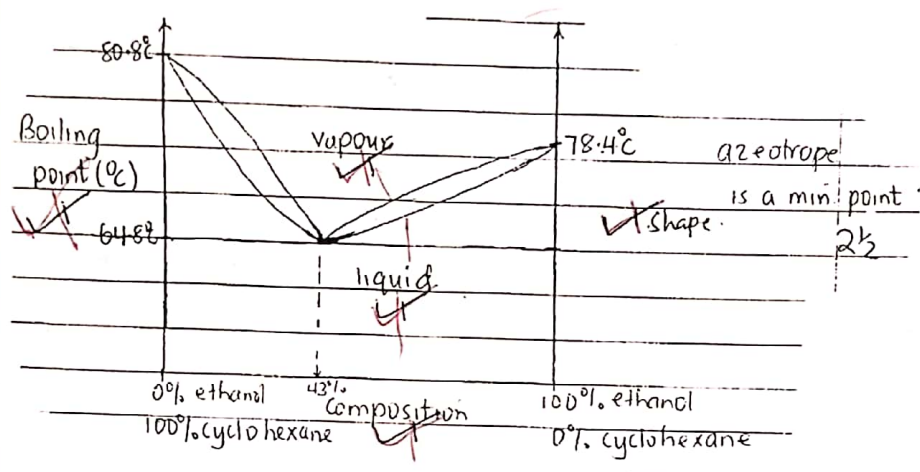
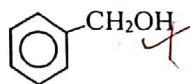
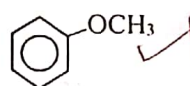
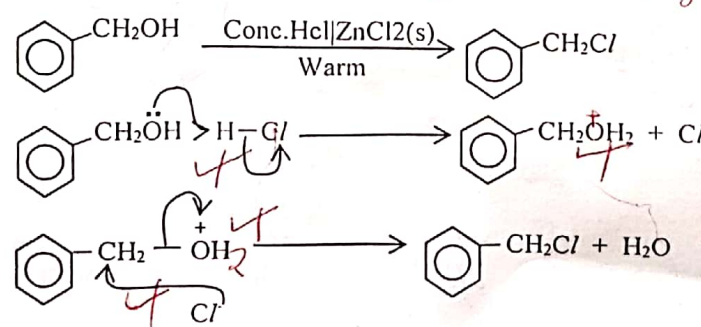


16.(a)(i)	Chlorine - 35	01
(ii)	$R.A.M = \frac{\sum \text{Isotopic mass} \times \text{abundance}}{100}$ $= \frac{35 \times 80 + 37 \times 20}{100}$ $= 35.4$	02
(b)	<p>Chlorine has two Isotopes ; Chlorine - 35 and Chlorine - 37. The diatomic gaseous molecule is formed by different isotopic combination to give molecular ions of mass 70, 72 and 74</p> $^{35}\text{Cl} + ^{35}\text{Cl} \rightarrow ^{70}\text{Cl}_2^+$ $^{35}\text{Cl} + ^{37}\text{Cl} \rightarrow ^{72}\text{Cl}_2^+$ $^{37}\text{Cl} + ^{37}\text{Cl} \rightarrow ^{74}\text{Cl}_2^+$ <p>Each molecular ion forms a peak in the mass spectrum</p>	04
(c)	<p>It is accurate A small amount of sample is used.</p>	02
		09 marks
17(a)	Graph paper.	
(i)	Freezing point of pure water = 0.00°C . (Intercept from graph)	01 mark
(ii)	<p>Slope = $\Delta T / \Delta M$</p> $= \frac{-0.05 + 0.80}{2.50 - 37.50}$ $= -0.0214286^\circ\text{Cg}^{-1}$ <p>Slope = $\frac{-100K_f}{\text{mass of solvent} \times M_x}$</p> <p>$-0.0214286 = \frac{-100 \times 1.86}{250 \times M_x}$</p> <p>$M_x = 347.20\text{g/mol.}$</p>	2½
(b)(i)	<p>Ionic compounds dissociate in water into ions. The number of non-volatile solute particles in solution increase thus increasing the freezing point depression. Since molar mass is inversely proportional freezing point depression, the observed value of molar mass is less than expected.</p>	1½
(ii)	<p>Hexane is a non polar solvent in which short chain carboxylic acids dimerise/associate through hydrogen bonding. This halves/ reduces the number of non-volatile solute particles doubling/ increase the observed value of molar mass</p>	1½

END

(ii)	<p>Positive deviation. ✓</p> <p>Addition of cyclohexane to ethanol weakens the hydrogen bonds between ethanol molecules. The escaping tendency of the molecules in solution increases thus the vapour pressure above the solution is greater than that expected for ideal solutions. At lower temperatures, the vapour pressure above solution is equal to the atmospheric pressure. ✓</p> <p><i>and the soln boils at lower temp</i></p>	2½
(b)(i)		2½
(ii)	<p>Distillate – azeotropic mixture (liquid mixture 43% ethanol)</p> <p>Residue – pure ethanol.</p>	02
(c)	<p>Solvent extraction ✓</p> <p>Distillation with a third component</p> <p><i>dilute HNO₃ / HCl followed by NaOH</i></p>	01
15.(a)(i)	<p>BaCO_3 - green precipitate insoluble in excess alkali.</p> <p>BaCO_3 - White precipitate insoluble in excess alkali.</p>	09 marks
(ii)	<p>Silver nitrate solution followed by dilute nitric acid KI – Yellow precipitate soluble in nitric acid.</p> <p>NaBr – Cream/pale yellow precipitate sparingly soluble in nitric acid.</p>	02
(b)(i)	<p>Iodide ions reduce copper (ii) ions to sparingly soluble copper (i) iodide and themselves oxidized to Iodine.</p> <p>$2\text{Cu}^{+}_{(aq)} + 4\text{I}^{-}_{(aq)} \rightarrow \text{Cu}_2\text{I}_{2(s)} + \text{I}_{2(aq)}$ ✓</p>	02½
(ii)	<p>Hydrogen sulphide partially ionizes in water or eqn.</p> <p>$\text{H}_2\text{S}_{(g)} + aq \rightleftharpoons 2\text{H}^{+}_{(aq)} + \text{S}^{2-}_{(aq)}$ ✓ <i>$\text{H}_2\text{S} + \text{H}_2\text{O} \rightleftharpoons 2\text{H}_3\text{O}^{+} + \text{S}^{2-}$</i></p> <p>In acidic medium, the high concentration of hydrogen ions suppresses the ionization of hydrogensulphide. Reducing the concentration of sulphide ions.</p> <p>The ionic product of sulphide ions and Nickel(ii) ions does not exceed the solubility product of Nickel(ii) Sulphide hence no precipitation.</p>	02½
09 marks		

7. (a)	Acid strength increases with increase in oxidation state of chlorine ($\text{HOCl} \angle \text{HClO}_2 \angle \text{HClO}_3 \angle \text{HClO}_4$).	01
(b)	Oxygen is more electronegative than chlorine hence has a negative inductive effect whose magnitude increase with number of oxygen atoms per acid molecule. The oxygen-hydrogen bond in each oxo acid becomes much weaker as oxidation state increases from chloric(i) to chloric (vii) acid and thus breaks more readily to release higher concentrations of hydrogen ions into solution.	03
(c)	(Pale) green solution turns brown. <i>accept brown solution</i>	$\frac{1}{2}$
(i)	$\text{ClO}_4^- + 8\text{H}^+ + 8\text{Fe}^{2+} \rightarrow \text{Cl}^- + 8\text{Fe}^{3+} + 4\text{H}_2\text{O}_{(l)}$ <i>(ignore states)</i>	01 $\frac{1}{2}$
(ii)		06 marks
8. (a)	Observation Yellow oily liquid (yellow oil) $(\text{CH}_3\text{CH}_2)_2\text{NH} \xrightarrow[\text{O}^\circ\text{C}]{\text{NaNO}_2(\text{aq}) / \text{Conc. HCl}} (\text{CH}_3\text{CH}_2)_2\text{N}-\text{N}=\text{O}$	$1\frac{1}{2}$
(b)	Observation White precipitate <i>(Dont PPT or solid). Don't penalize if NaCl is absent</i> $\text{C}_6\text{H}_5\text{COONa}_{(\text{aq})} + \text{HCl}_{(\text{aq})} \rightarrow \text{C}_6\text{H}_5\text{COOH}_{(\text{s})} + \text{NaCl}_{(\text{aq})}$ <i>- 1/2 for wrong state</i>	$1\frac{1}{2}$
(c)	Observation Orange red precipitate <i>(ignore)</i> $\text{C}_6\text{H}_5\text{N} \equiv \text{NCl}^- + \text{C}_6\text{H}_5\text{OH} \rightarrow \text{C}_6\text{H}_5\text{N}=\text{N}-\text{C}_6\text{H}_4\text{OH}$ <i>(pen 4)</i>	02
		05 marks
9. (a) (i)	$\text{SnCl}_2, \text{SnCl}_4, \text{CCl}_4$	$1\frac{1}{2}$
(ii)	$\text{SnCl}_{2(\text{s})} + 2\text{H}_2\text{O}_{(l)} \rightleftharpoons \text{Sn}(\text{OH})\text{Cl}_{(\text{s})} + \text{HCl}_{(\text{aq})}$ or <i>(ignore states)</i> $\text{SnCl}_{4(\text{l})} + \text{H}_2\text{O}_{(l)} \rightarrow \text{SnO}_{2(\text{s})} + 4\text{HCl}_{(\text{aq})}$ <i>ignore states</i>	01 01 01
(b) (i)	The orange solution turns green	$\frac{1}{2}$
(ii)	$3\text{Sn}^{2+}(\text{aq}) + \text{Cr}_2\text{O}_7^{2-}(\text{aq}) + 14\text{H}^+(\text{aq}) \rightarrow 3\text{Sn}^{4+}(\text{aq}) + 2\text{Cr}^{3+}(\text{aq}) + 7\text{H}_2\text{O}_{(l)}$	01 $\frac{1}{2}$
		05 $\frac{1}{2}$
SECTION B (54 MARKS)		
10. (a)	The ratio of distance between two parallel (identical) electrodes to their cross-section area. OR Cell constant = $\frac{l}{A}$ where; l = distance between parallel electrodes. A = cross-sectional area of identical electrodes.	01
(b) (i)	Cell constant = $\frac{l}{A}$ $= 0.12 / 1.25664 \times 10^{-3}$ $= 95.493 \text{ m}^{-1}$	01
(ii)	Electrolytic conductivity, $K^1 = \frac{1}{R} \times \frac{l}{A}$	

	$= \frac{1}{93.92} \times 95.493 = 1.01675 \Omega^{-1} m^{-1}$ $M_{\text{slow}} = \frac{K'}{C} = \frac{1.01675}{0.05 \times 1000}$ $0.02033 \Omega^{-1} m^{-2} mol^{-1} \text{ (with units)}$	03																				
(c) (i)	Barium chloride has higher conductivity than Magnesium chloride.	01																				
(ii)	The barium ion (Ba^{2+}) has a larger ionic radius hence a lower charge density than magnesium ion (Mg^{2+}). In solution, the Ba^{2+} ions is less hydrated thus has higher ionic mobility than the more hydrated Mg^{2+} ion.	03																				
		09 marks																				
11.(a)(i)	<table><tr><td>C</td><td>:</td><td>H</td><td>:</td><td>O</td></tr><tr><td>$\frac{77.78}{12}$</td><td>:</td><td>$\frac{7.41}{1}$</td><td>:</td><td>$\frac{14.81}{16}$</td></tr><tr><td>$\frac{6.481}{0.926}$</td><td>:</td><td>$\frac{7.410}{0.926}$</td><td>:</td><td>$\frac{0.926}{0.926}$</td></tr><tr><td>7</td><td>:</td><td>8</td><td>:</td><td>1</td></tr></table> <p style="text-align: center;">C_7H_8O</p>	C	:	H	:	O	$\frac{77.78}{12}$:	$\frac{7.41}{1}$:	$\frac{14.81}{16}$	$\frac{6.481}{0.926}$:	$\frac{7.410}{0.926}$:	$\frac{0.926}{0.926}$	7	:	8	:	1	02
C	:	H	:	O																		
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7	:	8	:	1																		
(ii)	$\frac{P H_2O}{P Q} = \frac{\text{Number of water in distilled}}{\text{Number of moles of Q in distilled}}$ $\frac{639}{760-639} = \frac{(100-53.18)/18}{53.18/MQ}$ <p>MQ = 108 (12 x 7 + 8 x 1 + 16)n = 108. n = 1 Q - C_7H_8O</p>	03																				
(b)(i)	 Phenylmethanol.	01																				
(ii)		$\frac{1}{2}$																				
(c) (i)	cloudy solution. Rej. two layers	$\frac{1}{2}$																				
(ii)	 <p>Accept water loss</p>	02																				
		09 marks																				

12.(a)	<table border="1"> <tr> <td>Na</td><td>Al</td><td>P</td></tr> <tr> <td>NaCl ✓</td><td>AlCl₃ ✓</td><td>PCl₃ or PCl₅ ✓</td></tr> <tr> <td>Ionic ✓</td><td>Covalent ✓</td><td>Covalent ✓</td></tr> </table>	Na	Al	P	NaCl ✓	AlCl ₃ ✓	PCl ₃ or PCl ₅ ✓	Ionic ✓	Covalent ✓	Covalent ✓	03½
Na	Al	P									
NaCl ✓	AlCl ₃ ✓	PCl ₃ or PCl ₅ ✓									
Ionic ✓	Covalent ✓	Covalent ✓									
b)	<p>NaCl > AlCl₃ > PCl₃ > PCl₅ ignore ✓ I ignore ✓ even student gives PCl₃ ✓</p> <p>NaCl has a giant ionic structure with strong electrostatic forces between sodium ions and Chloride ions which require a lot of heat energy to break.</p> <p>AlCl₃ is predominantly covalent due to high charge density of Aluminum ions which polarize the large Chloride ions thus reducing charge separation. Its molecules are held by weak intermolecular forces. PCl₃ and PCl₅ molecules are discrete and held by Van der Waals forces whose magnitude increase with molecular mass.</p>	2½									
(c)	<p>NaCl – dissociates in water into its free ions.</p> <p>$NaCl_{(aq)} \rightarrow Na^+_{(aq)} + Cl^-_{(aq)}$ ✓</p> <p>AlCl₃, PCl₃ and PCl₅ undergo hydrolysis to form Aluminium hydroxide, Phosphorous acid and phosphoric acid respectively with evolution of hydrogen chloride gas</p> <p>$AlCl_{3(s)} + 3H_2O_{(l)} \rightarrow Al(OH)_{3(s)} + 3HCl_{(g)}$ ✓</p> <p>$PCl_{3(l)} + 3H_2O_{(l)} \rightarrow H_3PO_3 + 3HCl_{(g)}$ ✓ or one of the Chlorides of P</p> <p>$PCl_{5(s)} + 4H_2O_{(l)} \rightarrow H_3PO_4 + 5HCl_{(g)}$ ✓</p>	03 02½									
13.(a)(i)	<p>CH₃CH = CCl₂ to CH₃CH = CHO.</p> <p>NaOH(aq) Reflux</p> <p>CH₃CH = CHO</p> <p>1. NaBH₄ 2. H₂O</p> <p>Alkene ✓</p>	09 marks									
(ii)	<p>CH₃C ≡ CCH₃ to CH₃C(OH) = COOH</p> <p>60°C H₂O/Conc. H₂SO₄ H₂SO₄ ✓</p> <p>CH₃CCH₂CH₃ ✓</p> <p>HCN or KCN/H⁺ or HCN/OH⁻ ✓</p> <p>CH₃C(OH)CN ✓</p> <p>05</p>	03 05									
(b).	<p>Ethanoic acid is reduced to ethanol by Lithium Aluminium tetrahydride in dry ether. Ethanol is mixed with excess concentrated sulphuric and heated to 180°C. it is dehydrated to ethane which is then reacted with bromine water to form 2 – bromoethanol.</p>	03 09 marks									
14 (a)(i)	<p><u>Azeotrope.</u></p> <p>Uniform mixture of two or more miscible liquids which boils at constant temperature to form a vapour of the same composition at a given pressure.</p> <p>or mixture that distills without change in composition at constant pressure ✓</p>	01									

4(a)(i)	Hydrogen bond or intermolecular hydrogen bonds	½
(ii)	R.F.M of HCOOH = 46 Observed R.F.M = 46X2 = 92	½
(b)	$\frac{\Delta P}{P^0} = \frac{\text{moles of solute}}{\text{moles of solvent}}$ $\frac{\Delta P}{122} = \frac{0.092/92}{156/78}$ $\Delta P = \frac{0.001 \times 122}{2} = 0.061 \text{ mmHg.}$ <p>Vapour pressure of solution = 122 - 0.061 = 121.939 mmHg.</p>	2½
(c)	The vapour pressure of solution is lower than the vapour pressure pure benzene. The non-volatile methanoic acid particles occupy the surface of the solvent reducing the escaping tendency of volatile benzene molecules into vapour phase	1½
		05 marks
5.(a)(i)	$\text{pH} = -\log_{10} [H_3O^+]$ $[H_3O^+] = 10^{-6.24}$ $= 5.7544 \times 10^{-7} \text{ mol dm}^{-3}$	1½
(ii)	$K_h = \frac{[H_3O^+][NH_3]}{[NH_4^+]}$ $\frac{1.0 \times 10^{-14}}{1.78 \times 10^{-5}} = \frac{(5.7544 \times 10^{-7})^2}{[NH_4^+]}$ $[NH_4^+] = 5.894134 \times 10^{-4} \text{ M.}$ $(NH_4)_2 SO_4(aq) \rightarrow 2NH_4^+(aq) + SO_4^{2-}(aq)$ <p>Concentration of $(NH_4)_2 SO_4 = \frac{1}{2} \times 5.894134 \times 10^{-4} \times 132$ = 0.0389 g dm⁻³</p>	03
(b)(i)	The pH increases (slightly above 7).	½
(ii)	The mixture of ammonia and ammonium sulphate constitutes a basic buffer with a low concentration of hydroxide ions due to partial ionization of ammonia.	½
		06 marks
6.(a)	Bond dissociation energy Heat energy absorbed to break/dissociate one mole of covalent bonds into free gaseous atoms.	01
(b)	$\Delta H_R = \sum B.E \text{ broken bonds} - \sum B.E \text{ bonds formed}$ $= [C=O + 2(N-H)] - [C=N + 2(O-H)]$ $= (1581 + 2(391)) - (615 + 2(463))$ $= +40 \text{ KJ mol}^{-1}$	03
		04 marks

WAKISSHA JOINT MOCK EXAMINATIONS
MARKING GUIDE
Uganda Advanced Certificate of Education
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CHEMISTRY P525/1

15



1. (a)	Measurement of colour intensity of the reaction mixture. Titration (of the reaction mixture with a standard solution of sodium thiosulphate/starch indicator)	01
(b) (i)	Rate of reaction increases Iron (ii) ions act as a catalyst by providing an alternative reaction path of lower activation energy.	1½
(ii)	Rate of reaction remains constant. Order of reaction with respect to each reactant is one.	1½
(iii)	Rate decreases/reduces. The kinetic energy of reactant ions is lowered, Frequency of effective collisions decreases and the value of the rate constant decreases	1½
		5 ½ marks
2. (a)	$\text{CH}_3\text{C}(=\text{O})\text{NH}_2 + \text{NaOBr}_{(\text{aq})} \longrightarrow \text{CH}_3\text{NH}_2$ Name of product Methylamine (amino methane)	1½
(b)	$\text{C}_6\text{H}_5\text{CH}_2\text{OH} \xrightarrow[\text{Heat}]{\text{Excess NaBr(s) / Conc. H}_2\text{SO}_4} \text{C}_6\text{H}_5\text{CH}_2\text{Br}$ phenylethane.	1½
(c)	$(\text{CH}_3)_2\text{C}=\text{CHCH}_3 \xrightarrow[\text{Heat}]{\text{Conc. KMnO}_4/\text{H}_2\text{SO}_4} \text{CH}_3\text{COCH}_3 + \text{CH}_3\text{COOH}$ Names CH_3COCH_3 Propanone CH_3COOH Ethanoic acid	02
		05marks
3. (a) (i)	MnO	½
(ii)	MnO ₂	½
(iii)	Mn ₂ O ₇	½
(b) (i)	$\text{MnO}_{(\text{s})} + \text{H}_2\text{SO}_{4(\text{aq})} \longrightarrow \text{MnSO}_{4(\text{aq})} + \text{H}_2\text{O}_{(\text{l})}$	01
(ii)	$\text{MnO}_{2(\text{s})} + 4\text{HCl}_{(\text{aq})} \longrightarrow \text{MnCl}_{2(\text{aq})} + \text{Cl}_{2(\text{g})} + 2\text{H}_2\text{O}_{(\text{l})}$	01
(iii)	$\text{Mn}_2\text{O}_7_{(\text{l})} + 2\text{OH}_{(\text{aq})} \longrightarrow 2\text{MnO}_4^{-}(\text{aq}) + \text{H}_2\text{O}_{(\text{l})}$ Or $\text{Mn}_2\text{O}_7_{(\text{l})} + 2\text{NaOH}_{(\text{aq})} \longrightarrow 2\text{NaMnO}_4(\text{aq}) + \text{H}_2\text{O}_{(\text{l})}$	01
		05 marks