

**JEE-MAIN EXAMINATION – APRIL 2025**(HELD ON TUESDAY 08<sup>th</sup> APRIL 2025)

TIME : 3:00 PM TO 6:00 PM

**CHEMISTRY****TEST PAPER WITH SOLUTION****SECTION-A**

51. In a first order decomposition reaction, the time taken for the decomposition of reactant to one fourth and one eighth of its initial concentration are  $t_1$  and  $t_2$  (s), respectively. The ratio  $t_1/t_2$  will :
- (1)  $\frac{4}{3}$       (2)  $\frac{3}{2}$   
 (3)  $\frac{3}{4}$       (4)  $\frac{2}{3}$

**Ans. (4)****Sol.** For 1<sup>st</sup> order reactionWhen  $C_t = C_0/4$  $t_1 = 2t_{50\%}$ .when  $C_t = C_0/8$  $t_2 = 3t_{50\%}$ 

$$\text{so } \frac{t_1}{t_2} = \frac{2}{3}$$

52. Match the **LIST-I** with **LIST-II**

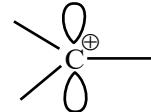
<b>LIST-I</b>		<b>LIST-II</b>	
A.	Carbocation	I.	Species that can supply a pair of electrons.
B.	C-Free radical	II.	Species that can receive a pair of electrons.
C.	Nucleophile	III.	$sp^2$ hybridized carbon with empty p-orbital.
D.	Electrophile	IV	$sp^2/sp^3$ hybridized carbon with one unpaired electron.

Choose the **correct** answer from the options given below :

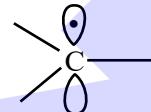
- (1) A-IV, B-II, C-III, D-I  
 (2) A-II, B-III, C-I, D-IV  
 (3) A-III, B-IV, C-II, D-I  
 (4) A-III, B-IV, C-I, D-II

**Ans. (4)**

**Sol.** (A) Carbocation  $\rightarrow sp^2$  hybridised carbon with empty P-orbital

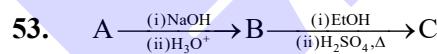


(B) Carbon free radical  $\rightarrow sp^2/sp^3$  hybridised carbon with one unpaired electron.



(C) Nucleophile  $\rightarrow$  species of that can supply a pair of electron.

(D) Electrophile  $\rightarrow$  species that can receive a pair of electron.

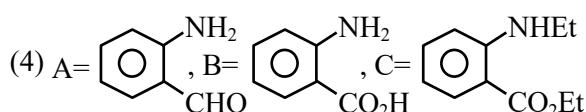
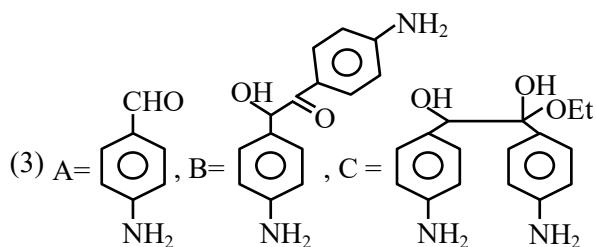
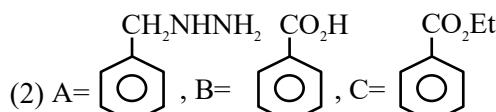
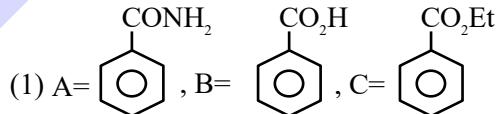


'A' shows positive Lassaigne's test for N and its molar mass is 121.

'B' gives effervescence with aq.  $NaHCO_3$ .

'C' gives fruity smell.

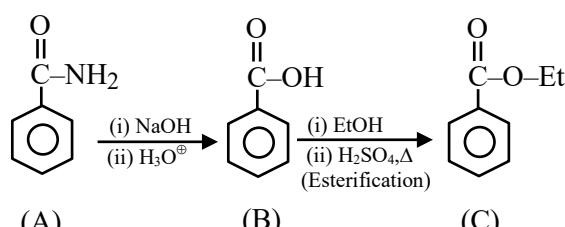
Identify A, B and C from the following.

**Ans. (1)**

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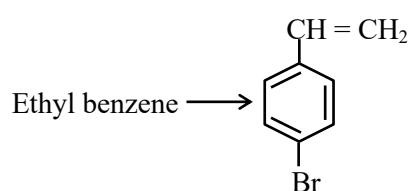
Sol.



Molar mass = 121

A  $\rightarrow$  Benzamide Shows positive Lassaigh's test.B  $\rightarrow$  Benzoic acid gives effervescence with aq.  $\text{NaHCO}_3$ .C  $\rightarrow$  Ester gives fruity smell.

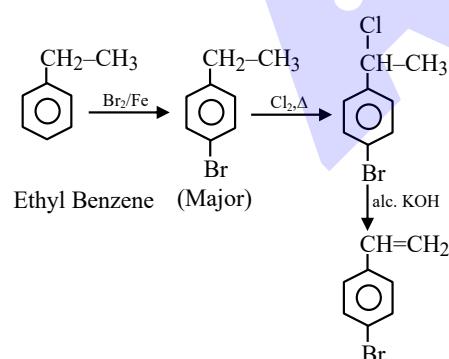
54. Choose the correct set of reagents for the following conversion.



- $\text{Br}_2/\text{Fe}$ ;  $\text{Cl}_2, \Delta$ ; alc. KOH
- $\text{Cl}_2/\text{Fe}$ ;  $\text{Br}_2/\text{anhy. AlCl}_3$ ; aq. KOH
- $\text{Br}_2/\text{anhy. AlCl}_3$ ;  $\text{Cl}_2, \Delta$ ; aq. KOH
- $\text{Cl}_2/\text{anhy. AlCl}_3$ ;  $\text{Br}_2/\text{Fe}$ ; alc. KOH

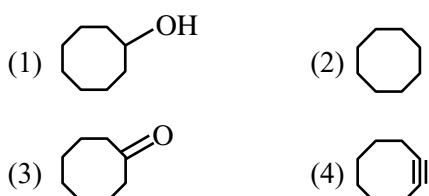
Ans. (1)

Sol.

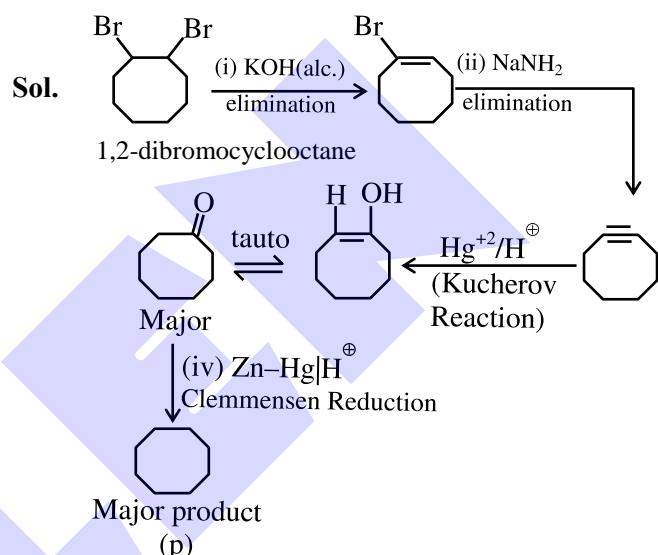


55. 1, 2-dibromocyclooctane  $\xrightarrow{\begin{array}{l} \text{(i) KOH (alc.)} \\ \text{(ii) NaNH}_2 \\ \text{(iii) } \text{Hg}^{2+}/\text{H}^+ \\ \text{(iv) Zn-Hg/H}^+ \end{array}} \text{P}$  (Major product)

'P' is



Ans. (2)



56. Given below are two statements :

**Statement I :** A homoleptic octahedral complex, formed using monodentate ligands, will not show stereoisomerism.

**Statement II :** cis- and trans- platin are heteroleptic complexes of Pd.

In the light of the above statements, choose the *correct* answer from the options given below.

- Both statement I and Statement II are false.
- Statement I is false but Statement II is true.
- Both statement I and Statement II are true.
- Statement I is true but Statement II is false.

Ans. (4)

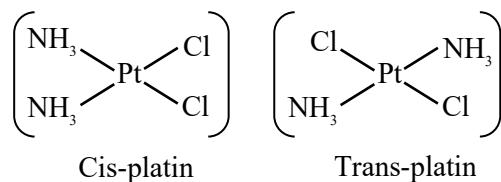


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**Sol.** Homoleptic complex of type  $[Ma_6]$  (Where  $a \Rightarrow$  monodentate ligand) cannot show geometrical as well as optical isomerism.

Cis-platin and trans-platin has formula  $[\text{Pt}(\text{NH}_3)_2\text{Cl}_2]$  which is a heteroleptic complex of platinum.



- 57.** The atomic number of the element from the following with lowest 1<sup>st</sup> ionisation enthalpy is :

- (1) 32                          (2) 35  
(3) 87                          (4) 19

**Ans. (3)**

**Sol.** Atomic no. 32  $\Rightarrow$  Ge

Atomic no. 35  $\Rightarrow$  Br

Atomic no. 87  $\Rightarrow$  Fr

Atomic no. 19  $\Rightarrow$  K

Lowest first I.E. among the given element will be of Fr [87].

Fr – [Rn] 7s<sup>1</sup>

- 58.** Which of the following binary mixture does not show the behaviour of minimum boiling azeotropes ?

- (1)  $\text{H}_2\text{O} + \text{CH}_3\text{COC}_2\text{H}_5$   
(2)  $\text{C}_6\text{H}_5\text{OH} + \text{C}_6\text{H}_5\text{NH}_2$   
(3)  $\text{CS}_2 + \text{CH}_3\text{COCH}_3$   
(4)  $\text{CH}_3\text{OH} + \text{CHCl}_3$

**Ans. (2)**

**Sol.** Binary mixture of  $\text{C}_6\text{H}_5\text{OH}$  and  $\text{C}_6\text{H}_5\text{NH}_2$  shows negative deviation from Raoult's law

So vapour pressure of solution is less than V.P of pure  $\text{C}_6\text{H}_5\text{OH}$  &  $\text{C}_6\text{H}_5\text{NH}_2$

So B.P. of solution is greater than boiling point of pure  $\text{C}_6\text{H}_5\text{OH}$  &  $\text{C}_6\text{H}_5\text{NH}_2$

So shows maximum Boiling azeotrope

- 59.**  $\text{HA(aq)} \rightleftharpoons \text{H}^+(\text{aq}) + \text{A}^-(\text{aq})$

The freezing point depression of a 0.1 m aqueous solution of a monobasic weak acid HA is 0.20°C.

The dissociation constant for the acid is

Given :

$$K_f(\text{H}_2\text{O}) = 1.8 \text{ K kg mol}^{-1}, \text{ molality} \equiv \text{molarity}$$

$$(1) 1.38 \times 10^{-3}$$

$$(2) 1.1 \times 10^{-2}$$

$$(3) 1.90 \times 10^{-3}$$

$$(4) 1.89 \times 10^{-1}$$

**Ans. (1)**

**Sol.**  $\Delta T_f = ik_f m$

$$0.2 = i \times 1.8 \times 0.1$$

$$i = \frac{20}{18} = \frac{10}{9}$$

$$\text{For } \text{HA}_{(\text{aq})} \rightleftharpoons \text{H}_{(\text{aq})}^+ + \text{A}_{(\text{aq})}^-$$

$$t=0 \quad 1$$

$$t=t_{\text{eq}} \quad 1-\alpha \quad \alpha \quad \alpha$$

$$i = 1 + \alpha$$

$$\frac{10}{9} = 1 + \alpha$$

$$\alpha = \frac{1}{9}$$

$$K_{\text{eq}} = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]} = \frac{Ca^2}{1-\alpha}$$

$$= \frac{0.1 \left( \frac{1}{9} \right)^2}{1 - \frac{1}{9}} = \frac{1}{720}$$

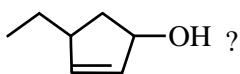
$$K_{\text{eq}} = 1.38 \times 10^{-3}$$



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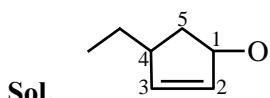
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60. What is the correct IUPAC name of



- (1) 4-Ethyl-1-hydroxycyclopent-2-ene
- (2) 1-Ethyl-3-hydroxycyclopent-2-ene
- (3) 1-Ethylcyclopent-2-en-3-ol
- (4) 4-Ethylcyclopent-2-en-1-ol

**Ans. (4)**



4-Ethylcyclopent-2-en-1-ol

61. The correct decreasing order of spin only magnetic moment values (BM) of  $\text{Cu}^+$ ,  $\text{Cu}^{2+}$ ,  $\text{Cr}^{2+}$  and  $\text{Cr}^{3+}$  ions is :

- (1)  $\text{Cu}^+ > \text{Cu}^{2+} > \text{Cr}^{3+} > \text{Cr}^{2+}$
- (2)  $\text{Cu}^{2+} > \text{Cu}^+ > \text{Cr}^{2+} > \text{Cr}^{3+}$
- (3)  $\text{Cr}^{2+} > \text{Cr}^{3+} > \text{Cu}^{2+} > \text{Cu}^+$
- (4)  $\text{Cr}^{3+} > \text{Cr}^{2+} > \text{Cu}^+ > \text{Cu}^{2+}$

**Ans. (3)**

**Sol.**  $\text{Cu}^+ : [\text{Ar}] 3d^{10}$ , Spin only magnetic moment = 0

B.M.

$\text{Cu}^{2+} : [\text{Ar}] 3d^9$ , Spin only magnetic moment =  $\sqrt{3}$

B.M.

$\text{Cr}^{2+} : [\text{Ar}] 3d^4$ , Spin only magnetic moment =  $\sqrt{24}$

B.M.

$\text{Cr}^{3+} : [\text{Ar}] 3d^3$ , Spin only magnetic moment =  $\sqrt{15}$

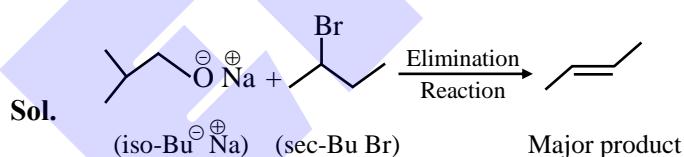
B.M.

Order of  $\mu$  :  $\text{Cr}^{2+} > \text{Cr}^{3+} > \text{Cu}^{2+} > \text{Cu}^+$

62. Which one of the following reactions will not lead to the desired ether formation in major proportion?  
(iso-Bu  $\Rightarrow$  isobutyl, sec-Bu  $\Rightarrow$  sec-butyl,  
nPr  $\Rightarrow$  n-propyl,  $t\text{Bu} \Rightarrow$  tert-butyl, Et  $\Rightarrow$  ethyl)



**Ans. (4)**

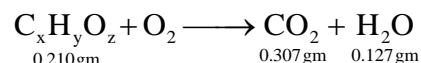


63. On combustion 0.210 g of an organic compound containing C, H and O gave 0.127 g  $\text{H}_2\text{O}$  and 0.307 g  $\text{CO}_2$ . The percentages of hydrogen and oxygen in the given organic compound respectively are:

- (1) 53.41, 39.6
- (2) 6.72, 53.41
- (3) 7.55, 43.85
- (4) 6.72, 39.87

**Ans. (2)**

- Sol.** In the combustion of organic compound, all "C" in  $\text{CO}_2$  and all "H" in  $\text{H}_2\text{O}$  comes from organic compound



Weight of "C" in  $\text{CO}_2 = \frac{12}{44} \times 0.307$

= 0.0837 gm

Weight of "H" in  $\text{H}_2\text{O} = \frac{2}{18} \times 0.127 = 0.0141\text{ g}$



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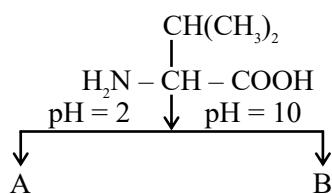
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$$\% \text{ 'H' in compound} = \frac{0.0141}{0.21} \times 100 = 6.719 \% \\ = 6.72 \%$$

Weight of "O" in compound  
 $= 0.210 - (0.0837 + 0.0141)$   
 $= 0.1122$

% of "O" in compound  $= \frac{0.1122}{0.21} \times 100$   
 $= 53.41 \%$

64.

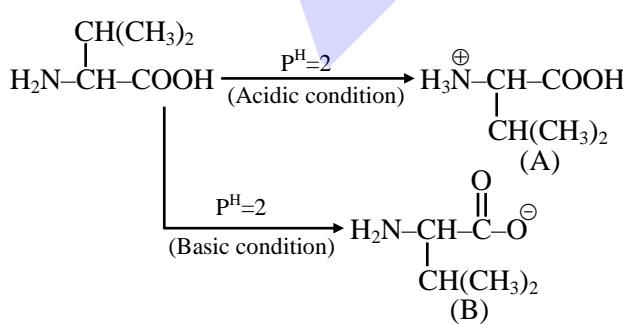


Choose the correct option for structures of A and B, respectively.

- (1)  $\text{H}_3\text{N}^+ - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{COOH}$  and  $\text{H}_2\text{N} - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{CO}\bar{\text{O}}$
- (2)  $\text{H}_2\text{N} - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{CO}\bar{\text{O}}$  and  $\text{H}_3\text{N}^+ - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{COOH}$
- (3)  $\text{H}_2\text{N} - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{COO}^\ominus$  and  $\text{H}_3\text{N}^+ - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{CO}\bar{\text{O}}$
- (4)  $\text{H}_3\text{N}^+ - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{CO}\bar{\text{O}}$  and  $\text{H}_3\text{N}^+ - \underset{\text{CH}(\text{CH}_3)_2}{\text{CH}} - \text{COOH}$

Ans. (1)

Sol.



65. Correct statements for an element with atomic number 9 are

A. There can be 5 electrons for which  $m_s = +\frac{1}{2}$  and

4 electrons for which  $m_s = -\frac{1}{2}$

B. There is only one electron in  $p_z$  orbital

C. The last electron goes to orbital with  $n = 2$  and  $l = 1$

D. The sum of angular nodes of all the atomic orbitals is 1.

Choose the correct answer from the options given below:

- (1) C and D Only
- (2) A and C Only
- (3) A, C and D Only
- (4) A and B Only

Ans. (2)

Sol. Element with atomic number 9 is Fluorine

$$\text{F}(9) = 1s^2 2s^2 2p^5$$



(A) 5 electrons can be up-spin  $\left[ m_s = +\frac{1}{2} \right]$  and

4 electrons can be down spin  $\left[ m_s = -\frac{1}{2} \right]$

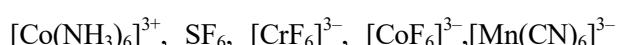
(B) Unpaired electron can be in anyone of  $p_x$ ,  $p_y$  or  $p_z$  orbital

(C) Last electron is in  $2p$  subshell with  $n = 2$ ,  $l = 1$

(D) Angular node for s-orbital = 0 while of each p-orbital = 1

Sum of all angular node = 3

66. The number of species from the following that are involved in  $sp^3d^2$  hybridization is



and  $[\text{MnCl}_6]^{3-}$

(1) 5

(2) 6

(3) 4

(4) 3

Allen Ans. (4)

NTA Ans. (3)



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Sol. In  $[\text{Co}(\text{NH}_3)_6]^{3+}$ ,  $\text{Co}^{3+}$  :  $[\text{Ar}]3\text{d}^6$ ,  $\text{NH}_3$  is S.F.L  
 Hybridisation state of  $\text{Co}^{3+}$  is  $\text{d}^2\text{sp}^3$   
 In  $\text{SF}_6$ , Hybridisation state of sulphur is  $\text{sp}^3\text{d}^2$   
 In  $[\text{CrF}_6]^{3-}$ ,  $\text{Cr}^{3+}$  :  $[\text{Ar}]3\text{d}^3$   
 Hybridisation state of  $\text{Cr}^{3+}$  is  $\text{d}^2\text{sp}^3$   
 $[\text{CoF}_6]^{3-}$ ,  $\text{Co}^{3+}$  :  $[\text{Ar}]3\text{d}^6$  F $^-$  is W.F.L  
 Hybridisation state of  $\text{Co}^{3+}$  is  $\text{sp}^3\text{d}^2$   
 $[\text{Mn}(\text{CN})_6]^{3-}$ ,  $\text{Mn}^{3+}$  :  $[\text{Ar}]3\text{d}^4$  CN $^-$  is S.F.L  
 Hybridisation state of  $\text{Mn}^{3+}$  is  $\text{d}^2\text{sp}^3$   
 $[\text{MnCl}_6]^{3-}$ ,  $\text{Mn}^{3+}$  :  $[\text{Ar}]3\text{d}^4$  Cl $^-$  is W.F.L  
 Hybridisation state of Cl $^-$  is  $\text{sp}^3\text{d}^2$   
 Total number of  $\text{sp}^3\text{d}^2$  hybridized molecules is 3

67. Match the **LIST-I** with **LIST-II**

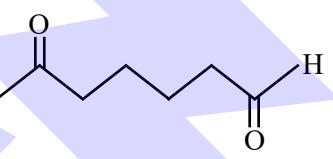
<b>LIST-I</b> (Reagent)		<b>LIST-II</b> (Functional Group detected)	
A.	Sodium bicarbonate solution	I.	double bond/unsaturation
B.	Neutral ferric chloride	II.	carboxylic acid
C.	ceric ammonium nitrate	III.	phenolic - OH
D.	alkaline $\text{KMnO}_4$	IV	alcoholic - OH

Choose the *correct* answer from the options given below :

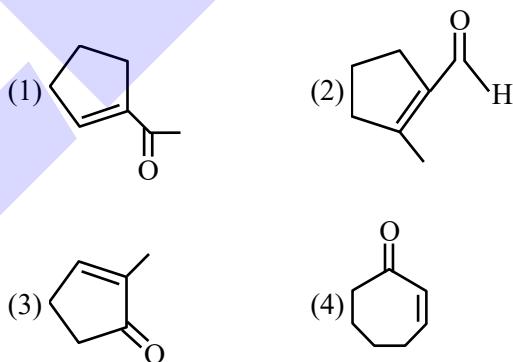
- (1) A-II, B-III, C-IV, D-I
- (2) A-II, B-III, C-I, D-IV
- (3) A-III, B-II, C-IV, D-I
- (4) A-II, B-IV, C-III, D-I

Ans. (1)

Sol. (1) Carboxylic acid gives effervescence with sodium bicarbonate solution  
 (2) Phenolic-OH gives violet coloured complex with Neutral  $\text{FeCl}_3$ .  
 (3) Alcoholic-OH gives Red colour with ceric ammonium Nitrate.  
 (4) When alkaline  $\text{KMnO}_4$  reacts with an unsaturated compound (Alkene or alkyne) the purple colour of  $\text{KMnO}_4$  solution disappears, indicating positive test for unsaturation.

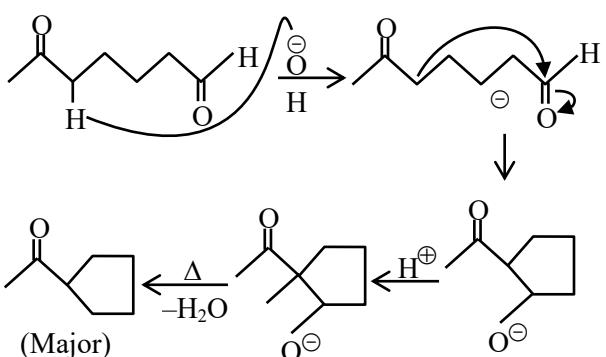
68. When  undergoes

intramolecular aldol condensation, the major product formed is :



Ans. (1)

Sol. Aldol condensation reaction



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69. Match the LIST-I with LIST-II

LIST-I (Complex/Species)		LIST-II (Shape & magnetic moment)	
A. $[\text{Ni}(\text{CO})_4]$	I.	Tetrahedral, 2.8 BM	
B. $[\text{Ni}(\text{CN})_4]^{2-}$	II.	Square planar, 0 BM	
C. $[\text{NiCl}_4]^{2-}$	III.	Tetrahedral, 0 BM	
D. $[\text{MnBr}_4]^{2-}$	IV	Tetrahedral, 5.9 BM	

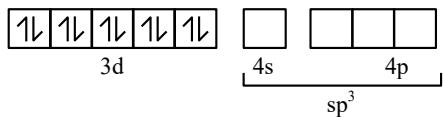
Choose the *correct* answer from the options given below :

- (1) A-III, B-IV, C-II, D-I
- (2) A-I, B-II, C-III, D-IV
- (3) A-III, B-II, C-I, D-IV
- (4) A-IV, B-I, C-III, D-II

**Ans. (3)**

**Sol.** (A)  $[\text{Ni}(\text{CO})_4]$ ,  $\text{Ni}^0$  :  $[\text{Ar}]3\text{d}^8 4\text{s}^2$

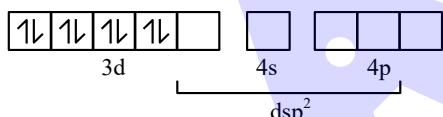
Valence orbitals of  $\text{Ni}^0$  in pre-hybridisation state :



Tetrahedral, Diamagnetic,  $\mu = 0$  B.M.

(B)  $[\text{Ni}(\text{CN})_4]^{2-}$ ,  $\text{Ni}^{+2}$  :  $[\text{Ar}]3\text{d}^8 4\text{s}^0$

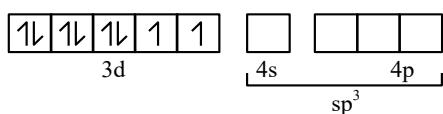
Valence orbitals of  $\text{Ni}^{+2}$  in pre-hybridisation state :



Square planar, Diamagnetic,  $\mu = 0$  B.M.

(C)  $[\text{NiCl}_4]^{2-}$ ,  $\text{Ni}^{+2}$  :  $[\text{Ar}]3\text{d}^8 4\text{s}^0$

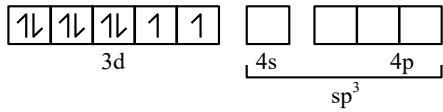
Valence orbitals of  $\text{Ni}^{+2}$  in ground state :



Tetrahedral, paramagnetic,  $\mu = \sqrt{8} = 2.8$  B.M.

(D)  $[\text{MnBr}_4]^{2-}$ ,  $\text{Mn}^{+2}$  :  $[\text{Ar}]3\text{d}^5$

Valence orbitals of  $\text{Mn}^{+2}$  in ground state :



Tetrahedral, paramagnetic,  $\mu = \sqrt{35} = 5.9$  B.M.

70. Given below are two statements :

**Statement I** :  $\text{H}_2\text{Se}$  is more acidic than  $\text{H}_2\text{Te}$ .

**Statement II** :  $\text{H}_2\text{Se}$  has higher bond enthalpy for dissociation than  $\text{H}_2\text{Te}$ .

In the light of the above statements, choose the *correct* answer from the options given below.

- (1) Both statement I and Statement II are false.
- (2) Both statement I and Statement II are true.
- (3) Statement I is true but Statement II is false.
- (4) Statement I is false but Statement II is true.

**Ans. (4)**

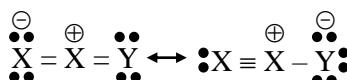
**Sol.** Acidic strength :  $\text{H}_2\text{Se} < \text{H}_2\text{Te}$



[276 KJ/mol] [238 KJ/mol]

## SECTION-B

71. Resonance in  $\text{X}_2\text{Y}$  can be represented as



The enthalpy of formation of

$\text{X}_2\text{Y} \left( \text{X} \equiv \text{X(g)} + \frac{1}{2} \text{Y} = \text{Y(g)} \rightarrow \text{X}_2\text{Y(g)} \right)$  is  $80 \text{ kJ mol}^{-1}$ .

The magnitude of resonance energy of  $\text{X}_2\text{Y}$  is \_\_\_\_\_  $\text{kJ mol}^{-1}$  (nearest integer value)

Given : Bond energies of  $\text{X} \equiv \text{X}$ ,  $\text{X} = \text{X}$ ,  $\text{Y} = \text{Y}$  and  $\text{X} = \text{Y}$  are 940, 410, 500 and  $602 \text{ kJ mol}^{-1}$  respectively.

valence X : 3, Y : 2

**Ans. (98)**



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**Sol.**  $\Delta H_{R.E} = \Delta H_f(\text{exp}) - \Delta H_f(\text{Theo})$   
 $\Delta H_f(\text{exp}) \text{ for } X_2Y_{(g)} = 80 \text{ kJ/mole}$   
for  $\Delta H_f(\text{Theo})$   
 $X_{2(g)} + \frac{1}{2} Y_{2(g)} \rightarrow X_2Y_{(g)} \Delta H_f = ?$   
 $\Delta H_{f(\text{Theo})} = \left( BE_{X=X} + \frac{1}{2} BE_{Y=Y} \right) - (BE_{X=X} + BE_{X=Y})$   
 $= \left( 940 + \frac{1}{2} \times 500 \right) - (410 + 602)$   
 $= 178 \text{ kJ/mole}$   
 $\Delta H_{R.E} = 80 - 178$   
 $= -98 \text{ kJ/mol}$   
 $|\Delta H_{R.E}| = 98$

72. The energy of an electron in first Bohr orbit of H-atom is  $-13.6 \text{ eV}$ . The magnitude of energy value of electron in the first excited state of  $\text{Be}^{3+}$  is \_\_\_\_\_ eV. (nearest integer value)

**Ans. (54)**

**Sol.**  $E_T = -13.6 \frac{z^2}{n^2} \text{ eV}$

For energy of H-atom, energy of 1<sup>st</sup> Bohr orbit  
 $E_1 = -13.6 \text{ eV}$  [z = 1, n = 1]

For  $\text{Be}^{3+}$  ion, energy of 1<sup>st</sup> E.S. [z = 4, n = 2]

$$\frac{E_H}{E_{\text{Be}^{3+}}} = \frac{z_1^2}{n_1^2} \times \frac{n_2^2}{z_2^2}$$

$$\frac{E_H}{E_{\text{Be}^{3+}}} = \frac{1}{1} \times \frac{4}{16}$$

$$E_{\text{Be}^{3+}} = -13.6 \times 4 = -54.4 \text{ eV}$$

$$|E_{\text{Be}^{3+}}| = 54.4 \text{ eV}$$

73. 20 mL of sodium iodide solution gave 4.74 g silver iodide when treated with excess of silver nitrate solution. The molarity of the sodium iodide solution is \_\_\_\_\_ M. (Nearest Integer value)  
(Given : Na = 23, I = 127, Ag = 108, N = 14, O = 16 g mol<sup>-1</sup>)

**Ans. (1)**

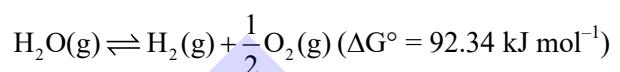
**Sol.**  $\text{NaI}_{(aq)} + \text{AgNO}_3{}_{(aq)} \rightarrow \text{AgI}_{(s)} + \text{NaNO}_3{}_{(aq)}$   
M, 20 ml excess 4.74g

$$\text{Moles of I}^- \text{ in NaI} = \text{Moles of (I}^- \text{) in AgI} = \frac{4.74}{235}$$

$$\text{Moles of NaI} = \frac{4.74}{235}$$

$$\text{Molarity [NaI]} = \frac{4.74}{235 \times 0.02} = 1.008$$

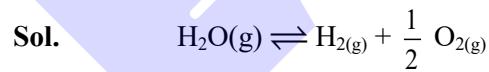
74. The equilibrium constant for decomposition of  $\text{H}_2\text{O}(g)$



is  $8.0 \times 10^{-3}$  at 2300 K and total pressure at equilibrium is 1 bar. Under this condition, the degree of dissociation ( $\alpha$ ) of water is \_\_\_\_\_  $\times 10^{-2}$  (nearest integer value).

[Assume  $\alpha$  is negligible with respect to 1]

**Ans. (5)**



$$t = 0 \quad 1 \text{ mole}$$

$$t = t_{eq} \quad 1-\alpha \quad \alpha \quad \frac{\alpha}{2}$$

$$n_T = 1 + \frac{\alpha}{2} \approx 1 \quad (\alpha \ll 1)$$

$$K_p = \frac{P_{\text{H}_2} \cdot P_{\text{O}_2}^{1/2}}{P_{\text{H}_2\text{O}}} = \frac{(\alpha \cdot P) \left( \frac{\alpha}{2} P \right)^{1/2}}{(1-\alpha)P}$$

$$P = 1$$

$$8 \times 10^{-3} = \frac{\alpha^{3/2}}{\sqrt{2}}$$

$$\alpha^{3/2} = 8\sqrt{2} \times 10^{-3}$$

$$\alpha^3 = 128 \times 10^{-6}$$

$$\alpha = \sqrt[3]{128} \times 10^{-2}$$

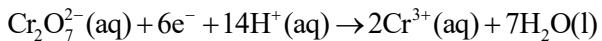
$$= 5.03 \times 10^{-2}$$



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75. Consider the following half cell reaction



The reaction was conducted with the ratio of

$$\frac{[\text{Cr}^{3+}]^2}{[\text{Cr}_2\text{O}_7^{2-}]} = 10^{-6}. \text{ The pH value at which the EMF}$$

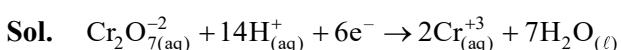
of the half cell will become zero is \_\_\_\_\_.

(nearest integer value)

[Given : standard half cell reduction potential

$$E_{\text{Cr}_2\text{O}_7^{2-}, \text{H}^+/\text{Cr}^{3+}}^{\circ} = 1.33\text{V}, \frac{2.303\text{RT}}{\text{F}} = 0.059\text{V} ]$$

**Ans. (10)**



$$E_R = E_R^{\circ} - \frac{0.059}{6} \log \frac{[\text{Cr}^{3+}]^2}{[\text{Cr}_2\text{O}_7^{2-}][\text{H}^+]^{14}}$$

$$0 = 1.33 - \frac{0.059}{6} \log \frac{10^{-6}}{[\text{H}^+]^{14}}$$

$$\frac{1.33 \times 6}{0.059} = \log \frac{10^{-6}}{[\text{H}^+]^{14}}$$

$$135.254 = -6 - 14 \log [\text{H}^+]$$

$$141.254 = 14 \text{ pH}$$

$$\text{pH} = \frac{141.254}{14} = 10.08$$



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