

EUNOIA JUNIOR COLLEGE

JC2 Preliminary Examination 2020 General Certificate of Education Advanced Level Higher 1

CHEMISTRY

8873/01

Paper 1 Multiple Choice

18 September 2020

1 hour

Additional Materials:

Multiple Choice Answer Sheet

Data Booklet

READ THESE INSTRUCTIONS FIRST

Write in soft pencil.

Do not use staples, paper clips, glue or correction fluid.

Write your name, civics group and registration number on the Answer Sheet in the spaces provided unless this has been done for you.

There are thirty questions on this paper. Answer all questions. For each question there are four possible answers A, B, C and D.

Choose the one you consider correct and record your choice in soft pencil on the separate Answer Sheet.

Read the instructions on the Answer Sheet very carefully.

Each correct answer will score one mark. A mark will not be deducted for a wrong answer. Any rough working should be done in this question paper.

The use of an approved scientific calculator is expected, where appropriate.

This document consists of 10 printed pages.

1	Use	of th	e Data Bool	(let l	s reievant to t	nis quesi	ion.			
	Wh	at do	the ions ³⁶ S	²⁻ a	nd ³⁷ C <i>l</i> ⁻ have	e in comr	non?			
	Α	Both	ions have r	nore	electrons tha	an neutro	ns.			
	В	Both	ions contal	n the	same numb	er of nucl	eons in the	ir nuclei.		
	C	Both	ions have a	an ol	uter electronic	configur	ation 2s² 2p	0 ⁶ .		
	D	Both	ions have 2	20 ne	eutrons in the	ir nuclei.				
2	Use	of th	e Data Boo	klet i	is relevant to	this ques	tion.			
	Wh	ich pa	article would	, on	losing an ele	ctron, ha	ve a half-fill	ed set of p	orbitals?	
	A	C-		В	N	С	N-	D	O ⁺	
3	Wh	ich of	the followin	ıg sp	ecies does n	ot have t	he same bo	ond angle a	s the rest?	
	A	oci	2	В	HCN	С	BeH ₂	D	XeF ₂	
4	Wh	nich of	f the followir	ng st	atements abo	out water	are correct	?		
		.1	Water can t	form	hydrogen bo	nds with	propanone.			
		2	Water has than in ice.	a hi	gher density	than ice	as hydroge	n bonds a	re stronger in	water
		3	Water has water than			int than p	entanol as	hydrogen l	oonds are wea	ker in
	A	1 ar	nd 2	В	1 and 3	С	2 and 3	D	1 only	
5	WI	hich o	f the followi	ng st	tatements abo	out graph	ite is incor i	rect?		
	A	All t	he bond and	ales	in it are 120°.					
	В				s bonded to ti		r carbon at	oms.		
	С		able to con		-	e to the	presence o	f delocalise	ed electrons th	at are
	D	it h	as low me	lting	-	to the w	eak instant	aneous di	pole-induced	dipole
					-					

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6 Use of the Data Booklet is relev	evant to this auestion.
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Which of the following contains twice the number of atoms as 16 g of oxygen gas, O₂?

- A 2 g of hydrogen gas
- B 2 g of helium gas
- C 16 g of nitrogen gas
- D 16 g of methane gas

7 The percentage by mass of water in a hydrated magnesium sulfate salt, MgSO₄. xH₂O is 51.1%.

What is the value of x?

- A 4
- **B** 5
- **C** 6
- D 7

8 Dithionite, S₂O₄²⁻ reacts with water under acidic conditions to form thiosulfate, S₂O₃²⁻ and hydrogen sulfite, HSO₃⁻.

$$2S_2O_4^{2-} + H_2O \longrightarrow S_2O_3^{2-} + 2HSO_3^{-}$$

Which of the following statements about the reaction and the species involved is correct?

- A H₂O is the reducing agent.
- B HSO, is the oxidising agent.
- C The oxidation number of sulfur decreases from +3 in $S_2O_4^{2-}$ to +2 in $S_2O_3^{2-}$.
- D The oxidation number of hydrogen increases from +1 in H₂O to +2 in HSO₃.
- 9 1 mol of zinc metal was added to 1 mol of VO₃. The zinc metal was found to be oxidised by VO₃ to form Zn²⁺.

Assuming that all the reactants added were reacted, what is the oxidation number of vanadium in the product?

- A +4
- **B** +3
- C +2
- D +1

10	The first eight ionisation energies	in kJ mol ⁻¹	of an unknown	element A are	given below.
10	THE HISLERING ROUND CHEINICS	. וניונו נאינון	, OI GII GINGIVIIII	OCCUPATION OF A COLO	giron boios

420

3100

4400

5900

8000 9600

11400

13300

Which of the following statements about A is correct?

- A Element A is in group 2 of the Periodic Table.
- B Element A can conduct electricity in the solid state.
- C The oxide of A is an amphoteric oxide.
- D The chloride of A hydrolyses in water to give an acidic solution.

11 Which statements concerning the elements in the third period, from sodium to chlorine, are correct?

- 1 The element that has the lowest melting point is phosphorus.
- 2 The element with the highest electrical conductivity is aluminium.
- 3 The element with the smallest anion is chlorine.
- A 2 and 3
- **B** 1 and 3
- C 1 and 2
- D 2 only

12 Use of the Data Booklet is relevant to this question.

Hypothetically, N₄ could be formed from nitrogen gas by the following reaction.

$$2N_2(g) \rightarrow N_4(g)$$

 ΔH

By considering the bonds broken and bonds formed, as well as the structure of N_4 given below, what would be the value of ΔH for the above reaction?



A +140 kJ mol⁻¹

B +928 kJ mol⁻¹

C -140 kJ mol⁻¹

928 kJ mol⁻¹

13 The enthalpy change of reaction between calcium and water, ΔH_r , can be measured in the laboratory.

$$Ca(s) + 2H_2O(l) \rightarrow Ca(OH)_2(s) + H_2(g)$$
 ΔH_r

What other data is needed in order to calculate the enthalpy change of formation of $Ca(OH)_2(s)$?

- A Lattice energy of calcium hydroxide
- B Enthalpy change of formation of calcium
- C Enthalpy change of combustion of hydrogen
- D First and second ionisation energies of calcium
- 14 A commercial pharmaceutical drug has a constant half-life of 2 h. The drug will lose its effectiveness in the human body once its concentration in the patient's bloodstream falls below 40 mg dm⁻³.

Given that when a patient first consumes the drug, the concentration of the drug in his bloodstream increases to 320 mg dm⁻³, how often should he take his prescription in order to maintain the effectiveness of this drug in his body?

- A every 2 h
- B every 4 h
- C every 6 h
- D every 8 h
- 15 The reactants D and E react in the presence of a dilute acid as follows:

$$D(aq) + E(aq) \rightarrow G(aq) + J(aq)$$

It is found that the rate equation for the reaction is rate = $k[\mathbf{D}]^2[H^+]$.

Which of the following statements is not correct?

- A The magnitude of *k* increases with temperature.
- **B** H⁺ lowers the activation energy of the reaction.
- C Doubling of [E] at constant temperature causes a doubling of the reaction rate.
- **D** Doubling of [H⁺] at constant temperature causes a doubling of the reaction rate.

- 16 In which of the following reactions is the underlined species behaving as a base?
 - $A N_2 + 3H_2 \Longrightarrow 2NH_3$
 - **B** $NH_4^+ + H_2O \rightleftharpoons NH_3 + H_3O^+$
 - $\textbf{C} \quad \text{CH}_3\text{CONH}_2 \, + \, \text{OH}^- \longrightarrow \text{CH}_3\text{COO}^- \, + \, \text{NH}_3$
- 17 Which of the following statements about the titration involving 25.0 cm³ of 0.001 mol dm⁻³ CH₃COOH(aq) and 0.002 mol dm⁻³ NaOH(aq) is correct?
 - A The initial pH is 3.
 - B The equivalence volume is 25.00 cm³.
 - C The solution obtained at equivalence point is acidic.
 - D After adding 6.00 cm³ of NaOH(aq), a buffer solution is obtained.
- 18 Consider a reaction for which the equilibrium constant is independent of temperature. Which of the following statements about this reaction is correct?
 - A The enthalpy change of reaction is zero.
 - B Its rate constant does not vary with temperature.
 - C There are equal amounts of reactants and products.
 - D The activation energies for both the forward and reverse reactions are zero.
- 19 Myoglobin, Mb, is an oxygen-carrier protein that exists in the muscle fibres of most mammals. Each Mb molecule will bind to one O₂ molecule, according to the following equation.

$$Mb(aq) + O_2(aq) \rightleftharpoons MbO_2(aq)$$

$$K_c = 1 \times 10^6 \text{ mol}^{-1} \text{ dm}^3$$

Given that the equilibrium concentration of O_2 is 6.5×10^{-6} mol dm⁻³, what is the percentage of MbO₂ in the Mb-MbO₂ mixture at equilibrium?

- A 50.5 %
- **B** 65.0 %
- C 86.7 %
- D 88.4 %

20 Compound L was reacted with hydrogen gas in the presence of platinum catalyst. How many sigma and pi bonds will be found in the product formed?

	number of sigma bonds	number of pi bonds
A	21	3
В	25	2
C	26	1
D	28	0

21 Compound M has a straight-chain structure with a molecular formula of C₅H₀O.

Which functional groups may be present in an isomer of compound M?

- 1 alcohol
- 2 aldehyde
- 3 ester
- A 1, 2 and 3
- B 1 and 2 only
- C 1 only
- D 2 only

22 (CH₃CH₂)₃CH can react with chlorine under *uv* light to produce monochloro-compounds.

How many possible constitutional isomers of monochloro-compounds can it produce?

- A 2
- **B** 3
- **C** 6
- D 7

23 Compound N reacts with 2 mol of HBr at room temperature in the following reaction:

Given that there are no adjacent carbon-carbon double bonds in **N**, what is the total number of *cis-trans* isomers that **N** can have?

- Δ 1
- **B** 2
- **C** 3
- D 4

24 The following compound was heated with ethanolic sodium hydroxide.

Which of the following represents the structure of the organic product?

A H₃C CH

25 2-methylbuta-1,3-diene, CH₂=C(CH₃)-CH=CH₂, is used as a monomer in the manufacture of synthetic rubbers.

Which compound will **not** produce this monomer on treatment with excess concentrated sulfuric acid at 170 °C?

- A (CH₃)₂C(OH)CH(OH)CH₃
- B HOCH₂CH(CH₃)CH₂CH₂OH
- C HOCH₂CH(CH₃)CH(OH)CH₃
- D HOCH₂C(CH₃)(OH)CH₂CH₃

- 26 P. Q and R are three isomeric alcohols.
 - P CH₃CH₂CH₂CH₂CH₂OH
 - Q CH₃CH₂CH(OH)CH₂CH₃
 - (CH₃)₂C(OH)CH₂CH₃ R

Two or more of these alcohols react with acidified potassium dichromate(VI). One of these alcohols, upon oxidation, produces an organic product S that reacts with sodium carbonate to give carbon dioxide.

Which row is correct?

	reacts with acidified potassium dichromate(VI)	forms S which reacts with sodium carbonate
A	P and Q only	P only
В	P and Q only	Q only
С	P, Q and R	P only
D	P, Q and R	Q only

- 27 Reaction I shows two ethanal molecules combining to form an aldol. Reaction II shows the reaction of the aldol when heated.
 - 2CH₃CHO → CH₃CH(OH)CH2CHO aldol
 - п CH₃CH(OH)CH₂CHO → CH₃CH=CHCHO + H₂O

Which of the following correctly describes reactions I and II?

II Ι elimination reduction A В substitution elimination addition reduction C elimination addition D

- 28 Which of the following molecules can react with both acidified potassium dichromate(VI) and sodium borohydride, NaBH₄?
 - A CH₃CH₂OH
- B CH₃CHO
- C CH₃COCH₃ D CH₃CH(OH)CH₃

29 Artemisinic acid is a useful intermediate for making the anti-malarial drug, artemisin.

Which statement about this compound is incorrect?

- A 15 mol of CO₂ is formed upon complete combustion of 1 mol of it.
- B It can exhibit cis-trans isomerism around a double bond.
- C In the presence of H⁺ ions, it can form esters with ethanol.
- D It can undergo condensation with ethylamine, in the presence of DCC.
- 30 The polymer having the repeat unit shown below occurs in bacteria as a cell storage material.

Which deductions about this substance can be made from this structure?

- 1 It is a condensation polymer.
- 2 Hydrogen bonding is formed between two polymer chains.
- 3 It can be made when a diol and a dicarboxylic undergo polymerisation.
- A 1 only
- B 1 and 2 only
- C 1 and 3 only
- D 2 and 3 only



EUNOIA JUNIOR COLLEGE JC2 Preliminary Examination 2020 General Certificate of Education Advanced Level Higher 1

CANDIDATE NAME						
CIVICS GROUP	1	9			INDEX NUMBER	
CHEMIS Paper 2 Stru		Questic	ons		8873/0 02 September 20 2 hou	20
Candidates a	nswer o	n the (Question	Paper		
Additional Ma	terials:	Data	a Bookle	t		

READ THESE INSTRUCTIONS FIRST

Write your name, civics group, index number on all the work you hand in. Write in dark blue or black pen.

You may use an HB pencil for any diagrams or graphs.

Do not use staples, paper clips, glue or correction fluid.

Section A

Answer all the questions.

Section B

Answer one question.

The use of an approved scientific calculator is expected, where appropriate.

A Data Booklet is provided.

At the end of the examination, fasten all your work securely together. The number of marks is given in brackets [] at the end of each question or part question.

For Examiner's Use				
Secti	ion A			
1	-1	8		
2	1	10		
3	1	10		
4	I	12		
5	1	8		
6	1	12		
Section B				
7 / 8	1	20		
Total	1	80		

Section A

For Examiner's Use

Answer all the questions in this section in the spaces provided.

1 (a) Beams of particles travelling at the same speed are subjected to an electric field. Fig 1.1 shows the experimental set-up in which protons are found to be deflected through an angle of $\pm x^{\circ}$.

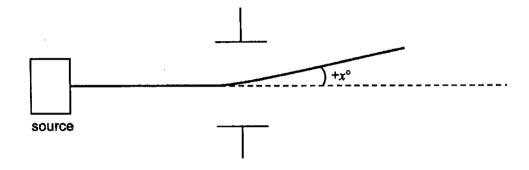


Fig 1.1

(i)	Indic	cate in Fig 1.1, the polarity of the plates and explain your answer.	

		[2	<u>'</u>]
(ii)		uming an identical set of experimental conditions, state and explain the anglidirection of deflection in terms of x° for the following particles.	e
	l:	⁴He nuclei	
			-
	*****	[2	<u>?]</u>
	H:	² H atom	

		[1	11

(b)	With reference to the electronic configurations, explain why the first ionisation energy of fluorine is smaller compared to that of neon.	For Examiner's Use
	Electronic configuration of fluorine:	
	Electronic configuration of neon:	
	·	
	[3]	
	[Total: 8]	

[Turn Over

2 Ascorbic acid (C₆H₈O₆), or more commonly known as Vitamin C, is an important component of our diet. Vitamin C pills are widely available in the market as dietary supplements, with about 1000 mg or 500 mg of Vitamin C in each pill.

For Examiner's Use

To determine the amount of Vitamin C in a pill, a titration involving the following steps can be carried out.

Step 1: IO₃ is reacted with I to produce a known amount of I₂:

$$IO_3^- + 6H^+ + 5I^- \longrightarrow 3I_2 + 3H_2O$$

- Step 2: A Vitamin C pill is dissolved in deionised water.
- Step 3: The I₂ produced in Step 1 is reacted with the Vitamin C solution from Step 2:

$$C_6H_8O_6 + I_2 \longrightarrow C_8H_6O_6 + 2I^- + 2H^+$$

Step 4: The unreacted I₂ from Step 3 will be titrated against S₂O₃²⁻:

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

(a) (i) Write half-equations to show the oxidation and reduction processes that occur in Step 1.

oxidation

reduction[2]

In Step 1, 5.00 g of KIO₃ (M_r = 214.0) was first dissolved in 10 cm³ of deionised water. The solution was then acidified and made up to 250 cm³. 25.0 cm³ of the resulting solution was pipetted out. 2.50 g of KI was added and stirred to ensure complete reaction.

(ii) Calculate the amount of KIO₃ in 25.0 cm³ of solution.

[1]

	(iii) Hence, calculate the amount of iodine formed in Step 1.	For Examiner's Use
	[3]	
(b)	Is Vitamin C, $C_6H_8O_6$, acting as a reducing agent or an oxidising agent in Step 3? Explain your answer in terms of the change in oxidation numbers.	
	[1]	
(c)	In Step 3, the solution of I_2 obtained in (a)(iii) was mixed with a solution of Vitamin C. The unreacted I_2 from Step 3 required 20.85 cm ³ of 0.400 mol dm ⁻³ $S_2O_3^{2-}$ for complete reaction in Step 4.	
	(i) Calculate the amount of I_2 that has reacted with $S_2O_3{}^{2-}$ in Step 4.	
	T41	
	[1]	
	(ii) Using your answer in (a)(iii), calculate the amount of I ₂ reacted with Vitamin C in Step 3.	
		:
	[1]	
	(iii) Hence, calculate the mass of Vitamin C in the pill.	
	[1]	
	[Total: 10]	

[Turn Over

3	Nitr	ogen (can form a few hyd	drides such as	NH₃ and N₂H₄.	;	For Examiner's Use
	(a)	Draw mole	/ a 'dot-and-cross' cule about each n	diagram to sh itrogen atom.	now the bonding in	N₂H₄. State the shape of the	Osa
					shape	[2]	
	(b)	The	boiling points of th	ree compound	ds are given in the t	able below.	
			!	compound	boiling point / °C		
				NH ₃	-33		
				N ₂ H ₄	114		
				C₂H₄	-103		:
		Expl	lain the difference	between the b	poiling points of:		
		(i)	N₂H₄ and C₂H₄				
		.,					
				,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		

			***************************************	•••••			
			***********	*******************************	.,		
				•••••	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	[2]	
		(II)	NH ₃ and N ₂ H ₄				
		(,	141 13 CITIC 1421 14				
				••••••			

			***************************************		••••••••••••	[1]	

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(c)	can	en NH ₃ is reacted with HC l , a white solid of NH ₄ C l with a melting point of 338 °C be obtained upon heating to dryness. The melting point of the NaC l solid obtained a similar reaction between NaOH and HC l is found to be much higher at 801 °C.	For Examiner's Use
	Ехр	lain why this is so with reference to their structure and bonding.	
	•••••		
	••••		
		· · · · · · · · · · · · · · · · · · ·	
	••••		
	••••		<u> </u>
		[3]	
(d)	(i)	NH ₃ reacts with BH ₃ to form ammonia borane, NH ₃ BH ₃ .	
		Draw a 'dot-and-cross' diagram to show the bonding in NH ₃ BH ₃ .	
		[1]	
	(ii)	Is ammonia borane a polar molecule? Explain your answer clearly.	
		[1]	
		[Total: 10]	

[Turn Over

4 (a) An experiment was carried out to determine the relative oxidising power of three unknown halogens X₂, Y₂ and Z₂. Table 4.1 shows the results of experiments in which the halogens X₂, Y₂ and Z₂ were added to separate aqueous solutions containing X⁻, Y⁻ and Z⁻ ions.

For Examiner's Use

	X ⁻ (aq)	Y⁻(aq)	Z- (aq)
Х2		no reaction	no reaction
Y ₂	brown solution formed		orange solution formed
Z ₂	brown solution formed	no reaction	

Table 4.1

(i)	Using the results given in Table 4.1, suggest the identity of X ₂ , Y ₂ and Z ₂ .
	X ₂ :
	Y ₂ :
	Z ₂ :
(ii)	Hence, state the most powerful oxidising agent among the halogens.
	[1]
(iii)	Explain, in terms of ease of gaining electrons, why the element you have identified in (a)(ii) is the most powerful oxidising agent.
	[2]

(b)	Describe what happens when separate samples of magnesium oxide and aluminium oxide are added to water. Give equations for any reactions that occur.	For Examiner's Use
	What is the effect of adding universal indicator to each resulting solution?	
		:
		:
	[3]	
(c)	Describe the reactions of the chlorides of magnesium and phosphorus with water. Include the approximate pH value of any resulting solutions and write equations for any reactions that occur.	
	[4]	
	[Total: 12]	
		1

[Turn Over

5 Ammonia, NH₃ is commonly used as a household cleaner to remove greasy dirt in ovens. It is also used to clean windows as it evaporates quickly and does not leave stains on glass.

For Examiner's Use

[2]

25.0 cm³ of 0.500 mol dm⁻³ NH₃ was titrated against 0.500 mol dm⁻³ HC*L* Fig. 5.1 shows the pH changes during the titration.

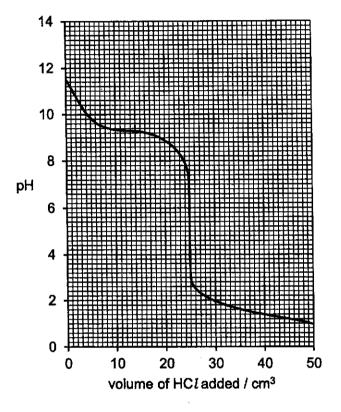


Fig. 5.1

(a) (i) Using the pH from the graph, calculate the concentration of hydroxide ions, in mol dm $^{-3}$, in the 25.0 cm 3 of 0.500 mol dm $^{-3}$ solution of NH $_3$.

(ii)	Explain how your answer in (a)(i) indicates that NH ₃ is a weak base.	
	***************************************	••••
	,	

For Examiner's Use

(b)	Duri	uring the addition of the first 15.0 cm^3 of HC L the mixture is behaving as a buffer.				
	(i)	State what is meant by	y the term buffe	er solution.		
			•••••	••••••		
					[1]	
	(ii)	Explain how the mixtur an equation in your an	_	s a buffer at thi	s stage of the titration. Include	
			••••••			
					[2]	
(c)	of th		nation from Fig.	5.1 and Table	or to determine the end-point 5.1, explain why the student's tor.	
٠		indicator	colour in acid	colour in alkali	pH range over which the colour change occurs	
		alizarin	yellow	orange	10.1–13.0	
		bromocresol green	yellow	blue	3.8–5.4	
		gentian violet	yellow	violet	0.0–1.6	
		phenolphthalein	colourless	pink	8.2–10.0	
			Tab	le 5.1		
-					[2]	
					[Total: 8]	

6	(a)	The two	most	common	types	of	nylon	used	in	textile	and	plastic	industries	are
		nylon-6 a	and nyl	on-6,6.										

For Examiner's Use

The structure of nylon-6,6 is shown below.

nylon-6,6

(i)	Define the term polymer.
	[1]
(ii)	Ropes made of nylon-6,6 are chosen for its high tensile strength. Ropes made from polyesters such as poly(ethylene terephthalate) (PET) are about 90% as strong as that made from nylon-6,6.
	With the aid of a diagram, showing relevant interactions, explain why nylon-6,6 is stronger than polyester.

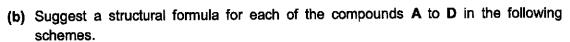
(iii) Rust on window grilles can be removed by spraying the grilles with the rust remover shown in Fig 6.1 before scrubbing with a brush.

For Examiner's Use

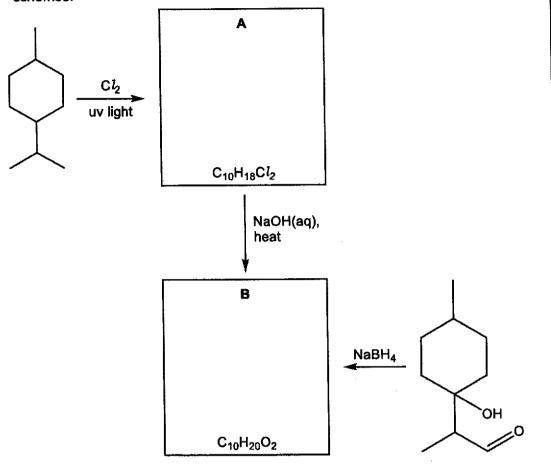


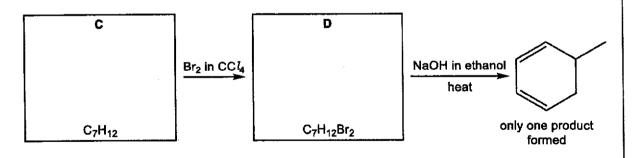
Fig 6.1

	Explain whether a brush with nylon bristles or high density poly(ethene) bristles should be used for the scrubbing.
	[2]
(iv)	State whether nylon-6,6 is classified as a thermoplastic or thermoset. Hence, explain whether it can be recycled.
	[2]



For Examiner's Use





[4]

[Total: 12]

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Section B

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Answer one question in this section in the space provided.

7 (a) Oleic acid is a monosaturated omega 9 fatty acid found in olive oil. It is said to potentially reduce the risk of coronary heart disease if taken daily. Oleic acid has the structure shown: CH₃(CH₂)₇CH=CH(CH₂)₇COOH (i) Describe the pi bonding in terms of orbital overlap. You may draw a diagram to illustrate your answer.[2] (ii) Oleic acid exists as a pair of isomers. Draw and label the structural formulae of the two isomers. [2] (iii) Describe a chemical test that would distinguish oleic acid from ethanoic acid.

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Turn Over

(b) Fig. 7.1 shows a bomb calorimeter, inside a controlled temperature water jacket, which is used to find an accurate value of the standard enthalpy change of combustion of oleic acid.

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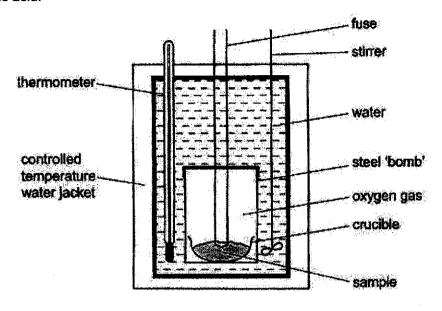


Fig. 7.1

(i)	Explain why the steel 'bomb' is flushed with a high pressure of oxygen at the start of the experiment.
	[1]
(ii)	There is minimal heat lost from the calorimeter because of the controlled temperature water jacket. Suggest how the controlled temperature water jacket achieves this.
	[1]

(iii) A 2.98 g sample of oleic acid (C₁₈H₃₄O₂) is burned in a bomb calorimeter with a heat capacity of 1.98 kJ K⁻¹. The temperature of the calorimeter increases by 59.3 °C. The process is known to be 95 % efficient.

For Examiner's Use

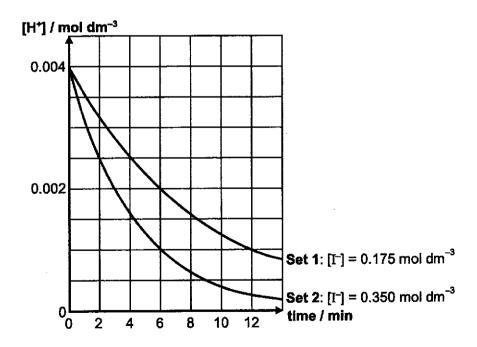
Calculate the standard enthalpy change of combustion of oleic acid.

[3]

(c) The Harcourt and Esson reaction is a reaction involving hydrogen peroxide and acidified potassium iodide.

$$H_2O_2 + 2I^- + 2H^+ \longrightarrow 2H_2O + I_2$$

To determine the order of reaction of each reactant, two sets of reaction mixtures containing varying concentrations of I⁻ were prepared. The concentration of hydrogen peroxide used for both experiments is 0.200 mol dm⁻³. The results are as follows.



(i) Given that the initial reaction rate for Set 2 is 1.00 × 10⁻³ mol dm⁻³ min⁻¹, calculate the initial rate of reaction for **Set 1**. Show your working clearly.

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[1]

(ii) Use the information given above to determine the order of reaction with respect to I⁻ and H⁺. Show your reasoning clearly.

[2]

(iii) The actual order of reaction with respect to H₂O₂ is one.

The rate of the reaction was measured as 4.4×10^{-5} mol dm⁻³ s⁻¹ when $[H_2O_2] = 0.002$ mol dm⁻³, $[H^+] = 0.2$ mol dm⁻³ and $[I^-] = 0.2$ mol dm⁻³

Determine the rate constant for this reaction and state its units

[2]

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(d)		frogen iodide can be formed by the reaction between H_2 and I_2 under strongly ited platinised asbestos. It exists as a homogeneous equilibrium system.	For Examiner's Use
		$H_2(g) + I_2(g) \rightleftharpoons 2HI(g) \cdots (1)$	
	(i)	Give the expression of the equilibrium constant, K_c of reaction (1).	
٠			
		[1]	
	(ii)	Calculate the value of K_c at 445 °C, if the equilibrium concentrations in mol dm ⁻³ of H ₂ (g), I ₂ (g) and HI (g) are 2.06, 13.40 and 36.98 respectively.	
		[1]	
	(ii)	The value for the equilibrium constant, K_c for reaction (1) at 1000 °C is 13. By comparing the values of K_c at 445 °C and 1000 °C, deduce whether the reaction between hydrogen and iodine is endothermic or exothermic.	
	٠		
		[2]	E
		[Total: 20]	
			1

8 (a) When 1 g of ethanol was burned under a container of water, it was found that 100 g of water was heated from 20 °C to 60 °C. The process was known to be only 65 % efficient.

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Use these data and values from the *Data Booklet* to calculate the enthalpy change of combustion of ethanol.

[3]

(b) Fig 8.1 shows an energy cycle involving ethanol.

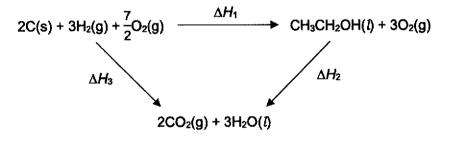


Fig 8.1

(i) Name the enthalpy change represented by ΔH_1 .

______[

(ii) Given that the enthalpy change of combustion of carbon and hydrogen are $-393.5 \text{ kJ moi}^{-1}$ and $-285.8 \text{ kJ moi}^{-1}$ respectively, use the energy cycle given above and your answer to (a) to calculate ΔH_1 .

[3]

(c) A series of reactions starting from 3-chloropropan-1-ol is shown.

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C1 OH
$$\stackrel{\text{I}}{\longrightarrow}$$
 A $\stackrel{\text{II}}{\longrightarrow}$ HO $\stackrel{\text{conc. H}_2SO_4}{\longrightarrow}$ C₆H₈O₄

m	State the r	eagents and	conditions	required	for reactions	I and II.
w	Otate the i	cagento and	CONGRETE	required	IOI ICACAONO	I WIN II

Step I:[1]

Step II:[1]

(ii) Draw the structural formulae of A and B.

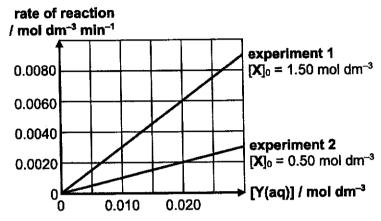
[2]

(d) The reaction between X and Y can be represented by the following equation:

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$$X(aq) + 2Y(aq) \longrightarrow Z_1(aq) + Z_2(g)$$

To study the kinetics of the above reaction, two experiments were carried out in the presence of excess **X**. The experimental results obtained are graphically represented below.



Note: $[X]_0$ = initial concentration of X(aq)

(i) With the aid of the graph, determine the order of reaction with respect to each of the two reactants, **X** and **Y**.

[2]

(ii) Hence, write the rate equation for the reaction and state the units for the rate constant k.

[2]

(e)	8.00 g of solid potassium chromate(VI), K ₂ CrO ₄ , is dissolved in acid to make a 100 cm ³
	solution. A dynamic equilibrium occurs in the mixture according to the equation below:

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$$2CrO_4^{2-}(aq) + 2H^{+}(aq) \rightleftharpoons Cr_2O_7^{2-}(aq) + H_2O(l)$$

(i) Write the K_c expression for the above equilibrium mixture.

[1]

(ii) At pH 5.75, the system reaches equilibrium and **one-fifth** of the original amount of chromate(VI) ions remain. Calculate the concentration of chromate(VI) ions at this pH.

[1]

(iii) Hence, calculate the value of Kc.

[3]

[Total: 20]

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2020 JC2 Prelim Exam H1 Chemistry 8873 Paper 1 Worked Solution

- A is incorrect as both ³⁸S²⁻ and ³⁷Ct have 20 neutrons but 18 electrons so that they have fewer (not more) electrons than neutrons.
 - B is incorrect as ³⁸S²⁻ has 36 nucleons but ³⁷Ct has 37 nucleons.
 - C is incorrect as ³⁶S²⁻ and ³⁷C*I*⁻ have outer electronic configuration of 3s²3p⁶.
 - **D** is correct as both ³⁶S²⁻ and ³⁷C*I*⁻ contain 20 neutrons.

⇒D

2

	Species	Before losing an electron	After losing an electron
A	C-	1s ² 2s ² 2p ³	1s ² 2s ² 2p ²
В	N	1s ² 2s ² 2p ³	1s ² 2s ² 2p ²
C	N-	1s ² 2s ² 2p ⁴	1s ² 2s ² 2p ³
D	O,	1s ² 2s ² 2p ³	1s ² 2s ² 2p ²

3

	Structure	No. of bp & lp; shape
A	CI CI	2bp, 2lp Bent
В	H—C <u>≔</u> N	2 bp, 0lp linear
С	HBeH	2 bp, 0lp linear
D	F—Xe—F	2 bp, 3lp linear

⇒A

- 4 1 is correct as there is a lone pair on the O atom on propanone which can form attractions to the H atom with a partial positive charge on water.
 - 2 is wrong as the lower density of ice is due to its open structure with large amount of spaces in between that arises due to hydrogen bonding. The strength of hydrogen bonds is not affected.
 - 3 is wrong as the lower boiling point is due to the much larger number of electrons in pentanol, which leads to significantly stronger instantaneous dipole-induced dipole interactions.

⇒D

5 A and B are true as can seen from the structure of graphite below:



C is true as the delocalised electrons can act as mobile charge carriers to conduct electricity within each layer.

D is false as graphite has high thermal stability due to the strong covalent bonds between the carbon atoms.

 $\Rightarrow D$

		Amt of molecule s	Amount of atoms
	16 g of O₂	0.5 mol	2 × 0.5 = 1 mol
Α	2 g of H ₂	1 mol	2 mol
	- g - 1.1.2	1	<u> </u>
В	2 g of He	0.5 mol	0.5 mol

7 Let the empirical formula of the salt be MgSO₄*xH₂O

$$\frac{18.0x}{24.3 + 32.1 + 4 \times 16.0 + 18.0x} \times 100\% = 51.1$$

$$x = 7$$

⇒D

£

8 A and B are false as this is a disproportionation reaction, and S₂O₄²⁻ is both the oxidising and reducing agent. C is true cos of the following.

Oxid no of S in S₂O₄^{t-} =
$$\frac{-2-4(-2)}{2}$$
 = +3;

oxid no of S in
$$S_2O_3^{2-} = \frac{-2-3(-2)}{2} = +2$$
.

D is false as the oxidation number of hydrogen is +1 in both H_sO and HSO_s^* . \Rightarrow C

9 Amt of zinc metal reacted = 65.4 = 0.02945 mol Zn → Zn² + 2e⁻

Amt of electrons lost by zinc metal = 2 × 0.02945 mol = amt of electrons gained by VO₃⁻

Amt of VO₁ reacted

$$= \frac{20}{1000} \times 1.5 = 0.03 \text{ mol}$$

Hence, 1 mol of VO_3 gained 2×0.02945

0.03 ≈ 2 mol of electrons ⇒ oxid no. of VO₃ decreased by 2.

Oxid no. of V in VO₃⁻ = -1 - 3(-2) = +5 Oxid no of Vin product = +5 - 2 = +3 \Rightarrow B

- 10 A is incorrect as there is a big jump between the first and second ionisation energies, indicating that the element is in group 1 and not group 2.
 - B is correct as group 1 metals can conduct electricity due to presence of sea of mobile valence electrons in its metallic lattice structure.
 - C is incorrect as group 1 oxides are basic.

D is incorrect as group 1 chlorides do not hydrolyse in water and produces a neutral solution.

⇒B

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- 11 1 The element that has the highest melting point is silicon, not aluminium.
 - 2 The element with the highest electrical conductivity is aluminium due to largest amt of delocalised valence electrons in its metallic lattice structure.
 - 3 The element with the smallest anion is chlorine due it having the largest nuclear charge amongst the anions that are all isoelectronic.

⇒ /

- 12 ΔH = BE(reactants)- BE(products) = 2(+944) - 6(+160) = +928 kJ mol⁻¹ ⇒ B
- 13 ΔH of reaction = ΣΔΗ_ε^θ of products ΣΔΗ_ε^θ of reactants. Since Ca(s) and H₂(g) are elements in their standard states, ΔH_ε^θ(Ca) = ΔH_ε^θ(H₂)=0 the other data required would be the ΔH_ε^θ(H₂O) which is equal to ΔH_ε^θ(H₂). Since both are represented by H₂ (g) + ½ O₂ (g) → H₂O (l).
 C

14 Given t_{1/2} = 2.0 hour

After 1st $t_{1/2}$, mass of $X = \frac{1}{2}(320) = 160 \text{ mg}$

After $2^{nd} t_{1/2}$, mass of $X = \frac{1}{2}(160) = 80 \text{ mg}$

After 3rd $t_{1/2}$, mass of $X = \frac{1}{2}(80) = 40 \text{ mg}$

Hence, drug X will still be effective after 3 half-lives (3 x 2.0 = 6.0 hour). \Rightarrow C

15 The order of reaction with respect to D is 2, hence tripling its concentration will lead to a (3)² = 9 times increased in reaction rate. Acid is the catalyst in the reaction, thus it introduces alternative reaction pathway with lower activation energy.

Based on Arrhenius equation

(k=Ae RT), an increase in temperature causes rate constant increase as it leads to increase in reaction rate.

⇒ C

- 16 A: H₂O is behaving as a base as it gains a proton from NH₄⁺.
 - B: NaOH is behaving as a base as the OH gains a proton from C_8H_5OH .
 - C: Na is not behaving as a base as it does not gain a proton. This reaction is a redox reaction, not an acid-base reaction.
 - D: CH₃CH₂NH₂ is behaving as a base as it gains a proton from H₃O⁺.

 ⇒ D

17 A is false as CH₃COOH is a weak acid, so the pH should be higher than -lg(0.001).

B is false as the equivalence volume shid be $\left(\frac{25.0}{1000} \times 0.001\right) + 0.002 \times 1000 = 12.50 \text{ cm}^3$

C is false as a basic salt, CH₃COO⁻Na⁺ is formed.

D is true as a mixture of CH₃COO⁻ and CH₃COOH will be obtained upon addition of 6.00 cm³ of NaOH.

⇒D

form 2 double bonds. ⇒ D

For option D, the product 25 OHCH₂C(CH₃)CHCH₃ or HOCHC(CH₃)CH₂CH₃ will be formed. 19 Let % of MbO₂ be y% & [MbO₂]_{eqm} be <u>xy</u>

> Primary alcohol L is oxidized by acidified potassium dichromate to form a carboxviic acid which reacts with sodium carbonate to form carbon dioxide gas. Secondary alcohol M is oxidised by acidified potassium dichromate to form a ketone. Tertiary alcohol N does not undergo oxidation.

27 Since the 2 aldehydes combine to form a single aldol, reaction 1 is an addition reaction. For reaction 2, water is removed from the original molecule thus it is an elimination reaction.

 \Rightarrow D

28

	Rxn with	Rxn
	K ₂ Cr ₂ O ₇ ?	with
		NaBH ₄ ?
CH₃CH₂OH	Yes	No
СН₃СНО	Yes	Yes
CH₃COCH₃	No	Yes
CH₃CH(OH)CH₃	Yes	No

ے B

- 29 A: There are 15 C atoms, 22 H atoms and 2 O atoms. So 15 mol of CO2 will be formed upon complete combustion of 1 mol of this cod.
 - B: Cls-trans isomerism requires each C on the double bond to have two different atoms/ groups. However the alkene is RR'- C = CH₂ and alkene in the ring can only exhibit cis configuration. Thus it CANNOT exhibit cis-trans isomerism
 - C: There is a carboxylic acid functional group that can undergo condensation reaction with ethanol upon heating in acidic conditions.

D:Carboxylic acid undergo can condensation with amine group in the presence of DCC to form an amide

30 The structure shows a condensation polymer (polyester). The polymer does not satisfy the requirements of hydrogen bonding. be made can HOCH(CH₃)CH₂COOH and not between a dicarboxylic acid and diol.

 $K_C = \frac{[MbO_2]}{[O_2][Mb]} = \frac{\frac{100}{100}}{[6.5 \times 10^{-6}](\frac{100 - y}{100})x}$ % of MbO2 in the Mb-MbO2 mixture, y = 86.7%

⇒ C

20 Both alkenes and carbonyl group will undergo reduction to form

18 When the enthalpy change of reaction is zero, it is neither endothermic nor

Hence,

not be changed as a result.

temperature will not affect its equilibrium

position, thus its equilibrium constant will

Let x be the initial concentration of Mb(aq).

Let % of Mb be (100-v)% & [Mb].com be

changes

 $-=1\times10^6 \, mol^{-1} dm^3$

exothermic.

 $(100 - y)_{x}$

100

21 An unsaturated 5-carbon alcohol would have molecular formula, C₅H₁₂O, e.g. CH3CH2CH2CH2CH2OH.

E has 4 less H atoms which implies that there is presence of 2 multiple bonds which could be C=C or C=O bond.

Hence, the possible isomers are CH2=CHCH=CHCH2OH, CH₂=CHCH₂COCH₃ and $CH_2=CHCH_2CH_2CHO$.

Hence, there can be alkene, alcohol, ketone and aldehyde present in the isomers of C₅H₈O. (1 and 2 only)

An ester is not possible as there is only 1 oxygen atom present in E. ⇒B

The chlorine atom can substitute once with Ha, Ha or Ha. (i.e. there are 3 types of hydrogen atoms - He, He and He, with different chemical environments) Hence, there can only be 3 possible structural isomers of monochloro-compounds.

 $\Rightarrow B$

⇒ C

23

There are 2 double bonds that can exhibit cis-trans. So there are 3 cis trans isomers. cis-cis, trans-trans and cistrans(take note trans-cis is the same as cis-trans since the molecule symmetrical)

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Answer Key

Qn	Ans	Qn	Ans	Qn	Ans
1	D	11	Α	21	В
2	С	12	В	22	В
3	A	13	С	23	С
4	ם	14	C	24	ם
5	۵	15	С	25	D
6	Α	16	D	26	Α
7	D	17	D	27	D
8	С	18	Α	28	В
9	В	19	С	29	В
10	В	20	D	30	Α

22



EUNOIA JUNIOR COLLEGE JC2 Preliminary Examination 2020 General Certificate of Education Advanced Level

CANDIDATE NAME					
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CHEMISTRY

8873/02

Paper 2 Structured Questions

02 September 2020 2 hours

Candidates answer on the Question Paper

Additional Materials: Data Booklet

READ THESE INSTRUCTIONS FIRST

Write your name, civics group, index number on all the work you hand in. Write in dark blue or black pen.

You may use an HB pencil for any diagrams or graphs.

Do not use staples, paper clips, glue or correction fluid.

Section A

Answer all the questions.

Section B

Answer one question.

The use of an approved scientific calculator is expected, where appropriate.

A Data Booklet is provided.

At the end of the examination, fasten all your work securely together. The number of marks is given in brackets [] at the end of each question or part question.

For Examiner's Use					
Sect	ion A				
1	1	8			
2	1	10			
3	1	10			
4	1	12			
5	1	8			
6	1	12			
Section B					
7 / 8	1	20			
Total	1	80			

This document consists of 23 printed pages and 1 blank page.

Section A

For Examiner's Use

Answer all the questions in this section in the spaces provided.

1 (a) Beams of particles travelling at the same speed are subjected to an electric field. Fig 1.1 shows the experimental set-up in which protons are found to be deflected through an angle of $+x^{\circ}$.

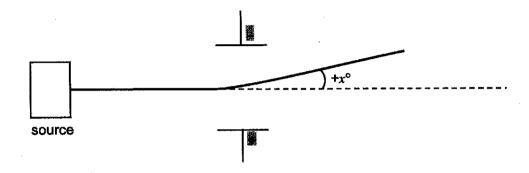


Fig 1.1

(i)	Indicate in Fig 1.1, the polarity of the plates and explain your answer.
	The plate on top is negatively charged as the positively charged proton is
	attracted to it.
	[2]
(ii)	Assuming an identical set of experimental conditions, state and explain the angle and direction of deflection in terms of x° for the following particles.
	I: ⁴ He nuclei
	$\frac{q}{m} = \frac{2}{4} = \frac{1}{2}$ Hence, angle of deflection is 1/2 that of the proton's, $+x/2^{\circ}\Box$
	(towards negative plate)
	[2]
	II: ² H atom
	0°. The atom has no net charge and hence remains undeflected.
	[4]

b)	With reference to the electronic configurations, explain why the first ionisation energy of fluorine is smaller compared to that of neon.
	Electronic configuration of fluorine: .1s².2s².2p⁵
	Electronic configuration of neon: 1s².2s².2p ⁶
	Ne has a greater nuclear charge compared to F, while the shielding effect
	experienced by both species is approximately the same as they both have same
	number of inner core electrons. The 2p electron in Ne experiences a greate
	effective nuclear charge than F as it is more strongly attracted to the nucleus
	Hence, more energy is required to remove 2p electron from Ne atoms.
	[3]
	[Total: 8]

[Turn Over

2 Ascorbic acid (C₆H₆O₈), or more commonly known as Vitamin C, is an important component of our diet. Vitamin C pills are widely available in the market as dietary supplements, with about 1000 mg or 500 mg of Vitamin C in each pill.

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To determine the amount of Vitamin C in a pill, a titration involving the following steps can be carried out.

Step 1: IO₃ is reacted with I to produce a known amount of I₂:

$$IO_3^- + 6H^+ + 5I^- \longrightarrow 3I_2 + 3H_2O$$

- Step 2: A Vitamin C pill is dissolved in deionised water.
- Step 3: The I₂ produced in Step 1 is reacted with the Vitamin C solution from Step 2:

$$C_6H_8O_6 + I_2 \longrightarrow C_6H_6O_6 + 2I^- + 2H^+$$

Step 4: The unreacted I₂ from Step 3 will be titrated against S₂O₃²⁻:

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

(a) (i) Write half-equations to show the oxidation and reduction processes that occur in Step 1.

oxidation $2I^- \longrightarrow I_2 + 2e^-$

reduction
$$2IO_3^- + 12H^+ + 10e^- \longrightarrow I_2 + 6H_2O$$
 [2]

In Step 1, 5.00 g of KIO₃ (M_r = 214.0) was first dissolved in 10 cm³ of deionised water. The solution was then acidified and made up to 250 cm³. 25.0 cm³ of the resulting solution was pipetted out. 2.50 g of KI was added and stirred to ensure complete reaction.

(ii) Calculate the amount of KIO₃ in 25.0 cm³ of solution.

Total amount of KIO₃ dissolved = $\frac{5.00}{214.0}$ = 0.02336 mol

Amount of KIO₃ in 25 cm³ of solution = $\frac{5.00}{214.0} \times \frac{25.0}{250} = 0.00234$ mol

[1]

(iii) Hence, calculate the amount of iodine formed in Step 1.

Amount of KI reacted =
$$\frac{2.50}{166.0}$$
 = **0.0151 mol**

Since $KIO_3 = 5KI$,

 $0.01506\,\mathrm{mol}\,\mathrm{of}\,\mathrm{KI}$ will need $0.003012\,\mathrm{mol}\,\mathrm{of}\,\mathrm{KIO_3}$ for complete reaction.

Hence, KIO₃ is the limiting reagent.

Amount if I_2 formed = $3 \times amount$ of $KIO_3 = 0.00701 mol$

[3]

(b) Is Vitamin C, C₆H₆O₈, acting as a reducing agent or an oxidising agent in Step 3? Explain your answer in terms of the change in oxidation numbers.

Reducing agent. It reduces iodine as can be seen from the decrease in oxidation

number from 0 in iodine to -1 in iodide. [1]

- (c) In Step 3, the solution of I₂ obtained in (a)(ii) was mixed with a solution of Vitamin C. The unreacted I₂ from Step 3 required 20.85 cm³ of 0.400 mol dm⁻³ S₂O₃²⁻ for complete reaction in Step 4.
 - (i) Calculate the amount of I₂ that has reacted with S₂O₃²⁻ in Step 4.

Amount of I₂ reacted with
$$S_2O_3^{2-} = \frac{20.85}{1000} \times 0.400 \times \frac{1}{2}$$

= **0.00417 mol**

[1]

(ii) Using your answer in (a)(ii), calculate the amount of I_2 reacted with Vitamin C in Step 3.

Amount of
$$I_2$$
 reacted with Vitamin C = 0.007009 - 0.004170
= 0.00284 mol

[1]

(III) Hence, calculate the mass of Vitamin C in the pill.

Amount Vitamin C reacted = Amount of
$$I_2$$
 reacted = 0.002839 mol
Mass of Vitamin C in pill = 0.002839 × 176.0
= **0.500 g**

[1]

[Total: 10]

For Examiner's Use

- 3 Nitrogen can form a few hydrides such as NH₃ and N₂H₄.
 - (a) (i) Draw a 'dot-and-cross' diagram to show the bonding in N₂H₄. State the shape of the molecule about each nitrogen atom.

Shape Trigonal pyramidal [2]

(b) The boiling points of three compounds are given in the table below.

compound	boiling point/ °C
NH ₃	-33
N₂H₄	114
C₂H₄	-103

Explain the difference between the boiling points of:

(i) N₂H₄ and C₂H₄

The intermolecular hydrogen bonds between N_2H_4 are stronger than the instantaneous dipole-induced dipole interactions between C_2H_4 . Hence, more energy is required to overcome the intermolecular forces of attraction between N_2H_4 than C_2H_4

(ii) NH₃ and N₂H₄

N₂H₄ can form an average of two hydrogen bonds per molecule as compare to

NH₃ which can only form one. More energy is required to break the <u>more</u>

extensive hydrogen bonds between N₂H₄, hence it has a higher boiling point.

.....[1]

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(c) When NH₃ is reacted with HCl, a white solid of NH₄Cl with a melting point of 338 °C can be obtained upon heating to dryness. The melting point of the NaCl solid obtained after a similar reaction between NaOH and HCl is found to be much higher at 801 °C.

Explain why this is so.

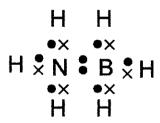
NH₄Cl and NaCl both have giant ionic structures, with strong electrostatic forces of attractions between oppositely charged ions. The strength of ionic bonds is measured by the magnitude of lattice energy.

The charges of the ions in both solids are the same, but NH₄* has a larger ionic radius than Na*. Since $|\Delta H_{lag}| \propto \frac{q_+ \times q_-}{r_+ + r_-}$, the magnitude of lattice energy of NH₄Cl is smaller, and ionic bonds are weaker. Hence, NH₄Cl has a lower melting

point than NaCL [3]

(d) (i) NH₃ reacts with BH₃ to form ammonia borane, NH₃BH₃.

Draw a 'dot-and-cross' diagram to show the bonding in NH₃BH₃.



[1]

(ii) Is ammonia borane a polar molecule? Explain your answer clearly.

Yes. There is a net dipole moment as the dipole moments due to the polar

N-H and N-B bonds do not cancel out. [1]

[Total: 10]

Turn Over

4 (a) An experiment was carried out to determine the relative oxidising power of three unknown halogens X₂, Y₂ and Z₂. Table 4.1 shows the results of experiments in which the halogens X₂, Y₂ and Z₂ were added to separate aqueous solutions containing X⁻, Y⁻ and Z⁻ ions.

For Examiner's Use

	X- (aq)	Y ⁻ (aq)	Z⁻ (aq)
X ₂		no reaction	no reaction
Y ₂	brown solution formed		orange solution formed
Z ₂	brown solution formed	no reaction	

Table 4.1

(i)	Using the results given in Table 4.1, suggest the identity of X_2 , Y_2 and Z_2 .
	X ₂ : I ₂ or iodine
	Y ₂ : Ct ₂ or chlorine (accept F ₂ or fluorine)
	Z_2 : Br_2 or bromine [2]
(ii)	
	Chlorine, C1 ₂ (accept Y ₂) [1]
(iii	Explain, in terms of ease of gaining of electrons, why the element you have identified in (a)(ii) is the most powerful oxidising agent.
	Chlorine has the smallest atomic radius, hence the tendency to accept
	electrons is highest
	[2]

(b)	Describe what happens when separate samples of magnesium oxide and aluminium oxide are added to water. Give equations for any reactions that occur.
	What is the effect of adding universal indicator to each resulting solution?
	Aluminium oxide does not dissolve in or react with water. Universal indicator is
	green colour.
	MgO is sparingly soluble in water to give a weakly alkaline solution. Universal
	indicator is blue in colour. MgO(s) + H ₂ O(/) ⇒ Mg(OH) ₂ (ag)
	[3]
(c)	Describe the reactions of the chlorides of magnesium and phosphorus with water. Include the approximate pH value of any resulting solutions and write equations for any reactions that occur.
	When MgC ½ dissolves in water, the Mg²+ gets hydrated.
	The hydrated Mg²⁺ undergoes slight hydrolysis to give H₃O⁺. The resultant solution
	has a <u>pH ≈ 6.5</u> .
	MgC l_2 + 6H ₂ O → [Mg(H ₂ O) ₆] ²⁺ + 2C l^-
	$[Mg(H_2O)_6]^{2+} + H_2O \Rightarrow [Mg(OH)(H_2O)_5]^{+} + H_3O^{+}$
	PC <i>l</i> ₅ hydrolyses in water to give H₃O⁺ as shown.
	The resultant solution has a <u>pH ≈ 2</u> .
	PC 1 ₅ + 4H ₂ O → H ₃ PO ₄ + 5HC 1
	[4]
	[Total: 12]

[Turn Over

5 Ammonia, NH₃ is commonly used as a household cleaner to remove greasy dirt in ovens. It is also used to clean windows as it evaporates quickly and does not leave stains on glass.

For Examiner's Use

 25.0 cm^3 of $0.500 \text{ mol dm}^{-3} \text{ NH}_3$ was titrated against $0.500 \text{ mol dm}^{-3} \text{ HC} L$ Fig. 5.1 shows the pH changes during the titration.

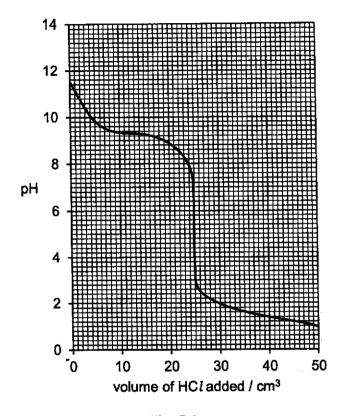


Fig. 5.1

(a) (I) Using the pH from the graph, calculate the concentration of hydroxide ions, in mol dm⁻³, in the 25.0 cm³ of 0.500 mol dm⁻³ solution of NH₃.

$$pOH = 14 - 11.5$$

= 2.5
 $OH^{-} = 10^{-2.5} = 0.00316 \text{ mol dm}^{-3}$

[2]

(ii) Explain how your answer in (a)(i) indicates that NH_3 is a weak base.

The concentration of OH⁻ produced from the dissociation of NH₃ is lower than

the initial concentration of NH₃, hence only partial dissociation has occured.

. [1]

(b)	During the addition of the first 15.0 cm³ of HCL the mixture is behaving as a buffer.								
	(i)	(i) State what is meant by the term buffer solution.							
	A buffer solution is one which is able to resist changes in pH upon addition of								
	a small amount of acid or base.								
	(ii) Explain how the mixture is behaving as a buffer at this stage of the titration. Include an equation in your answer.								
		$NH_3 + H^+ \rightarrow NH_4^{\pm} N$	early all the H	ions added	are removed by the large				
	reservoir of NH ₃ present. Hence, the concentration of H* does not incre								
	significantly, and the pH is kept approximately constant.								
(c)	of th		nation from Fig.	. 5.1 and Table	or to determine the end-point 5.1, explain why the student's tor.				
indicator colour in colour in pH range over which									
		alizarin	yellow	orange	10.1-13.0				
		bromocresol green	yellow	blue	3.8-5.4				
		gentian violet	yellow	violet	0.0–1.6				
		phenolphthalein	colourless	pink	8.2–10.0				

Table 5.1

The region of rapid pH change of this titration occurs below pH 8, hence is not within the working range of phenolphthalein.

A more suitable indicator will be bromocresol green. [2]

[Turn Over

6 (a) The two most common types of nylon used in textile and plastic industries are nylon-6 and nylon-6,6.

For Examiner's Use

The structure of nylon-6,6 is shown below.

(i) Define the term polymer.

A polymer is a macromolecule made from monomers with average molecular mass of at least 1000 or at least 100 repeat units.

(ii) Ropes made of nylon-6,6 are chosen for its high tensile strength. Ropes made from polyesters such as poly(ethylene terephthalate) (PET) are about 90% as strong as that made from nylon-6,6.

With the aid of a diagram, showing relevant interactions, explain why nylon-6,6 is stronger than polyester.

There are stronger hydrogen bonds in between nylon polymer chains as compared to permanent dipole permanent dipole interactions in polyester chain.

(iii) Rust on window grilles can be removed by spraying the grilles with the rust remover shown in Fig 6.1 before scrubbing with a brush.

For Examiner's Use



Fig 6.1

Explain whether a brush with nylon bristles or high density poly(ethene) bristles should be used for the scrubbing.

A brush with high density poly(ethene) bristles should be used.

The acid in the rust remover would cause hydrolysis of the amide linkage in nylon bristles while poly(ethene) bristles are unaffected.

[2]

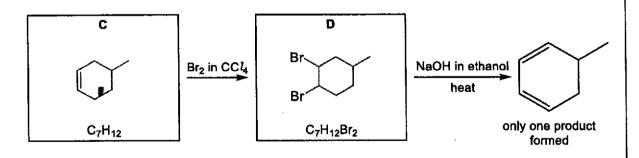
(iv) State whether nylon-6,6 is classified as a thermoplastic or thermoset. Hence, explain whether it can be recycled.

Nylon-6,6 is a thermoplastic. It is recyclable as it melts upon heating as the hydrogen bonds between the chains can be broken.

(b) Suggest a structural formula for each of the compounds A to D in the following schemes.

For Examiner's Use

$$\begin{array}{c|c} & & & \\ & & & \\ \hline & & & \\ & & & \\ \hline & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ \hline & & & \\ & & & \\ & & & \\ \hline & & & \\ & & & \\ & & & \\ \hline & & & \\ & & & \\ & & & \\ \hline & & & \\ & & & \\ & & & \\ & & & \\ \hline & & & \\ & & \\ & & & \\ & & & \\ & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & &$$



[4]

[Total: 12]

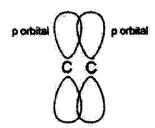
Section B

Answer one question in this section in the space provided.

7 (a) Oleic acid is a monosaturated omega 9 fatty acid found in olive oil. It is said to potentially reduce the risk of coronary heart disease if taken daily.

Oleic acid has the structure shown:

(i) Describe the pi bonding in terms of orbital overlap. You may draw a diagram to illustrate your answer.



The C-C π bond is formed by the sideway overlap between the 2p orbital of

the carbon atom and 2p orbital of another carbon. [2]

(ii) Oleic acid exists as a pair of isomers.

Draw and label the structural formulae of the two isomers.

[2]

(iii) Describe a chemical test that would distinguish oleic acid from ethanoic acid.

Add Br₂ in CCL to each compound in the dark at r.t.p.

Oleic acid: Orange-red bromine solution is decolourised.

Ethanoic acid: Orange-red bromine solution is **not decolourised**. [2]

Turn Over

(b) Fig. 7.1 shows a bomb calorimeter, inside a controlled temperature water jacket, which is used to find an accurate value of the standard enthalpy change of combustion of oleic acid.

For Examiner's Use

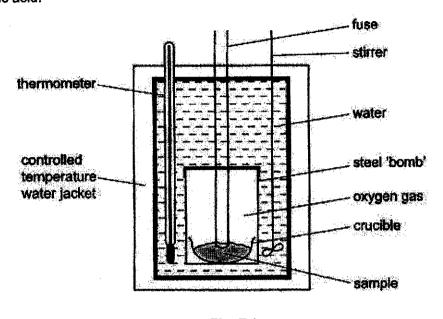


Fig. 7.1

(i) Explain why the steel 'bomb' is flushed with a high pressure of oxygen at the start of the experiment.

It is to ensure that there is **sufficient O₂** present in the steel bomb for **complete**

combustion. [1]

(ii) There is minimal heat lost from the calorimeter because of the controlled temperature water jacket. Suggest how the controlled temperature water jacket achieves this.

The temperature of water in the jacket is raised with an electric heater so that

it just matches the average temperature of the calorimeter. [1]

(iii) A 2.98 g sample of oleic acid (C₁₈H₃₄O₂) is burned in a bomb calorimeter with a heat capacity of 1.98 kJ K⁻¹. The temperature of the calorimeter increases by 59.3 °C. The process is known to be 95 % efficient.

For Examiner's Use

Calculate the standard enthalpy change of combustion of oleic acid.

Heat absorbed by calorimeter = 1.98 × 59.3 = 117 kJ

Amount of oleic acid used =
$$\frac{2.98}{282.0}$$
 = 0.0106 mol

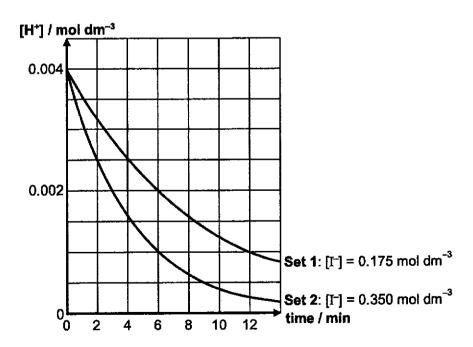
$$\Delta H_c^{\circ}$$
(oleic acid) = $-\frac{117 \times (\frac{100}{95})}{0.0106}$
= $-11700 \text{ kJ mol}^{-1}$

[3]

(c) The Harcourt and Esson reaction is a reaction involving hydrogen peroxide and acidified potassium iodide.

$$H_2O_2 + 2I^- + 2H^+ \longrightarrow 2H_2O + I_2$$

To determine the order of reaction of each reactant, two sets of reaction mixtures containing varying concentrations of I⁻ were prepared. The concentration of hydrogen peroxide used for both experiments is 0.200 mol dm⁻³. The results are as follows.



(i) Given that the initial reaction rate for Set 2 is 1.00 × 10⁻³ mol dm⁻³ min⁻¹, calculate the initial rate of reaction for **Set 1**. Show your working clearly.

For Examiner's Use

Taking gradient of tangent at t = 0,

set 1 =
$$0.004 \div 8$$

= 5×10^{-4} mol dm⁻³ min⁻¹

[1]

(ii) Use the information given above to determine the order of reaction with respect to I⁻ and H⁺. Show your reasoning clearly.

When $[\dot{H}^+]$ decreases from 0.004 mol dm⁻³ to 0.002 mol dm⁻³, time taken is 3 min When $[H^+]$ decreases from 0.002 mol dm⁻³ to 0.001 mol dm⁻³, time taken is 3 min Since t_{32} is constant at 3 min, order of reaction wrt H^+ is $\underline{1}$.

	[H ₂ O ₂] / mol dm ⁻³	[l ⁻] / mol dm ⁻³	[H ⁺] / mol dm ⁻³	initial rate / mol dm ⁻³ min ⁻¹
Set 1	0.200	0.175	0.004	0.0005
Set 2	0.200	0.350	0.004	0.001

Comparing Set 1 and Set 2, when concentration of $[I^-]$ is doubled, the rate is also <u>doubled</u>. Since rate is directly proportional to $[I^-]$, order of reaction wrt $[I^-]$ is $\underline{1}$.

OR Deduce constant half life to deduce order of reaction to I-

[2]

(iii) The actual order of reaction with respect to H₂O₂ is one.

The rate of the reaction was measured as 4.4×10^{-5} mol dm⁻³ s⁻¹ when $[H_2O_2] = 0.002$ mol dm⁻³, $[H^+] = 0.2$ mol dm⁻³ and $[I^-] = 0.2$ mol dm⁻³

Determine the rate constant for this reaction and state its units

rate =
$$k [H_2O_2] [l^-] [H^+]$$

4.4 × 10⁻⁵ = $k(0.002)(0.2)(0.2)$
 $k = 0.550 \text{ mol}^{-2} \text{ dm}^6 \text{ s}^{-1}$

[2]

	Hydrogen iodide can be formed by the reaction between H2 and I2 under stro	ngly
	heated platinised asbestos. It exists as a homogeneous equilibrium system.	

$$H_2(g) + I_2(g) \rightleftharpoons 2HI(g) \cdots (1)$$

(i) Give the expression of the equilibrium constant, K_c of reaction (1).

$$K_c = \frac{[HI]^2}{[H_2][I_2]}$$

[1]

(ii) Calculate the value of K_c at 445 °C, if the equilibrium concentrations in mol dm⁻³ of $H_2(g)$, $I_2(g)$ and $H_2(g)$ are 2.06, 13.40 and 36.98 respectively.

$$K_c = \frac{36.98^2}{2.06 \times 13.40} = 49.5$$

[1]

(ii) The value for the equilibrium constant, K_c for reaction (1) at 1000 °C is 13. By comparing the values of K_c at 445 °C and 1000 °C, deduce whether the reaction between hydrogen and iodine is endothermic or exothermic.

K_c at 1000 °C is less than K_c at 445 °C.

At a higher temperature, there is a smaller proportion of HI at equilibrium.

This implies that backward reaction is favoured to absorb the heat.

Hence, the forward reaction is exothermic.

_____[2]

[Total: 20]

8 (a) When 1 g of ethanol was burned under a container of water, it was found that 100 g of water was heated from 20 °C to 60 °C. The process was known to be only 65 % efficient.

For Examiner's Use

Use these data and values from the *Data Booklet* to calculate the enthalpy change of combustion of ethanol.

$$(CH_3CH_2OH + 3O_2 \longrightarrow 2CO_2 + 3H_2O)$$

Heat absorbed by water (65% efficient) = $mc\Delta T$ = (100)(4.18)(60 - 20) = 16720 J

Heat evolved by combustion of ethanol (100%) = $\frac{16720}{0.65}$ = 25.72 kJ

Enthalpy change of combustion of ethanol = $-\frac{25.72}{\frac{1}{46}}$ = $-\frac{1180 \text{ kJ mo}}{1}$

[3]

(b) Fig 8.1 shows an energy cycle involving ethanol.

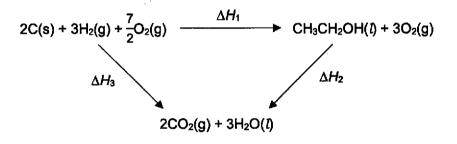


Fig 8.1

(i) Name the enthalpy change represented by ΔH_1 .

(standard) enthalpy change of formation of ethanol

(ii) Given that the enthalpy change of combustion of carbon and hydrogen are $-393.5 \text{ kJ mol}^{-1}$ and $-285.8 \text{ kJ mol}^{-1}$ respectively, use the energy cycle given above and your answer to (a) to calculate ΔH_1 .

$$\Delta H_1 = \Delta H_3 - \Delta H_2$$

= [2(-393.5) + 3(-285.8)] - (-1180)
= -1644.4 + 1330
= -464 kJ mol⁻¹

[3]

(c) A series of reactions starting from 3-chloropropan-1-ol is shown.

For Examiner's Use

C1 OH
$$\stackrel{\text{I}}{\longrightarrow}$$
 A $\stackrel{\text{II}}{\longrightarrow}$ HO $\stackrel{\text{conc. H}_2SO_4}{\longrightarrow}$ C₆H₈O₄

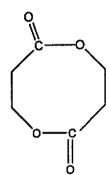
(i) State the reagents and conditions required for reactions I and II.

Step I: KMnO₄(aq) or K₂Cr₂O₇(aq), H₂SO₄(aq) heat under reflux [1]

Step II: NaOH(aq), heat under reflux, followed by addition of H₂SO₄ (aq) [1]

(ii) Draw the structural formulae of A and B.

CI COOH



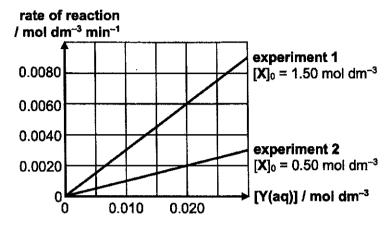
[2]

(d) The reaction between X and Y can be represented by the following equation:

For Examiner's Use

$$X(aq) + 2Y(aq) \longrightarrow Z_1(aq) + Z_2(g)$$

To study the kinetics of the above reaction, two experiments were carried out in the presence of excess **X**. The experimental results obtained are graphically represented below.



Note: $[X]_0$ = initial concentration of X(aq)

(i) With the aid of the graph, determine the order of reaction with respect to each of the two reactants, X and Y.

Consider either experiment 1 or experiment 2.

The graph obtained is a straight line having a positive gradient and passing through the origin. This implies that the rate is directly proportional to [Y]. Hence order of reaction with respect to Y is 1.

Consider the two rates of reaction when [Y] = $0.020 \text{ mol dm}^{-3}$. For expt. 1 where [X]₀ = 1.50 mol dm^{-3} , rate = $0.0060 \text{ mol dm}^{-3} \text{ min}^{-1}$.

For expt. 2 where $[X]_0 = 0.50 \text{ mol dm}^{-3}$, rate = 0.0020 mol dm⁻³ min⁻¹.

Hence when [X] is tripled, rate is corresponding tripled.

This implies that rate is directly proportional to [X]. Therefore order of reaction with respect to X is 1.

[2]

(ii) Hence write down the rate equation for the reaction and state the units for the rate constant k.

rate = k[X][Y]

units for rate constant, k: mol-1 dm3 min-1

[2]

(e) 8.00 g of solid potassium chromate(VI), K₂CrO₄, is dissolved in acid to make a 100 cm³ solution. A dynamic equilibrium occurs in the mixture according to the equation below:

For Examiner's Use

$$2CrO_4^{2-}(aq) + 2H^{\dagger}(aq) \rightleftharpoons Cr_2O_7^{2-}(aq) + H_2O(l)$$

(i) Write the Kc expression for the above equilibrium mixture.

$$K_c = \frac{[Cr_2O_7^{2-}]}{[CrO_4^{2-}]^2[H^+]^2}$$

[1]

(ii) At pH 5.75, the system reaches equilibrium and **one-fifth** of the original amount of chromate(VI) ions remain. Calculate the concentration of chromate(VI) ions at this pH.

Initial [CrO₄²⁻] =
$$\frac{8}{194.2} \div \frac{100}{1000} = 0.4119 \text{ mol dm}^{-3}$$

[CrO₄²⁻] at eqm = $\frac{1}{5}$ (0.4119) = 0.0824 mol dm⁻³

[1]

(iii) Hence, calculate the value of Kc.

$$[Cr_2O_7^{2-}]$$
 at eqm = $\frac{4}{5}(0.4119) \div 2 = 0.1648 \text{ mol dm}^{-3}$
 $[H^+] = 10^{-5.75} = 1.778 \times 10^{-6} \text{ mol dm}^{-3}$

Substituting into Kc,

$$K_c = \frac{[0.1648]}{[0.08239]^2[1.778 \times 10^{-6}]^2} = 7.67 \times 10^{12} \text{mol}^{-3} \text{dm}^9$$

[3]

[Total: 20]

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