

Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME					
CENTRE NUMBER			CANDIDATE NUMBER		

CHEMISTRY
Paper 4 A Le

9701/43

Paper 4 A Level Structured Questions

October/November 2017

2 hours

Candidates answer on the Question Paper.

Additional Materials: Data Booklet

READ THESE INSTRUCTIONS FIRST

Write your Centre number, candidate number and name 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.

DO **NOT** WRITE IN ANY BARCODES.

Answer all questions.

Electronic calculators may be used.

You may lose marks if you do not show your working or if you do not use appropriate units.

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.



International Examinations

Answer all the questions in the spaces provided.

$${\rm 2NO} \, + \, {\rm Br_2} \, \rightleftharpoons \, {\rm 2NOBr}$$

(a)	Using oxidation numbers, explain why this reaction is a redox reaction.

(b) Nitrosyl bromide contains a trivalent nitrogen atom.

Draw the 'dot-and-cross' diagram for NOBr. Show outer electrons only.

[2]

(c) The rate of the reaction was measured at various concentrations of the two reactants, NO and Br₂, and the following results were obtained.

experiment	[NO]/moldm ⁻³	[Br ₂]/moldm ⁻³	initial rate /moldm ⁻³ s ⁻¹
1	0.03	0.02	3.4×10^{-3}
2	0.03	0.04	6.8 × 10 ⁻³
3	0.09	0.04	6.1 × 10 ⁻²
4	0.12	0.06	to be calculated

The general form of the rate equation for this reaction is as follows.

rate =
$$k[NO]^a[Br_2]^b$$

(i)	What is meant by the term <i>order of reaction</i> with respect to a particular reagent?	
. ,		
		1

	(ii)	Use the data in the table to deduce the values of <i>a</i> and <i>b</i> in the rate equation. Show your reasoning.						
((iii)	Use the data in the table to calculate the initial rate for experiment 4.	. <u>~</u>]					
		initial rate = mol dm ⁻³ s ⁻¹	[1]					
(1	iv)	Use the results of experiment 1 to calculate the rate constant, k , for this reaction. Include the units of k .						
		rate constant, <i>k</i> = units	[2]					
	(v)	By considering the rate equation, explain why the rate decreases with decreasing temperature.	ng					
(d)	The	reaction between X and Y was studied.						
		$2X + Y \rightarrow Z$						
	The	following sequence of steps is a proposed mechanism for the reaction.						
		$\begin{array}{lll} \text{step 1} & 2X \rightarrow V \\ \text{step 2} & V + Y \rightarrow Z \end{array}$						
	The	general form of the rate equation for this reaction is as follows.						
		$rate = k[X]^m[Y]^n$						
	Step	o 1 is the slower step in the mechanism.						
	Dec	Suce the values of m and n in the rate equation.						
	m =	n =						
			[1]					
		[Total: 1	2]					

2 (a) The table lists values of solubility products, $K_{\rm sp}$, of some Group 2 carbonates.

	solubility product in water at 298 K, $K_{\rm sp}/{\rm mol^2dm^{-6}}$		
MgCO ₃	1.0 × 10 ⁻⁵		
CaCO ₃	5.0 × 10 ⁻⁹		
SrCO ₃	1.1 × 10 ⁻¹⁰		

		3100 ₃ 1.1 × 10
		the data in the table to describe the trend in the solubility of the Group 2 carbonates down group.
(b)	(i)	Write an equation to show the equilibrium for the solubility product for ${\rm MgCO_3}$. Include state symbols.
		[1]
	(ii)	With reference to your equation in (i), suggest what is observed when a few cm 3 of concentrated Na $_2$ CO $_3$ (aq) are added to a saturated solution of MgCO $_3$. Explain your answer.
		[2]
(0)	Hod	
(6)	USE	e the data in the table to calculate the solubility of MgCO ₃ in water at 298 K, in g dm ⁻³ .
		solubility of $MgCO_3 = \dots gdm^{-3}$ [2]

(d) (i)	Magnesium nitrate decomposes at a lower temperature than barium nitrate.
	Explain why.
	[2]
(ii)	A sample of barium nitrate was heated strongly until no further change occurred. A white solid was formed.
	Write an equation for the action of heat on barium nitrate.
	[1]
(iii)	When water was added to the white solid produced in (d)(ii) , an alkaline solution was produced. Adding sulfuric acid to this solution produced a white precipitate.
	Write equations to explain these observations.
	[2]
	[Total: 11]

3	(a)	Def	fine the term standard cell potential.					
						[2]		
	(b)	(i)		d electrode potenti			I use to measure the clude the necessary	
							[4]	
		(ii)	The E° for a Pb ²⁺ (aq)/Pb(s) electrode is -0.13 V.					
							f the concentration of ppropriate box in the	
				more negative	no change	less negative		
			Explain	your answer.			1	
							[2]	

(c) Car batteries are made up of rechargeable lead-acid cells. Each cell consists of a negative electrode made of Pb metal and a positive electrode made of PbO₂. The electrolyte is H₂SO₄(aq).

When a lead-acid cell is in use, Pb²⁺ ions are precipitated out as PbSO₄(s) at the negative electrode.

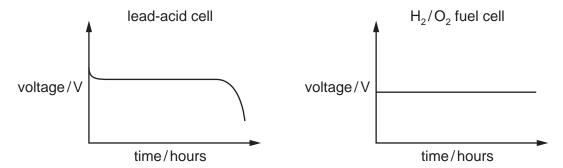
$$Pb(s) + SO_4^{2-}(aq) \rightarrow PbSO_4(s) + 2e^{-}$$

(i) Calculate the mass of Pb that is converted to PbSO₄ when a current of 0.40A is delivered by the cell for 80 minutes.

(ii) Complete the half-equation for the reaction taking place at the positive electrode.

$$PbO_{2}(s) + SO_{4}^{2-}(aq) + \dots + PbSO_{4}(s) + \dots$$
 [1]

(d) The diagrams show how the voltage across two different cells changes with time when each cell is used to provide an electric current.



Suggest a reason why

•	the voltage of the lead-acid cell changes after several hours,
•	the voltage of the fuel cell remains constant.

[Total: 13]

[2]

4 (a	(a)	Des	scribe and explain how the density and melting point of cobalt compare to those of calcium	n.
		den	sity of cobalt	
		ехр	lanation	
		mel	ting point of cobalt	
		ехр	lanation	
				[3]
	(b)	Tra	nsition metals can form complexes.	
		Wh	at is meant by the term transition metal complex?	
				[1]
		4 0		
	(C)	(1)	Cobalt can form the compounds $[Co(NH_3)_5Br]SO_4$ and $[Co(NH_3)_5SO_4]Br$. These two compounds are structural isomers.	
			Define the term <i>structural isomer</i> .	
				[1]
		(ii)	Draw a three-dimensional diagram to show the structure of the ion [Co(NH ₃) ₅ Br] ²⁺ .	
		` ,	Name its shape.	
			[Co(NH ₃) ₅ Br] ²⁺	
			shape	[1]
	((iii)	State the type of bonding between the cobalt ion and NH_3 groups in the $[Co(NH_3)_5Br]^{2+}$ ion $[Co(NH_3)_5Br]^{2+}$ in $[Co(NH_3)_5Br]^{2+}$	n.
				[4]

	(iv)	State the oxidation number of c	obalt in					
		• [Co(NH ₃) ₅ Br] ²⁺		oxidation	number	of Co =		
		• [Co(NH ₃) ₅ SO ₄] ⁺		oxidation	number	of Co =		
								[1]
(d)	each	tions of the compounds [Co(NF other by simple chemical tests ime that any species bonded to	5.					rom
		plete the table with two differ cound. Give the expected obse				to positive	ely identify e	ach
		test		rvation with I ₃) ₅ Br]SO ₄ (a			vation with) ₅ SO ₄]Br(aq)	
			l .					[2]
(e)	The	two compounds [Co(NH ₃) ₅ Br]S0	O ₄ and [Co(NH ₃) ₅ SO ₄]E	Br are dif	ferent colo	urs.	
	Expl	ain why the colours of the two c	compounds	are differer	nt.			
								[2]
(f)	Som	e transition metals and their co	mnounde a	ct as cataly	vete The	catalveie (ean he classi	fiec
(1)		eterogeneous or homogeneous		ct as cataly	/3t3. THE	Catalysis	an be classi	IICC
	Complete the table by placing one tick (\checkmark) in each row to indicate the type of catalysis in each reaction.							
			hetero	ogeneous	homo	geneous		
		Fe in the Haber process						
		Fe ²⁺ in the I ⁻ /S ₂ O ₈ ²⁻ reaction	on					
		NO ₂ in the oxidation of SC	2					
		V ₂ O ₅ in the Contact proces	ss					
			'				1	[2]

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[Total: 14]

Compound **P** contains several functional groups.

$$Cl$$
 CN
 CN

(a)	Name the functional groups present in P .	
		[2]
(b)	Compound P can be polymerised.	
	Draw a section of the polymer of P showing two repeat units. Name the type of polymerisation.	
	type of polymerisation	 [2]

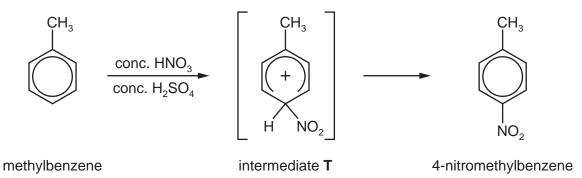
(c) Complete the following table to show the structures of the products formed and the *type of organic reaction* when **P** reacts with the four reagents.

reagent	structure(s) of product(s)	type of organic reaction
excess Br ₂ (aq)		
excess hot, concentrated, acidified MnO ₄ ⁻ (aq)		
excess hot HC1(aq)		
excess H ₂ /Pt catalyst		

[8]

[Total: 12]

6 (a) 4-nitromethylbenzene can be prepared via an electrophilic substitution reaction as shown.



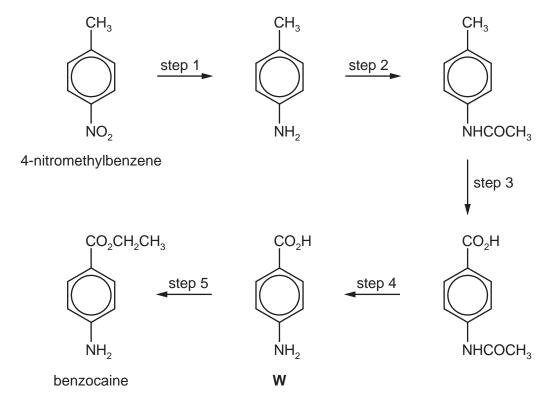
(i) This reaction also forms an isomer of 4-nitromethylbenzene as a by-product.

Draw the structure of this by-product.

(ii)	Write an equation for the reaction between ${\rm HNO_3}$ and ${\rm H_2SO_4}$ that forms the electrophile fo this reaction.
	[1
(iii)	Describe how the structure and bonding of the six-membered ring in intermediate 1 differs from that in methylbenzene.
	[3

[1]

(b) Benzocaine is used as a local anaesthetic. It can be synthesised from 4-nitromethylbenzene by the route shown.



(i) Give the systematic name of compound W.

		ĮΙ
(ii)	Suggest the reagents and conditions for steps 1–5.	

step 1step 2

step 3

step 4

step 5[6]

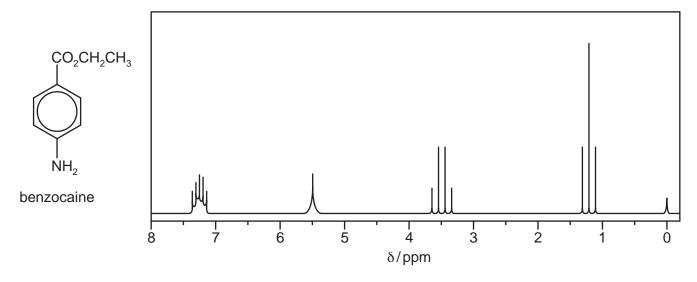
(c) Suggest how the basicity of benzocaine would compare to that of ethylamine.

Explain your answer.

(d)	A sample of benzocaine,	shown below,	was	analysed	by	proton	NMR	and	carbon-13	NMR
	spectroscopy.									

(i) Predict the number of peaks that would be seen in the carbon-13 NMR spectrum.

(ii) Benzocaine was dissolved in ${
m CDC} l_3$ and the proton NMR spectrum of this solution was recorded



Suggest why ${\rm CDC} l_3$ and not ${\rm CHC} l_3$ is used as the solvent when obtaining a proton NMR spectrum.

(iii) Use the *Data Booklet* and the spectrum in **(d)(ii)** to complete the table for the proton NMR spectrum of benzocaine. The actual chemical shifts, δ , for the four absorptions have been added.

δ/ppm	group responsible for the peak	number of ¹ H atoms responsible for the peak	splitting pattern
1.2			
3.5			
5.5			
7.1–7.4			multiplet

(iv) Explain the splitting pattern for the absorption at $\delta 1.2\,\text{ppm}.$

	15	5	
(v)) The proton NMR spectrum of benzocair	ine dissolved in D ₂ O was recorded.	
	Suggest how this spectrum would differ Explain your answer.	er from the spectrum in (d)(ii).	
			[1]
(e) Be	senzocaine can also be used to synthesise	e the dyestuff S by the following route.	
	CO ₂ CH ₂ CH ₃ step 1		
	benzocaine	R	
		step 2 NaOH(aq), OH	

(i) Suggest the reagents used for step 1.

.....[1]

S

(ii) Suggest structures for compounds **R** and **S** and draw them in the boxes. [2]

[Total: 25]

7	(a)	Complete the following electronic structures.

• the iron atom, Fe 1s²2s²2p⁶

• the iron(III) ion, Fe
$$^{3+}$$
 1s 2 2s 2 2p 6

[1]

(b) Solutions of iron(III) salts are acidic due to the equilibrium shown.

$$[Fe(H_2O)_6]^{3+}(aq) \iff [Fe(H_2O)_5(OH)]^{2+}(aq) + H^+(aq) \qquad K_a = 8.9 \times 10^{-4} \, mol \, dm^{-3}$$

Calculate the pH of a $0.25 \, \text{mol dm}^{-3} \, \text{FeC} \, l_3$ solution.

(c) The table shows numerical values of the stability constants for the following equilibrium where M can be one of the metal ions listed and L one of the ligands which replaces one H₂O molecule.

$$[M(H_2O)_6]^{n+}(aq) + L^{-}(aq) \rightleftharpoons [M(H_2O)_5L]^{(n-1)+}(aq) + H_2O(I)$$

metal ion, M	ligand, L	stability constant, K _{stab}
Fe ³⁺	F-	1.0 × 10 ⁶
Fe ³⁺	C1-	2.5×10^{1}
Fe ³⁺	SCN-	9.0×10^{2}
Hg ²⁺	C1-	5.0 × 10 ⁶

)	What is meant by the term <i>stability constant</i> , K_{stab} ?	
		•••
		[1]
		1,1

(ii) Use the data in the table to predict the formula of the complex formed in the greatest amount when

 a solution containing equal concentrations of both F⁻ and SCN⁻ ions is added to Fe³⁺(aq),

• a solution containing equal concentrations of both Fe^{3+} and Hg^{2+} ions is added to $Cl^{-}(aq)$.

Ethanedioate ions, $^{-}O_{2}CCO_{2}^{-}$, are bidentate ligands. The abbreviation ed^{2-} can be used to represent ethanedioate ions.

(d) The complex $[Fe(ed)_2Cl_2]^{3-}$ can be formed according to the equation shown.

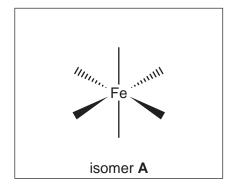
$$[Fe(H_2O)_4Cl_2]^+(aq) + 2ed^{2-}(aq) \rightleftharpoons [Fe(ed)_2Cl_2]^{3-}(aq) + 4H_2O(l)$$

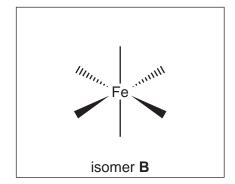
Write the expression for the equilibrium constant, K_{stab} , and state its units.

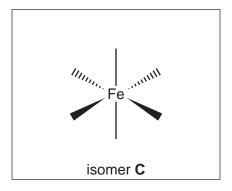
$$K_{\text{stab}} =$$

units[2]

- (e) $[Fe(ed)_2Cl_2]^{3-}$ shows geometrical and optical isomerism.
 - (i) Complete the three-dimensional diagrams to show the three stereoisomers of $[Fe(ed)_2Cl_2]^{3-}$. You may use $_{-O}^{-}$ to represent ed^{2-} .







[3]

(ii) Give the letters of two isomers of $[Fe(ed)_2Cl_2]^{3-}$ which are geometrical isomers of each other.

.....[1]

(iii) Give the letters of the two isomers of $[Fe(ed)_2Cl_2]^{3-}$ which show optical isomerism.

.....[1]

(iv) Give the letter of the isomer which has **no** dipole moment.

......[1]

[Total: 13]

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