

Guangyang Secondary School
Chemistry 6092
Secondary Four Express
Preliminary Examination

MARK SCHEME

Paper 1

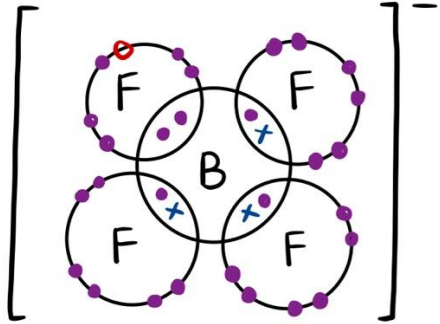
Q1	Q2	Q3	Q4	Q5	Q6	Q7	Q8	Q9	Q10
D	D	B	C	B	D	A	B	C	C
Q11	Q12	Q13	Q14	Q15	Q16	Q17	Q18	Q19	Q20
D	B	A	C	A	A	C	D	B	D
Q21	Q22	Q23	Q24	Q25	Q26	Q27	Q28	Q29	Q30
C	A	C	D	D	D	A	C	A	B
Q31	Q32	Q33	Q34	Q35	Q36	Q37	Q38	Q39	Q40
D	C	B	C	D	A	B	D	D	C

Paper 2

SECTION A

Question	Solutions	Marks	Remarks
1 (a)	barium chloride, zinc sulfate	1	<i>both answer for 1m</i>
(b)	sulfur dioxide	1	
(c)	silver chloride, zinc carbonate	1	<i>both answer for 1m</i>
(d)	ammonium carbonate	1	

Question	Solutions	Marks	Remarks
2 (a) (i)	<i>Labelled arrow of electron highly deflected towards positive plate</i>	1	
(ii)	Electron is negatively charged with a very small / negligible mass	1	<i>both points 1m</i>
(b) (i)	Unlike neutrons, protons are positively-charged and will be repelled by the positively-charged nucleus .	1	<i>both points 1m</i>
(ii)	No change in chemical properties as the new isotope / atom formed has the same electronic configuration (or same number of valence electrons) as the original element . <i>Reject no change in chemical properties as they are isotopes.</i>	1	<i>award marks even if the term isotope is not used</i>

Question	Solutions	Marks	Remarks
3 (a)	Typical: Bonding in the compound is covalent and involves the sharing of electrons which is expected as boron and fluorine are both non-metals .	1	
	Not typical: The valence electron shell of boron only has 6 valence electrons, instead of a complete octet electronic configuration .	1	
(b)	 key: • e⁻ of F atom - x e⁻ of B atom ○ e⁻ of Cs atom 1m for correct electronic structure 1m for square brackets and charge number Accept if donation of electron pair includes electron from caesium atom.	2	deduct 1m if key not included
(c)	Boron trifluoride has a simple molecular structure while caesium tetrafluoroborate has a giant ionic crystal lattice structure. There are weak intermolecular forces of attraction between boron trifluoride molecules, which only require a small amount of energy to overcome. There are strong electrostatic forces of attraction between caesium cations and tetrafluoroborate anions. Thus, a larger amount of energy is required to overcome these forces of attraction within the ionic compound. Hence, boron trifluoride exists as a gas and caesium tetrafluoroborate exists as a solid at rtp.	3	1m: both structures 1m: bonding present in both substances 1m: comparison of energy required

Question	Solutions	Marks	Remarks
4 (a)	$\text{CaCO}_3 + \text{H}_2\text{SO}_4 \rightarrow \text{CaSO}_4 + \text{H}_2\text{O}$ $2\text{KOH} + \text{H}_2\text{SO}_4 \rightarrow \text{K}_2\text{SO}_4 + 2\text{H}_2\text{O}$ $\text{no. of moles of KOH} = \frac{25.00}{1000} \times 0.320$ $= 0.00800 \text{ mol}$ mole ratio of KOH : H₂SO₄ $2 : 1$ $\text{no. of moles of excess H}_2\text{SO}_4 = 0.008 \div 2$ $= 0.00400$ mol $\text{total no. of moles of H}_2\text{SO}_4 = \frac{50.00}{1000} \times 0.200$ $= 0.0100 \text{ mol}$ $\text{no. of moles of H}_2\text{SO}_4 \text{ reacted with CaCO}_3$ $= 0.0100 - 0.00400$ $= 0.00600 \text{ mol}$ mole ratio of H₂SO₄ : CaCO₃ $1 : 1$ $\text{no. of moles of CaCO}_3 = 0.00600 \text{ mol}$ $\text{mass of CaCO}_3 = 0.00600 \times (40 + 12 + 48)$ $= 0.6 \text{ g}$ $\% \text{ purity of CaCO}_3 = \frac{0.6}{0.835} \times 100\%$ $= 71.9 \% (3\text{sf})$	1 1 1 1	1m for excess H₂SO₄ that reacted with KOH 1m for amount of H₂SO₄ that reacted with CaCO₃ 1m for mass of CaCO₃ 1m for percentage purity of CaCO₃
(b) (i)	Point Q was the equivalence point where nitric acid was completely neutralised by sodium hydroxide. At R, sodium hydroxide is in excess. The presence of <u>free moving Na⁺ and OH⁻ ions from excess sodium hydroxide act as mobile charge carriers</u> to conduct electricity. Thus, electrical conductivity of solution increases.	1	To state the presence of ions from EXCESS sodium hydroxide

(ii)	Similar to curve for experiment, except for an initial pH value of 4. Same volume of NaOH used and same height at end of reaction.	1	
(iii)	As ethanoic acid is a <u>weak acid</u>, it ionises partially in water to produce hydrogen ions. Nitric acid, on the other hand, is a <u>strong acid</u> which ionises completely to produce hydrogen ions. Since ethanoic acid has a <u>lower concentration of hydrogen ions</u>, it has a <u>higher pH</u>	1 1	1m for difference in strength of acids 1m pH value to concentration of hydrogen ions

Question	Solutions	Marks	Remarks
5 (a)	Iron(II) ion is a catalyst as it is chemically unchanged in the overall reaction / is regenerated at the end of step 2.	1	
(b) (i)	Aqueous sodium hydroxide could be added to the mixture of iron(II) ions and aqueous iodine, to form insoluble iron(II) hydroxide. Aqueous sodium hydroxide is added in excess until no further change is seen. Filtration can be carried out to remove the solid from the mixture.	1 1	Accept use of aqueous ammonia 1m not awarded if reagent is not added in excess
(ii)	A black solid is formed and the solution turns into a lighter shade of brown. Iodine is more reactive than astatine, and will displace astatine from sodium astatide solution.	1 1	No mark awarded if explanation is incorrect.

Question	Solutions	Marks	Remarks
6 (a) (i)	Z, W, X, Y	1	
(ii)	$3X(s) + 2W(NO_3)_3(aq) \rightarrow 3X(NO_3)_2(aq) + 2W(s)$	1	
(b)	Zinc	1	
(c) (i)	No gas is produced for metal Z. Metal Z, which is less reactive than copper, must be an unreactive metal and does not react with steam.	1	
	X is more reactive than W and will produce more gas than W during the fixed time interval.	1	
(ii)	Fused/anhydrous calcium chloride	1	
(d)	Metal V is below Y in the Periodic Table. Since metal V is more reactive than Y, the electrostatic forces of attraction between the nucleus and valence electrons for V must be weaker.	1	<i>1m on linking reactivity to forces of attraction</i>
	This implies that the valence electron is further from the positively charged nucleus, and V has more electron shells than Y.	1	<i>1m on concluding the no. of electron shells</i>

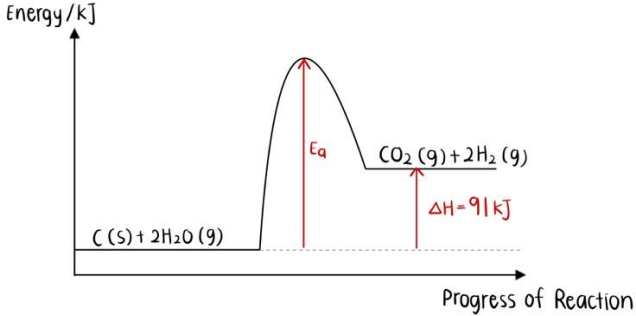
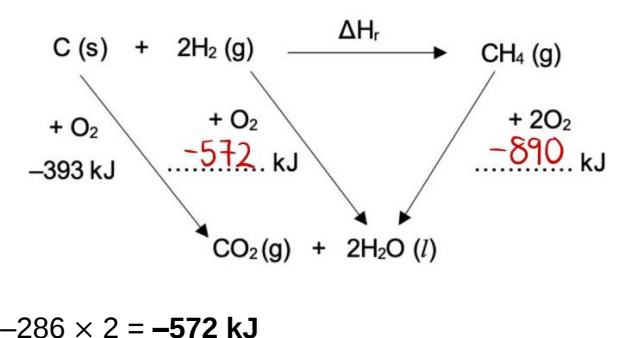
Question	Solutions	Marks	Remarks
7 (a) (i)	Carbon monoxide and oxides of nitrogen react with each other to form nitrogen gas and carbon dioxide through redox reactions .	1	<i>type of reaction and products must be stated clearly for mark to be awarded</i>
	Unburnt hydrocarbons combust / burn / react with oxygen in air to form carbon dioxide and water vapour	1	
(ii)	When the temperature of the car engine is higher, nitrogen gas will react with the oxygen present in the air to form more oxides of nitrogen .	1	
(b)	Ammonia is oxidised as the oxidation state of nitrogen increases from -3 in NH_3 to 0 in N_2	1	
	Nitrogen dioxide is reduced as the oxidation state of nitrogen decreases from $+4$ in NO_2 to 0 in N_2 . Since oxidation and reduction occurs simultaneously , this is a redox reaction.	1	
			<i>Deduct 1m if last sentence on both oxidation and reduction occurring is not stated</i>

Question	Solutions	Marks	Remarks
8 (a)	C_2H_2	1	
(b) (i)	$2C_4H_6 + 11O_2 \rightarrow 8CO_2 + 6H_2O$ <i>accept $C_4H_6 + \frac{11}{2} O_2 \rightarrow 4CO_2 + 3H_2O$ but must be in mixed fractions</i>	1	
(ii)	Add a few drops of aqueous bromine to separate test-tubes of butyne and butane. Butyne: Aqueous bromine decolourises Butane: Aqueous bromine remains brown.	1	<i>Test and results for both butyne and butane must be included for 1m</i>
(iii)	Addition reaction $ \begin{array}{ccccccc} H & & H & & H & & \\ & & & & & & \\ C & = & C & - & C & - & C - H \\ & & & & & & \\ OH & H & H & & H & & \end{array} $ OR $ \begin{array}{ccccccc} H & & H & & H & & \\ & & & & & & \\ C & = & C & - & C & - & C - H \\ & & & & & & \\ H & & OH & & H & & H \end{array} $	1 1	<i>or any structure that makes sense (such as double addition where only single bonds and 2 hydroxyl groups are present)</i>
(c)	$ \begin{array}{ccccccc} & H & & & H & & \\ & & & & & & \\ H & - C & - & C \equiv C & - & C & - H \\ & & & & & & \\ & H & & & H & & \end{array} $	1	
(d)	No, the bromine atoms are not on adjacent carbon atoms and cannot be broken to form an alkyne.	1	

(e)	$ \begin{array}{c} \text{CH}_3 \\ \\ \text{HO}-\text{C}-\text{CH}_2-\text{C} \begin{array}{l} \text{O} \\ \\ \text{OH} \end{array} \\ \\ \text{H} \end{array} $ $ \begin{array}{c} \text{CH}_2\text{CH}_3 \\ \\ \text{HO}-\text{C}-\text{CH}_2-\text{C} \begin{array}{l} \text{O} \\ \\ \text{OH} \end{array} \\ \\ \text{H} \end{array} $	1	
		1	

Question	Solutions	Marks	Remarks
9 (a)	<u>Experiment 1:</u> anode: $4\text{OH}^- (\text{aq}) \rightarrow 2\text{H}_2\text{O} (\text{l}) + \text{O}_2 (\text{g}) + 4\text{e}^-$ cathode: $\text{Ag}^+ (\text{aq}) + \text{e}^- \rightarrow \text{Ag} (\text{s})$ Observations & Explanations: Effervescence observed at the platinum electrode. Colourless, odourless gas evolved relights a glowing splint. Hydroxide ions are preferentially discharged to form water and oxygen gas , due to the greater ease of discharge than nitrate ions. Silver solid deposited on steel electrode. Silver ions are preferentially discharged to form silver solid , due to the greater ease of discharge than hydrogen ions.	1	1m correct half equations at Expt 1
		1	1m correct observations and explanation at Expt 1 anode
		1	1m correct observations and explanation at Expt 1 and 2 cathode
	<u>Experiment 2:</u> anode: $\text{Ag} (\text{s}) \rightarrow \text{Ag}^+ (\text{aq}) + \text{e}^-$ cathode: $\text{Ag}^+ (\text{aq}) + \text{e}^- \rightarrow \text{Ag} (\text{s})$ Observations & Explanations: Silver anode decreases in size and mass. Silver electrode is reactive and is preferentially oxidised to form silver ions, over hydroxide and nitrate ions. (Similar to Expt 1) Silver solid deposited on steel electrode. Silver ions are preferentially discharged to form silver solid, due to the greater ease of discharge than hydrogen ions.	1	1m correct half equations at Expt 2
	<u>The runtime for experiment 2 is longer</u> as the silver ions in the electrolyte are constantly replenished by the oxidation of the silver electrode , unlike experiment 1 which will end once all the silver ions in the electrode is reduced.	1	1m difference in run times

(b)	The steel electrode will decrease in mass and become smaller.	1	
	Iron is more reactive than silver and will be preferentially oxidised to form iron ions in the simple cell.	1	

Question	Solutions	Marks	Remarks
10 (a)	 1m for correct diagram (endothermic) with correct reactants and products, and enthalpy change 1m for presentation including labelling of axes, and activation energy	2	
(b) (i)	 $-286 \times 2 = -572 \text{ kJ}$	1 1	1m for each value
(ii)	By Hess Law, $\Delta H_r + (-890) = -393 + (-572)$ $\Delta H_r = -393 + (-572) - (-890)$ $= -75 \text{ kJ}$	1 1	1m for arranging enthalpy changes correctly (CW=ACW)

(c) (i)		substance	sample	1	
		methane	2		
		ethanol	1		
		butanol	3		
(ii)	The higher the temperature rise, the greater the enthalpy change of combustion of substance. Butanol, with the highest enthalpy change of combustion, will result in the highest temperature rise, followed by ethanol and methane. In the combustion of all three fuels, the energy released from forming bonds in carbon dioxide and water (C=O and O-H) is more than the energy absorbed to break the C-H and O=O bonds present in the molecules. Since butanol contains the largest number of C and H atoms, most number of C=O and O-H bonds are formed and most amount of energy is released from the reaction.			1	1m for linking temperature rise to enthalpy change
				1	1m for comparing energy absorbed and energy released
				1	1m for stating that butanol is the largest molecule and will result in the most energy released from most number of products formed.
(iii)	-2670 kJ/mol			1	proportional to temperature rise, compared to methane and ethanol
(d)	The black powder is soot which is formed from the incomplete combustion of butanol, due to limited amount of oxygen gas.			1	both points 1m

[illegible]

(d)	At lower temperature, the initial rate of reaction will decrease. At a lower temperature, the particles have less kinetic energy. They move slower and collide more less frequently. In addition, there are fewer reacting particles with energy greater than or equal to the activation energy. Hence, there is a decrease in the frequency of effective collisions, resulting in a slower reaction. <i>deduct 1m if frequency of effective collision is not seen</i>	1 1 1	
(e)	No gas will be produced as silver is unreactive and does not react with acid.	1	

Question	Solutions	Marks	Remarks
12 (a)	No, the concentration ClO_2 of has a greater effect on the rate of reaction than the concentration of OH^- ions .	1	
	In experiment 1 and 2, when the concentration of ClO_2 doubles from 0.020 to 0.040 mol/dm ³ , the initial rate of reaction increased by four times from 0.00358 to 0.01432 mol/dm ³ s.	1	
	In experiment 1 and 3, when the concentration of OH^- doubles 0.015 to 0.030 mol/dm ³ , the initial rate of reaction only increased by two times from 0.00358 to 0.0716 mol/dm ³ s.	1	
	Hence, the effect of concentration of each reactant is not equal.		
(b)	$0.00358 \times 9 \times 5 = \mathbf{0.1611 \text{ mol/dm}^3\text{s}}$	1	
(c)	When catalyst is present, it provides an alternative pathway with a lower activation energy for the reaction to occur.	1	
	Hence, more reactant particles have energy greater than or equal to activation energy .	1	
	Hence, there is an increase in the frequency of effective collisions , resulting in a faster reaction.	1	
(d)	mole ratio of $\text{ClO}_2 : \text{OH}^-$ 1 : 1		
	Since the same volume of both solutions is used, ClO_2 , with a lower concentration, is the limiting reactant.		
	no. of moles of $\text{ClO}_2 = 0.020 \times \frac{50}{1000}$ = 0.00100 mol	1	
	mole ratio of $\text{ClO}_2 : \text{H}_2\text{O}$ 2 : 1		
	no. of moles of $\text{H}_2\text{O} = 0.00100 \div 2$ = 0.0005 mol	1	
	mass of $\text{H}_2\text{O} = 0.0005 \times 18$ = 0.009 g	1	