#### Solution

#### CHEMISTRY

#### JEE main - Chemistry

#### **CHEMISTRY (Section-A)**

1.

# (c) Tritium ${}_{1}^{3}H$

# **Explanation:**

Among the isotopes of hydrogen, only tritium is radioactive and emits low energy  $\beta^-$  particles.

2.

# **(c)** Te

# Explanation:

Metalloids have intermediate properties between metals and non-metals Sc, Pb, Bi are metals Te is a metalloid.

# 3.

# (b) CH(CN)<sub>3</sub>

# **Explanation:**

Due to the resonance stabilisation of the conjugate base, CH(CN)<sub>3</sub> is the strongest acid amongst the given compounds.

$$\begin{array}{c} \mathrm{CH}(\mathrm{CN})_{3}\rightleftharpoons\mathrm{H}^{+}+\overline{\mathrm{C}}(\mathrm{CN})_{3}\\\\ \overset{\mathrm{NC}}{\longrightarrow}\overline{\mathrm{C}}\overset{\mathrm{C}}{=}\overset{\mathrm{C}}{=}\overset{\mathrm{N}}{N} \longleftrightarrow \overset{\overset{\mathrm{N}}{\longrightarrow}\overset{\mathrm{C}}{\longrightarrow}\overset{\mathrm{C}}{=}\overset{\mathrm{C}}{=}\overset{\mathrm{C}}{=}\overset{\mathrm{N}}{N}\\\\ \overset{\mathrm{NC}}{\longrightarrow}\overset{\mathrm{N}}{\longrightarrow}\overset{\mathrm{C}}{\longleftarrow}\overset{\mathrm{N}}{\longrightarrow}\overset{\mathrm{N}}{\underset{N}=}\overset{\mathrm{C}}{\subset}\overset{\mathrm{C}}{=}\overset{\mathrm{C}}{=}\overset{\mathrm{N}}{N} \end{array}$$

The conjugate bases of CHBr<sub>3</sub> and CHI<sub>3</sub> are stabilised by inductive effect of halogens. This is why, they are less stable. Also, the conjugate base of CHCl<sub>3</sub> involves back-bonding between 2p and 3p orbitals.

4.

(b) A-B has the stiffest bond

# Explanation:

A-B bond has the highest intermolecular potential energy among the given molecules. Hence, it is the strongest bond and has maximum bond enthalpy.

# 5. **(a)** 9.0 L

# Explanation:

 $pH = 1 :: [H^+] = 10^{-1} = 0.1 M$   $pH = 2 :: [H^+] = 10^{-2} = 0.01 M$ For dilution of HCl,  $M_1V_1 = M_2V_2$   $0.1 \times 1 = 0.01 \times V_2$   $V_2 = 10 L$ Volume of water to be added = 10 -1 = 9 L

6.

**(c)** 2

#### **Explanation:**

 $\begin{array}{l} \mathrm{Mn}O_4^- + 5\mathrm{e}^- \longrightarrow \mathrm{Mn}^{2+} \\ \mathrm{i.} \ \mathrm{FeC}_2\mathrm{O}_4 \longrightarrow \mathrm{Fe}^{3+} + 2\mathrm{CO}_2 + 3\mathrm{e}^- \\ 1 \ \mathrm{mole} \ \mathrm{of} \ \mathrm{FeC}_2\mathrm{O}_4 \ \mathrm{reacts} \ \mathrm{with} \ \frac{3}{5} \ \mathrm{mole} \ \mathrm{of} \ \mathrm{acidified} \ \mathrm{KMnO}_4 \\ \mathrm{ii.} \ \mathrm{Fe}_2(\mathrm{C}_2\mathrm{O}_4)_3 \longrightarrow \mathrm{Fe}^{3+} + \mathrm{CO}_2 + 6\mathrm{e}^- \end{array}$ 

1 mole of Fe<sub>2</sub>(C<sub>2</sub>O<sub>4</sub>)<sub>3</sub> reacts with  $\frac{6}{5}$  moles of KMnO<sub>4</sub>

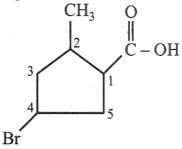
(iii)  $FeSO_4 \longrightarrow Fe^{3+} + e^{-1}$ 

1 mole of FeSO<sub>4</sub> react with  $\frac{1}{5}$  moles of KMnO<sub>4</sub>

- (iv)  $Fe_2(SO_4)_3$  does not oxidise
- $\therefore$  Total moles required =  $\frac{3}{6} + \frac{6}{5} + \frac{1}{5} = 2$

#### 7.

(c) 4-Bromo-2-methylcyclopentane carboxylic acid **Explanation:** 



4-Bromo-2-methylcyclopentane carboxylic acid

#### 8. (a) I<sup>-</sup>

#### Explanation:

If an electronegative element is in its lowest possible oxidation state in a compound or in a free state. It can function as a powerful reducing agent, e.g. I<sup>-</sup>

9. (a) 1-phenyl-2-butene

# Explanation:

 $\begin{array}{c} PhCH_2 \\ H \end{array} C = C \begin{pmatrix} CH_3 \\ H \end{pmatrix} \begin{pmatrix} PhCH_2 \\ H \end{pmatrix} C = C \begin{pmatrix} H \\ CH_3 \\ H \end{pmatrix} \begin{pmatrix} CH_3 \\ H \end{pmatrix} \begin{pmatrix} PhCH_2 \\ H \end{pmatrix} C = C \begin{pmatrix} H \\ CH_3 \\ H \end{pmatrix} \begin{pmatrix} CH_3 \\ H \end{pmatrix} \begin{pmatrix} PhCH_2 \\ H \end{pmatrix} \begin{pmatrix} CH_3 \\ H \end{pmatrix} \begin{pmatrix} CH_3 \\ H \end{pmatrix} \begin{pmatrix} PhCH_2 \\ H \end{pmatrix} \begin{pmatrix} CH_3 \\ H \end{pmatrix} \begin{pmatrix} C$ 

10.

(b) 38.0 torr and 0.589

# **Explanation:**

Total V.P. of solution =  $P_A^{\circ} X_A + P_B^{\circ} X_B$ Given,  $P_A^{\circ}$  = 74.7 torr,  $P_B^{\circ}$  = 22.3 torr  $n_{benzene}$  = 1.5 mol,  $n_{toluene}$  = 3.5 mol  $n_{solution}$  = 1.5 + 3.5 = 5 mol  $x_A = \frac{n_{benzene}}{n_{solution}} = \frac{1.5}{5} = 0.3$   $x_B = \frac{n_{toluene}}{n_{solution}} = \frac{3.5}{3} = 0.7$ Total V.P. of solution = (0.3 × 74.7 + 0.5 × 22.3) torr = (22.4 + 15.6) torr = 38 torr Mole fraction of benzene in vapour form =  $\frac{22.4}{38} = 0.589$ 

#### 11.

# Explanation:

(c) 1.51

The molality of 20% (mass/mass) aqueous solution of KI can be calculated by following formula.  $m = \frac{w_2 \times 1000}{Mw_2 \times w_1}$ 

20% aqueous solution of KI means that 20 gm of KI is present in 80 gm solvent.  $m = \frac{20}{166} \times \frac{1000}{80} = 1.506 \approx 1.51 \text{ mol/kg}$ 

12.

(d) 
$$C_2 = \sqrt{2}C_1$$

#### Explanation:

For the concentration cell,  $E_{cell}^{o} = 0$ Anode:  $Cu(s) \longrightarrow Cu^{2+} (aq)_A$ Cathode:  $Cu^{2+} (aq)_e \longrightarrow Cu(s)$ Overall:  $Cu^{2+} (aq)_C \longrightarrow Cu^{2+} (aq)_A$   $(C_2M) \longrightarrow Cu^{2+} (aq)_A$ As  $\Delta G = -nFE$ If  $\Delta G = -ve$ , then  $E_{cell}$  is +ve.  $E_{cell} = 0 - \frac{RT}{2F} \ln \frac{C_1}{C_2}$  $E_{cell} = \frac{RT}{2F} \ln \frac{C_2}{C_1}$ 

So, 
$$C_2 > C_1$$

Thus,  $C_2 = \sqrt{2}C_1$  relation is correct.

# 13.

# **(c)** 1

# **Explanation:**

Time of 75% reaction is twice the time taken for 50% reaction if it is first-order reaction w.r.t P. From the graph, [Q] decreases linearly with time, thus it is zeroth-order reaction w.r.t. Q

 $\frac{dx}{dt} = bk[P]^a [Q]^b$ 

Order w.r.t P = a = 1

Order w.r.t Q = b = 0

Thus, overall order of the reaction = 1 + 0 = 1

# 14. (a) -2.69 V; the reaction will not occur

# **Explanation:**

 $\mathrm{Mn}^{2+}$  + 2e<sup>-</sup>  $\rightarrow$  Mn; E<sup>o</sup> = -1.18 V; ...(i)

 $Mn^{3+} + e^- \rightarrow Mn^{2+}$ ;  $E^o = +1.51$  V; ...(ii)

Now multiplying equation (ii) by two and subtracting from equation (i):

 $3Mn^{2+} \rightarrow Mn^+ + 2Mn^{3+}$ ;

 $E^{0} = -1.18 - (+ 1.51) = -2.69 V$ 

-ve value of EMF (i.e.,  $\Delta G$  = +ve) shows that the reaction is non-spontaneous.

15.

(c) Superoxide and  $-\frac{1}{2}$ 

# **Explanation:**

In KO<sub>2</sub>, the nature of oxygen species and the oxidation state of oxygen atom are superoxide (superoxide ion is  $O_2^-$ )and -1/2 respectively.

Let x be oxidation state of oxygen. The oxidation state of K is +1. Hence + 1 + 2(x) =  $0 \Rightarrow x = -\frac{1}{2}$ 

16.

**(b)**  $[Co(H_2O)_5Cl] Cl_2 \cdot H_2O$ 

# **Explanation:**

- Molarity (M) =  $\frac{\text{Number of moles of solute}}{\frac{1}{2}}$
- Volume of solution ( in L ) : Number of moles of complex
- Molarity  $\times$  volume ( in mL)

$$= \frac{1000}{1000} = \frac{0.1 \times 100}{1000} = 0.01 \text{mole}$$

Number of moles of ions precipitate

$$=rac{1.2 imes 10^{22}}{6.02 imes 10^{23}}=0.02$$
 moles

... Number of Cl<sup>-</sup> present in ionisation sphere

 $\frac{\text{Number of Net } r}{\text{Number of moles of ions precipitated}} = \frac{0.02}{0.01} = 2$ Number of moles of complex

∴ 2 Cl<sup>-</sup> are present outside the square brackets, i.e. in ionisation sphere. Thus, the formula of complex is  $[Co(H_2O)_5Cl]Cl_2 \cdot H_2O$ 

#### 17. (a) 2-chloro-2-methylbutane

# **Explanation:**

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The optically inactive compound must contains achiral carbon atom(s). Option (d) contains achiral carbon atom

$$\mathrm{CH}_3 - \overset{\mathrm{Cl}}{\overset{|}{\underset{\mathrm{H}}{\mathrm{CHO}}}} - \mathrm{CHO}$$

2-Chloropropanol (chiral)

$$\mathrm{CH}_3-\mathrm{CH}_2-\overset{\mathrm{Cl}}{\underset{\mathrm{H}}{\overset{\mathrm{C}}{\mathrm{H}}}}-\mathrm{CH}_3$$

2-Chlorobutane (chiral)

$$\mathrm{CH}_3-\mathrm{CH}_2-\mathrm{CH}_2-\overset{|}{\operatorname*{CH}}_2-\mathrm{CH}_3$$

Cl

2-Chloropentane (chiral)  $\mathrm{CH}_3$ 

$$\mathrm{CH}_3 - \mathrm{CH}_2 - egin{array}{c} ert \ \mathrm{CH}_3 & -\mathrm{CH}_2 \ ert \ \mathrm{Cl} \ ert \ \mathrm{Cl} \end{array}$$

2-Chloro-2-methylbutane (achiral)

# 18.

(d) (P) - (1), (Q) - (3), (R) - (4), (S) - (2)

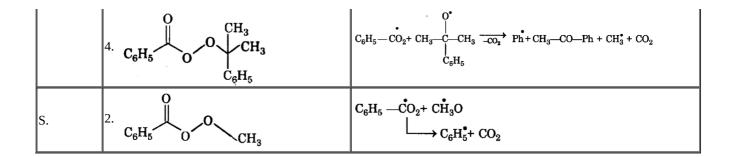
# **Explanation:**

Allylic radical are more stable than alkyl radical, so when there is a possibility of formation of allyl radical, it will undergo fragmentation

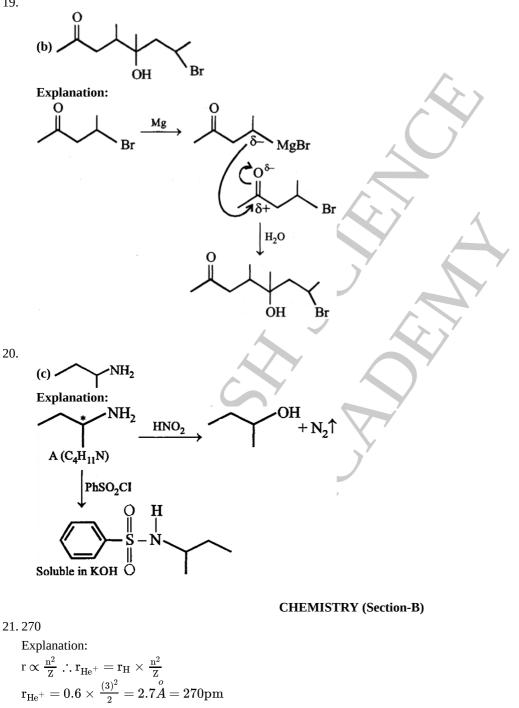
through formation of allyl radical, i.e. fragmentation produces stable radical.

On the basis of stability of radical, fragmentation can be done as

Column I	Column II	Explaination
Р.	<sup>1.</sup> C <sub>6</sub> H <sub>5</sub> H <sub>2</sub> C O CH <sub>3</sub>	$C_6H_5$ — $CH_2$ + $CO_2$ + $CH_3O$
Q.	3. $C_6H_5H_2C$ O $CH_3$ O CH <sub>3</sub> CH <sub>2</sub> C <sub>6</sub> H <sub>5</sub>	$\begin{array}{c} O^{\bullet} \\ C_{8}H_{5} - \overset{\bullet}{C}H_{2} + CO_{2} + Ph - CH_{2} - \overset{\bullet}{C} - CH_{3} \longrightarrow Ph - \overset{\bullet}{C}H_{2} + CH_{3} - CO - CH_{3} \\ \\ CH_{3} \end{array}$
R.		



19.



22.1350.0

Explanation:

$$\frac{t_1}{t_2} = \frac{\frac{1}{K} \ln \frac{a_0}{0.4a_0}}{\frac{1}{K} \ln \frac{a_0}{0.1a_0}}$$

$$\frac{540}{t_2} = \frac{\ln \frac{10}{4}}{\ln 10} \Rightarrow \frac{540}{t_2} = \frac{\log 10 - \log 4}{\log 10}$$

$$\frac{540}{t_2} = \frac{1 - 0.6}{1} \Rightarrow \frac{540}{t_2} = 0.4 \Rightarrow t_2 = \frac{540}{0.4} = 1350 \text{ sec.}$$

23.2.0

Explanation:  
$$\begin{bmatrix} {}^{+1}_{Ag}(NH_3)_2 \end{bmatrix}^{+} \begin{bmatrix} {}^{+1}_{Ag}(CN)_2 \end{bmatrix}^{-1}$$

# 24.4

Explanation:  $R = N_A \times k$ = 6.023 × 10<sup>23</sup> × 1.380 × 10<sup>-23</sup> = 8.312 which has 4 significant figures

# 25.6.0

Explanation:  $\Delta G^{\circ} = -RT \ell nK$   $-nFE_{cell}^{\circ} = -RT \times 2.303 (log_{10}K)$   $\frac{E_{cell}^{\circ}}{0.06} \times n = log K ...(i)$   $Pd^{+2}(aq.) + 2e^{-} \rightleftharpoons Pd(s), E_{cat, redn}^{\circ} = 0.83$   $Pd(s) + 4Cl^{-}(aq.) \rightleftharpoons PdCl_{4}^{2-}, (aq) + 2e^{-}, E_{anode, oxidn}^{\circ} = 0.65$ Net Reaction  $\rightarrow Pd^{2+}$  (aq.)  $4Cl^{-}(aq.) \rightleftharpoons PdCl_{4}^{2-}(aq.)$   $E_{cell}^{\circ} = E_{cat, redn}^{\circ} - E_{anode, oxidn}^{\circ}$   $E^{o}_{cell} = 0.83 - 0.65 \Rightarrow E^{o}_{cell} = 0.18 ...(ii)$ Also n = 2 using equation (i), (ii) & (iii), (iii)  $\Rightarrow logK = 6$