intermediate but not the actual species was quite common.



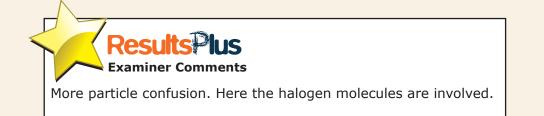
The intermediates shown here are particularly well drawn and the candidate gains a further mark for noting the (greater) stability of the tertiary carbocation. However, while the candidate is aware that the methyl groups are important in determining the mechanism, their role in stabilising the carbocation is not made clear.

Question 21 (b) (iii)

This was a straightforward question with many correct answers. Although the question was worth just one mark, many candidates gave elaborate discussions involving electronegativity differences often without touching on the key point. There were a surprising number of responses which referred to the H—X rather than the C—X bond.

(iii) If C4H9I is used instead of C4H9Br in reaction D, the rate of formation of C4H9NH2 increases. Explain why the rate of reaction increases. (1) The C-I bond is weaker than the C-Br bond, therefore will break easier, so that the reaction takes place quicker.

(iii) If C₄H₉I is used instead of C₄H₉Br in reaction **D**, the rate of formation of C₄H₉NH₂ increases. Explain why the rate of reaction increases. (1) <u>Because Brz is more reactive flam Iz</u> Is substituted to meedice



Examiner Comments

This answer is very much to the point.

(iii) If C4H9I is used instead of C4H9Br in reaction D, the rate of formation of C4H9NH2 increases. Explain why the rate of reaction increases.
(1)
8 Sodine is a larger molecule, with large bord length and hence weaker than the broomcal kane, so it reacts paoily. Thus reafe of reaction increases.

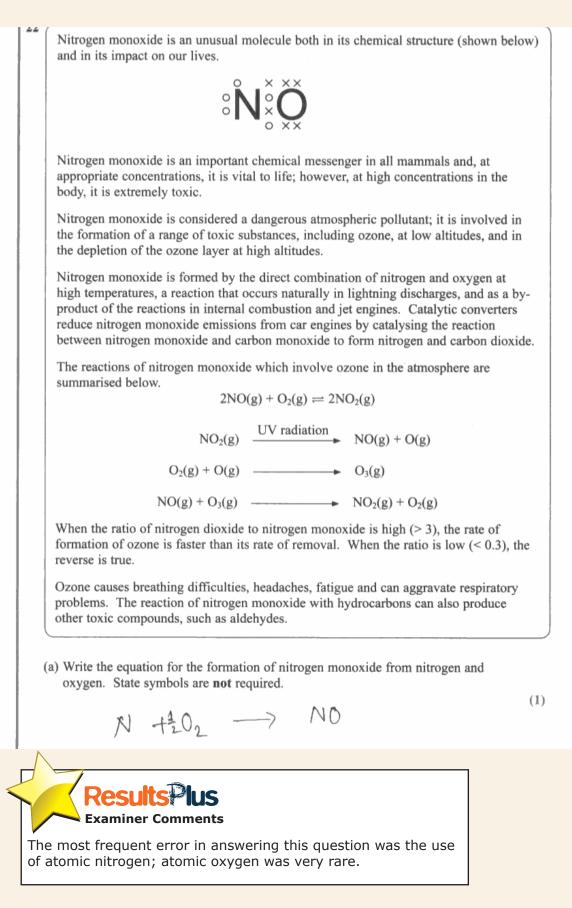


Question 21 (c)

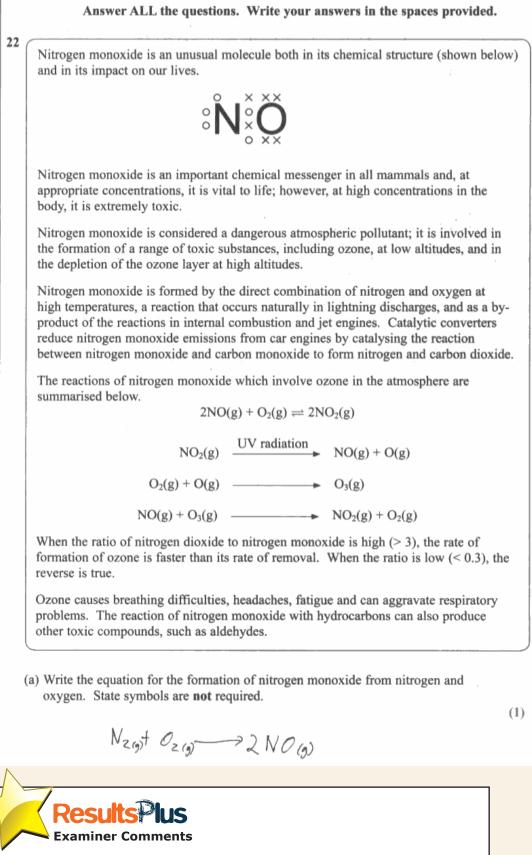
While there were many excellent answers to this question, there was some confusion about how a refrigerant cools and a failure of common sense in choosing the properties that might be desirable in a refrigerant.

Question 22 (a)

While most candidates were able to write the correct balanced equation for this reaction, a significant minority could not.



SECTION C



Despite the instruction in the question. the candidate includes the state symbols. This does not affect the mark but is unnecessary.

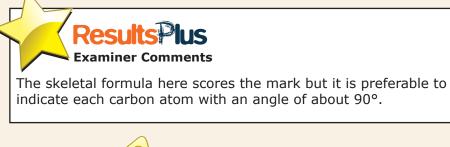
Question 22 (b) (i)

Most candidates were familiar with the terms 'free radical' and homolytic fission.

Question 22 (c) (i-iii)

Fully correct answers to this sequence of questions were quite rare. Few candidates appreciated that unburnt fuel in vehicle exhaust emissions was the most likely source of hydrocarbons in the atmosphere and difficulties were exacerbated by poor use of language. Most candidates realised that the conversion of a hydrocarbon to an aldehyde was an oxidation reaction but there were also references to the (partial) oxidation of alcohols indicating that the question had not been fully understood. The skeletal formula question produced a range of answers. Common errors were the skeletal formula of ethanal and full structural formula of propanal.

(c) (i)	Suggest the most likely source of the hydrocarbons that react with nitrogen monoxide to form toxic compounds.	(1)
(ii)	Suggest the type of reaction that is involved when a hydrocarbon is converted into an aldehyde.	·····
	nio un unonguo.	(1)
	Oxidation	
(iii) Draw the skeletal formula of the aldehyde with three carbon atoms.	(1)
	*	
	i i i	





Many candidates find skeletal formulae difficult so it is well worth the time ensuring that you understand the principles and practising drawing examples.

(c) (i) Suggest the most likely source of the hydrocarbons that react with nitrogen monoxide to form toxic compounds. (1) (ii) Suggest the type of reaction that is involved when a hydrocarbon is converted into an aldehyde. (1)axidation (iii) Draw the skeletal formula of the aldehyde with three carbon atoms. (1)H - C - C - C - H**Results**Plus **Examiner Comments** Skeletal is bold in the question but this is a full structural formula. (c) (i) Suggest the most likely source of the hydrocarbons that react with nitrogen monoxide to form toxic compounds. (1)alcohols. (ii) Suggest the type of reaction that is involved when a hydrocarbon is converted into an aldehyde. (1)ovidation substitution. (iii) Draw the skeletal formula of the aldehyde with three carbon atoms. (1)**Examiner Comments** The inclusion of the hydrogen in the skeletal formula means the mark cannot be awarded.

Question 22 (c) (iv)

There were many excellent answers to this question but some candidates failed to take account of the mark allocation and gave lengthy and elaborate responses while others ignored the word 'explain'.

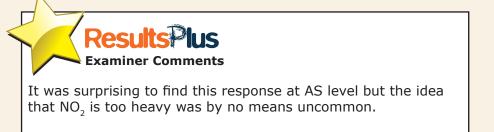
(iv) By considering the equation $NO(g) + O_3(g) \longrightarrow NO_2(g) + O_2(g)$ explain the effect of the reaction of hydrocarbons with nitrogen monoxide on the breakdown of ozone. (1)More depletion of ozone, more breakdown the ozone. **Examiner Comments** This candidate has ignored the key word 'explain' in the question and simply given the outcome of the reaction between nitrogen monoxide and hydrocarbons. **Results**Plus **Examiner Tip** Make sure that you identify the command word in a question e.g. 'state', 'identify', 'define', 'explain'. These words indicate the required scope of the answer. (iv) By considering the equation $NO(g) + O_3(g) \longrightarrow NO_2(g) + O_2(g)$ explain the effect of the reaction of hydrocarbons with nitrogen monoxide on the breakdown of ozone. (1)Nitrogen monoxide attacks the ozone layer and produces more NO2 gas, which is a greenhouse gas and causes global warning. *lesults*Plus **Examiner Comments**

This response not only ignores the essential point required by the question but also brings in irrelevant reference to global warming.

Question 22 (d)

Most candidates referred to the effect of UV radiation but often without mentioning that its intensity would be greater at higher altitudes. Those candidates who discussed the effect of the reduced pressure on the equilibrium were more likely to gain both marks. A surprising number of candidates believed that nitrogen dioxide was too heavy a molecule to reach high altitudes.

(d) Suggest why the proportion of nitrogen dioxide might be reduced at high altitudes. (2) UV radiction and when it and the inter No" + 0" free radiculy radiculy
Results Pus Examiner Comments A typical answer which makes good use of the material in the passage but fails to address the high altitude issue.
Results Plus Examiner Tip There were two marks for this item but the answer only makes one point.
(d) Suggest why the proportion of nitrogen dioxide might be reduced at high altitudes. (2) NHNO2 is heavy, so it would normally drop back to a lower level, hence at high altitudes ND2 is reduced.



Question 22 (e)

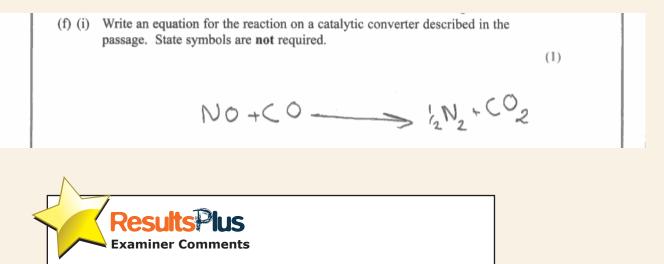
Most candidates understood the role of ozone in absorbing UV radiation and why this is important but there were many examples of careless use of terminology. Common examples were the idea that UV radiation is reflected back into space or prevented from entering the atmosphere. Some candidates failed to identify the radiation as UV, referring to 'radiation' or 'light' or UV and IR; there were many references to global warming being caused by UV radiation.

In order to prevent too much in sight coming into the atmosphere. as as this Too much in light may heat up the carths atmosphere causing global warning

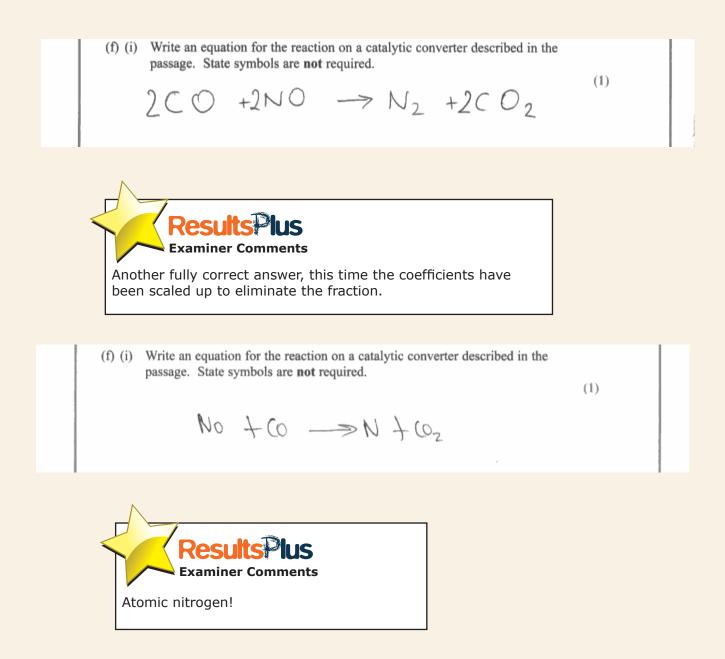


Question 22 (f) (i)

Some candidates failed to balance this equation and there was more use of nitrogen atoms (as in 22(a)).



A correct answer; using $\frac{1}{2}N_2$ is perfectly acceptable.



Question 22 (f) (ii)

Almost all candidates had the right idea in this question; there very few Maxwell-Boltzmann distribution curves. However, as elsewhere in the paper, marks were often lost due to a lack of precision. The main problems were incorrect or missing labels although there were also some endothermic systems drawn. Most examiners commented on the poor quality of the diagrams,

Question 22 (f) (iii)

This question was very well answered with the final marking point the one most likely to be lost.

Question 22 (g)

Candidates usually appreciated the importance of aircraft flying close to the ozone layer although it was often expressed with insufficient clarity, candidates frequently stating to the effect that 'aircraft fly higher than cars'. The second mark proved more difficult to gain with vague ideas such as the time that it would take the NO to reach the upper atmosphere being fairly typical. There was also much vague reference to 'emissions' and 'pollutants' and 'greenhouse gases'.

(g) Jet aircraft are considered a greater threat to the ozone layer than road vehicles. Suggest an explanation for this. (2)expell their the waits products directly in the air at allitudes This nears ozone is found' quicke by Novet. more ozone is affected, whereas in road rehicles the NO will have to travel to the upper abrosphere by the most will have dissipated. (Total for Question 22 = 20 marks) **Examiner Comments** The idea of NO being 'dissipated' is sufficient to gain the second mark. Suggest an explanation for this. (2)innant release Nubagen m p in the outmosp n , so are more likely to ema O 0 36 (Total for Question 22 = 20 marks) **Examiner Comments** The ideas in this response lack sufficient precision ('higher up in the atmosphere than car engines') and development (what exactly does the last sentence mean?) **Examiner Tip** Do ensure that what you have written fully

expresses what you want to say.

(2) because the Jet aircraft travels closer to the Earthas atmosphere so the harmful gases or greenhouse gases attack the ozone utyer faster, also Jet aircrafts may produce larger amounts (Total for Question 22 = 20 marks)	(g) Jet aircraft are considered a greater threat to the ozone layer than road vehicl Suggest an explanation for this.	es.
the Earthas atmosphere so the harmful gases or granhause gases attack the ozone vayer faster, also Jet aircrafts may produce larger amounts	Support an explanation for anot	(2)
grænhausegases attack the ozone layer faster, also Jet arrorafts may produæ larger amounts	because the Jet aircraft travels closer	<u>+0</u>
	the Earth's atmosphere so the harmful	gases or
	granhause gases attack the ozone layer	faster,
(Total for Ouestion $22 = 20$ marks)	also Jet aircrafts may produce larger.	amounts
	(Total for Question 22 =	= 20 marks)



Paper Summary

The number of marks for a question indicates the number of scoring points the examiner will be looking for and the number of lines shows the **maximum** likely length of a reasonable response. Use this information in framing your answers.

Do read your answers carefully and check that you have answered the question set by the examiner.

The 'theory' papers will all include questions that test practical knowledge and understanding.

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