

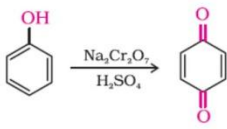


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Marking scheme – 2017 (Compartment)

CHEMISTRY (043)/ CLASS XII

Set 56/1

Q.No	Value Points	Marks
1	Frenkel defect	1
2	Liquid –liquid colloidal systems ; example- milk (or any other)	½, ½
3	Dichloridobis(ethane-1,2-diamine)cobalt(III) ion	1
4	 / Benzoquinone is formed	1
5	N,N-dimethylbutan-1-amine	1
6	$Ag^+ (aq) + e^- \rightarrow Ag(s)$ Because it has higher reduction potential	1 1
7	Hypophosphorous acid is a good reducing agent as it contains two P-H bonds. There is no P-H bond in orthophosphoric acid , so it is not a reducing agent Example : It reduces $AgNO_3$ to metallic silver/ chemical equation	1 1
8	a) Due to high activation energy b) Rate = $k [A_2]^0 [B_2]^0$	1 1
OR		
8	$R \rightarrow P$ $Rate = -\frac{d[R]}{dt} = k[R]$ or $\frac{d[R]}{[R]} = -kdt$ Integrating this equation, we get $\ln [R] = -kt + I$ (4.8) When $t = 0$, $R = [R]_0$, where $[R]_0$ is the initial concentration of the reactant. Therefore, equation (4.8) can be written as $\ln [R]_0 = -k \times 0 + I$ $\ln [R]_0 = I$ Substituting the value of I in equation (4.8) $\ln[R] = -kt + \ln[R]_0$ (4.9) Rearranging this equation $\ln \frac{[R]}{[R]_0} = -kt$ or $k = \frac{1}{t} \ln \frac{[R]_0}{[R]}$ $k = \frac{2.303}{t} \log \frac{[R]_0}{[R]}$	1 1
9	i. Because it has incompletely filled d orbitals in one of its oxidation state (Cu^{2+})	1

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	ii. Cr^{2+} (d^4) changes to Cr^{3+} (d^3) while Fe^{2+} (d^6) changes to Fe^{3+} (d^5). In aqueous medium d^3 is more stable than d^5 .	1
10.	a) $\text{CH}_3\text{-CH}(\text{Br})\text{-CH}_3 \xrightarrow{\text{alc. KOH}} \text{CH}_3\text{-CH=CH}_2 \xrightarrow{\text{HBr, Peroxide}} \text{CH}_3\text{-CH}_2\text{-CH}_2\text{-Br}$ b) .	1
		1
11	In bcc, $z=2$; $d = (z \times M) / a^3 \times N_A$ (i) Putting values of M in equation (i) $M = 7.2 \text{ g/cm}^3 \times (288 \times 10^{-10} \text{ cm})^3 \times N_A / 2$ $= 51.8 \text{ g/mol}$ (or any other correct method)	1 1 1
12	$\Delta rG^\circ = -nFE^\circ_{\text{cell}}$, $n=6$ $= -6 \times 96500 \text{ C/mol} \times 0.34 \text{ V}$ $= -196860 \text{ J/mol}$ or -196.860 kJ/mol $E^\circ_{\text{cell}} = 0.059 \text{ V} / n \times \log K_c$ $\log K_c = 0.34 \text{ V} \times 6 / 0.059 \text{ V} = 34.5762$	$\frac{1}{2}$ 1 $\frac{1}{2}$ 1
13	$t = \frac{2.303}{k} \log [R]_0 / [R]$ $t_{99\%} = \frac{2.303}{k} \log 100/1 = \frac{2.303}{k} \times 2$ ----- (i) $t_{90\%} = \frac{2.303}{k} \log 100/10 = \frac{2.303}{k}$ ----- (ii) Dividing equation (i) by (ii) $\frac{t_{99\%}}{t_{90\%}} = \frac{\frac{2.303}{k} \times 2}{\frac{2.303}{k}}$ $t_{99\%} = 2 t_{90\%}$	1 1 1
14	i) The colloidal particles scatter light in all directions in space. ii) The zig-zag movement of particles of the dispersed phase due to unbalanced bombardment of the colloidal particles by the molecules of dispersion medium. iii) As the adsorption is an exothermic process, it decreases with increase in temperature.	1 1 1
15	a) i) The impurities are more soluble in the melt than in the solid state of the metal. ii) The more basic / reactive metal gets deposited at the cathode and the less basic / reactive ones go to the anode mud. b) i) Ni ii) Ti/Zr	1 1 $\frac{1}{2}, \frac{1}{2}$
16	A: Na_2CrO_4 ; B: $\text{Na}_2\text{Cr}_2\text{O}_7$ $4 \text{ FeCr}_2\text{O}_4 + 8 \text{ Na}_2\text{CO}_3 + 7 \text{ O}_2 \rightarrow 8 \text{ Na}_2\text{CrO}_4 + 2 \text{ Fe}_2\text{O}_3 + 8 \text{ CO}_2$ $2\text{Na}_2\text{CrO}_4 + 2 \text{ H}^+ \rightarrow \text{Na}_2\text{Cr}_2\text{O}_7 + 2 \text{ Na}^+ + \text{H}_2\text{O}$	$\frac{1}{2}, \frac{1}{2}$ 1 1
OR		
16	a) i) Due to d-d transition ii) Due to higher oxidation state of Mn in Mn_2O_7 / Due to high polarizing power of Mn(VII). b) $\mu = \sqrt{4(4+2)} = 4.90 \text{ B.M}$	1 1 1

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17	Hybridisation : d^2sp^3 Magnetic character : Paramagnetic 	1 1 1
18.	a) i) Due to -I effect of X, the ring gets partially deactivated ii) They fail to form Hydrogen bonds with water/ more energy is required to break hydrogen bonds in water and less energy is released when new attractions are set up. b) 2-Bromo-2-methylbutane < 2-Bromopentane < 1-Bromopentane	1 1 1
19.	a) Due to resonance, phenoxide ion is more stable than phenol whereas there is no resonance in alkoxide ion / explained with the help of resonating structures. b) . (i) $CH_3-CH_2-\ddot{O}-H + H^+ \rightarrow CH_3-CH_2-\overset{+}{O}-H$ (ii) $CH_3CH_2-\ddot{O}: + CH_3-CH_2-\overset{+}{O}-H \rightarrow CH_3CH_2-\overset{+}{O}-CH_2CH_3 + H_2O$ (iii) $CH_3CH_2-\overset{+}{O}-CH_2CH_3 \rightarrow CH_3CH_2-O-CH_2CH_3 + H^+$	1 ½ 1 ½
20.	i) A: CH_3-CH_2CN ; B: $CH_3-CH_2-CH_2NH_2$; C: $CH_3-CH_2-CH_2-NH-COCH_3$ ii) A: $Ar-\overset{+}{N}_2\overset{-}{B}F_4$; B: ; C:	½ x 3 ½ x 3
21	a) Because they are excreted in urine and cannot be stored in body; Vitamin C / B ₁ / B ₂ / B ₆ b) i) Essential amino acids are those which cannot be synthesized in the body and are supplied through diet whereas non-essential amino acid can be synthesized in the body ii) In fibrous proteins, the polypeptide chains run parallel and are held together by hydrogen or disulphide bonds while in globular, polypeptide chains coil around to give a spherical shape	½, ½ 1 1
22	i) Phenol / 0.2 % phenol is antiseptic while 1% is disinfectant. ii) Aspartame iii) Cationic detergents are quaternary ammonium salts of amines with acetates, chlorides or bromides as anions/ Cationic part has a long chain hydrocarbon which is involved in cleansing action.	1 1 1
23	a) <i>Poly β-hydroxybutyrate – co-β-hydroxyvalerate / (PHBV)</i> Monomers : $CH_3-\overset{OH}{ }CH-CH_2-COOH$, $CH_3-CH_2-\overset{OH}{ }CH-CH_2-COOH$ Repeating unit : $\left(O-\underset{CH_3}{ }CH-CH_2-\overset{O}{ }C-O-\underset{CH_2CH_3}{ }CH-CH_2-\overset{O}{ }C \right)_n$	½ ½, ½ ½

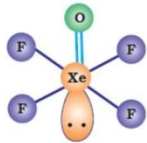
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	b) PHBV is used in speciality packaging, orthopaedic devices and in controlled release of drugs. (any two)	½, ½
	c) Concern for environment, caring (or any other)	½, ½
24	a) Vapour pressure of the solvent decreases in the presence of non-volatile solute (glucose) hence boiling point increases	2
	b) $p_{CO_2} = K_H X_{CO_2}$	½
	$X_{CO_2} = p_{CO_2} / K_H$ $= 2.53 \times 10^5 \text{ Pa} / 1.67 \times 10^8 \text{ Pa} = 1.51 \times 10^{-3}$	1
	$n_{H_2O} = 500 \text{ g} / 18 \text{ g/mol} = 27.77 \text{ mol}$	
	Let $n_{CO_2} = n \text{ mol}$	
	$X_{CO_2} = n / (27.77 + n) = 1.51 \times 10^{-3}$	½
	$n_{CO_2} = 1.51 \times 10^{-3} \times 27.77 \text{ mol} = 0.042 \text{ mol}$	1
OR		
24	a) i) The solutions which obey Raoult's law over the entire range of concentration.	1
	ii) It is the excess pressure that must be applied to a solution to prevent osmosis.	1
	b) $\Delta T_b = i K_b m$	
	Here, $m = w_b \times 1000 / M_b \times w_a$	1
	$\Delta T_b = [3 \times 0.512 \text{ K kg mol}^{-1} \times 1000 \times 10 \text{ g}] / [111 \text{ g mol}^{-1} \times 200 \text{ g}]$	1
	$= 0.69 \text{ K}$	1
25	a) A: NO_2 ; B: N_2O_4	½, ½,
	$\text{NaNO}_3 + \text{conc. H}_2\text{SO}_4 \rightarrow \text{NaHSO}_4 + \text{HNO}_3$ (or any other nitrate)	1
	$\text{Cu} + 4 \text{HNO}_3 \rightarrow \text{Cu}(\text{NO}_3)_2 + 2 \text{NO}_2 + 2 \text{H}_2\text{O}$	1
	$2 \text{NO}_2 \xrightarrow{\text{cool}} \text{N}_2\text{O}_4$	1
	b) .	
		1
OR		
25	a) i) Stability of higher oxidation state decreases down the group from S to Te/ Stability of lower oxidation state increases down the group from S to Te.	1
	ii) ClO_3^- is more stable than $\text{ClO}^- / \text{ClO}_2^-$ is a weak conjugate base than $\text{ClO}^- / \text{ClO}_2^-$ Due to higher oxidation state of chlorine in HClO_3	1
	iii) Fluorine and oxygen are most electronegative and very reactive.	1
	b)	
	i) .	1
	$4 \text{NaCl} + \text{MnO}_2 + 4 \text{H}_2\text{SO}_4 \rightarrow \text{MnCl}_2 + 4 \text{NaHSO}_4 + 2 \text{H}_2\text{O} + \text{Cl}_2$	1
	ii).	1
	$6 \text{XeF}_4 + 12 \text{H}_2\text{O} \rightarrow 4 \text{Xe} + 2 \text{XeO}_3 + 24 \text{HF} + 3 \text{O}_2$	1
26	a) i) Due to steric and +I effect of two methyl groups in propanone.	1
	ii) Because it is a deactivating group / Due to electron withdrawing carboxylic group resulting in decreased electron density at o- and p- position.	1
	iii) Due to resonance, electrophilicity of carbonyl carbon is reduced.	1
	b) i) Add NaOH and I_2 to both the compounds and heat, acetophenone forms yellow ppt of iodoform.	1
	ii) Add NaHCO_3 solution to both the compounds, benzoic acid will give effervescence and liberate CO_2 .	1
	(Or any other suitable test)	
OR		
26	a) A: CH_3CHO ; B: $\text{CH}_3\text{-CH(OH)-CH}_2\text{-CHO}$; C: $\text{CH}_3\text{-CH=CH-CHO}$;	1×4

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D: $\text{CH}_3\text{-CH}(\text{CH}_3)\text{-OH}$ b) $\text{CH}_3\text{-O-CH}_3 < \text{CH}_3\text{CHO} < \text{CH}_3\text{-CH}_2\text{-OH} < \text{CH}_3\text{-COOH}$	1
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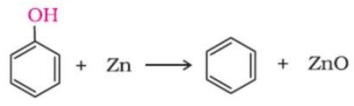
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Marking scheme – 2017 (Compartment)
CHEMISTRY (043)/ CLASS XII
Set 56/2

Q.No	Value Points	Marks
1	Dispersion medium- liquid/ water ; Dispersed phase – liquid/ oil	$\frac{1}{2}$, $\frac{1}{2}$
2	Tetraamminechloridonitrito-N -cobalt(III) ion	1
3	N,N-dimethylbutan-1-amine	1
4	Schottky Defect	1
5	 <p align="right">/ Benzene is formed</p>	1
6	<p>a) Due to high activation energy</p> <p>b) $\text{Rate} = k [A_2]^0 [B_2]^0$</p> <p align="center">OR</p> <p>R → P</p> $\text{Rate} = -\frac{d[R]}{dt} = k[R]$ <p>or $\frac{d[R]}{[R]} = -kdt$</p> <p>Integrating this equation, we get</p> $\ln [R] = -kt + I \quad (4.8)$ <p>When $t = 0$, $R = [R]_0$, where $[R]_0$ is the initial concentration of the reactant.</p> <p>Therefore, equation (4.8) can be written as</p> $\ln [R]_0 = -k \times 0 + I$ $\ln [R]_0 = I$ <p>Substituting the value of I in equation (4.8)</p> $\ln [R] = -kt + \ln [R]_0 \quad (4.9)$ <p>Rearranging this equation</p> $\ln \frac{[R]}{[R]_0} = -kt$ <p>or $k = \frac{1}{t} \ln \frac{[R]_0}{[R]}$</p> $k = \frac{2.303}{t} \log \frac{[R]_0}{[R]}$	1 1 1
7	<p>i) Silver can exhibit +2 oxidation state wherein it will have incompletely filled d-orbital.</p> <p>ii) Much higher third ionisation energy of Mn where the required change is from d^5 to d^4</p>	1 1
8	<p>a) $\text{CH}_3\text{-CH(Br)-CH}_3 \xrightarrow{\text{alc. KOH}} \text{CH}_3\text{-CH=CH}_2 \xrightarrow{\text{HBr, Peroxide}} \text{CH}_3\text{-CH}_2\text{-CH}_2\text{-Br}$</p> <p>b) .</p>	1

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		1
9	Hypophosphorous acid is a good reducing agent as it contains two P-H bonds. There is no P-H bond in orthophosphoric acid, so it is not a reducing agent Example : It reduces AgNO ₃ to metallic silver/ chemical equation	1 1
10.	Ag ⁺ (aq) + e ⁻ → Ag(s) Because it has higher reduction potential	1 1
11	i) Phenol / 0.2 % phenol is antiseptic while 1% is disinfectant. ii) Aspartame iii) Cationic detergents are quaternary ammonium salts of amines with acetates, chlorides or bromides as anions/ Cationic part has a long chain hydrocarbon which is involved in cleansing action.	1 1 1
12	a) Because they are excreted in urine and cannot be stored in body; Vitamin C / B ₁ / B ₂ / B ₆ b) i) Essential amino acids are those which cannot be synthesized in the body and are supplied through diet whereas non-essential amino acid can be synthesized in the body ii) In fibrous proteins, the polypeptide chains run parallel and are held together by hydrogen or disulphide bonds while in globular, polypeptide chains coil around to give a spherical shape	½, ½ 1 1
13	i) A: CH ₃ -CH ₂ CN ; B: CH ₃ -CH ₂ -CH ₂ NH ₂ ; C: CH ₃ -CH ₂ -CH ₂ -NH-COCH ₃ ii) A: Ar-N ⁺ ₂ B ⁻ F ₄ ; B: ; C:	½ × 3 ½ × 3
14	a) i) Due to -I effect of X, the ring gets partially deactivated ii) They fail to form Hydrogen bonds with water/ more energy is required to break hydrogen bonds in water and less energy is released when new attractions are set up. b) 2-Bromo-2-methylbutane < 2-Bromopentane < 1-Bromopentane	1 1 1
15	i). ii). Step 1: Formation of protonated alcohol. Step 2: Formation of carbocation: It is the slowest step and hence, the rate determining step of the reaction. Step 3: Formation of ethene by elimination of a proton. 	1 ½ ½ 1
16	Hybridisation : d ² sp ³ Spin : Low spin	1 1

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		1
17	i) The impurities are more soluble in the melt than in the solid state of the metal. ii) Different components of a mixture are differently adsorbed on the surface of adsorbent. iii) The more basic / reactive metal gets deposited at the cathode and the less basic / reactive ones go to the anode mud.	1 1 1
18.	A: Na_2CrO_4 ; B: $\text{Na}_2\text{Cr}_2\text{O}_7$ $4 \text{FeCr}_2\text{O}_4 + 8 \text{Na}_2\text{CO}_3 + 7 \text{O}_2 \rightarrow 8 \text{Na}_2\text{CrO}_4 + 2 \text{Fe}_2\text{O}_3 + 8 \text{CO}_2$ $2\text{Na}_2\text{CrO}_4 + 2 \text{H}^+ \rightarrow \text{Na}_2\text{Cr}_2\text{O}_7 + 2 \text{Na}^+ + \text{H}_2\text{O}$	$\frac{1}{2}$, $\frac{1}{2}$ 1 1
OR		
18	a) i) Due to d-d transition ii) Due to higher oxidation state of Mn in Mn_2O_7 / Due to high polarizing power of Mn(VII). b) $\mu = \sqrt{4(4+2)} = 4.90 \text{ B.M}$	1 1 1
19.	i) The colloidal particles scatter light in all directions in space. ii) The zig-zag movement of particles of the dispersed phase due to unbalanced bombardment of the colloidal particles by the molecules of dispersion medium. iii) As the adsorption is an exothermic process, it decreases with increase in temperature.	1 1 1
20.	$t = \frac{2.303}{k} \log \frac{[R]_0}{[R]}$ $t_{99\%} = \frac{2.303}{k} \log \frac{100}{1} = \frac{2.303}{k} \times 2 \text{ ---- (i)}$ $t_{90\%} = \frac{2.303}{k} \log \frac{100}{10} = \frac{2.303}{k} \text{ ---- (ii)}$ Dividing equation (i) by (ii) $\frac{t_{99\%}}{t_{90\%}} = \frac{\frac{2.303}{k} \times 2}{\frac{2.303}{k}}$ $t_{99\%} = 2 t_{90\%}$	1 1 1
21	In bcc, $z=2$; $d = (z \times M) / a^3 \times N_A$ (i) Putting values of M in equation (i) $M = 7.2 \text{ g/cm}^3 \times (288 \times 10^{-10} \text{ cm})^3 \times N_A / 2$ $= 51.8 \text{ g/mol}$ (or any other correct method)	1 1 1
22	$\Delta rG^\circ = -nFE^\circ_{\text{cell}}$, $n=6$ $= -6 \times 96500 \text{ C/mol} \times 0.30 \text{ V}$ $= -173700 \text{ J/mol} = -173.7 \text{ kJ/mol}$ $E^\circ_{\text{cell}} = 0.059 \text{ V} / n \times \log K_c$ $\log K_c = 0.30 \text{ V} \times 6 / 0.059 \text{ V} = 30.5$	$\frac{1}{2}$ 1 $\frac{1}{2}$ 1
23	a) <i>Poly</i> β -hydroxybutyrate – <i>co</i> - β -hydroxyvalerate / (PHBV) 	$\frac{1}{2}$ $\frac{1}{2}$, $\frac{1}{2}$

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	<p>Repeating unit :</p> $\left(\text{O}-\underset{\text{CH}_3}{\text{CH}}-\text{CH}_2-\overset{\text{O}}{\parallel}{\text{C}}-\text{O}-\underset{\text{CH}_2\text{CH}_3}{\text{CH}}-\text{CH}_2-\overset{\text{O}}{\parallel}{\text{C}} \right)_n$ <p>b) PHBV is used in speciality packaging, orthopaedic devices and in controlled release of drugs.(any two) ½, ½</p> <p>c) Concern for environment , caring (or any other) ½, ½</p>	
24	<p>a) i) Due to steric and +I effect of two methyl groups in propanone. ii) Because it is a deactivating group / Due to electron withdrawing carboxylic group resulting in decreased electron density at o- and p- position. iii) Due to resonance, electrophilicity of carbonyl carbon is reduced.</p> <p>b) i) Add NaOH and I₂ to both the compounds and heat, acetophenone forms yellow ppt of iodoform. ii) Add NaHCO₃ solution to both the compounds, benzoic acid will give effervescence and liberates CO₂.</p> <p align="right">(Or any other suitable test)</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
OR		
24	<p>a) A: CH₃CHO ; B: CH₃-CH(OH)-CH₂-CHO ; C: CH₃-CH=CH-CHO ; D: CH₃-CH(CH₃)-OH</p> <p>b) CH₃-O-CH₃ < CH₃CHO < CH₃-CH₂-OH < CH₃-COOH</p>	<p>1×4</p> <p>1</p>
25	<p>a) Vapour pressure of the solvent decreases in the presence of non – volatile solute (glucose) hence boiling point increases</p> <p>b) $p_{\text{CO}_2} = K_H X_{\text{CO}_2}$ $X_{\text{CO}_2} = p_{\text{CO}_2} / K_H$ $= 2.53 \times 10^5 \text{ Pa} / 1.67 \times 10^8 \text{ Pa} = 1.51 \times 10^{-3}$ $n_{\text{H}_2\text{O}} = 500\text{g} / 18 \text{ g/mol} = 27.77 \text{ mol}$ Let $n_{\text{CO}_2} = n \text{ mol}$ $X_{\text{CO}_2} = n / (27.77 + n) = 1.51 \times 10^{-3}$ $n_{\text{CO}_2} = 1.51 \times 10^{-3} \times 27.77 \text{ mol} = 0.042 \text{ mol}$</p>	<p>2</p> <p>½</p> <p>1</p> <p>½</p> <p>1</p>
OR		
25	<p>a) i) The solutions which obey Raoult's law over the entire range of concentration. ii) It is the excess pressure that must be applied to a solution to prevent osmosis.</p> <p>b) $\Delta T_b = i K_b m$ Here , $m = w_b \times 1000 / M_b \times w_A$ $\Delta T_b = [3 \times 0.512 \text{ K kg mol}^{-1} \times 1000 \times 10 \text{ g}] / [111 \text{ g mol}^{-1} \times 200\text{g}]$ $= 0.69\text{K}$</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
26	<p>a) A: NO₂ ; B: N₂O₄ $\text{NaNO}_3 + \text{conc. H}_2\text{SO}_4 \longrightarrow \text{NaHSO}_4 + \text{HNO}_3$ (or any other nitrate) $\text{Cu} + 4 \text{HNO}_3 \longrightarrow \text{Cu}(\text{NO}_3)_2 + 2 \text{NO}_2 + 2 \text{H}_2\text{O}$ $2\text{NO}_2 \xrightarrow{\text{cool}} \text{N}_2\text{O}_4$</p> <p>b) .</p>	<p>½, ½,</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
OR		
26	<p>a) i) Stability of higher oxidation state decreases down the group from S to Te/ Stability of lower oxidation state increases down the group from S to Te. ii) ClO₃⁻ is more stable than ClO⁻ / ClO₃⁻ is a weak conjugate base than ClO⁻ / Due to higher oxidation state of chlorine in HClO₃</p>	<p>1</p> <p>1</p>

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iii) Fluorine and oxygen are most electronegative and very reactive.	1
b)	
i) .	1
$4\text{NaCl} + \text{MnO}_2 + 4\text{H}_2\text{SO}_4 \rightarrow \text{MnCl}_2 + 4\text{NaHSO}_4 + 2\text{H}_2\text{O} + \text{Cl}_2$	
ii).	1
$6\text{XeF}_4 + 12 \text{H}_2\text{O} \rightarrow 4\text{Xe} + 2\text{XeO}_3 + 24 \text{HF} + 3 \text{O}_2$	

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CHEMISTRY (043)/ CLASS XII

Set 56/3

Q.No	Value Points	Marks
1	<p>2,4,6-Tribromophenol is formed</p>	1
2	Dichloridobis(ethane-1,2-diamine)cobalt(III) ion	1
3	AgBr	1
4	N,N-dimethylbutan-1-amine	1
5	Dispersed phase - liquid/ water ; Dispersion medium – liquid/ oil	1
6	<p>a) $\text{CH}_3\text{-CH}(\text{Br})\text{-CH}_3 \xrightarrow{\text{alc. KOH}} \text{CH}_3\text{-CH=CH}_2 \xrightarrow{\text{HBr, Peroxide}} \text{CH}_3\text{-CH}_2\text{-CH}_2\text{-Br}$</p> <p>b) .</p>	1
7	i) Due to absence of unpaired electrons ii) Due to high $\Delta_a H^\ominus$ and low $\Delta_{\text{hyd}} H^\ominus$.	1 1
8	Hypophosphorous acid is a good reducing agent as it contains two P-H bonds. There is no P-H bond in orthophosphoric acid , so it is not a reducing agent Example : It reduces AgNO_3 to metallic silver/ chemical equation	1 1
9	$\text{Cu}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Cu}(\text{s})$ Because it has higher reduction potential	1 1
10.	i) Zero Order ii) Pseudo-first Order	1 1
11	a) i)The impurities are more soluble in the melt than in the solid state of the metal. ii)The more basic / reactive metal gets deposited at the cathode and the less basic / reactive ones go to the anode mud. b) i)Ni ii) Ti/Zr	1 1 $\frac{1}{2}, \frac{1}{2}$
12	i) High energy of activation is needed ii) Blood being a colloidal solution, it gets coagulated by alum (an electrolyte). iii) Dust particles along with water suspended in air scatter blue light which reaches our eyes.	1 1 1
13	A: Na_2CrO_4 ; B: $\text{Na}_2\text{Cr}_2\text{O}_7$ $4 \text{FeCr}_2\text{O}_4 + 8 \text{Na}_2\text{CO}_3 + 7 \text{O}_2 \rightarrow 8 \text{Na}_2\text{CrO}_4 + 2 \text{Fe}_2\text{O}_3 + 8 \text{CO}_2$ $2\text{Na}_2\text{CrO}_4 + 2 \text{H}^+ \rightarrow \text{Na}_2\text{Cr}_2\text{O}_7 + 2 \text{Na}^+ + \text{H}_2\text{O}$	$\frac{1}{2}, \frac{1}{2}$ 1 1
OR		
13	a) i)Due to d-d transition ii)Due to higher oxidation state of Mn in Mn_2O_7 / Due to high polarizing power of Mn(VII). b) $\mu = \sqrt{4(4+2)} = 4.90 \text{ B.M}$	1 1 1
14	$\Delta_r G^\ominus = -nFE^\ominus_{\text{cell}}$, n=6	$\frac{1}{2}$

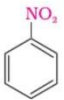
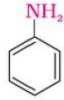
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	$= -6 \times 96500 \text{ C/mol} \times 2.02\text{V}$ $= -1169580 \text{ J/mol or } -116.958 \text{ kJ/mol}$ $E^\circ_{\text{cell}} = 0.059\text{V} / n \times \log Kc$ $\log Kc = 2.02 \text{ V} \times 6 / 0.059\text{V} = 205.42$	1 ½ 1
15	In bcc, $z=2$; $d = (z \times M) / a^3 \times N_A$ (i) Putting values of M in equation (i) $M = 7.2\text{g/cm}^3 \times (288 \times 10^{-10} \text{ cm})^3 \times N_A / 2$ $= 51.8 \text{ g/mol}$ (or any other correct method)	1 1 1
16	a) i) Due to -I effect of X, the ring gets deactivated ii) They fail to form Hydrogen bonds with water/ more energy is required to break hydrogen bonds in water and less energy is released when new attractions are set up. b) 2-Bromo-2-methylbutane < 2-Bromopentane < 1-Bromopentane	1 1 1
17	i) A: $\text{CH}_3\text{-CH}_2\text{CN}$; B: $\text{CH}_3\text{-CH}_2\text{-CH}_2\text{NH}_2$; C: $\text{CH}_3\text{-CH}_2\text{-CH}_2\text{-NH-COCH}_3$ ii) A: $\text{Ar}-\overset{+}{\text{N}}_2\text{BF}_4^-$; B:  ; C: 	½ × 3 ½ × 3
18.	a) Because they are excreted in urine and cannot be stored in body; Vitamin C / B ₁ / B ₂ / B ₆ b) i) Essential amino acids are those which cannot be synthesized in the body and are supplied through diet whereas non-essential amino acid can be synthesized in the body ii) In fibrous proteins, the polypeptide chains run parallel and are held together by hydrogen or disulphide bonds while in globular, polypeptide chains coil around to give a spherical shape	½, ½ 1 1
19.	i) Phenol / 0.2 % phenol is antiseptic while 1% is disinfectant. ii) Aspartame iii) Cationic detergents are quaternary ammonium salts of amines with acetates, chlorides or bromides as anions/ Cationic part has a long chain hydrocarbon which is involved in cleansing action.	1 1 1
20.	i) $[\text{Cr}(\text{H}_2\text{O})_6] \text{Cl}_3$ ii) Hexaaquachromium(III) chloride iii) Paramagnetic and high spin	1 1 ½, ½
21	$t_{1/2} = \frac{0.693}{k}$ $k = \frac{0.693}{693 \text{ s}}$ $= 0.001 \text{ s}^{-1}$ $k = \frac{2.303}{t} \log [R]_0/[R]$ $t = \frac{2.303}{k} \log [R]_0/[R]$ $= \frac{2.303}{0.001} \log 100/10$ $k = 2303 \text{ s}$	½ 1 ½ 1
22	a) Due to resonance, phenoxide ion is more stable than phenol whereas there is no resonance in alkoxide ion / explained with the help of resonating structures. b) .	1

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	<p>(i) $\text{CH}_3\text{-CH}_2\text{-}\ddot{\text{O}}\text{-H} + \text{H}^+ \rightarrow \text{CH}_3\text{-CH}_2\text{-}\overset{\text{H}}{\underset{\text{H}}{\text{O}^+}}\text{-H}$</p> <p>(ii) $\text{CH}_3\text{CH}_2\text{-}\ddot{\text{O}}\text{:} + \text{CH}_3\text{-CH}_2\text{-}\overset{\text{H}}{\underset{\text{H}}{\text{O}^+}} \rightarrow \text{CH}_3\text{CH}_2\text{-}\overset{\text{H}}{\underset{\text{H}}{\text{O}^+}}\text{-CH}_2\text{CH}_3 + \text{H}_2\text{O}$</p> <p>(iii) $\text{CH}_3\text{CH}_2\text{-}\overset{\text{H}}{\underset{\text{H}}{\text{O}^+}}\text{-CH}_2\text{CH}_3 \rightarrow \text{CH}_3\text{CH}_2\text{-O-CH}_2\text{CH}_3 + \text{H}^+$</p>	<p>½</p> <p>1</p> <p>½</p>
23	<p>a) <i>Poly β-hydroxybutyrate – co-β-hydroxyvalerate / (PHBV)</i></p> <p><i>Monomers:</i> $\text{CH}_3\text{-}\overset{\text{OH}}{\text{CH}}\text{-CH}_2\text{-COOH}$, $\text{CH}_3\text{-CH}_2\text{-}\overset{\text{OH}}{\text{CH}}\text{-CH}_2\text{-COOH}$</p> <p><i>Repeating unit:</i></p> $\left(\text{O}-\underset{\text{CH}_3}{\text{CH}}-\text{CH}_2-\overset{\text{O}}{\underset{\text{O}}{\text{C}}}-\text{O}-\underset{\text{CH}_2\text{CH}_3}{\text{CH}}-\text{CH}_2-\overset{\text{O}}{\underset{\text{O}}{\text{C}}} \right)_n$ <p>b) PHBV is used in speciality packaging, orthopaedic devices and in controlled release of drugs. (any two)</p> <p>c) Concern for environment , caring (or any other)</p>	<p>½</p> <p>½ , ½</p> <p>½</p> <p>½ , ½</p> <p>½ , ½</p>
24	<p>a) A: NO_2 ; B: N_2O_4</p> <p>$\text{NaNO}_3 + \text{conc. H}_2\text{SO}_4 \rightarrow \text{NaHSO}_4 + \text{HNO}_3$ (or any other nitrate)</p> <p>$\text{Cu} + 4 \text{HNO}_3 \rightarrow \text{Cu}(\text{NO}_3)_2 + 2 \text{NO}_2 + 2 \text{H}_2\text{O}$</p> <p>$2\text{NO}_2 \xrightarrow{\text{cool}} \text{N}_2\text{O}_4$</p> <p>b) .</p>	<p>½ , ½ ,</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
OR		
24	<p>a) i) Stability of higher oxidation state decreases down the group from S to Te/ Stability of lower oxidation state increases down the group from S to Te.</p> <p>ii) ClO_3^- is more stable than ClO^- / ClO_3^- is a weak conjugate base than ClO^- / Due to higher oxidation state of chlorine in HClO_3</p> <p>iii) Fluorine and oxygen are most electronegative and very reactive.</p> <p>b)</p> <p>i) .</p> <p>$4\text{NaCl} + \text{MnO}_2 + 4\text{H}_2\text{SO}_4 \rightarrow \text{MnCl}_2 + 4\text{NaHSO}_4 + 2\text{H}_2\text{O} + \text{Cl}_2$</p> <p>ii).</p> <p>$6\text{XeF}_4 + 12 \text{H}_2\text{O} \rightarrow 4\text{Xe} + 2\text{XeO}_3 + 24 \text{HF} + 3 \text{O}_2$</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
25	<p>a) i) Due to steric and +I effect of two methyl groups in propanone.</p> <p>ii) Because it is a deactivating group / Due to electron withdrawing carboxylic group resulting in decreased electron density at o- and p- position.</p> <p>iii) Due to resonance, electrophilicity of carbonyl carbon is reduced.</p> <p>b) i) Add NaOH and I_2 to both the compounds and heat, acetophenone forms yellow ppt of iodoform.</p> <p>ii) Add NaHCO_3 solution to both the compounds, Benzoic acid will give effervescence and liberates CO_2.</p> <p align="right">(Or any other suitable test)</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>

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	OR	
25	a) A: CH ₃ CHO ; B: CH ₃ -CH(OH)-CH ₂ -CHO ; C: CH ₃ -CH=CH-CHO ; D: CH ₃ -CH(CH ₃)-OH b) CH ₃ -O-CH ₃ < CH ₃ CHO < CH ₃ -CH ₂ -OH < CH ₃ -COOH	1×4 1
26	a) Vapour pressure of the solvent decreases in the presence of non – volatile solute (glucose) hence boiling point increases b) $p_{CO_2} = K_H X_{CO_2}$ $X_{CO_2} = p_{CO_2} / K_H$ $= 2.53 \times 10^5 \text{ Pa} / 1.67 \times 10^8 \text{ Pa} = 1.51 \times 10^{-3}$ $n_{H_2O} = 500\text{g} / 18 \text{ g/mol} = 27.77 \text{ mol}$ Let $n_{CO_2} = n \text{ mol}$ $X_{CO_2} = n / (27.77 + n) = 1.51 \times 10^{-3}$ $n_{CO_2} = 1.51 \times 10^{-3} \times 27.77 \text{ mol} = 0.042 \text{ mol}$	2 ½ 1 ½ 1
	OR	
26	a) i) The solutions which obey Raoult's law over the entire range of concentration. ii) It is the excess pressure that must be applied to a solution to prevent osmosis. b) $\Delta T_b = i K_b m$ Here, $m = w_b \times 1000 / M_b \times w_A$ $\Delta T_b = [3 \times 0.512 \text{ K kg mol}^{-1} \times 1000 \times 10 \text{ g}] / [111 \text{ g mol}^{-1} \times 200\text{g}]$ $= 0.69\text{K}$	1 1 1 1 1

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