

Marking scheme Chemistry Prelims 2022

Paper 1:

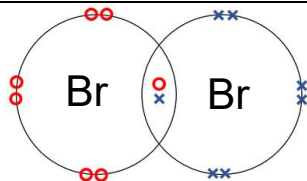
1	2	3	4	5	6	7	8	9	10
D	A	C	A	A	B	A	B	C	A
11	12	13	14	15	16	17	18	19	20
A	B	D	B	C	B	C	D	C	A
21	22	23	24	25	26	27	28	29	30
D	D	C	A	B	C	D	D	B	B
31	32	33	34	35	36	37	38	39	40
C	C	B	B	D	D	A	C	D	C

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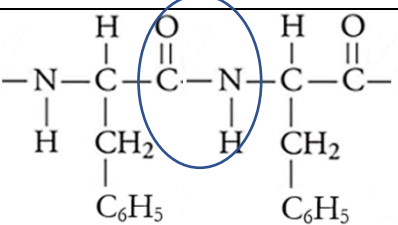
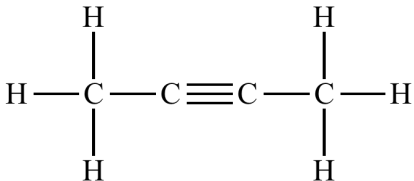
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Paper 2:

A1	(a) Fe (b) Pt (c) Zn (d) C (e) C / N (f) K / Ca	1m each
A2(a)	The molecules can <u>roll and glide over one another</u> and can <u>move throughout the liquid</u> . They are <u>arranged closed together</u> in a <u>disorderly arrangement</u> .	1m 1m
A2(b)		1m
A2(c)	Bromine is <u>more reactive</u> than iodine and gains electrons more readily. Hence it will <u>displace</u> iodine from KI to give KBr and I ₂ . $\text{Br}_2 (\text{l}) + 2 \text{I}^- (\text{aq}) \rightarrow 2 \text{Br}^- (\text{aq}) + \text{I}_2 (\text{s/aq})$	1m 1m 1m
A3(a)	(i) <u>no change</u> /(solution remains pink) (ii) grey solid; forms in green solution	1m 1/2m, 1/2m
A3(b)	$\text{CuO} + 2 \text{E} \rightarrow \text{E}_2\text{O} + \text{Cu}$	1m formulae 1m balanced
A3(c)	Electrolysis	1m
A4(a)	Total amount of energy absorbed during bonding breaking $= (413 \times 8) + (390 \times 6) + (3 \times 498) = 7138 \text{ kJ}$ Total amount of energy released during bonding forming $= (413 \times 2) + (891 \times 2) + (464 \times 12) = 8176 \text{ kJ}$ Total amount of energy released is <u>greater</u> than total amount of energy absorbed OR Enthalpy change of energy is <u>negative</u>	1m 1m 1m
A4(b)	Platinum is a catalyst. It provides an <u>alternative pathway with a lower activation energy</u> . The <u>proportion of particles with sufficient energy to overcome the activation energy increases</u> . As a result, the reacting molecules/ particles (reject ions) collide with each other more often causing the <u>frequency of effective collisions to increase</u> and therefore the rate of reaction.	1m 1m 1m

A4(c)	The reaction is <u>reversible</u> so hydrogen cyanide can decompose back to its reactants as it is formed.	1m
A5(a)	F: ammonium sulfate G: barium sulfate L: calcium carbonate	1m 1m 1m
A5(b)	<u>Blue precipitate formed which dissolves in excess ammonia solution to give a dark blue solution</u>	1m 1m
A6(a)	1. Purple solution turned colourless. 2. Effervescence observed.	1m 1m
A6(b)	<u>The oxidation state of C increases from +3 in $C_2O_4^{2-}$ to +4 in CO_2.</u> <u>$C_2O_4^{2-}$ is oxidised to form CO_2. $C_2O_4^{2-}$ is the reducing agent.</u> <u>OR The oxidation state of Mn decreases from +7 in MnO_4^- to +2 in Mn^{2+}.</u> <u>MnO_4^- is reduced to Mn^{2+}. $C_2O_4^{2-}$ is the reducing agent.</u>	1m 1m
A6(c)	No. of moles of $KMnO_4$ used = $250/1000 \times 0.5 = 0.125$ mol No. of moles of CO_2 produced = $0.125 \times (10/2) = 0.625$ mol Volume of CO_2 produced = $0.625 \times 24 = 15.0$ dm ³	1m 1m 1m
A7(a)	(i) electrons (ii) ions	1m 1m
A7(b)	Apparatus A: <u>A1</u> Apparatus B: <u>B1</u>	1/2m 1/2m
A7(c)	Copper(II) ions (Cu^{2+}) gain electrons more easily than hydrogen ions (H^+) to form copper solid. OR Copper(II) ions (Cu^{2+}) are preferentially discharged to form copper solid. $Cu^{2+} (aq) + 2e \rightarrow Cu (s)$	1m 1m
A7(d)	Electrode A2: $4 OH^- (aq) \rightarrow 2 H_2O (l) + O_2 (g) + 4e$ Electrode B2: $Cu (s) \rightarrow Cu^{2+} (aq) + 2e$	1m 1m
A8(a)	Carboxyl, Amine, Amide, Ester (any two answers)	2m
A8(b)	$ \begin{array}{c} O \\ \\ OH - C - CH_2 - \overset{\text{NH}_2}{\underset{H}{ }{C}} - COOH \end{array} $	1m
A8(ci)	Methyl propanoate	1m
A8(cii)	concentrated sulfuric acid	1m
A8(di)	A macromolecule is a very large molecule that is made up of many small molecules.	1m

A8(dii)		1m 1m circle linkage
B9(a)	Coagulation <u>increases size of particles</u> , allow it to be <u>trapped by the filters/cannot pass through</u> the pores. Coagulation <u>increases mass of particles</u> , hence is <u>heavier and settles to the bottom</u> of the tank for removal.	1m 1m
B9(b)	Filtration removes <u>undissolved, insoluble</u> particles while BAC removes <u>dissolved, soluble</u> particles. Filtration <u>separates by size</u> , while BAC specifically <u>attracts organic compounds</u> and not other compounds.	1m 1m
B9(c)	Calcium oxide is a basic oxide that can neutralise acids in water to increase pH.	1m
B9(d)	Error 1: <u>quote any 2 ions</u> in Table B9.4 which will form precipitate; state they will <u>form precipitate / insoluble salt</u> OR Error 2: <u>Precipitate of lead(II) hydroxide and aluminium hydroxide</u> will react away / <u>dissolve in excess</u> (sodium) hydroxide solution	1m 1m OR 1m 1m
B9(e)	one spot for copper(II) at 4.1 cm one spot for iron(III) at 2.4 cm (-1m for no labelling)	1m 1m
B9(f)	No. The components would have <u>different solubilities</u> in a different solvent, hence the <u>R_f values would be different</u> , resulting in a different chromatogram.	1m 1/2m 1/2m
B10(a)	They have the same functional group <u>carbon-carbon triple bond</u> ; C≡C They have the same general formula <u>C_nH_{2n-2}</u> . The two members differ by a <u>-CH₂</u> unit.	1m each (Any two)
B10(bi)	Any value within the range of 36-43.	1m
B10(bii)	As the <u>molecular sizes increase</u> , there are <u>more intermolecular forces of attraction</u> to overcome. <u>More energy</u> is required to overcome these forces of attraction during boiling.	1m 1m
B10(c)		1m
B10(cii)	<u>Isomers</u> . OR They have the <u>same molecular formula but different structural formulae</u> .	1m

B10(d)	$2 \text{C}_4\text{H}_6 + 11 \text{O}_2 \rightarrow 8 \text{CO}_2 + 6 \text{H}_2\text{O}$	2m
B11E (a)	$\text{CuCO}_3 \rightarrow \text{CuO} + \text{CO}_2$	1m
B11E (b)	Down Group II, the <u>reactivity of the metal increases</u> . The more reactive the metal, the <u>greater the thermal stability of the metal carbonate</u> , hence the <u>temperature for decomposition</u> of the metal carbonate <u>increases</u> .	1m 1m 1m
B11E (ci)	When sulfuric acid is added to CaCO_3 , insoluble CaSO_4 , CO_2 and water is produced. The <u>CO_2</u> produced gives the <u>effervescence</u> . The <u>insoluble CaSO_4</u> formed forms a <u>protective layer</u> over the CaCO_3 , preventing further reaction.	1m 1m
B11E (cii)	No of moles of sulfuric acid = $0.200 \times 25/1000 = 0.00500 \text{ mol}$ No of moles of $\text{MgCO}_3 = 1/[24 + 12 + (16 \times 3)] = 0.0119 \text{ mol}$ Mole ratio $\text{H}_2\text{SO}_4 : \text{MgCO}_3$ is 1:1 0.005 mol of H_2SO_4 requires 0.005 mol of MgCO_3 but there is 0.0119 mol, hence MgCO_3 is in excess. Since there is excess MgCO_3 , there should be solid MgCO_3 left behind and not disappear.	1m 1m 1m for explanation 1m for observation
B11O (ai)	Gas syringe	1m
B11O (aii)	Insert a lighted splint into the gas. The gas will extinguish the lighted splint with a pop sound.	1m 1m
B11O (aiii)	Observation 1 – sulfuric acid is <u>strong</u> acid and hence <u>ionises completely</u> to produce a <u>higher concentration of H^+ ions</u> . Observation 2 – Citric acid is a <u>tribasic</u> acid while sulfuric acid is <u>dibasic</u> . The <u>number of mole of H^+ ions</u> produced is <u>greater</u> with citric acid.	1m 1m
B11O (bi)	Ionic AND Covalent	1m
B11O (bii)	It is formed when the hydrogen ion in citric acid is replaced by the potassium ion.	1m
B11O (biii)	Heat the solution until it is <u>saturated</u> and allow it to <u>cool</u> for crystals to form. <u>Filter</u> to collect crystals as <u>residue</u> . <u>Wash</u> the crystals with a little <u>cold</u> deionised water and <u>dry</u> by pressing between sheets of filter paper.	1m 1m 1m