CHEMISTRY - CET 2021 - VERSION CODE – B2 SOLUTIONS

- 1. Zeta potential is
 - (A) Potential required to bring about coagulation of a colloidal sol.
 - (B) Potential required to give the particle a speed of 1 cm S^{-1}
 - (C) Potential difference between fixed charged layer and the diffused layer having opposite charges
 - (D) Potential energy of the colloidal particles.

Ans (C)

The potential difference between fixed charged layer and the diffused layer having opposite charges is known as zeta potential

2. Which of the following compound on heating given N_2O ?

(A) $Pb(NO_3)_2$ (B) NH_4NO_3 (C) NH_4NO_2 (D) $NaNO_3$ **Ans** (B) $NH_4NO_3 \xrightarrow{\Delta} N_2O + 2H_2O$

3. Which of the following property is true for the given sequence $NH_3 > PH_3 > AsH_3 > SbH_3 > BiH_3$?

(A) Reducing property	(B) Thermal stability
(C) Bond angle	(D) Acidic character
Ans GRACE	
(B), (C)	

Thermal stability and bond angle decreases down the group due to increase in bond length.

4. The correct order of boiling point in the following compounds is

(A) $HF > H_2O > NH_3$ (B) $H_2O > HF > NH_3$ (C) $NH_3 > H_2O > HF$ (D) $NH_3 > HF > H_2O$ Ans (B)

Boiling point is directly proportional to molecular mass and strength of hydrogen bond.

5. XeF₆ on partial hydrolysis gives a compound X, which has square pyramidal geometry 'X' IS (A) XeO₃ (B) XeO₄ (C) XeOF₄ (D) XeO₂F₂ Ans (C) XeF₆ + H₂O \longrightarrow XeOF₄ + 2HF

6. A colourless, neutral, paramagnetic oxide of Nitrogen 'P' on oxidation gives reddish brown gas Q. Q on cooling gives colourless gas R. R on reaction with P gives blue solid S. Identify P, Q, R, S, respectively (A) N₂O NO NO₂ N₂O₅
(B) N₂O NO₂ N₂O₄ N₂O₃

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(C) NO NO<sub>2</sub> N<sub>2</sub>O<sub>4</sub> N<sub>2</sub>O<sub>3</sub>

Ans (C)

NO \underset{Q}{\overset{[O]}{\longrightarrow}} \underset{Q}{\overset{Ooling}{\longrightarrow}} \underset{R}{\overset{Cooling}{\longrightarrow}} \underset{R}{\overset{O_4}{\xrightarrow{[H]}}} \underset{S}{\overset{(H)}{\longrightarrow}} \underset{S}{\overset{O_4}{\xrightarrow{[H]}}} \underset{S}{\overset{O_4}{\overset{O_4}{\xrightarrow{[H]}}} \underset{S}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}{\overset{O_4}
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7. Which of the following does not represent property stated against it?

(A) $CO^{+2} < Fe^{+2} < Mn^{+2}$ – Ionic size

- (B) Ti < V < Mn Number of oxidation states
- (C) $Cr^{+2} < Mn^{+2} < Fe^{+2}$ Paramagnetic behaviour
- (D) Sc > Cr > Fe Density

Ans GRACE





(C) and (D) Correct option are both C and D $Cr [Ar]3d^54s^1$ $Cr^{2+}3d^44s^0 = 4$ unpaired electrons $Mn [Ar]3d^54s^2$ $Mn^{2+}3d^54s^0 = 5$ unpaired electrons $Fe [Ar]3d^64s^2$ $Fe^{2+}3d^64s^0 = 4$ unpaired electrons Since both Cr^{2+} and Fe^{2+} contain 4 unpaired electrons, the correct option is C. Option D is also correct because density order given is not correct.

- 8. Which one of the following is correct for all elements from Sc to Cu?
 - (A) The lowest oxidation state shown by them is +2
 - (B) 4S orbital is completely filled in the ground state
 - (C) 3d orbital is not completely filled in the ground state
 - (D) The ions in +2 oxidation states are paramagnetic.

Ans (D)

Paramagnetism is due to the presence of one or more unpaired electrons.

- 9. When the absolute temperature of ideal gas is doubled and pressure is halved, the volume of gas
 - (A) will be half of original volume
- (B) will be 4 times the original volume
- (C) will be 2 times the original volume (D) will be 1/4 times the original volume

Ans (B)

From the combined gas equation,

$$\frac{p_1 V_1}{T_1} = \frac{p_2 V_2}{T_2}$$
$$\frac{p_1 V_1}{T_1} = \frac{\frac{1}{2} p_1 V_2}{2T_1}$$

$$\mathbf{V}_1 = \frac{1}{4} \mathbf{V}_2$$

$$\mathbf{V}_2 = 4\mathbf{V}_1$$

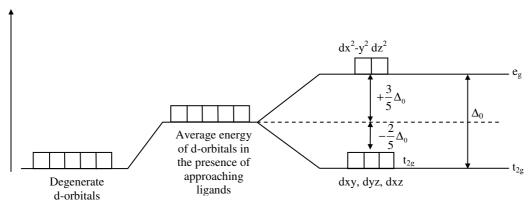
10. Which of the following pairs has both the ions coloured in aqueous solution? [Atomic numbers of Sc = 21, Ti = 22, Ni = 28, Cu = 29, Mn = 25]

(A)
$$Sc^{3+}$$
, Mn^{2+} (B) Ni^{2+} , Ti^{4+} (C) Ti^{3+} , Cu^{+} (D) Mn^{2+} , Ti^{3+}
Ans (D)
 $Sc = 21$, $Sc^{3+} = [Ar] 3d^{0}4s^{0} = no$ unpaired electrons
 $Ti = 22$, $Ti^{3+} = [Ar] 3d^{1}4s^{0} = 1$ unpaired electrons
 $Ti^{4+} = [Ar] 3d^{0}4s^{0}$ no unpaired electrons
 $Ni = 28$, $Ni^{2+} = [Ar] 3d^{8}4s^{2} = 2$ unpaired electrons

Cu = 29, $Cu^+ = [Ar]3d^{10}4s^0 = no$ unpaired electrons

- Mn = 25, $Mn^{2+} = [Ar] 3d^54s^2 = 5$ unpaired electrons
- 11. For the crystal field splitting in octahedral complexes,
 - (A) the energy of the e_g orbitals will decrease by $(3/5)\Delta_0$ and that of the t_{2g} will increase by $(2/5)\Delta_0$
 - (B) the energy of the e_g orbitals will increase by $(3/5)\Delta_0$ and that of the t_{2g} will decrease by $(2/5)\Delta_0$
 - (C) the energy of the e_g orbitals will increase by $(3/5)\Delta_0$ and that of the t_{2g} will increase by $(2/5)\Delta_0$
- (D) the energy of the e_g orbitals will decrease by $(3/5)\Delta_0$ and that of the t_{2g} will decrease by $(2/5)\Delta_0$ Strategic Academic Alliance with





- 12. Peroxide effect is observed with the addition of HBr but not with the addition of HI to unsymmetrical alkene because
 - (A) H-I bond is strong that H-Br and is not cleaved by the free radical
 - (B) H-I bond is weaker than H-Br bond so that iodine free radicals combine to form iodine molecules
 - (C) Bond strength of HI and HBr are same but free radicals are formed in HBr
 - (D) All of these.

Ans (B)

Due to free radical mechanism (secondary free radical)

- 13. The IUPAC name of [Co(NH₃)₅(CO₃)] Cl is
 - (A) Pentaamminecarbonatocobalt (III) Chloride
 - (B) Carbonatopentamminecobalt (III) Chloride
 - (C) Pentaamminecarbonatocobaltate (III) Chloride
 - (D) Pentaammine cobalt (III) Carbonate Chloride

Ans (A)

Pentaamminecarbonatocobalt(III) chloride

14. Homoleptic complexes among the following are

	(A) $K_3[Al(C_2O_4]_3,$	(B) $[\operatorname{CoCl}_2(\operatorname{en})_2]^+$	(C) $K_2[Zn(OH)_4]$	
(A) A only	7	(B) (A) and (B) only		
(C) (A) and (C) only		(D) (C) only		
Ans (C)				

Complex compounds having only one type of ligands are homoleptic complexes

15. The correct order for wavelengths of light absorbed in the complex ions

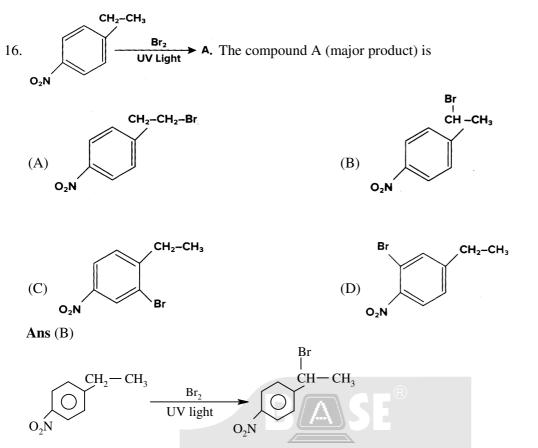
$$\begin{bmatrix} \text{CoCl}(\text{NH}_{3})_{5} \end{bmatrix}^{2^{+}}, \begin{bmatrix} \text{Co}(\text{NH}_{3})_{6} \end{bmatrix}^{3^{+}} \text{ and } \begin{bmatrix} \text{Co}(\text{CN})_{6} \end{bmatrix}^{3^{-}} \text{ is} \\ \text{(A)} \begin{bmatrix} \text{CoCl}(\text{NH}_{3})_{5} \end{bmatrix}^{2^{+}} > \begin{bmatrix} \text{Co}(\text{NH}_{3})_{6} \end{bmatrix}^{3^{+}} > \begin{bmatrix} \text{Co}(\text{CN})_{6} \end{bmatrix}^{3^{-}} \\ \text{(B)} \begin{bmatrix} \text{Co}(\text{NH}_{3})_{6} \end{bmatrix}^{3^{+}} > \begin{bmatrix} \text{Co}(\text{CN})_{6} \end{bmatrix}^{3^{-}} > \begin{bmatrix} \text{CoCl}(\text{NH}_{3})_{5} \end{bmatrix}^{2^{+}} \\ \text{(C)} \begin{bmatrix} \text{Co}(\text{CN})_{6} \end{bmatrix}^{3^{-}} > \begin{bmatrix} \text{CoCl}(\text{NH}_{3})_{5} \end{bmatrix}^{2^{+}} > \begin{bmatrix} \text{Co}(\text{CN})_{6} \end{bmatrix}^{3^{-}} \\ \text{(D)} \begin{bmatrix} \text{Co}(\text{NH}_{3})_{6} \end{bmatrix}^{3^{+}} > \begin{bmatrix} \text{CoCl}(\text{NH}_{3})_{5} \end{bmatrix} > \begin{bmatrix} \text{Co}(\text{CN})_{6} \end{bmatrix}^{3^{-}} \\ \end{bmatrix}$$





Ans (A)

Wavelength of light absorbed is inversely proportional to strength of the ligand.



In presence of uv light, substitution takes place in side chain by free radical halogenation. Halogenation takes place to the carbon which has less number of hydrogen.

17. Bond enthalpies of A_2 , B_2 and AB are in the ratio 2 : 1 : 2. If bond enthalpy of formation of AB is -100 KJ mol^{-1} . The bond enthalpy of B_2 is

4

(A) 100 KJ mol^{-1} (B) 50 KJ mol^{-1} (C) 200 KJ mol^{-1} (D) 150 KJ mol^{-1} Ans (C)

Let bond enthalpy of A_2 be x, B_2 be $\frac{x}{2}$, AB be x

$$A_{2} + B_{2} \rightarrow 2AB$$

$$x + \frac{x}{2} \qquad 2x$$

$$\Delta H_{form} = \frac{1}{2}A_{2} + \frac{1}{2}B_{2} \rightarrow AB$$

$$-100 = \left(\frac{x}{2} + \frac{x}{4}\right) - x$$

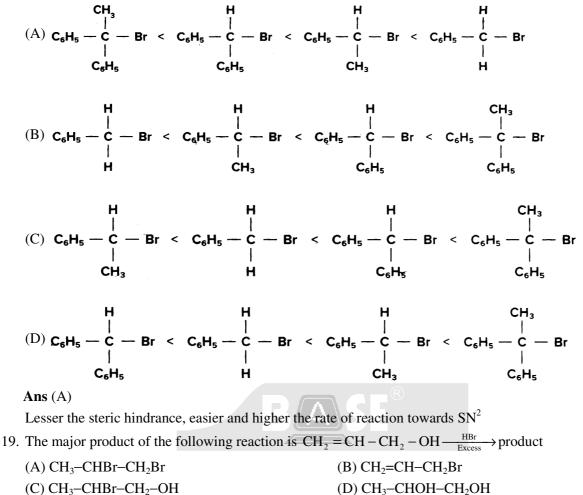
$$x = 400$$

$$B_{2} = \frac{x}{2} = 200 \text{ kJ}$$

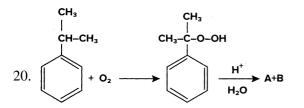




18. The order of reactivity of the compounds $C_6H_5CH_2Br$, $C_6H_5CH(C_6H_5)Br$, $C_6H_5CH(CH_3)Br$ and $C_6H_5C(CH_3)(C_6H_5)Br$ in S_N2 reaction is



$$CH_2 = CH - CH_2OH \xrightarrow{HBr} CH_3 - CH_2H_2Br$$



The product 'A' gives white precipitate when treated with bromine water. The product 'B' is treated with Barium hydroxide to give the product C. The compound C is heated strongly to form product D. The product D is

- (A) 4-Methylpent-3-en-2-one
- (C) 3-Methylpent-3-en-2-one

Ans (A)

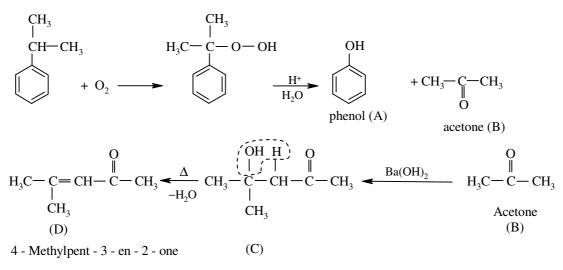
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(B) But-2 enal

(D) 2-Methylbut-2-enal



21. For the reaction $A(g) + B(g) \rightleftharpoons C(g) + D(g); \Delta H = -QKJ$

The equilibrium constant cannot be disturbed by

- (A) Addition of A (B) Addition of D
- (C) Increasing of pressure (D) Increasing of temperature

Ans GRACE

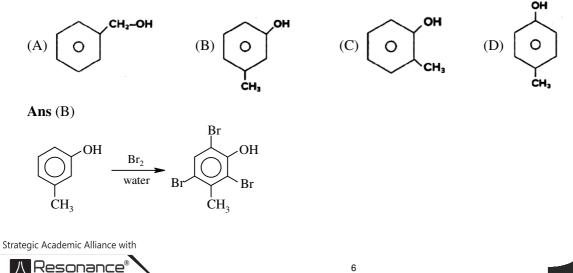
Equilibrium constant is directly proportional to temperature

So, Options (A), (B), (C) are correct.

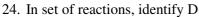
22. An organic compound 'X' on treatment with PCC in dichloromethane gives the compound Y. Compound 'Y' reacts with I2 and alkali to form yellow precipitate of triiodomethane. The compound X is

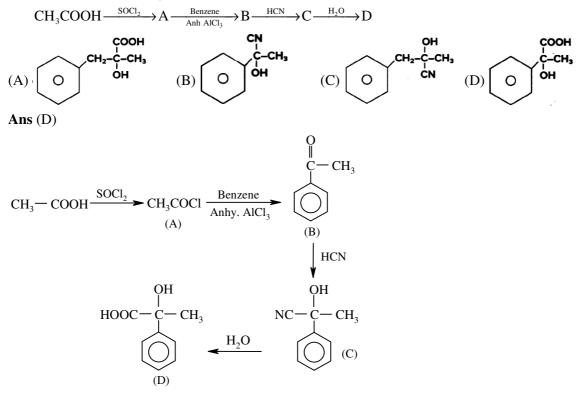
(A) CH₃CHO
(B) CH₃COCH₃ (C) CH₃CH₂OH (D) CH₃COOH
Ans (C)
CH₃CH₂OH
$$\xrightarrow{PCC}_{CH_2Cl_2}$$
 CH₃ $\xrightarrow{O}_{T} \stackrel{I}{C}_{-}$ H $\xrightarrow{I_2}_{NaOH}$ CHI₃
X Iodoform (Yellow precipitate)

23. A compound 'A' (C7H8O) is insoluble in NaHCO3 solution but dissolve in NaOH and gives a characteristic colour with neutral FeCl₃ solution. When treated with Bromine water compound 'A' forms the compound B with the formula C₇H₅OBr₃. 'A' is







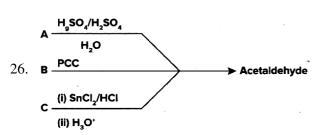


25. K_a values for acids H₂SO₃, HNO₂, CH₃COOH and HCN are respectively 1.3×10^{-2} , 4×10^{-4} , 1.8×10^{-5} and 4×10^{-10} , which of the above acids produces stronger conjugate base in aqueous solution? (A) H₂SO₃ (B) HNO₂ (C) CH₃COOH (D) HCN Ans (D)

7

 $K_a(HCN) = 4 \times 10^{-10}$ $HCN \Longrightarrow H^+ + CN^-$

Weaker the acid, stronger the conjugate base.



A, B and C respectively are

- (A) ethanol, ethane nitrile and ethyne
- (C) ethyne, ethanol and ethane nitrile

$$HC \equiv CH \xrightarrow{H_2SO_4} CH_2 = CHOH \xrightarrow{tautomerises} CH_3 - C - H$$
(A)

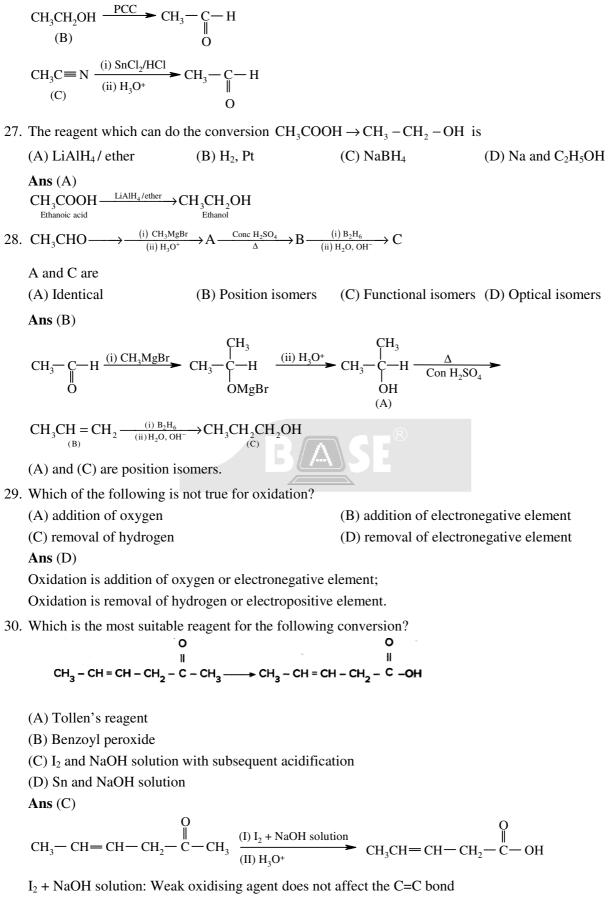
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(B) ethane nitrile, ethanol and ethyne

(D) ethyne, ethane nitrile and ethanol







31. $C_6H_5CH_2Cl \xrightarrow{alc.NH_3} A \xrightarrow{2CH_3Cl} B$

The product B is

(A) N, N-Dimethyl phenyl methanamine

(C) N-Benzyl-N-methyl methanamine

Ans (A)

(B) N, N-Dimethyl benzenamine

(D) phenyl-N, N-dimethyl methanamine

$$C_6H_5CH_2Cl \xrightarrow{alc. NH_3} C_6H_5CH_2NH_2 \xrightarrow{2CH_3Cl} C_6H_5CH_2 \xrightarrow{-2HCl} C_6H_5CH_2 \xrightarrow{-N-CH_3CH_3} C_6H_5CH_3 \xrightarrow{-N-CH_3CH_3} C_6H_5CH_3 \xrightarrow{-N-CH_3CH_3} C_6H_5CH_3 \xrightarrow{-N-CH_3CH_3} C_6H_5CH_3 \xrightarrow{-N-CH_3CH_3} C_6H_3CH_3 \xrightarrow{-N-CH_3} C_6H_3 C_6H_3 \xrightarrow{-N-CH_3} C_6H_3 C$$

Phenyl-N,N-dimethylmethanamine

- 32. The method by which aniline cannot be prepared is
 - (A) Nitration of benzene followed by reduction with Sn and con HCl
 - (B) Degradation of benzamide with bromine in alkaline solution
 - (C) Reduction of nitrobenzene with H₂/Pd is ethanol
 - (D) Potassium salt of phthalimide treated with chlorobenzene followed by the hydrolysis with aqueous NaOH solution

Ans (D)

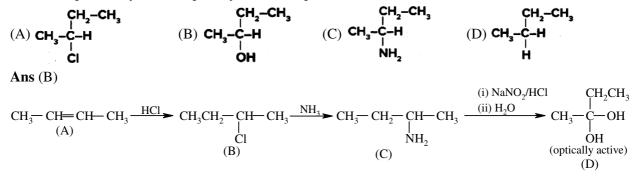
Gabriel phthalimide synthesis not preferred for preparing aniline because amine formation involves nucleophilic substitution of alkyl halides by the anion formed by phthalimide. But aryl halides do not undergo nucleophilic substitution.

- 33. Permanent hardness cannot be removed by
 - (A) Using washing soda
 - (C) Clark's method

Ans (C)

Clark's method is for the removal of temporary hardness.

34. A hydrocarbon A (C_4H_8) on reaction with HCl gives a compound B (C_4H_9Cl) which on reaction with 1 mol of NH₃ gives compound C($C_4H_{10}N$). On reacting with NaNO₂ and HCl followed by treatment with water, compound C yields an optically active compound D. The D is



- 35. RNA and DNA are chiral molecules, their chirality is due to the presence of
 - (A) D-Sugar component (B) L-Sugar component
 - (C) Chiral bases (D) Chiral phosphate ester unit

Ans (A)

Deoxyribose and ribose sugars are D-chiral sugars in DNA and RNA.

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CET2021Chemistry(B2)

- 36. The property of the alkaline earth metals that increases with their atomic number is
 - (A) Ionisation enthalpy (B) Electronegativity
 - (C) Solubility of their hydroxide in water (D) Solubility
- (D) Solubility of their sulphate in water

Ans (C)

The solubility of alkaline earth metal hydroxides increases with increase in atomic number.

37. Primary structure in a nucleic acid chain contains bases as G A T G C _____ The chain which is complementary to this chain is

38. In the detection of II group acid radical, the salt containing chloride is treated with concentrated sulphuric acid, the colourless gas is liberated. The name of the gas is

(A) Hydrogen chloride gas	(B) Chlorine gas
(C) Sulphur dioxide gas	(D) Hydrogen gas
Ans (A)	
$MCl + H_2SO_4 \longrightarrow MHSO_4 + HCl_{(g)}$	$ \mathbf{R} $

39. The number of six membered and five membered rings in Buckminster Fullerene respectively is (A) 20, 12 (B) 12, 20 (C) 14, 18 (D) 14, 11

Ans (A)

- 12:5 Membered ring
- 20:6 Membered ring
- 40. In Chrysoberyl, a compound containing Beryllium, Aluminium and oxygen, oxide ions form cubic close packed structure. Aluminium ions occupy 1/4th of tetrahedral voids and Beryllium ions occupy 1/4th of octahedral voids. The formula of the compound is
 - (A) $BeAlO_4$ (B) $BeAl_2O_4$ (C) Be_2AlO_2 (D) $BeAlO_2$ Ans (B) CCP = N = 4Octahedral void = $N = 1 \times 4 = 4 \times \frac{1}{4} = 1$: Be

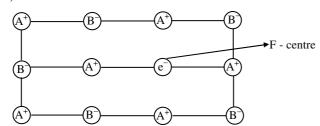
Tetrahedral void = $2N = 2 \times 4 = 2 \times \frac{1}{4} = 2$: Al

- 41. The correct statement regarding defects in solids is
 - (A) Frenkel defect is a vacancy defect
 - (B) Schottky defect is a dislocation defect
 - (C) Trapping of an electron in the lattices leads to the formation of F-centre
 - (D) Schottky defect has no effect on density.





Ans (C)



42. A metal crystallises in BCC lattice with unit cell edge length of 300 pm and density 6.15 g cm⁻³. The molar mass of the metal is

(A) 50 g mol⁻¹ (B) 60 g mol⁻¹ (C) 40 g mol⁻¹ (D) 70 g mol⁻¹ Ans (A) $d = \frac{ZM}{a^3N_A}$ $6.15 = \frac{2 \times M}{(3 \times 10^{-8})^3 \times 6.02 \times 10^{23}}$ $M = \frac{6.15 \times 27 \times 10^{-24} \times 6.02 \times 10^{23}}{2}$ M = 50 g/mol

43. Henry's law constant for the solubility of N_2 gas in water at 298 K is 1.0×10^5 atm. The mole fraction of N_2 in air is 0.8. The number of moles of N_2 from air dissolved in 10 moles of water at 298 K and 5 atm pressure is

(A) 4.0×10^{-4} (B) 4.0×10^{-5} (C) 5.0×10^{-4} (D) 4.0×10^{-6} Ans (A) $p_{N_2} = x_{N_2} \times p_T$ $= 0.8 \times 5$ = 4.0 atm Dalton's law of partial pressures Henry's law, $p_{N_2} = x_{N_2} \times K_H$

 $4 = \frac{n_{N_2}}{10} \times 10^5$:. $n_{N_2} = 4 \times 10^{-4}$

44. A pure compound contains 2.4 g of C, 1.2×10^{23} atoms of H, 0.2 moles of oxygen atoms. Its empirical formula is

(A) C₂HO (B) C₂H₂O₂ (C) CH₂O (D) CHO Ans (D) Number of moles of $C = \frac{2.4}{12} = 0.2$ Number of moles of $H = \frac{1.2 \times 10}{6.0 \times 10} = 0.2$ Number of moles of O = 0.2 mol Empirical formula is CHO.



CET2021Chemistry(B2)

45. Choose the correct statement

- (A) K_H value is same for a gas in any solvent
- (B) Higher the K_H value more the solubility of gas
- (C) K_H value increases on increasing the temperature of the solution
- (D) Easily liquefiable gases usually has lesser K_H values

Ans (C) and (D)

 $K_{\rm H} \propto$ temperature

46. The K_H value (K bar) of Argon (I), Carbon dioxide (II) formaldehyde (III) and methane (IV) are respectively 40.3, 1.67, 1.83×10^{-5} and 0.413 at 298 K. The increasing order of solubility of gas in liquid is

(A) I < II < IV < III(B) III < IV < II < I(C) I < III < II < IV(D) I < IV < II < IIIAns (A)

K_H is inversely proportional to solubility of gas in liquid.

47. The vapour pressure of pure liquids A and B are 450 and 700 mm of Hg at 350 K respectively. If the total vapour pressure of the mixture is 600 mm of Hg, the composition of the mixture in the solution is

(A) $x_A = 0.4$, $x_B = 0.6$ (B) $x_A = 0.6$, $x_B = 0.4$ (C) $x_A = 0.3$, $x_B = 0.7$ (D) $x_A = 0.7$, $x_B = 0.3$ Ans (A) $p_{tot} = p_A^0 x_A + p_B^0 x_B$ $600 = 450x_A + 700(1 - x_A)$ $600 = 450x_{\rm A} + 700 - 700x_{\rm A}$ $100 = -250x_{A}$ $x_{A} = \frac{100}{250} = \frac{2}{5}$ = 0.4 $x_{\rm B} = 1 - x_{\rm A} = 1 - 0.4$ = 0.648. Consider the following electrodes
$$\begin{split} P &= Zn^{2+} (0.0001 \text{ M})/Zn & Q &= Zn^{2+} (0.1 \text{ M})/Zn \\ R &= Zn^{2+} (0.01 \text{ M})/Zn & S &= Zn^{2+} (0.001 \text{ M})/Zn \end{split}$$
 $E_{7n/7n^{2+}}^{0} = -0.76$ V Electrode potentials of the above electrodes in volts are in the order (A) P > S > R > Q(B) S > R > Q > P(C) Q > R > S > P(D) P > Q > R > SAns (C) Electrode potential is directly proportional to concentration of Zn^{+2} ions. 49. The number of angular and radial nodes in 3p orbital respectively are (A) 3, 1 (C) 2, 1 (B) 1, 1 (D) 2, 3 Ans (B) Number of angular nodes = lNumber of radial nodes = n - l - 1 $\begin{bmatrix} s & p & d & f \\ l = 0 & 1 & 2 & 3 \end{bmatrix}$ \therefore For 3p orbital, angular nodes = 1 For 3p orbital, radial nodes = 3 - 1 - 1 = 1







50. The resistance of 0.01 m KCl solution at 298 K is 1500 Ω . If the conductivity of 0.01 m KCl solution at 298 K is 0.146×10^{-3} S cm⁻¹. The cell constant of the conductivity cell in cm⁻¹ is

(C) 0.301 (A) 0.219 (B) 0.291 (D) 0.194 Ans (A) $\kappa = \frac{1}{R} \left(\frac{l}{a} \right)$ $\therefore \frac{l}{a} = \kappa R$ $= 0.146 \times 10^{-3} \times 1500$ $= 219 \times 10^{3} = 0.219 \text{ cm}^{-1}$ 51. $H_{2(g)} + 2AgCl_{(s)} \rightleftharpoons 2Ag_{(s)} + 2HCl_{(aq)}$ E_{cell}^{0} at 25° C for the cell is 0.22 V. The equilibrium constant at 25 °C is (A) 2.8×10^7 (B) 5.2×10^8 (C) 2.8×10^5 (D) 5.2×10^4 Ans (A) $E_{cell}^0 = \frac{0.059}{n} \log K_C$ $0.22 = \frac{0.059}{2} \log K_{c}$ $\log K_{\rm c} = \frac{0.22 \times 2}{0.059} = 7.457$ $K_{\rm C} = 2.8 \times 10^7$

52. For a reaction $A + 2B \rightarrow$ Products, when concentration of B alone is increased half life remains the same. If concentration of A alone is doubled, rate remains the same. The unit of rate constant for the reaction is

(A) S^{-1} (B) $L \mod^{-1} S^{-1}$ (C) $\mod L^{-1} S^{-1}$ (D) atm^{-1}

Ans (A)

Order w.r.t. B = 1 as $t_{1/2}$ remains same when concentration is increased.

Order w.r.t. A = 0 as rate does not change when concentration is doubled.

 \therefore rate = k[A]⁰[B]¹

 \therefore It is a first order reaction.

Hence, unit is S^{-1}

(A) Alkali metals

53. The third ionisation enthalpy is highest in

(B) Alkaline earth metals

(D) Pnictogens

(C) Chalcogens

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Ans (B)
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Alkaline earth metals have ns² configuration.

After the removal of 2^{nd} electron, they get stable noble gas configuration.

54. If the rate constant for a first order reaction is k, the time (t) required for the completion of 99% of the reaction is given by

(A)
$$t = \frac{4.606}{k}$$
 (B) $t = \frac{2.303}{k}$ (C) $t = \frac{0.693}{k}$ (D) $t = \frac{6.909}{k}$





CET2021Chemistry(B2)

Ans (A)

$$t = \frac{2.303}{k} \log \frac{100}{1} = \frac{2.303}{k} \times 2$$

 $t = \frac{4.606}{k}$

55. The rate of a gaseous reaction is given by the expression $k[A][B]^2$. If the volume of vessel is reduced to one half of the initial volume, the reaction rate as compared to original rate is

(B) 1-Ethyl-4-Fluoro-3-nitrobenzene

(D) 5-Ethyl-2-Fluoronitrobenzene

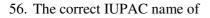
(A)
$$\frac{1}{16}$$
 (B) $\frac{1}{8}$ (C) 8 (D) 16

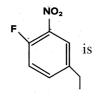
Ans (C)

 $\mathbf{r} = \mathbf{K}[\mathbf{A}] \left[\mathbf{B}\right]^2$

When volume is reduced to half, pressure increases by two times.

 $r = k[2A]^{1} [2B]^{2} = 2 \times 4 = 8$ times





(A) 4-Ethyl-1-Fluoro-2-nitrobenzene

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(C) 3-Ethyl-6-Fluoronitrobenzene
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Ans (A)

Based on lowest sum of locants,



4-Ethyl-1-fluoro-2-nitrobenzene

- 57. Higher order (>3) reactions are rare due to
 - (A) Shifting of equilibrium towards reactants due to elastic collisions
 - (B) Loss of active species on collision
 - (C) Low probability of simultaneous collision of all reacting species
 - (D) Increase in entropy as more molecules are involved

Ans (C)

Collisions between more than three reactant molecules is rare.

58. Arrange benzene, n-hexane and ethyne in decreasing order of their acidic behaviour

- (A) Benzene > n-hexane > ethyne
- (B) n-hexane > Benzene > ethyne
- (C) ethyne > n-hexane > Benzene
- (D) ethyne > Benzene > n-hexane





Ans (D)

%

НС	C≡CH	>		>	CH ₃ (CH ₂) ₄ CH ₃
	sp		sp ²		sp ³
s-character	50%		33.3%		25%

Higher the percentage s-character, higher the acidic character.

59. A colloidal solution is subjected to an electric field than colloidal particles more towards anode. The amount of electrolytes of BaCl₂, AlCl₃ and NaCl required to coagulate the given colloid is in the order (A) $NaCl > BaCl_2 > AlCl_3$ (B) $BaCl_2 > AlCl_3 > NaCl$

(C)
$$AlCl_3 = NaCl = BaCl_2$$
 (D) $AlCl_3 > BaCl_2 > NaCl$

Based on Hardy-Schulze rule, higher the valence of flocculating ion, higher the flocculating power.

Flocculating power $\propto \frac{1}{\text{Flocculating value}}$

Order of the amount of electrolyte required is $NaCl > BaCl_2 > AlCl_3$

- 60. Which of the following is an incorrect statement?
 - (A) Hydrogen bonding is stronger than dispersion forces
 - (B) Sigma bonds are stronger than π -bonds
 - (C) Ionic bonding is non-directional
 - (D) σ -electrons are referred to as mobile electrons

Ans (D)

 π -electrons are referred to as mobile electrons (with respect to graphite).

* * *



