

# Our country, our future 525/1

## **S6 CHEMISTRY**

### Exam 11

#### PAPER 1

**DURATION: 2 HOUR 45 MINUTES** 

For Marking guide contact and consultations: Dr. Bbosa Science 0776 802709,

#### Instructions

- This paper consists of two sections A and B
- Section A is compulsory
- Attempt only six questions in section B
- Answers must be written in the spaces provided only.
- Where applicable
   Gas constant, R, = 8.31 JK-1mol-1
   Molar volume at s.t.p is 22.4liters

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1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17

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## SECTION A

1(a)	Define the term "standard heat of reaction."	
<i>a</i> >		
(b)	Given the following data:	
	Enthalpy of formation of magnesium chloride	$= -641.62 \text{kJmol}^{-1}$
	Lattice energy of Magnesium chloride	$= +2495.6 \text{ kJmol}^{-1}$
	Hydration energy of chloride ions	$= -378.0 \text{ kJmol}^{-1}$
	Enthalpy of solution of hydrogen chloride gas	$= -74.8 \text{ kJmol}^{-1}$
	Heat of formation of hydrogen chloride gas	$= -92.32 \text{ kJmol}^{-1}$
	Hydration energy of magnesium ions	$= -1926 \text{ kJmol}^{-1}$
	Calculate the enthalpy of the reaction;	
	$Mg(s) + 2HC1 (a q) \longrightarrow MgCl^2 (aq) + H_2 (g)$	(4 marks)


	gaseous oxide of sulphur $SO_n$ diffused through a porous qual vaolume methane at the same temperature diffused	_
(a) D	etermine the value of n in the oxide SO <sub>n</sub> ,	(2 ½ marks)
• • • • • •		
•••••		
•••••		
•••••		
(b)	The oxide in (a) above reacts with benzene via an elec	trophilic substitution mechanism.
	(i) Stale whether the oxide in (a) above acts as an elect	rophile or nucleophile. Give
	a reason for your answer.	(1½ marks)
	a reason for your answer.	,
	······································	,
	······································	
		and outline the mechanism.
		and outline the mechanism.
		and outline the mechanism.

each of the following pairs he reagent is reacted with ea
(2marks each)
(2 marks)
(2 marks)

(c)	Na <sub>2</sub> C <sub>2</sub> O <sub>4</sub> (aq) and Na <sub>2</sub> SO <sub>3</sub> (aq)	
	Reagent	
	Oobservations.	
••••		
	Soron and nitrogen form chlorides in oxidation state +3	
	Draw the structure and name of the shape adopted by each of the chloride	
(b) E	Explain why the molecules formed by booron in (a) above is non-polar.	(2marks)
(c) V	Write an equation for the reaction of each compound in 9a) above with water.	(2marks)

	••••
	••••
4. The standard redox potentials of Fe $^{3+}$ (aq)/Fe $^{2+}$ and Sn $^{2+}$ (aq)/Sn $^{4+}$ (aq) are -+0.76 and -0.15V respectively.	5 V
(a) Write an equation for the half-cell reaction at the (2 marks)	s)
	••••
	••••
(ii) anode (2mark	s) 
(b) State two conditions under which the electrode potentials above are meas (2marks)	 sured
	••••
	••••
	••••
(c) Deduce if the overall reaction is feasible or not. Give a reason for your answer. (2 ma	ırks)

5. (a)Define the term freezing point depression constant of a solvent.	(1mark)
(b) In an experiment, a 5 per cent solution of glucose , $C_6H_{12}O_6$ in water found give freezing point depression as a 3.3% aqueous solution of $C_nH_{2n}O_n$ .	ve the same
(i) Determine the molecular formula of $C_nH_{2n}O_n$ •	(3 marks)
(ii) The compound in b(i) above forms a crystalline white precipitate with saturable hydrogen sulphite solution but no observable change with ammoniacal silver not lead to reacts with ethanoic acid in the presence of concentrated sulphuric a product with a sweet fruity smell.	itrate solution.
Write the structural formulae and I U PAC name of one of the isomers of C	$C_nH_{2n}O_n$ . (1 mark)
	•••••
6. The first ionization energies of group II metals and the melting point of the	neir

ionization energies of group II metals and the melting points of their chl€l MetalMgCaSr[ First ionisation energy ((KJmol -1 )738590549505 Melting point of chloride CC) 708 772 873 967 Explain. why the ionisation energy decreases from Mg to Ba. (2 (a) marks)

melting points of the chlorides of the metals increase With increase	se in acornž
mbera	(2 marks;
g of ammoniagas were into of water at In a closed sy stern The mixture equilibrium was established and the	e was mainta
uali brium mixture found to contain I .67 <sup>i</sup> 7í34g 01 unionised arnrnonta	a,
Write an equation for the ionisation of ammonia in water.	(1 mark)
Calculate the degree of Ionisation of ammonia in water at 25 °C.	(l mark)
	mbera  The mixture of ammoniagas were into of water at In a closed system. The mixture equilibrium was established and the maintain I .67 i7i34g 01 unionised arnrontain I .67 i7i34g 01 unionised arrontain I .67 i7i34g 01 unionised arrontain I .67 i7i34g 01 unionised arr

above	(1½ ms
	ASSESS PROCESS
basic dissociation constant of methylarnme is 4.38 x 10 <sup>4</sup> mol dill <sup>3</sup> at 2	
(i)Compare the basic strength of ammonia and methylamine.	( '/9. mark
(ii)Give a reason for your answer in c(i) above.	rrla
A solid mixture of manganese (IV) oxide, potassium chlorate (V) and	<del>-</del>
(1) onto, pousseum ontoine (1) une	•
was fused an II-on crucible. Hot water was added to the product to form	
was fused an Il-on crucible. Hot water was added (o the produci to form	(
was fused an II-on crucible. Hot water was added (o the produci to form Name the green solution.	( <sup>-</sup> p, ma
	_
Name the green solution.	
(i)Name the green solution.	
Name the green solution.	
Name the green solution.	

(ii) Explain your observation in b(i) above.	(2 r

## SECTaON g

Attempt any six questions from this section.

20cm of an organic compound Q belonging to C II series were mixed with 200cm ox vgcm. The \_rruxt-urc was sparked and the residual gaseous product absorbed in Lo concentŸated potassium hydroxide solution. The final volume of gas at room 'temperatu re was found to be 30.0cm3.

	minc the molecular formula of Q.	(3 marks)
		******
	eact with bromine water and also decolourises acidified potassium ganate(VII) (Mole ratio $Q:Br_2=1$	
	eact with bromine water and also decolourises acidified potassium ganate(VII) (Mole ratio Q: $Br_2=1$ Identify Q.	( <sup>1</sup> /2 mar- <b>I</b> 0
	ganate(VII) (Mole ratio Q: $Br_2 = 1$	( <sup>1</sup> /2 mar-I
	ganate(VII) (Mole ratio Q: $Br_2 = 1$	
(1)	ganate(VII) (Mole ratio Q: $Br_2 = 1$ Identify Q.	( <sup>1</sup> /2 mar-I0
(1)	ganate(VII) (Mole ratio Q: $Br_2 = 1$ Identify Q.	

(i (ii)	Identify P and R. Suggesl a	ii-ablc mcchanlsyxı for i he reactionlociwccn	(l mark 'l?and hydroxyla
	in Ûhc presence	e of arı acid.	(3 111 ar
İ i İ)	Name the final	product for the reaction in c(ii) above.	mark

(a) The soluibility product Ksp of calcium fluoride can be determined by conductivity rneasurement at [25  $^{0}$  C. The electrolytic conductivities of water amer a salurated

[Define tY1e terrr1 solutbility	prod
L	1
Calculate the solubility -pr	roduci of calcium fluoride at 2 5 °C. State any
assumption s) made.	(5 marks)

solution of ca]cturvn fluoricle arc2 75 x 10  $^4$  ç) cm 554 x 1 0  $^{-5}\Omega^{-1}$  c/respectively. Given

(b)

	O. 1M potassium fluoride solution at 25 ° c	(: 2 mark)
	(11 Explain your answer In b(i) above.	(2 má: ks)
(c)	State one other factor that can affect the solu bility of calcium f	fluoride. ( rnarl<)
$(X = \mathbf{F})$	(VII) elements react with hydrogen gas to form hydrides of the feet, Cli and I).	formula HX
(a)De	Sodium carbonate solution.	(1 <sup>1</sup> /9 marks)
	(ii) Sulphuric acid	marks)

Phe boiling points and bone on the table below.	d dissociation	n encrgics (	of the hydric	des (HX) are
Hydride (HX)	HF	HCl	HBr	HI
Bond energy $((KImol^{-1})$	562	431	366	298
	+19.9	-85.0	-67.0	-35.0
	on In:			
he table below. State and explain the variation	on In: dri(hns.	PIF 1.0		(2 m
he table below.  State and explain the variation  Boiling points of the hy	on In: dri(hns.	PIF 1.0		(2 m
he table below.  State and explain the variation  Boiling points of the hy	on In: dri(hns.	PIF 1.0		(2 m

 $\vee Vrite$  equations to show how the following compounds can bc synthesized.

aIodoetharyc	to	Iodomethane.	(3½ n
	***************************************		***************************************
(b) Phenol	4 -bromobc	nzencsulphonic acid.	(3 marks)
			(21/2
enzoic aci	$\bigcirc c$		(2½ marks)

ain ea	ach of the following observations.	
	ch of the following observations. ethanol are misciblel in all proportions however hexan-l-ol is immi	
		scible with wate (2 <sup>i</sup> 2 marl

a mixture of water and bromobenzene boils below 100 ° c at atmospheric

	esolution added to aqueous nickel (II) sulphate solution a
	moniasolution added to aqueous nickel (I l) sulphate solution, a green soÀüble in excess ammonia to íorrn a blue solution. (3 marks)
[ Ivdr	rooen -peroxide can act as an oxidisinø or reducing agent.
(a)	State the condition(s) under which hydrogen peroxide acts as.  (i) an oxidising agent.  (ii) mar
	(i) all olderoning agoing

Write half reactions to show how hydrogen peroxic	dc acts as a:
• • •	
(i)reducing agent.	(i tnavk
• •	

O)	Write the redox reaction that occurs during	the titration (1 n	nark)
(ii) <b>G</b>	ive a reason why the titration of hydrogen peroxide is do but not a solution of Iron (II) ions. (1 marks)	ne with potassiur	m man <sup>a</sup> anate (V
(iii)	Calculate the volume strength of the hydrogen peroxid $2 [1202 (aq) \longrightarrow 2H_20(l)]$ molar gas	e used in the abovolume at s. t. p	ve experiment.  22400cm <sup>cs</sup> )  (3 5/2 marks
(iii)			22400cm <sup>3</sup> )
(iii)			22400cm <sup>3</sup> )
(iii)			22400cm <sup>3</sup> )
(iii)			22400cm <sup>3</sup> )

In an experiment if) deterrnme the volume strength of Hydrogen peroxide (volume of oxygenges

at s l. liberated by the decomposition of of sob Itian), of I lydrogenperoxi

	roethcne and hcxane-1, 6-dioic acid and hexane-1 6-diaminc are many polymers namely; polyvjnyl chloride and nylon respectively.	nonorners of two
	Define the tervn "Polymer	(1 m
(b)	Write the structural formula of;	
(=)	(i)chloroethcne	( <sup>I</sup> /O mar
	(ii) hexane- I ,6-dioic acid and hexane- 1 ,6-diarnine.	/ > rnar
(c)	Write the structural formula of each of the polymers in (b) above an polymerisation involved.	nd state the tv pc

e)	Name one natural polymer formed by the same reaction as •  (i) Polyvinyl chloride.	(½ mark)
	(ii)nylon, 6, 6.	ma rk)
Ethy	yl bromide reacts with sodium hvdroxide according to the equation $CH_3CH_2Br(l) + NaOH(aq) \longrightarrow CH_3CH_3CH_2OH(aq) + Na/3r(aq)$	
cíhv	Rai e KIC/13C//È/3rl <sup>X</sup> iNaC)//l <sup>Y</sup> where x and y are the respect of the vibrornide and sodium hydroxide •were mixed and refluxed in a fl	tive orders of reactio).02.
cíhv samj	rate equation IS (Yiven by Since Rai e KIC/13C//È/3rl XiNaC)//l Ywherc x and y are the respect	tive orders of reactio).02.
cíhv samp again Time (minu 50751	Rai e KIC/13C//È/3rl <sup>X</sup> iNaC)//l <sup>Y</sup> where x and y are the respect of the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in a fluxed where withdrawn at regular time Intervals and poured in excess of the vibration of the v	tive orders of reactio).02.
cíhv samp again Time (minu 50751	Rai e KIC/13C//È/3rl <sup>X</sup> iNaC)//l <sup>Y</sup> where x and y are the respect of the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in a fluxed where withdrawn at regular time Intervals and poured in excess of the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide •were mixed and poured in excellent the vibrornide and sodium hydroxide and s	tive orders of reactio).02.  ask.20cm ess cold water thentitrate
cíhv samp agair Time (minu 50751 )32.01	Rai e KIC/13C//È/3rl <sup>X</sup> iNaC)//l <sup>Y</sup> where x and y are the respect of the vibrornide and sodium hydroxide •were mixed and refluxed in a fluxed in a fluxed where withdrawn at regular time Intervals and poured in excess to 1.05M hydrochloric acid as shown below.  Intes)o25 1 1 100200 11	tive orders of reactio).02.  ask.20cm ess cold water thentitrate  250 4.0

	Rate =	Cl[2Brl <sup>x</sup> [Na0H			
_					
(i!)	Deduce the va	llue of x in ratc equa	ntion if t_hc ove	rall order of the	reaction IS (l mark)
	_	rate of reaction v		• •	turc if the