# **JEE MAIN 2024** Paper with Solution

Chemistry | 30<sup>th</sup> January 2024 \_ Shift-2





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### SECTION – A

- **1.** Which among the following purification methods is based on the principle of "Solubility" in two different solvents?
  - (1) Column Chromatography
  - (3) Distillation

(2) Sublimation

(4) Differential Extraction

Ans. 4

Different layers are formed which can be separated in funnel. (Theory based).

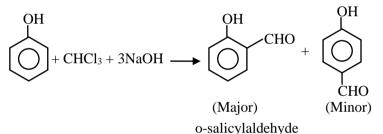
2. Salicylaldehyde is synthesized from phenol, when reacted with

$$(1) H Cl, NaOH$$

$$(2) CO2, NaOH$$

$$(3) CCl4, NaOH$$

(4) HCCl<sub>3</sub>, NaOH



Reimer Tiemann Reaction

**3.** Given below are two statements:

**Statement – I :** High concentration of strong nucleophilic reagent with secondary alkyl halides which do not have bulky substituents will follow  $S_N^2$  mechanism.

**Statement – II :** A secondary alkyl halide when treated with a large excess of ethanol follows  $S_N^{-1}$  mechanism. In the light of the above statements, choose the most appropriate from the options given below:

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

#### Ans. 4

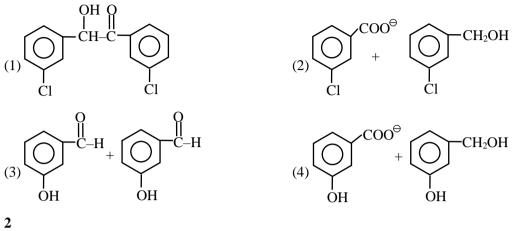
Statement 1 : High conc<sup>n</sup>. of strong nu<sup>-</sup> reagent with 2° Alkyl Halide which do not have bulky substituents will follow  $S_N^2$  mechanism.

Hence statement I is true.

Statement 2 : 2° Alkyl Halide reacts with excess of ethanol undergo  $S_N^{-1}$  reaction.

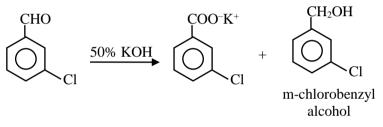
Hence statement 2 is true

#### 4. m-chlorobenzaldehyde on treatment with 50% KOH solution yields



#### Ans. 2

Cannizaro reaction (Disproportination reaction)



5. Given below are two statements: One is labelled as Assertion A and the other is labelled as Reason R: Assertion A:  $H_2Te$  is more acidic than  $H_2S$ .

Reason R: Bond dissociation enthalpy of  $H_2Te$  is lower than  $H_2S$ .

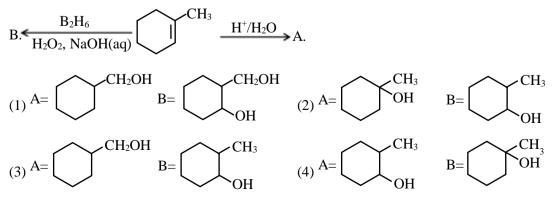
In the light of the above statements, choose the most appropriate from the options given below:

- (1) Both A and R are true but R is NOT the correct explanation of A.
- (2) Both A and R are true and R is the correct explanation of A.
- (3) A is false but R is true.
- (4) A is true but R is false.

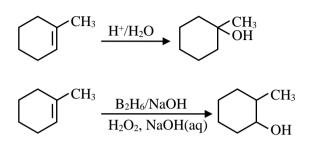
#### Ans. 2

Due to lower bond dissociation energy of  $H_2Te$  it ionizes to give  $H^+$  more easily as compare to  $H_2S$ 

6. Products A and B formed in the following set of reactions are



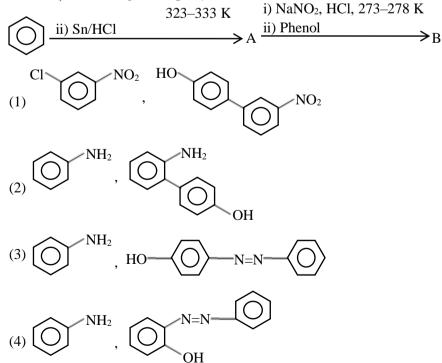
Ans. 2



- 7. IUPAC name of following compound is : CH<sub>3</sub>--CH--CH<sub>2</sub>--CN NH<sub>2</sub>
  - (1) 2-Aminopentanenitrile
  - (2) 2-Aminobutanenitrile
  - (3) 3-Aminobutanenitrile
  - (4) 3-Aminopropanenitrile

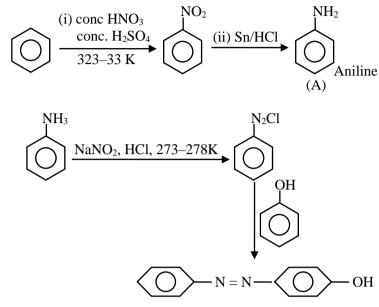
Ans. 3

- 4 3 2 1 C-C-C-CN 3-Amino butanenitrile
- 8. The products A and B formed in the following reaction scheme are respectively i) con. HNO<sub>3</sub>/con. H<sub>2</sub>SO<sub>4</sub>





Ans. 3



9. The molecule / ion with square pyramidal shape is

(1)  $[Ni(CN)_4]^{2-}$ (2) PCl<sub>5</sub> (3) BrF<sub>5</sub> (4) PF<sub>5</sub> 3

$$F \xrightarrow{F} F$$

square pyramidal

- The orange colour of  $K_2Cr_2O_7$  and purple colour of  $KMnO_4$  is due to 10.
  - (1) Charge transfer transition in both.
  - (2) d  $\rightarrow$  d transitions in KMnO<sub>4</sub> and charge transfer transitions in K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>.
  - (3)  $d \rightarrow d$  transitions in K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> and charge transfer transitions in KMnO<sub>4</sub>.
  - (4)  $d \rightarrow d$  transitions in both

Ans. 1

orange colour of K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> and purple colour of KMnO<sub>4</sub> is due to ligand to metal charge transfer (LMCT)

Alkaline oxidative fusion of MnO<sub>2</sub> gives "A" which on electrolytic oxidation in alkaline solution produces B. 11. A and B respectively are

(2)  $MnO_4^{2-}$  and  $MnO_4^{-}$ (1)  $Mn_2O_7$  and  $MnO_4^-$ 

(4)  $MnO_4^{2-}$  and  $Mn_2O_7$ (3)  $Mn_2O_3$  and  $MnO_4^{2-}$ 

#### Ans. 2

Alkaline oxidative fusion of  $MnO_2$  gives  $MnO_4^{2-}$ 

 $2MnO_2 + 4OH^- + O_2 \rightarrow 2 MnO_4^{2-} + 2H_2O$ 

electrolytic oxidation of  $MnO_4^{2-}$  in alkaline solution gives  $MnO_4^{-}$ 

 $MnO_4^{2-} \rightarrow MnO_4^{-} + e^{-}$ 

12. If a substance 'A' dissolves in solution of a mixture of 'B ' and 'C' with their respective number of moles as  $n_A, n_B$  and  $n_C$ . Mole fraction of C in the solution is

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(1) 
$$\frac{n_{\rm C}}{n_{\rm A} \times n_{\rm B} \times n_{\rm C}}$$
 (2)  $\frac{n_{\rm C}}{n_{\rm A} + n_{\rm B} + n_{\rm C}}$  (3)  $\frac{n_{\rm C}}{n_{\rm A} - n_{\rm B} - n_{\rm C}}$  (4)  $\frac{n_{\rm B}}{n_{\rm A} + n_{\rm B}}$ 

Ans. (2)

mole fraction of C(X<sub>c</sub>) =  $\frac{n_C}{n_A + n_B + n_C}$ 

**13.** Given below are two statements:

**Statement** – I: Along the period, the chemical reactivity of the elements gradually increases from group 1 to group 18.

**Statement – II :** The nature of oxides formed by group 1 elements is basic while that of group 17 elements is acidic.

In the light of the above statements, choose the most appropriate from the options given below:

- (1) Both Statement I and Statement II are True
- (2) Statement I is True But Statement II is False
- (3) Statement I is False but statement II is true
- (4) Both Statement I and Statement II are False

#### Ans. 3

Chemical reactivity along the period decreases

So, statement-I is wrong

Group-I elements form basic nature oxide while group 17 elements form acidic nature oxide

- **14.** The coordination geometry around the manganese in decacarbonyldimanganese (0) is
  - (1) Octahedral(3) Square pyramidal
- (2) Trigonal bipyramidal(4) Square planar

Ans. 1

15.

$$\begin{array}{c|c} CO & CO & CO \\ CO & 1 & CO \\ CO & Mn & Mn & CO \\ CO & 1 & CO \\ CO & CO & CO \end{array}$$

Octahedral around Mn

Given below are two statements:

**Statement** – **I** : Since Fluorine is more electronegative than nitrogen, the net dipole moment of  $NF_3$  is greater than  $NH_3$ .

**Statement – II :** In  $NH_3$ , the orbital dipole due to lone pair and the dipole moment of NH bonds are in opposite direction, but in  $NF_3$  the orbital dipole due to lone pair and dipole moments of N-F bonds are in same direction. In the light of the above statements, choose the most appropriate from the options given below:

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

#### Ans.

2

Both statement are false

16. The correct stability order of carbocations is

(1) 
$$(CII_3)_3 C^+ > CII_3 - \overset{+}{C}II_2 > (CII_3)_2 \overset{+}{C}II > \overset{+}{C}II_3$$
  
(2)  $\overset{+}{C}H_3 > (CH_3)_2 \overset{+}{C}H > CH_3 - \overset{+}{C}H_2 > (CH_3)_3 \overset{+}{C}$   
(3)  $(CH_3)_3 \overset{+}{C} > (CH_3)_2 \overset{+}{C}H > CH_3 - \overset{+}{C}H_2 > \overset{+}{C}H_3$   
(4)  $\overset{+}{C}H_3 > CH_3 - \overset{+}{C}H_2 > CH_3 - \overset{+}{C}H_2 (CH_3)_3 C^+$   
 $\downarrow CH_3$ 

#### Ans. 3

Stability order carbocation

$$CH_{3}-\overset{\dagger}{C}-CH_{3} > CH_{3}-\overset{\dagger}{C}-CH_{3} > \overset{\dagger}{C}-CH_{3} > \overset{\dagger}{C}H_{3}$$

$$CH_{3}$$

$$(9 \alpha-H) \qquad (6 \alpha-H) \qquad (3 \alpha-H)$$

- 17. The solution from the following with highest depression in freezing point/lowest freezing point is
  - (1) 180 g of acetic acid dissolved in water
  - (2) 180 g of acetic acid dissolved in benzene
  - (3) 180 g of benzoic acid dissolved in benzene
  - (4) 180 g of glucose dissolved in water

#### Ans. 1

Highest depression in freezing point (Colligative Properties) is inversely proportional to experimentally molar mass of non volatile solute

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\* on dissociation exp. Molar mass dec.

\* On association exp. Molar mass Inc.

**18.** A and B formed in the following reactions are:

$$CrO_2Cl_2 + 4NaOH \rightarrow A + 2NaCl + 2H_2O$$
$$A + 2HCl + 2H_2O_2 \rightarrow B + 3H_2O$$

(1) 
$$A = Na_2CrO_4, B = CrO_5$$

(2) 
$$A = Na_2Cr_2O_4, B = CrO_4$$

(3) 
$$A = Na_2Cr_2O_7, B = CrO_3$$

(4) 
$$A = Na_2Cr_2O_7, B = CrO_5$$

#### Ans. 1

 $CrO_{2}Cl_{2}+4NaOH \rightarrow Na_{2}CrO_{4}+2NaCl+2H_{2}O$ (A)  $Na_{2}CrO_{4}+2HCl+2H_{2}O_{2}\rightarrow CrO_{5}+3H_{2}O+2NaCl$ 

**19.** Choose the correct statements about the hydrides of group 15 elements.

A. The stability of the hydrides decreases in the order  $NH_3 > PH_3 > AsH_3 > SbH_3 > BiH_3$ .

B. The reducing ability of the hydride increases in the order  $NH_3 < PH_3 < AsH_3 < SbH_3 < BiH_3$ .

C. Among the hydrides, NH<sub>3</sub> is strong reducing agent while BiH<sub>3</sub> is mild reducing agent.

D. The basicity of the hydrides increases in the order  $NH_3 < PH_3 < AsH_3 < SbH_3 < BiH_3$ .

Choose the most appropriate from the options given below:

(1) B and C only (2) C and D only (3) A and B only (4) A and D only **3**  $NH_3 \mid M$ -H bond length  $\uparrow$ 

PH<sub>3</sub> Thermal stability  $\downarrow$ ASH<sub>3</sub> Reducing nature  $\uparrow$ SbH<sub>3</sub> BiH<sub>3</sub>

20. Reduction potential of ions are given below:

 $ClO_{4}^{-} IO_{4}^{-} BrO_{4}^{-}$   $E^{\circ} = 1.19 V E^{\circ} = 1.65 V E^{\circ} = 1.74 V$ The correct order of their original

The correct order of their oxidizing power is.

(1)  $ClO_{4}^{-} > IO_{4}^{-} > BrO_{4}^{-}$ (2)  $BrO_{4}^{-} > IO_{4}^{-} > ClO_{4}^{-}$ (3)  $BrO_{4}^{-} > ClO_{4}^{-} > IO_{4}^{-}$ (4)  $IO_{4}^{-} > BrO_{4}^{-} > ClO_{4}^{-}$ 

#### Ans. 2

Ans.

Oxidizing power  $\propto$  Reduction potential

 $ClO_4^ IO_4^ BrO_4^-$ 

 $E_{red}^{o}$  1.19V 1.65V 1.74V

order of standard reduction potential

 $E^{\circ} ClO_{4}^{-} < IO_{4}^{-} < BrO_{4}^{-}$ 

Thus order of oxidizing power

 $ClO_{4}^{-} < IO_{4}^{-} < BrO_{4}^{-}$ 

#### SECTION - B

**21.** Number of complexes which show optical isomerism among the following is

$$\operatorname{cis} - [\operatorname{Cr}(\operatorname{ox})_{2}\operatorname{Cl}_{2}]^{3^{-}}, [\operatorname{Co}(\operatorname{en})_{3}]^{3^{+}}, \operatorname{cis} - [\operatorname{Pt}(\operatorname{en})_{2}\operatorname{Cl}_{2}]^{2^{+}}, \operatorname{cis} - [\operatorname{Co}(\operatorname{en})_{2}\operatorname{Cl}_{2}]^{+},$$
  
trans  $- [\operatorname{Pt}(\operatorname{en})_{2}\operatorname{Cl}_{2}]^{2^{+}}, \operatorname{trans} - [\operatorname{Cr}(\operatorname{ox})_{2}\operatorname{Cl}_{2}]^{3^{-}}$ 

Ans. 4

Those complex which has no COS and POS are optically active  $Cis-[Cr(ox)_2Cl_2]^{3-}$ ,  $[Co(en)_3]^{3+}$ ,  $Cis-[Pt(en)_2Cl_2]^{+2}$ ,  $[Co(en)_2Cl_2]^+$ 

22. NO<sub>2</sub> required for a reaction is produced by decomposition of N<sub>2</sub>O<sub>5</sub> in CCl<sub>4</sub> as by equation  $2N_2O_{5(g)} \rightarrow 4NO_{2(g)} + O_{2(g)}$ 

The initial concetration. of  $N_2O_5$  is  $3 \text{mol}L^{-1}$  and it is  $2.75 \text{mol}L^{-1}$  after 30 minutes.

The rate of formation of NO<sub>2</sub> is  $x \times 10^{-3} \text{ mol } \text{L}^{-1} \text{ min}^{-1}$ , value of x is (nearest integer) \_\_\_\_\_.

Ans. 17

$$\frac{1}{2} \frac{d[N_2O_5]}{dt} = -\frac{1}{4} \frac{d[NO_2]}{dt}$$
$$\frac{d[NO_2]}{dt} = -2 \frac{d[N_2O_5]}{dt}$$
$$\frac{d[NO_2]}{dt} = -2 \frac{(3-2.75)}{30}$$
$$\frac{d[NO_2]}{dt} = \frac{2 \times 0.25}{30}$$
$$= 1.667 \times 10^{-2}$$
$$= 16.67 \times 10^{-3}$$
$$x \approx 17 \times 10^{-3}$$

**23.** Two reactions are given below:

$$2\operatorname{Fe}_{(s)} + \frac{3}{2}\operatorname{O}_{2(g)} \to \operatorname{Fe}_{2}\operatorname{O}_{3(s)}, \Delta \operatorname{H}^{\circ} = -822 \,\mathrm{kJ} \,/ \,\mathrm{mol}$$
$$\operatorname{C}_{(s)} + \frac{1}{2}\operatorname{O}_{2(g)} \to \operatorname{CO}_{(g)}, \Delta \operatorname{H}^{\circ} = -110 \,\mathrm{kJ} \,/ \,\mathrm{mol}$$

Then enthalpy change for following reaction  $3C_{(s)} + Fe_2O_{3(s)} \rightarrow 2Fe_{(s)} + 3CO_{(g)}$  is \_\_\_\_\_ kJ/mol.

#### Ans. 492

By Hess law  $2 \text{ Fe} + \frac{3}{2} O_2 \rightarrow \text{Fe}_2 O_3$   $\Delta H^\circ = -822 \text{ KJ } \dots (1)$   $C + \frac{1}{2} O_2 \rightarrow CO$   $\Delta H^\circ = -110 \text{ KJ/mol } \dots (2)$ Fe<sub>2</sub>O<sub>3</sub>  $\rightarrow 2 \text{ Fe} + 3/2 O_2$   $\Delta H^\circ = 822 \text{ KJ } \dots (3)$ eq. (2) multiply by 3  $3C + \frac{3}{2} O_2 \rightarrow 3 \text{ CO}$   $\Delta H^\circ = -330 \text{ KJ } \dots (4)$ eq. (3)+(4)  $3C + \text{Fe}_2 O_3 \rightarrow 2 \text{Fe} + 3 \text{CO}$   $\Delta H = 492 \text{ KJ/mol}$ 

**24.** The total number of correct statements, regarding the nucleic acids is

- A. RNA is regarded as the reserve of genetic information
- B. DNA molecule self-duplicates during cell division
- C. DNA synthesizes proteins in the cell
- D. The message for the synthesis of particular proteins is present in DNA
- E. Identical DNA strands are transferred to daughter cells.

#### Ans. 3

А	В	С	D	E
(False)	(True)	(False)	(True)	(True)

25. The pH of an aqueous solution containing 1 M benzoic acid  $(pK_a = 4.20)$  and 1 M sodium benzoate is 4.5. The volume of benzoic acid solution in 300 mL of this buffer solution is \_\_\_\_\_ mL. (given : log 2 = 0.3)

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Ans. 100

$$pH=pK_{a}+log \frac{(V_{salt})}{(V_{acid})}$$

$$4.5=4.2+log \frac{(V_{salt})}{(V_{acid})}$$

$$log \frac{(V_{salt})}{(V_{acid})} = 0.3$$

$$\frac{(V_{salt})}{(V_{acid})} = 2 \qquad \dots \dots (Eq. 1)$$

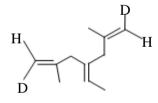
$$V_{salt} + V_{acid} = 300 \text{ ml} \qquad \dots \dots (Eq. 2)$$

$$By Eq. (1) \& (2)$$

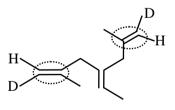
$$V_{salt} = 200 \text{ ml}$$

$$V_{acid} = 100 \text{ ml}$$

26. Number. of geometrical isomers possible for the given structure is/are \_\_\_\_\_.







(E, E) (E, Z) (Z, Z) (Z, E)

27. Total number of species from the following which can undergo disproportionation reaction is \_\_\_\_\_\_.  $H_2O_2, ClO_3^-, P_4, Cl_2, Ag, Cu^{+1}, F_2, NO_2, K^+$ 

Ans. 6

Intermediate oxidation state of element can undergo disproportionation reaction  $H_2O_2$ ,  $ClO_3^-$ ,  $P_4$ ,  $Cl_2$ ,  $Cu^+$ ,  $NO_2$ 

28. Number of metal ions characterized by flame test among the following is \_\_\_\_\_\_.

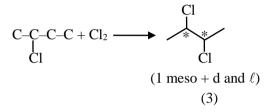
#### Ans. 4

Metal ions (Ca<sup>2+</sup>, Sr<sup>2+</sup>, Ba<sup>2+</sup>Cu<sup>2+</sup>) responds flame test

**29.** 2-chlorobutane  $+Cl_2 \rightarrow C_4H_8Cl_2$  (isomers)

Total number of optically active isomers shown by  $C_4H_8Cl_2$ , obtained in the above reaction is \_\_\_\_\_.

#### Ans. 3



**30.** Number of spectral lines obtained in He<sup>+</sup> spectra, when an electron makes transition from fifth excited state to first excited state will be

#### Ans. 10

Fifth excited state  $n_2=6$ first excited state  $n_1=2$ 

Number of spectra line 
$$\frac{\left[\left(n_{2}-n_{1}\right)\right]\left[\left(n_{2}-n_{1}\right)+1\right]}{2}$$

$$=\frac{(6-2)(6-2+1)}{2}$$
$$=\frac{4\times 5}{2}=\frac{20}{2}=10$$







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