# ISC Solved Paper 2018 <br> Chemistry <br> Class-XII 

(Maximum Marks : 80)
(Time allowed : Three hours)

Candidates are allowed an additional 15 minutes for only reading the paper. They must NOT start writing during this time.

All questions are compulsory
Question 1 is of 20 marks having four sub parts, all of which are compulsory. Question numbers 2 to 8 carry 2 marks each, with any two questions having internal choice.
Question numbers 9 to 15 carry 3 marks each, with any two questions having an internal choice.
Question numbers 16 to 18 carry 5 marks each, with an internal choice.
All working, including rough work, should be done on the same sheet as, and adjacent to the rest of the answer.
The intended marks for questions or parts of questions are given in brackets [J.
Balanced equations must be given wherever possible and diagrams where they are helpful.
When solving numerical problems, all essential working must be shown.
In working out problems, use the following data:.
Gas constant $R=1.987 \mathrm{cal} \mathrm{deg}^{-1} \mathrm{~mol}^{-1}=8.314 \mathrm{JK}^{-1} \mathrm{~mol}^{-1}=0.0821 \mathrm{dm}^{3} \mathrm{~atm} .11 \mathrm{~atm}=/ \mathrm{dm}^{3} \mathrm{~atm}=101.3 \mathrm{~J}$. 1 Faraday $=96500$ coulombs. Avogadro's number $=6.023 \times 10^{23}$

1. (a) Fill in the blanks by choosing the appropriate word/words from those given in the brackets: [ $4 \times 1$ ] (square pyramidal, electrical, $74,26, s p^{3} d^{2}$, $s p^{3} d$, chemical, 68, 32, tetrahedral, yellow, white, iodoform, Lucas)
(i) A Galvanic cell converts $\qquad$ energy into $\qquad$ energy.
*(ii) The percentage of unoccupied spaces in bcc and fcc arrangements are $\qquad$ and $\qquad$ respectively
(iii) Propan-2-ol on reaction with iodine and sodium hydroxide gives $\qquad$ precipitate and the reaction is called $\qquad$ test.
*(iv) The geometry of $\mathrm{XeOF}_{4}$ molecule is $\qquad$ and the hybridisation of xenon atom in the molecule is $\qquad$ .
(b) Complete the following statements by selecting the correct alternative from the choices given :
$[4 \times 1]$
(i) During the course of an $S_{N} 1$ reaction, the intermediate species formed is :
(1) a carbocation
(2) a free radical
(3) a carbanion
(4) an intermediate complex
*(ii) Purification of aluminium by electrolytic refining is called :
(1) Serpeck's process
(2) Hoope's process
(3) Hall's process
(4) Baeyer's process
(iii) An aqueous solution of urea freezes at $-0.186^{\circ} \mathrm{C}, \mathrm{K}_{f}$ for water $=1.86 \mathrm{~K} \mathrm{~kg} \mathrm{~mol}^{-1}, \mathrm{~K}_{b}$ for water $=0.512 \mathrm{~K} \mathrm{~kg} \mathrm{~mol}^{-1}$. The boiling point of urea solution will be :
(1) 373.065 K
(2) 373.186 K
(3) 373.512 K
(4) 373.0512 K
(iv) In the dehydration of alcohols to alkenes by heating with concentrated sulphuric acid, the initiation step is :
(1) formation of carbocation
(2) formation of an ester
(3) protonation of alcohol molecule
(4) elimination of water
(c) Match the following:
$[4 \times 1]$
(i) Rate constant
(a) Dialysis
(ii) Biodegradable polymer
(b) Glycine
(iii) Zwitter ion
(c) Arrhenius
equation
(iv) Purification of colloids
(d) PHBV
(d) Answer the following questions:
[4×2]
(i) (1) Why does the density of transition elements increase from Titanium to Copper ?
(at, no. $\mathrm{Ti}=22, \mathrm{Cu}=29$ )
(2) Why is zinc not regarded as a transition element?
(at, no. $\mathrm{Zn}=30$ )
(ii) Identify the compounds $\mathrm{A}, \mathrm{B}, \mathrm{C}$ and D .

(iii) Calculate the osmotic pressure of a solution prepared by dissolving 0.025 g of $\mathrm{K}_{2} \mathrm{SO}_{4}$ in 2.0 litres of water at $25^{\circ} \mathrm{C}$ assuming that $\mathrm{K}_{2} \mathrm{SO}_{4}$ is completely dissociated, (mol, wt. of $\mathrm{K}_{2} \mathrm{SO}_{4}=$ $174 \mathrm{~g} \mathrm{~mol}^{-1}$ )
(iv) What type of isomerism is shown by the following coordination compounds :
[ $\left.\mathrm{Pt} \mathrm{Cl}_{2}\left(\mathrm{NH}_{3}\right)_{4}\right] \mathrm{Br}_{2}$ and $\left[\mathrm{Pt} \mathrm{Br}_{2}\left(\mathrm{NH}_{3}\right)_{4}\right] \mathrm{Cl}_{2}$.
Write their 1UPAC names.
Ans. (a) (i) chemical to electrical.
(iii) yellow and Iodoform..
(b) (i) (1) a carbocation.
(iii) (4) 373.0512 K
(iv) (1) Formation of carbocation.
(c) (i) Rate constant $\rightarrow$ Arrhenius equation.
(ii) Biodegradable polymer $\rightarrow \mathrm{PHBV}$.
(iii) Zwitter ion $\rightarrow$ Glycine.
(iv) Purification of colloids $\rightarrow$ Dialysis.
(d) (i) (1) In transition elements the electrons are being added to the inner shells ( $d$ or $f$ sub-shells) where they are very effective at screening the outer electrons from the nuclear charge. This trend in density can be explained by the small and irregular decrease in metallic radii coupled with the relative increase in atomic mass.
(2) Zinc is not a transition metal because its stable ion $\mathrm{Zn}^{2+}$ has completely filled $d$-orbitals. Transition metals are compounds that form at least one stable ion with incomplete $d$-structure. The electronic configuration of Zn and $\mathrm{Zn}^{2+}$ is $[\mathrm{Ar}] 3 \mathrm{~d}^{10} 4 \mathrm{~s}^{2}$ and $[\mathrm{Ar}] 3 \mathrm{~d}^{10}$, respectively.
(ii)

$\xrightarrow{\mathrm{Br}_{2} / \mathrm{KOH}} \mathrm{CH}_{3} \mathrm{NH}_{2}$
Methyl amine
(iii) Osmotic pressure can be calculated as $\pi=\frac{m R T}{(M V)}$, where $m$ is the mass of the compound, R is the Gas constant, T is the temperature, and V is the volume. $m=$ $0.025 \mathrm{gm}, \mathrm{R}=8.31 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}, \mathrm{~T}=25^{\circ} \mathrm{C}$ or $298 \mathrm{~K}, \mathrm{M}=174 \mathrm{gm} \mathrm{mol}^{-1}$, and $\mathrm{V}=2 \mathrm{~L}$ or $2000 \times 10^{-6} \mathrm{~m}^{3}$.
$\pi=\frac{0.025 \times 8.31 \times 298}{174 \times 0.002}$
Thus, $\pi=177.9 \mathrm{~N} \mathrm{~m}^{-2}$, or $\pi=1779$ dyne $\mathrm{cm}^{-2}$, or $\pi=1779 \mathrm{~Pa}$.
(iv) They will exhibit geometric isomerism (cisand trans-).
The IUPAC names of $\left[\mathrm{PtCl}_{2}\left(\mathrm{NH}_{3}\right)_{4}\right]$
$\mathrm{Br}_{2}$ and $\left[\mathrm{PtBr}_{2}\left(\mathrm{NH}_{3}\right)_{4}\right] \mathrm{Cl}_{2}$ are
Tetraamminedichloroplatinum
(IV)bromide and

Tetraamminedibromoplatinum (IV)
chloride, respectively.
$\left[\mathrm{PtCl}_{2}\left(\mathrm{NH}_{3}\right)_{4}\right] \mathrm{Cl}$ shows geometrical isomerism :



Trans
Cis
$\left[\mathrm{PtBr}_{2}\left(\mathrm{NH}_{3}\right)_{4}\right] \mathrm{Br}$ shows geometrical isomerism :


Cis
Trans
2.(a) (i) Write the rate law expression for the reaction $A+B+C \rightarrow D+E$, if the order of reaction is first, second and zero with respect to $A, B$ and C, respectively.
(ii) How many times the rate of reaction will increase if the concentration of $A, B$ and $C$ are doubled in the equation given in (i) above?

OR
(b) The rate of reaction becomes four times when the temperature changes from 293 K to 313 K. Calculate the energy of activation ( $\mathrm{E}_{a}$ ) of the reaction assuming that it does not change with temperature. $\left(\mathrm{R}=8.314 \mathrm{~K}^{-1} \mathrm{~mol}^{-1}\right)$.

Ans. (a) (i) $\frac{d x}{d t}=k[\mathrm{~A}]^{1}[\mathrm{~B}]^{2}[\mathrm{C}]^{0}$
(ii) When the concentration doubles the reaction rate will increase by a factor of 8 as the order of the reaction is 3 .
(b) Given $\frac{k_{2}}{k_{1}}=4, \mathrm{~T}_{2}=313 \mathrm{~K}, \mathrm{~T}_{1}=293 \mathrm{~K}$, we have $\log \left(\frac{k_{2}}{k_{1}}\right)=\frac{\mathrm{E}_{a}}{2.303 \mathrm{R}}\left(\frac{\mathrm{T}_{2}-\mathrm{T}_{1}}{\mathrm{~T}_{2} \mathrm{~T}_{1}}\right)$,

$$
\text { or } \log 4=\frac{\mathrm{E}_{a}}{2.303 \times \frac{8.314 \mathrm{~J}}{\mathrm{~mol} / \mathrm{K}}} \times \frac{313 \mathrm{~K}-293 \mathrm{~K}}{313 \mathrm{~K} \times 293 \mathrm{~K}}, \text { or }
$$

Activation energy $\mathrm{E}_{a}=52.86 \frac{\mathrm{~kJ}}{\mathrm{~mol}}$
*3. (a) How do antiseptics differ from disinfectants?
(b) State the role of the following chemicals in
the food industry :
(i) Sodium benzoate
(ii) Aspartame

Ans. (a) Antiseptics are applied to the living skin or tissue to prevent infection whereas disinfectants are applied to surfaces, equipments or other nonliving things. They kill microorganisms such as bacteria, viruses, and fungi using chemicals called biocides.
(b) (i) Sodium benzoate is the sodium salt of benzoic acid. It is used as a preservative in the food industry and can be found in acidic foods such as salad dressings, carbonated drinks (as carbonic acid), sauces and juices (citric acid), and pickles (acetic acid).
(ii) Aspartame is used as an intense sweetener in the food industry. It is approximately 200 times sweeter than sugar and is used in soft drinks, low calorie sugar free foods.
4. An aromatic organic compound [A] on heating with $\mathrm{NH}_{3}$ and $\mathrm{Cu}_{2} \mathrm{O}$ at high pressure gives [B|. The compound [B] on treatment with ice cold solution of $\mathrm{NaNO}_{2}$ and HCl gives [C], which on heating with $\mathrm{Cu} / \mathrm{HCI}$ gives compound $[\mathrm{A}]$ again. Identify the compounds $[\mathrm{A}],[\mathrm{BJ}$ and $[\mathrm{C}]$. Write the name of the reaction for the conversion of (B] to [C].
Ans. $\mathrm{A} \xrightarrow[\mathrm{NH}_{3}, \mathrm{Cu}_{2} \mathrm{O}]{\Delta, \mathrm{High}} \mathrm{B} \xrightarrow[\mathrm{HCl}]{\mathrm{NaNO}_{2}} \mathrm{C} \xrightarrow[\mathrm{Cu} / \mathrm{HCl}]{\Delta} \mathrm{A}$

*5. Write the names of the monomers for each of the following polymers:
(a) Bakelite
(b) Nylon-2-nylon-6
[2]
6. Name the purine bases and pyrimidine bases present in RNA and DNA.
Ans. RNA : Purine base-Adenine and Guanine, Pyrimidine base-Uracil, Cytosine.
DNA : Purine base-Adenine and Guanine, Pyrimidine base-Thymine and Cytosine.
7. (a) How will you obtain the following? (Give balanced equation.)
(i) Picric acid from phenol.
(ii) Ethyl chloride from diethyl ether.

OR
(b) How will you obtain the following? (Give balanced equation.)
(i) Anisole from phenol
(ii) Ethyl acetate from ethanol

Ans. (a) (i) Picric acid from phenol


Phenol
2, 4, 6 - trinitrophenol (Picric acid) water
(ii)


Diethyl ether
Ethyl chloride
*Out of Syllabus
(b) (i) Anisole from phenol

(ii) Ethylacetate from ethanol

$$
\underset{\substack{\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH} \\ \text { Ethanol } \\ \text { Acetic acid } \\ \mathrm{CH}_{3} \mathrm{COOH}} \underset{\text { Ethylaetate }}{\mathrm{CH}_{3} \mathrm{COOC}_{2} \mathrm{H}_{5}}+\underset{\text { Water }}{\mathrm{H}_{2} \mathrm{O}}}{ }
$$

8. $40 \%$ of a first order reaction is completed in 50 minutes. How much time will it take for the completion of $80 \%$ of this reaction?
Ans. $40 \%$ of first order reaction is completed in 50 min .
$k=\left(\frac{2.303}{t}\right) \log \frac{a}{(a-x)}$, when
$x=0.4 a$, and $t=50 \mathrm{~min}, k=\left(\frac{2.303}{50}\right) \log \frac{1}{0.6}$,
$k=0.010218 \min ^{-1}$.
$t=\left(\frac{2.303}{0.010218}\right) \log \left(\frac{a}{a-0.8 a}\right)$
$t=157.5 \mathrm{~min}$
9. (a) The freezing point of a solution containing 5.85 g of NaCl in 100 g of water is $-3.348^{\circ} \mathrm{C}$. Calculate van't Hoff factor' $i$ ' for this solution. What will be the experimental molecular weight of NaCl ?
$\left(\mathrm{K}_{f}\right.$ for water $=1.86 \mathrm{~K} \mathrm{~kg} \mathrm{~mol}^{-1}$, at wt. $\mathrm{Na}=23$, $\mathrm{Cl}=35.5$ )

## OR

(b) An aqueous solution containing 12.48 g of barium chloride $\left(\mathrm{BaCl}_{2}\right)$ in 1000 g of water, boils at $100.0832^{\circ} \mathrm{C}$. Calculate the degree of dissociation of barium chloride, ( $\mathrm{K}_{b}$ for water $=0.52 \mathrm{~K} \mathrm{~kg} \mathrm{~mol}^{-1}$, at. wt. $\mathrm{Ba}=137, \mathrm{Cl}=35.5$ )
Ans. (a) Freezing point of water $-3.348^{\circ} \mathrm{C}$ when solute is added. Freezing point of water under normal conditions is $0^{\circ} \mathrm{C}$. Change in the freezing point $\Delta \mathrm{T}_{f}=0-(-3.348)^{\circ} \mathrm{C}$, or $\Delta \mathrm{T}_{f}=3.348^{\circ} \mathrm{C}$, or $\Delta \mathrm{T}_{f}=$ 3.348 K. The van't Hoff factor $i$ is calculated as $i$ $=\frac{\Delta \mathrm{T}_{f}}{\left(\mathrm{~K}_{f} m\right)}$, where $\mathrm{K}_{f}=1.86 \mathrm{~K} \mathrm{~kg} \mathrm{~mol}^{-1}$, molality $m$ is calculated as $m=$ mass of solute/(Molecular weight of solute $\times$ mass of solvent). Mass of solute $=5.85 \mathrm{~g}$, Molecular weight of NaCl is 58.5 $\mathrm{g} \mathrm{mol}^{-1}$ i.e., $(23+35.5) \mathrm{g}$ mol-1, and mass of solvent is 100 g or 0.1 kg . Thus, $m=\frac{5.85}{(58.5 \times 0.1)}$, or $m=1 \mathrm{~mol} \mathrm{~kg}^{-1}$. Using the above values, van' $t$ Hoff factor $i=1.8$.
(b) Change in the boiling point $\Delta \mathrm{T}_{b}=100.0832$ $-100{ }^{\circ} \mathrm{C}$, or $\Delta \mathrm{T}_{b}=0.0832^{\circ} \mathrm{C}$, or $\Delta \mathrm{T}_{b}=0.0832$ K. The van't Hoff factor $i$ is calculated as $i=$ $\frac{\Delta \mathrm{T}_{b}}{\left(\mathrm{~K}_{b} m\right)}$, where $\mathrm{K}_{b}=0.52 \mathrm{~K} \mathrm{~kg} \mathrm{~mol}^{-1}$, molality $m$ is calculated as $m=$ Mass of solute/(Molecular weight of solute $\times$ mass of solvent). Mass of solute $=12.48 \mathrm{~g}$, Molecular weight of $\mathrm{BaCl}_{2}$ is $208 \mathrm{~g} \mathrm{~mol}^{-1}$ i.e., $(137+2 \times 35.5) \mathrm{g} \mathrm{mol}^{-1}$, and mass of solvent is 1000 g or 1 kg . Thus, $m=$ $12.48 /(208 \times 1)$. Using the above values, van't Hoff factor $i=2.67$.
The degree of dissociation $\alpha$ is related to the van't Hoff factor $i$, as $i=1+\alpha(n-1)$, or $\alpha=$ $\frac{(i-1)}{(n-1)}$ where $n$ is the number of ions, and $n=$ 3 , as there are 3 ions in $\mathrm{BaCl}_{2}$. Using the above values, degree of dissociation $\alpha=\frac{(2.67-1)}{(3-1)}$, or
$\alpha=0.835$.
*10. Examine the defective crystal given below and answer the question that follows :

| $\mathbf{A}^{+}$ | $\mathbf{B}^{-}$ | $\mathbf{A}^{+}$ | $\mathbf{B}^{-}$ | $\mathbf{A}^{+}$ |
| :---: | :---: | :---: | :---: | :---: |
| $\mathbf{B}^{-}$ |  | $\mathbf{B}^{-}$ | $\mathbf{A}^{+}$ | $\mathbf{B}^{-}$ |
| $\mathbf{A}^{+}$ | $\mathbf{B}^{-}$ | $\mathbf{A}^{+}$ |  | $\mathbf{A}^{+}$ |
| $\mathbf{B}^{-}$ | $\mathbf{A}^{+}$ | $\mathbf{B}^{-}$ | $\mathbf{A}^{+}$ | $\mathbf{B}^{-}$ |

State if the above defect is stoichiometric or nonstoichiometric. How does this defect affect the density of the crystal? Also, write the term for this type of defect.
*11. Give reason for each of the following :
(a) For ferric hydroxide sol the coagulating power of phosphate ion is more than chloride ion
(b) Medicines are more effective in their colloidal form.
(c) Gelatin is added to ice creams.
12. (a) For the complex-ion $\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{3-}$, state : [3]
(i) the type of hybridisation.
(ii) the magnetic behaviour.
(iii) the oxidation number of the central metal atom.
(b) Write the IUPAC name of $\left[\mathrm{Co}(\mathrm{en})_{2} \mathrm{Cl}_{2}\right]^{+}$ion and draw the structures of its geometrical burners.
Ans. (a) (i) The hybridisation of $\left[\mathrm{Fe}\left(\mathrm{CN}_{6}\right)\right]^{3-}$ is $d^{2} s p^{3}$ because of strong field ligand, $\mathrm{CN}_{-}$.
(ii) The magnetic behaviour is paramagnetic with an octahedral geometry and one electron.
(iii) The oxidation number of the central metal atom $\mathrm{Fe}=+3$.
(b) Cis and Trans isomers of Dichlorobis ethylenediamine cobalt (III) chloride.


Trans


Cis
The IUPAC name of $\left[\mathrm{Co}(\mathrm{en})_{2} \mathrm{Cl}_{2}\right]^{+}$ion is Dichlorobis (ethylenediamine) Cobalt (III) Chloride.
13. (a) Explain why :
(i) $\mathrm{Mn}^{2+}$ is more stable than $\mathrm{Fe}^{2+}$ towards oxidation to +3 state.
(At. no. of Mn-25, Fe-26)
(ii) Transition elements usually form coloured ions.
(iii) Zr and Hf exhibit similar properties.
(At. no. of $\mathrm{Zr}=40, \mathrm{Hf}=72$ )
OR
(b) Complete and balance-the following chemical equations :
(i)

$\qquad$ $+$ $\qquad$ $+$
$\qquad$ $+$ $\qquad$
(ii) $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}+\mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{H}_{2} \mathrm{~S} \rightarrow$ $\qquad$ $+$ $\qquad$
$+$ $\qquad$ $+$
(iii) $\mathrm{KMnO}_{4}+\mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{FeSO}_{4} \rightarrow$ $\qquad$ $+$ $\qquad$
$+$ $\qquad$ $+$ $\qquad$
$\qquad$
Ans. (a) (i) The electronic configuration of $\mathrm{Mn}^{2+}$ and $\mathrm{Fe}^{2+}$ are $[\mathrm{Ar}]^{18} 3 \mathrm{~d}^{5}$ and $[\mathrm{Ar}]^{18} 3 \mathrm{~d}^{6}$, respectively. $\mathrm{Mn}^{2+}$ has a stable state $3 \underline{\mathrm{~d}}^{5}$ configuration and is thus resistant towards oxidation to $\mathrm{Mn}^{3+}$. On the other hand,
$\mathrm{Fe}^{2+}$ has a $3 \underline{d}^{6}$ configuration and by losing one electron, it can change to a more stable $3 \underline{\mathrm{~d}}^{5}$ configuration. Therefore, $\mathrm{Fe}^{2+}$ is easily oxidised to $\mathrm{Fe}^{3+}$ oxidation state.
(ii) Transition metals have partially filled $d$-orbitals. When they start bonding with other ligands, the $d$-orbitals split and become non-degenerate which means they have different energy levels.
The electrons are able to absorb certain wavelengths of electromagnetic radiation to get to higher energy level orbitals and during the downward transition emit light which gives them a coloured appearance.
(iii) Zr and Hf exhibit similar properties due to lanthanide contraction. Electrons present in the $f$-subshell are not too effective in shielding, due to which the size gets constricted with atomic number and so the size of Zr and Hf have nearly the same size and properties.
(b) (i) $2 \mathrm{KMnO}_{4}+10 \mathrm{KI}+8 \mathrm{H}_{2} \mathrm{SO}_{4} \longrightarrow 5 \mathrm{I}_{2}+2$
$\mathrm{MnSO}_{4}+6 \mathrm{~K}_{2} \mathrm{SO}_{4}+8 \mathrm{H}_{2} \mathrm{O}$
(ii) $\mathrm{K}_{2} \mathrm{Cr}_{2} \mathrm{O}_{7}+4 \mathrm{H}_{2} \mathrm{SO}_{4}+3 \mathrm{H}_{2} \mathrm{~S} \longrightarrow \mathrm{~K}_{2} \mathrm{SO}_{4}+$ $\mathrm{Cr}_{2}\left(\mathrm{SO}_{4}\right)_{3}+3 \mathrm{~S}+7 \mathrm{H}_{2} \mathrm{O}$
(iii) $2 \mathrm{KMnO}_{4}+8 \mathrm{H}_{2} \mathrm{SO}_{4}+10 \mathrm{FeSO}_{4} \longrightarrow 5$
$\mathrm{Fe}_{2}\left(\mathrm{SO}_{4}\right)_{3}+2 \mathrm{MnSO}_{4}+\mathrm{K}_{2} \mathrm{SO}_{4}+8 \mathrm{H}_{2} \mathrm{O}$
14. (a) Arrange the following in the increasing order of their basic strength :

$$
\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{NH}_{2}, \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{NH}_{2},\left(\mathrm{C}_{2} \mathrm{H}_{5}\right)_{2} \mathrm{NH}
$$

(b) Give a balanced chemical equation to convert methyl cyanide to ethyl alcohol,
(c) What happens when benzene diazonium chloride reacts with phenol in weak alkaline medium?
(Give balanced equation)
Ans. (a) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{NH}_{2}<\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{NH}_{2}<\left(\mathrm{C}_{2} \mathrm{H}_{5}\right)_{2} \mathrm{NH}$
$\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{NH}_{2}$ will be least basic because of the delocalisation of the lone pair of electron present on the N -atom over the benzene ring due to the -R effect of the $\mathrm{C}_{6} \mathrm{H}_{5}$ group. On the other hand, the alkyl group $\mathrm{C}_{2} \mathrm{H}_{5}$ will tend to increase the electron density on the N -atom making it more easily available for the donation to a proton.
(b) Methylcyanide to Ethylalcohol
$\underset{\text { methylcyanide }}{\mathrm{CH}_{3} \mathrm{CN}+2 \mathrm{H}_{2} \mathrm{O} \xrightarrow[-\mathrm{NH}_{4} \mathrm{Cl}]{\mathrm{H}^{+} / \mathrm{HCl}} \mathrm{CH}_{3} \mathrm{COOH} \xrightarrow[\text { acetic acid }]{4[\mathrm{H}]} \mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}}$ ethylalcohol
(c) Benzene diazonium chloride reacts with phenol in weak alkaline medium.
 Chloride

*15. Name the sulphide ore of Copper. Describe how pure copper is extracted from this ore.
16. (a) (i) Calculate the emf and $\Delta G^{\circ}$ for the cell reaction at $25^{\circ} \mathrm{C}$ :
$\mathbf{Z n}(\mathbf{s})\left|Z n_{(a q)}^{2+}\right|\left|C d_{(a q)}^{2+}\right| \mathbf{C d}_{(s)}$

$$
(0.1 \mathrm{M})(0.01 \mathrm{M})
$$

Given $\mathrm{E}_{\mathrm{Zn}}{ }^{2+} / \mathrm{Zn}=-0.763$ and $\mathrm{E}_{\mathrm{Cd}}^{\circ}{ }^{2+} / \mathrm{Cd}=-$ 0.403 V
(ii) Define the following terms:
(1) Equivalent conductivity
(2) Corrosion of metals

OR
(b) (i) The specific conductivity of a solution containing 5 g of anhydrous $\mathrm{BaCl}_{2}$ (mol. wt. $=$ 208) in $1000 \mathrm{~cm}^{3}$ of a solution is found to be $0.0058 \mathrm{ohm}^{-1} \mathrm{~cm}^{-1}$. Calculate the molar and equivalent conductivity of the solution.
(ii) What is an electrochemical series? How is it useful in predicting whether a metal can liberate hydrogen from acid or not?
Ans. (a) (i) $\mathrm{E}_{\text {cell }}^{\circ}=E_{\text {cathode }}^{\circ}-E_{\text {anode }}^{o \text { Right }}$
$\mathrm{E}_{\text {cell }}^{\circ}=-0.403-(-0.763) \mathrm{V}$, or $\mathrm{E}_{\text {cell }}^{\circ}=0.36$ V.
$\Delta \mathrm{G}^{\circ}=-n \mathrm{FE}_{\text {cell }}^{\circ}$
$\Delta \mathrm{G}^{\circ}=-(2 \mathrm{~mol})\left(96485 \mathrm{C} \mathrm{mol}^{-1}\right)\left(0.36 \mathrm{~J} \mathrm{C}^{-1}\right)$ $\Delta G^{\circ}=-69.469 \mathrm{~kJ}$
(ii) (1) Equivalent conductivity is defined as conducting power of all the ions produced by dissolving 1 g equivalent of an electrolyte in solution. Equivalent conductivity $\Lambda_{e}=1000 \mathrm{k} / \mathrm{c}$, where $k$ is the specific conductivity and c is the concentration in molarity.
(2) Corrosion of metals is the deterioration of a metal as a result of chemical reactions between the metal and the surrounding environment which converts a refined metal to a more chemically stable form such as oxide, hydroxide, or sulphide.
(b) (i) Molar conductivity $\Lambda_{m}=\frac{k}{c}$, where $k$ is the specific conductivity and $c$ is the concentration in molarity. $k=0.0058 \mathrm{ohm}^{-1} \mathrm{~cm}^{-1}$. Mass of $\mathrm{BaCl}_{2} m_{a}$
$=5 \mathrm{~g}$, volume of solvent $\mathrm{V}=1 \mathrm{~L}$, molecular weight of $\mathrm{BaCl}_{2} \mathrm{M}=208 \mathrm{~g} \mathrm{~mol}^{-1}$. Thus, $c$ $=\frac{\left(\frac{m_{a}}{M}\right)}{V}$, and hence, $\Lambda_{m}=\frac{0.0058}{0.024}$, or $\Lambda_{m}$ $=24.167 \mathrm{ohm}^{-1} \mathrm{~m}^{-1} \mathrm{M}^{-1}$.
Equivalent conductivity $\Lambda_{e}=1000 \Lambda_{m}$, hence $\Lambda_{e}=24.167 \times 10^{3} \mathrm{ohm}^{-1} \mathrm{~m}^{-1} \mathrm{M}^{-1}$
(ii) Electrochemical series is a series of chemical elements in order of their standard electrode potential. Hydrogen electrode is considered as having zero electrode potential. Electro-positive elements are the ones having greater tendency than Hydrogen to lose electrons to their solution. Thus they lie above Hydrogen in the series. Those that gain electrons from their solutions are electro-negative elements lie below Hydrogen in the series. Thus electro-positive elements tend to replace Hydrogen from acids.
*17. (a) (i) Explain why :
(1) Nitrogen does not form pentahalides.
(2) Helium is used for filling weather balloons.
(3) $I C l$ is more reactive than $I_{2}$.
(ii) Draw the structures of the following:
(1) $\mathrm{HClO}_{4}$
(2) $\mathrm{H}_{3} \mathrm{PO}_{3}$

## OR

*(b) (i) Explain why :
(1) Mercury loses its meniscus in contact with ozone.
(2) Halogens are coloured and the colour deepens on moving down in the group from fluorine to iodine.
(3) Hydride of sulphur is a gas while hydride of oxygen is a liquid.
(ii) Complete and balance the following reactions:
(1) $\mathrm{NaCl}+\mathrm{MnO}_{2}+\mathrm{H}_{2} \mathrm{SO}_{4} \rightarrow+{ }_{+}^{+}+$ ${ }^{+}+$
(2) $\mathrm{KMnO}_{4}+\mathrm{SO}_{2}+\mathrm{H}_{2} \mathrm{O} \rightarrow+\ldots+\ldots+$
18. (a) (i) Give balanced equations for the following reactions :
(1) Benzaldehyde reacts with hydrazine.
(2) Acetic acid reacts with phosphorous pentachloride.
(3) Acetone reacts with sodium bisulphite.
(ii) Give one chemical test each to distinguish between the following pairs of compounds :
(1) Ethanol and acetic acid
(2) Acetaldehyde and benzaldehyde

OR
(b) (i) Write chemical equations to illustrate the following name reactions:
(1) Clemmensen's reduction
(2) Rosenmund's reduction
(3) HVZ reaction
(ii) Explain why :
(1) Acetaldehyde undergoes aldol condensation, but formaldehyde does not.
(2) Acetic acid is a weaker acid as compared to formic acid.

Ans. (a) (i) (1) Benzaldehyde with hydrazine


Benzaldehyde hydrazine
Water

Acetone Sodium Bisulphate Sodium 3-hydroxy Propane-1-sulphonate
(ii) (1) Litmus test : Ethanol does not show any change, while with acetic acid, blue Litmus turns to red.
(2) Acetaldehyde gives Iodoform test with $\mathrm{I}_{2}$ and alkalis, while benzaldehyde does not.
(b) (i) (1) Clemmensen's Reduction :

Reduction of ketones to alkanes using zinc amalgam and HCl .

(2) Rosenmund's reduction:

The catalytic hydrogenation of acid chlorides to form aldehydes.

(3) Hell - Volhard - Zelinsky Reaction : Carboxylic acid on reacting with bromine and catalytic amount of phosphorus leads to $\alpha$-bromination of carboxylic acid.

(ii) (1) Acetaldehyde $\left(\mathrm{CH}_{3} \mathrm{CHO}\right)$ undergoes aldol condensation due to the presence of $\alpha$-hydrogen attached to the $\alpha$-carbon but formaldehyde (HCHO) has no $\alpha$-carbon attached to the functional group hence no $\alpha$-hydrogen is available for the reaction to occur.
(2) In Acetic acid, $\mathrm{CH}_{3}$ is an electron donating group, thus it increases
electron density towards $\mathrm{O}-\mathrm{H}$ bond making it difficult to remove the $\mathrm{H}^{+}$ to a base. Formic acid $(\mathrm{HCOOH})$, has
no electron donating group, thus $\mathrm{O}-\mathrm{H}$ bond is more polarised and can easily donate $\mathrm{H}^{+}$ion making it more acidic.

