Marking Scheme <u>Chemistry - 2014</u> Outside Delhi- SET (56 /1)

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1	It first increases then decreases or graphical representation.	1
2	Zn acts as reducing agent.	1
3	2	1
4	çı	1
	2–Chlorobutane or or first molecule of the pair.	
5	Proteins	1
6.	Diazotization	1
7.	Glucose & Fructose	1
8.	СНО	1
	1	
	[O]	
	\searrow	
	CH_3	
9.	Given; $d = 2.8 \text{g/cm}^3$; $Z = 4$; $a = 4 \times 10^{-8} \text{ cm}$ $N_A = 6.022 \times 10^{23} \text{ per mol}$	
	$d = \frac{Z \times M}{a^3 \times N_A}$ or $M = \frac{d \times a^3 \times N_A}{Z}$	1/2
	$2.8 \text{ g cm}^{-3} (4 \text{ x } 10^{-8} \text{ cm})^3 \text{ x } 6.022 \text{ x} 10^{23}$	
	\Rightarrow M = $\frac{1}{4}$	1/2
	$M = 2.8 \text{ x } 16 \text{ x } 10^{-1} \text{ x } 6.022 = 26.97 \text{ g/mol}$	1
10	(i) Metal excess defect / Metal excess defect due to anionic vacancies filled by free electrons	1
	/ Due to F – centers.	
	(ii) Schottky defect.	1
	Or	

10	(i) Tetrahedral void is surrounded by 4 constituent particles (atoms / molecules / ions).	
	Octahedral void is surrounded by 6 constituent particles (atoms / molecules / ions).	1
	OR	
	radius ratio (r $^+$ /r $^-$) for Tetrahedral void is 0.225 & radius ratio for octahedral voids is 0.414	
	(ii) A regular three dimensional arrangement of points in space is called a crystal lattice.	
	Unit cell is the smallest portion of a crystal lattice which, when repeated in three directions,	1
	generates an entire lattice. / unit cell is the miniature of crystal lattice / microscopic edition of the	
	crystal lattice.	
11	Kohlrausch law of independent migration of ions. The law states that limiting molar	1
	conductivity of an electrolyte can be stated as the sum of the individual contributions of the anion	
	and cation of the electrolyte.	
	On dilution, the conductivity (κ) of the electrolyte decreases as the number of ions per unit	1
	volume of solution decreases.	
12	(i) Zero order reaction	1
	(ii) slope = -k	1
13	In this method, the impure metal is made to act as anode. A strip of the same metal in pure form is	1
	used as cathode. They are put in a suitable electrolytic bath containing soluble salt of the same	
	metal. Pure metal is deposited at the cathode and impurities remain in the solution.	
	For example: electro refining of Cu, Ag, Au (any one)	1
14	(i) $P_4 + H_2O \longrightarrow$ no reaction or if attempted in any form, award one mark	1
	(ii) $XeF_4 + O_2F_2 \longrightarrow XeF_6 + O_2$.	1
15	F F F F	1+1
16	Reimer-Tiemann reaction	
	$\begin{array}{c c} OH & \overline{O} Na^{+} & OH \\ \downarrow & \downarrow & \downarrow & \downarrow \\ \downarrow & \downarrow & \downarrow & \downarrow \\ \downarrow & \downarrow &$	1
	$\bigcirc \xrightarrow{\text{CHCl}_3 + \text{ aq NaOH}} \bigcirc \xrightarrow{\text{CHO}} \xrightarrow{\text{H}^+} \bigcirc \xrightarrow{\text{CHO}} \xrightarrow{\text{CHO}}$	
	Williamson synthesis	
	$R-X + R'-O' Na \longrightarrow R-O'-R' + Na X$	1
17	$HBr \rightarrow H^+ + Br^-$	

	¥¥	1		
	$CH_3 - CH_2 - O - H + H^+ \rightarrow CH_3 - CH_2 - O - H$	1/2		
	H			
	$CH_3 - CH_2 - O - H \rightarrow CH_3 - CH_2 + H_2O$			
	$CH_3^+CH_2 \xrightarrow{Br^-} CH_3^-CH_2^-Br$			
	Or			
	$\mathbf{Br} + \mathbf{CH}_{2} - \mathbf{OH}_{2}^{+} \longrightarrow \mathbf{Br} - \mathbf{CH}_{2} + \mathbf{H}_{2}\mathbf{O}$			
	$(\text{where } \mathbf{K} = -\mathbf{C}\mathbf{H}_3)$			
18	(i) Phenol & Formaldehyde	1		
	(ii) 2–Chloro–1,3–butadiene (or Chloroprene)			
19	(a) Given $E^{\circ}Cell = +2.71V$ & $F = 96500C \text{ mol}^{-1}$ $n = 2$ (from the given reaction)			
	$\Delta r G^{O} = -n x F x E^{O} Cell$			
	$\Delta r G^{O} = -2 \times 96500 \text{ C mol}^{-1} \times 2.71 \text{ V}$			
	= - 523030 J / mol or - 523.030 kJ / mol	1		
	(b) Hydrogen – oxygen fuel Cell / Fuel cell.	1		
20	$SO_2 Cl_2 \rightarrow SO_2 + Cl_2$			
	At $t = 0s$ 0.4 atm 0 atm 0 atm			
	At $t = 100s$ (0.4 – x) atm x atm x atm			
	Pt = 0.4 - x + x + x			
	Pt = 0.4 + x			
	0.7 = 0.4 + x			
	x = 0.3			
	$k = \frac{2.303}{t} \log \frac{p_i}{2p_i - p_t}$	1		
	$k = \frac{2.303}{t} \log \frac{0.4}{0.8 - 0.7}$	1		
	$k = \frac{2.303}{100s} \log \frac{0.4}{0.1}$	1		
	$k = \frac{2.303}{100s} \times 0.6021 = 1.39 \times 10^{-2} \text{ s}^{-1}$			
21	1 These are liquid-liquid colloidal systems or the dispersion of one liquid in another liquid.			
	Types: (i) Oil dispersed in water (O/W type) Example; milk and vanishing cream			
	(ii) Water dispersed in oil (W/O type) Example; butter and cream.			

	(Any one example of each type)		
22	2 (i) As N can't form 5 covalent bonds / its maximum covalency is four.		
	(ii) This is due to very small size of Oxygen atom / repulsion between electrons is large in		
	relatively small 2p sub-shell.		
	(iii) In H ₃ PO ₂ there are 2 P–H bonds, whereas in H ₃ PO ₃ there is 1 P–H bond		
23	(i) Tetraamminedichloridochromium (III) chloride.		
	(ii) Optical isomerism		
	(iii) In [NiCl ₄] ²⁻ ; Cl ⁻ acts as weak ligand therefore does not cause forced pairing, thus electrons		
	will remain unpaired hence paramagnetic.		
	In [Ni(CO) ₄]; CO acts as strong ligand therefore causes forced pairing, thus electrons will		
	become paired hence diamagnetic.		
24	(a)		
	(i) $\langle \rangle$ -CH ₂ Cl		
	(11) \sim $CH_2CH_2CH_3$		
	Br		
	(b) (i) CH ₃ –I		
	(ii) CH ₃ –Cl		
25	(i) As primary amines form inter molecular H – bonds, but tertiary amines don't form H – bonds.	1	
	(ii) Aniline forms salt with Lewis acid AlCl _{3.}	1	
	(iii) This is because of the combined effect of hydration and inductive effect (+I effect).	1	
	Or		
25	(i) $C_6H_5NO_2 \xrightarrow{Sn+HCl} C_6H_5NH_2 \xrightarrow{NaNO_2+HCl}; 273K \rightarrow C_6H_5N_2^+Cl^- \xrightarrow{H_2O} C_6H_5OH$	1/2+1/2+1/2	
	A B C		
	H_2O/H^+ NH_3 $Br_2 + KOH$		
	(ii) CH ₃ CN \longrightarrow CH ₃ COOH $$ CH ₃ CONH ₂ \longrightarrow CH ₃ NH ₂	1/2+1/2+1/2	
	A B C		
26	(i) Peptide linkage is an amide formed between –COOH group and –NH ₂ group (-CO-NH-)	1	
	(ii) Specific sequence of amino acids in a polypeptide chain is said to be the primary structure	1	
	of the protein.		

	(iii) When a protein in its native form, is subjected to change in temperature or change in pH,			
	protein loses its biological activity. This is called denaturation of protein			
27	(i) (a) dedicated towards work/ kind/ compassionate (any two).	1		
	(b) Dutiful / caring / humane in the large interest of public health in rural area.	1⁄2		
	(any other suitable value)			
	(ii) Narcotic analgesics	1/2		
	(iii) Aspartame / Saccharin / Alitame / Sucrolose.(any one)	1		
28	(a)			
	(i) Molarity is defined as number of moles of solute dissolved in one litre of solution.	1		
	(ii) It is equal to elevation in boiling point of 1 molal solution.	1		
	(b) For isotonic solutions: π urea = π glucose	1/2		
	$\frac{W_{urea}}{M} = \frac{W_{Glucose}}{M}$ (As volume of solution is same)	1⁄2		
	Murea X V s M Glucose X V s			
	$W_{urea} = W_{Glucose} = 15g = W_{Glucose}$			
	$\frac{1}{M_{\text{urea}}} = \frac{1}{M_{\text{Glucose}}} \text{Or} \frac{1}{60 \text{g mol}^{-1}} = \frac{1}{180 \text{g mol}^{-1}}$	1		
	W _{Glucose} = $\frac{15g \times 180g \text{ mol}^{-1}}{60g \text{ mol}^{-1}} = 45g$	1		
	OR			
28	(a) It shows positive deviation	1		
20	It is due to weaker interaction between acetone and ethanol than ethanol-ethanol interactions	1		
	It is due to weaker interaction between accone and emanor than emanor-emanor interactions.	1		
	(b) Given: $W_{p} = 10g W_{q} = 100g W_{t} = 90g M_{p} = 180g/mol $ & $d = 1.2g/m J$			
	(b) Given: $w_B = 10g w_S = 100g$, $w_A = 90g w_B = 100g/mor a u = 1.2g/m L$			
	- Wt % x density x 10			
	$M = \frac{M}{Mol.wt.}$	1/2		
		72		
	$M = \frac{10 \times 1.2 \times 10}{180} = 0.66 \text{ M} \text{ or } 0.66 \text{ mol/L}$	1		
		1		
	$m = \frac{W_B \times 1000}{W_B \times 1000}$	1/		
	$M_B \times W_A (ing)$	72		
	10 x 1000			
	10 X 1000	1		
	$m = \frac{1}{180 \times 90}$	1		
	$m = \frac{1}{180 \times 90}$ = 0.61m or 0.61mol/kg (or any other suitable method)	1		

29	(a) (i) $\operatorname{Cr}_2\operatorname{O7}^{2-} + 2 \operatorname{OH}^- \longrightarrow 2\operatorname{Cr}\operatorname{O4}^{2-} + \operatorname{H}_2\operatorname{O}$			
	(ii) $MnO_4^- + 4H^+ + 3e^- \longrightarrow MnO_2 + 2H_2$	0	1	
	(b) (i) Zn / Zn^{2+} has fully filled d orbitals.			
	(ii) This is due to smaller ionic sizes / higher	ionic charge and availability of d orbitals.	1	
	(iii) because Mn $^{+2}$ is more stable(3d ⁵) than	$Mn^{3+}(3d^4).\ Cr^{+3}$ is more stable due to $t_2g^3\ /\ d^3$	1	
	configuration.			
	C	Dr		
29	(i)			
	Lanthanoids	Actinoids		
	Atomic / ionic radii does not show much	Atomic / ionic radii show much variation /		
	variation $/+3$ is the most common oxidation	Besides +3 oxidation state they exibit	1	
	state, in few cases $+2 \& +4$	+4,+5,+6,+7 also.		
	They are quite reactive	Highly reactive in finely divided state	1	
	(Any two Points)			
	(ii) Cerium (Ce ⁴⁺)			
	(iii) $MnO_4^- + 8H^+ + 5e^- \longrightarrow Mn^{2+} + 4H_2O$ (iv) Mn^{3+} is more paramgnetic			
	Because Mn^{3+} has 4 unpaired electrons (3d ⁴) therefore more paramagnetic whereas Cr^{3+} has			
	unpaired electrons (3d ³).			
30	(a) (i)			
	CH ₃ CN			
	H ² OH			
			1	
	(ii) CH ₃ CH=N–OH			
	(iii)			
	2 CH_{2} - CHO $\stackrel{\text{dil. NaOH}}{\longleftarrow}$ CH ₂ - CH-CH ₂ - CH	CHO	1	
	(b) (i) Add neutral FeCl ₃ in both the solutions, phenol forms violet colour but benzoic acid does			
	$(1) \mathbf{T}_{\mathbf{r}} \mathbf{H}_{\mathbf{r}} \mathbf{h}_{\mathbf{r}$		1	
	(ii) Tollen's reagent test : Add ammoniacal solution of silver nitrate (Tollen's reagent) in			
	both the solutions propanal gives silver mirror whereas propanone does not.			
	(or any other correct test)			



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