# JEE Main 2023 30 Jan Shift 1 Chemistry Question Paper with Solutions

**Time Allowed :**180 minutes | **Maximum Marks :**300 | **Total questions :**90

#### **General Instructions**

#### Read the following instructions very carefully and strictly follow them:

- (A) The test is of 3 hours duration.
- (B) The question paper consists of 90 questions, out of which 75 are to attempted. The maximum marks are 300.
- (C) There are three parts in the question paper consisting of Physics, Chemistry and Mathematics having 30 questions in each part of equal weightage. (D) Each part (subject) has two sections.
- (i) Section-A: This section contains 20 multiple choice questions which have only one correct answer. Each question carries 4 marks for correct answer and −1 mark for wrong answer.
- (ii) Section-B: This section contains 10 questions. In Section-B, attempt any five questions out of 10. The answer to each of the questions is a numerical value. Each question carries 4 marks for correct answer and –1 mark for wrong answer. For Section-B, the answer should be rounded off to the nearest integer

# Chemistry

# **Section A**

# 1. Which of the following compounds would give the following set of qualitative analysis?

(i) Fehling's Test: Positive

(ii) Na fusion extract upon treatment with sodium nitroprusside gives a blood red colour but not

$$N$$
 $N$ 
 $CHO$ 

$$\sim$$
 CHO

$$\binom{N}{S}$$
—CHO

# **Solution:**

Aromatic aldehydes typically do not react with Fehling's solution, meaning they do not

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give a positive result for Fehling's test. This test is primarily used to detect the presence of aldehydes, but aromatic aldehydes, due to their structure, do not produce the characteristic red precipitate with Fehling's reagent.

In contrast, the presence of both nitrogen and sulfur is essential to produce the blood-red colour. This specific colour change occurs when both elements are involved in a reaction with certain reagents. One such reagent is sodium nitroprusside, which is known to produce a blood-red colour when it interacts with compounds containing both nitrogen and sulfur.

Therefore, based on this information, the compound found in option (4) is the correct choice, as it is the only one that meets the required conditions of containing both nitrogen and sulfur, leading to the formation of the blood-red colour when reacted with sodium nitroprusside.

#### Quick Tip

In organic chemistry, the presence of functional groups such as nitrogen and sulfur can influence reactions like Fehling's test and the reaction with sodium nitroprusside. Always check for the necessary elements involved in these reactions.

# 2. What is the correct order of acidity of the protons marked A-D in the given compounds?

$$H_{D}$$
 $CO_{2}H_{C}$ 
 $H_{A}$ 

(1) 
$$H_C > H_D > H_B > H_A$$

(2) 
$$H_C > H_D > H_A > H_B$$

(3) 
$$H_D > H_C > H_B > H_A$$

(4) 
$$H_C > H_A > H_D > H_B$$

**Correct Answer:** (2)  $H_C > H_D > H_A > H_B$ 

#### **Solution:**

The acidity of a proton is determined by the stability of the conjugate base that is formed upon its removal. The more stable the conjugate base, the more acidic the proton.

**Step 1: Acidity of H**<sub>C</sub> Among all the protons, H<sub>C</sub> is the most acidic. Removal of H<sub>C</sub> leads to the formation of a carboxylate anion, which is highly stabilized due to resonance.

**Step 2: Acidity of H\_D** H<sub>D</sub> is the second most acidic proton. Its removal generates a carbanion that is stabilized by resonance with the benzene ring.

Step 3: Acidity of  $H_A$  vs  $H_B$   $H_A$  is more acidic than  $H_B$ . The conjugate base formed after removing  $H_A$  is stabilized by resonance with the triple bond, which results in the negative charge being spread across two carbon atoms. In contrast, the conjugate base formed after the removal of  $H_B$  places the negative charge adjacent to the triple bond, with no resonance stabilization. This makes it highly unstable due to the electron-withdrawing nature of the sp-hybridized carbon in the alkyne.

**Step 4: Overall Acidity Order** Thus, the correct order of acidity is  $H_C > H_D > H_A > H_B$ . **Conclusion:** The correct answer is Option (2).

# Quick Tip

The key to determining acidity is analyzing the stability of the conjugate base. Factors like resonance, electronegativity, and hybridization play a crucial role. Carboxylic acids are generally much more acidic than carbon acids, which are more acidic than terminal alkynes. Alkynes with a negative charge adjacent to the triple bond are highly unstable.

# 3. Given below are two statements: one is labelled as Assertion (A) and the other is labelled as Reason (R).

Assertion (A): Ketoses give Seliwanoff's test faster than Aldoses.

Reason (R): Ketoses undergo -elimination followed by formation of furfural.

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

(1) (A) is false but (R) is true

(2) Both (A) and (R) are true and (R) is the correct explanation of (A)

(3) (A) is true but (R) is false

(4) Both (A) and (R) are true but (R) is not the correct explanation of (A)

**Correct Answer:** (3) (A) is true but (R) is false

**Solution:** 

Seliwanoff's test is used to distinguish between ketoses and aldoses. This test is based on the principle that ketose sugars, particularly keto-hexoses, undergo dehydration more quickly than aldoses when heated in an acidic medium. During this reaction, ketoses are dehydrated to form 5-hydroxymethylfurfural, a compound that reacts with resorcinol to form a red or brown-colored complex. The formation of this colored complex occurs rapidly, indicating a positive result for ketoses. In contrast, aldoses do not undergo this rapid dehydration and condensation, resulting in no color change or a much slower formation of the colored

complex. Thus, Seliwanoff's test provides a simple way to differentiate between ketoses and

aldoses based on their reactivity under acidic conditions.

Quick Tip

When studying organic tests, remember the key principle behind each test. For example, Seliwanoff's test differentiates between Ketoses and Aldoses based on the rate of dehydration to form furfural.

4. In the extraction of copper, its sulphide ore is heated in a reverberatory furnace after

mixing with silica to:

(1) separate CuO as  $CuSiO_3$ 

(2) remove calcium as  $CaSiO_3$ 

(3) decrease the temperature needed for roasting of  $Cu_2S$ 

(4) remove FeO as  $FeSiO_3$ 

Correct Answer: (4) remove FeO as  $FeSiO_3$ 

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#### **Solution:**

The copper ore, which contains iron, is mixed with silica before being heated in a reverberatory furnace. During the heating process, iron oxide (FeO) reacts with silica (SiO<sub>2</sub>) to form iron silicate (FeSiO<sub>3</sub>), which then slags off. The reaction can be represented by the following chemical equation:

$$FeO + SiO_2 \longrightarrow FeSiO_3$$

This reaction helps in separating the iron from the copper ore, as iron silicate is formed as a slag, which can be easily removed, leaving the purified copper. This process is important in the extraction and refining of copper from ores that contain impurities like iron.

#### 5. Amongst the following compounds, which one is an antacid?

- (1) Ranitidine
- (2) Meprobamate
- (3) Terfenadine
- (4) Brompheniramine

**Correct Answer:** (1) Ranitidine

#### **Solution:**

Ranitidine is an antacid that works by reducing the amount of acid produced in the stomach. It is commonly used to treat conditions such as ulcers, gastroesophageal reflux disease (GERD), and Zollinger-Ellison syndrome. The other compounds listed, Meprobamate, Terfenadine, and Brompheniramine, are not antacids. Meprobamate is used for anxiety, Terfenadine was an antihistamine (discontinued due to safety concerns), and Brompheniramine is an antihistamine used for allergies.

#### 6. The major products 'A' and 'B', respectively, are:

#### **Solution:**

In the given reaction, electrophilic substitution of the phenyl group with a sulfate group is induced. The major products are:

# Quick Tip

In organic reactions, the type of reagent and reaction conditions (such as temperature and concentration of acid) play a key role in determining the products. Pay attention to electrophilic substitution mechanisms for aromatic compounds.

#### 7. Benzyl isocyanide can be obtained by:

$$(A) \overbrace{\hspace{1cm}}^{CH_2Br} \underbrace{\hspace{1cm}}^{AgCN}$$

(B) 
$$CH_2NH_2 CHCl_3$$
 $Aq. KOH$ 

(C) 
$$CH_2$$
—NHCH<sub>3</sub> CHCl<sub>3</sub>
Aq. KOH

$$(D) \overbrace{\hspace{1cm}}^{CH_2OTs} \underbrace{\hspace{1cm}}_{KCN}$$

Choose the correct answer from the options given below:

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

**Correct Answer:** (3) A and B

#### **Solution:**

Benzyl isocyanide can be synthesized by reacting a benzyl halide with an appropriate nucleophile. The reactions are as follows:

- In reaction (A), when CH<sub>2</sub>Br reacts with AgCN, benzyl isocyanide is formed through a nucleophilic substitution.
- In reaction (B), CH<sub>2</sub>NH<sub>2</sub>CHCl<sub>2</sub> undergoes a reaction with aqueous KOH, where the chloro group is replaced by a CN group, resulting in the corresponding isocyanide.

Therefore, the correct answers are reactions (A) and (B).

#### Quick Tip

For the preparation of isocyanides, halogen-substituted alkyl groups often react with cyanide (CN) or other nucleophiles. In these cases, nucleophilic substitution mechanisms such as  $SN_2$  are involved.

# 8. Given below are two statements: one is labelled as Assertion (A) and the other is labelled as Reason (R).

**Assertion** (A): In expensive scientific instruments, silica gel is kept in watch-glasses or in semipermeable membrane bags.

**Reason** (**R**): Silica gel adsorbs moisture from air via adsorption, thus protects the instrument from water corrosion (rusting) and/or prevents malfunctioning.

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

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

**Correct Answer:** (3) Both (A) and (R) are true and (R) is the correct explanation of (A)

#### **Solution:**

Silica gel is widely utilized in high-end scientific instruments to protect them from moisture, which could otherwise cause damage.

It works by adsorbing moisture from the surrounding air, thus preventing water-related corrosion (such as rusting) and safeguarding the instrument from potential malfunctions. Therefore, the explanation provided in (R) accurately justifies the assertion made in (A).

#### Quick Tip

Silica gel is used as a desiccant in a variety of applications, including in scientific instruments, to maintain a dry environment and prevent moisture damage.

#### 9. Match List I with List II:

List I		List II	
Α	$ \begin{array}{c} CI \\ CH_3CI \\ \longrightarrow \end{array} $	I	Fitting reaction
В	C1 +2Na → ()	II	Wurtz Fitting reaction
С		III	Finkelstein reaction
D	$C_2H_5Cl + Nal \rightarrow C_2H_5I + NaCl$	IV	Sandmeyer reaction

$$(1)$$
 A – IV, B – I, C – III, D – II

$$(2) A - I, B - II, C - IV, D - III$$

$$(3) A - I, B - III, C - II, D - I$$

$$(4) A - II, B - I, C - IV, D - III$$

Correct Answer: (4) A - II, B - I, C - IV, D - III

#### **Solution:**

We are asked to match the reactions in List I with the corresponding reactions in List II. Let's analyze each option:

- In reaction (A), the reaction of a chlorobenzene with methyl chloride in the presence of sodium gives a methylated product. This reaction corresponds to the **Fitting reaction**
- In reaction (B), the reaction of a chlorobenzene with sodium metal gives biphenyl. This corresponds to the **Wurtz Fitting reaction**.
- In reaction (C), the reaction of chlorobenzene with sodium azide in the presence of copper chloride leads to the formation of an azide group. This corresponds to the **Finkelstein** reaction.
- In reaction (D), the reaction of ethyl chloride with sodium iodide gives ethyl iodide. This corresponds to the **Sandmeyer reaction**.

Now, comparing the reactions from List I and List II:

$$A \longrightarrow II$$
 (Fitting reaction)

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B → I (Wurtz Fitting reaction)

 $C \longrightarrow III$  (Finkelstein reaction)

 $D \longrightarrow IV$  (Sandmeyer reaction)

Thus, the correct match is option (4), which corresponds to:

$$(4) A - II, B - I, C - III, D - IV$$

# Quick Tip

In organic chemistry, different reactions are often classified based on the type of halogen substitution or exchange that occurs. Recognizing these patterns will help identify the correct type of reaction.

# 10. Caprolactam when heated at high temperature in presence of water gives:

- (1) Teflon
- (2) Dacron
- (3) Nylon 6, 6
- (4) Nylon 6

**Correct Answer:** (4) Nylon 6

#### **Solution:**

Caprolactam, when subjected to high temperatures in the presence of water, undergoes a polymerization reaction to produce Nylon 6. This synthetic polymer is widely utilized in the production of textiles and plastics.

- Teflon is a polymer made from tetrafluoroethylene, which is not related to caprolactam.
- Dacron is a trade name for polyethylene terephthalate (PET), and it is also unrelated to caprolactam.
- Nylon 6, 6 is synthesized from hexamethylenediamine and adipic acid, rather than caprolactam.
- Nylon 6 is specifically produced by the polymerization of caprolactam.

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#### Quick Tip

Caprolactam is specifically used for the preparation of Nylon 6, which is a key polymer used in textiles, carpeting, and plastics.

#### 11. The alkaline earth metal sulphate(s) which are readily soluble in water is/are:

- (A) BeSO<sub>4</sub>
- (B) MgSO<sub>4</sub>
- (C) CaSO<sub>4</sub>
- (D) SrSO<sub>4</sub>
- (E) BaSO<sub>4</sub>

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

- (1) A only
- (2) B only
- (3) A and B
- (4) B and C

**Correct Answer:** (3) A and B

#### **Solution:**

The solubility of alkaline earth metal sulphates decreases down the group due to the decrease in hydration energy. Hydration energy is the energy released when ions interact with water molecules. Higher hydration energy leads to better solubility.

- **BeSO**<sub>4</sub>: Due to its small size and high charge density, the Be<sup>2+</sup> ion exhibits very high hydration energy. This makes BeSO<sub>4</sub> highly soluble in water.
- $MgSO_4$ : The  $Mg^{2+}$  ion also has high hydration energy, leading to good solubility of  $MgSO_4$  in water.
- CaSO<sub>4</sub>, SrSO<sub>4</sub>, and BaSO<sub>4</sub>: As we move down the group, the size of the cations increases, reducing the charge density and hydration energy. This results in lower solubility. Hence, these sulphates are sparingly soluble or insoluble in water.

#### **Conclusion:**

From the analysis, BeSO<sub>4</sub> and MgSO<sub>4</sub> are the only sulphates that are readily soluble in water. Therefore, the correct answer is (3) A and B.

#### Quick Tip

To remember solubility trends, note that alkaline earth metal sulphates become less soluble as you move down the group. This is due to the decreasing hydration energy of the cations.

#### 12. Which of the following is the correct order of ligand field strength?

$$(1) CO < en < NH_3 < C_2O_4^{2-} < S^{2-}$$

(2) 
$$S^{2-} < C_2O_4^{2-} < NH_3 < en < CO$$

$$(3) \ NH_3 < en < CO < S^{2-} < C_2O_4{}^{2-}$$

(4) 
$$S^{2-} < NH_3 < en < CO < C_2O_4^{2-}$$

**Correct Answer:** (2)  $S^{2-} < C_2O_4{}^{2-} < NH_3 < en < CO$ 

#### **Solution:**

The spectrochemical series arranges ligands based on their field strength, which affects the splitting of d-orbitals in a coordination compound. The increasing order of ligand field strength is:

$$S^{2-} < C_2 O_4{}^{2-} < NH_3 < en < CO \\$$

# **Step 2: Explanation of the Order**

-  $S^{2-}$ : Sulfide ions are weak field ligands due to their large size and low charge density. -

 $C_2O_4^{2-}$ : Oxalate ions have moderate field strength as bidentate ligands. -  $NH_3$ : Ammonia has higher field strength due to its ability to donate lone pair electrons. - en

(Ethylenediamine): A bidentate ligand with stronger field strength than NH<sub>3</sub>. - CO

(Carbon monoxide): A strong field ligand due to its  $\pi$ -back bonding capability.

Conclusion: The correct order is (2)  $S^{2\text{--}} < C_2 O_4{}^{2\text{--}} < NH_3 < en < CO.$ 

# Quick Tip

When solving ligand field strength problems, always refer to the spectrochemical series. Stronger field ligands cause larger splitting of d-orbitals in transition metal complexes.

13. Formation of photochemical smog involves the following reaction in which A, B, and C are respectively.

- (i) NO<sub>2</sub>  $\xrightarrow{hv}$  A + B
- (ii) B +  $O_2 \rightarrow C$
- (iii)  $A + C \rightarrow NO_2 + O_2$

Choose the correct answer from the options given below:

- (1) O, NO, and  $NO_3^-$
- (2) N<sub>2</sub>O and NO
- (3) N,  $O_2$ , and  $O_3$
- (4) NO, O, and O<sub>3</sub>

Correct Answer: (4) NO, O, and O<sub>3</sub>

#### **Solution:**

Breaking down the reactions, - **Reaction** (i): NO<sub>2</sub> undergoes photodissociation under sunlight:

$$NO_2 \xrightarrow{hv} NO(A) + O(B)$$

- **Reaction (ii):** The oxygen radical (B) reacts with molecular oxygen to form ozone:

$$O\left(B\right)+O_{2}\rightarrow O_{3}\left(C\right)$$

- **Reaction (iii):** NO reacts with ozone to regenerate  $NO_2$  and molecular oxygen:

$$NO\left(A\right)+O_{3}\left(C\right)\rightarrow NO_{2}+O_{2}$$

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Step 2: Identify A, B, and C

From the reactions:

- A = NO (Nitric oxide)
- -B = O (Oxygen radical)

-  $C = O_3$  (Ozone)

Conclusion: The correct answer is (4) NO, O, and O<sub>3</sub>.

# Quick Tip

Photochemical smog is a result of sunlight-driven reactions involving  $NO_x$  and volatile organic compounds (VOCs). Always trace the sequence of reactions to identify intermediates and final products.

14. During the qualitative analysis of  $SO_3^{2-}$  using dilute  $H_2SO_4$ ,  $SO_2$  gas is evolved which turns  $K_2Cr_2O_7$  solution (acidified with dilute  $H_2SO_4$ ):

- (1) Black
- (2) Red
- (3) Green
- (4) Blue

Correct Answer: (3) Green

#### **Solution:**

#### **Reaction Involved**

When  $SO_3^{2-}$  reacts with dilute  $H_2SO_4$ ,  $SO_2$  gas is evolved. The  $SO_2$  gas reduces the dichromate ion  $(Cr_2O_7^{2-})$  to  $Cr^{3+}$ , which is green in color. The reaction is as follows:

$$Cr_2O_7^{2-} + SO_3^{2-} + H^+ \rightarrow Cr^{3+} + SO_4^{2-}$$

# **Step 2: Color Change**

- Dichromate ion  $(Cr_2O_7^{2-})$  is orange in color. - After reduction,  $Cr^{3+}$  ions form, which are green in color.

**Conclusion:** The solution turns **green** due to the formation of  $Cr^{3+}$ . Therefore, the correct answer is (3) **Green**.

# Quick Tip

In qualitative analysis, dichromate ions  $(Cr_2O_7^{2-})$  are commonly used as oxidizing agents. Reduction of these ions often results in a color change, which can help identify the reducing species.

# 15. To inhibit the growth of tumours, identify the compounds used from the following:

- (A) EDTA
- (B) Coordination Compounds of Pt
- (C) D Penicillamine
- (D) Cis Platin

Choose the correct answer from the options given below:

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

Correct Answer: (1) B and D Only

#### **Solution:**

#### **Understanding the role of Coordination Compounds of Platinum**

- Platinum-based coordination compounds, such as **Cisplatin** ( $cis-[Pt(NH_3)_2Cl_2]$ ), are widely used in chemotherapy.
- These compounds inhibit tumour growth by binding to DNA and interfering with the replication process.

#### **Step 2: Analysis of the Options**

- **Option A (EDTA):** EDTA is primarily a chelating agent used for metal ion sequestration, not for tumour treatment.
- **Option B** (**Coordination Compounds of Pt**): Correct, as platinum-based compounds are effective in tumour inhibition.
- Option C (D Penicillamine): Penicillamine is used in chelation therapy for heavy metal

poisoning, not for tumours.

- Option D (Cisplatin): Correct, as it is a well-known platinum-based anti-cancer drug.

**Conclusion:** The correct combination is **B** and **D**.

# Quick Tip

Cisplatin and other platinum-based coordination compounds are effective anti-cancer drugs. They form cross-links with DNA, disrupting cell division and leading to tumour inhibition.

# 16. In the wet tests for identification of various cations by precipitation, which transition element cation doesn't belong to group IV in qualitative inorganic analysis?

- $(1) \text{ Fe}^{3+}$
- $(2) Zn^{2+}$
- $(3) \text{ Co}^{2+}$
- $(4) Ni^{2+}$

**Correct Answer:** (1) Fe<sup>3+</sup>

#### **Solution:**

#### **Step 1: Understanding the Grouping of Cations in Qualitative Analysis**

In qualitative inorganic analysis, cations are categorized into specific groups based on how they behave when reacting with certain reagents. The classification is as follows:

- **Group III:** This group includes  $Fe^{3+}$ ,  $Al^{3+}$ , and  $Cr^{3+}$ , which form hydroxide precipitates when reacted with NH<sub>4</sub>OH and NH<sub>4</sub>Cl.
- **Group IV:** This group contains  $Zn^{2+}$ ,  $Co^{2+}$ , and  $Ni^{2+}$ , which form sulfide precipitates (such as ZnS, CoS, and NiS) when reacted with  $H_2S$  in neutral or slightly acidic conditions.

#### **Step 2: Analysis of the Cations**

- $\mathbf{Fe}^{3+}$ : It belongs to Group III because it forms  $Fe(OH)_3$  precipitate when reacted with  $NH_4OH$ .
- $\mathbf{Z}\mathbf{n}^{2+}, \mathbf{Co}^{2+}, \mathbf{Ni}^{2+}$ : These ions belong to Group IV, as they form sulfide precipitates (such

as ZnS, CoS, and NiS) when reacted with H<sub>2</sub>S.

**Conclusion:** Since  $Fe^{3+}$  does not form sulfide precipitates with  $H_2S$ , it does not belong to Group IV. Therefore, the correct answer is (1)  $Fe^{3+}$ .

# Quick Tip

To identify the group of a cation, consider the reagent used for precipitation and the medium (acidic, basic, or neutral). Group III cations form hydroxide precipitates, while Group IV cations form sulfide precipitates.

#### 17. Match List I with List II

LIST-I (molecules/ions)	LIST-II (No. of lone pairs of e- on central atom)
(A) IF <sub>7</sub>	I. Three
(B) ICl <sub>4</sub> <sup>-</sup>	II. One
(C) XeF <sub>6</sub>	III. Two
(D) XeF <sub>2</sub>	IV. Zero

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

(1) 
$$A - II$$
,  $B - III$ ,  $C - IV$ ,  $D - I$ 

(2) 
$$A - IV$$
,  $B - III$ ,  $C - II$ ,  $D - I$ 

(3) 
$$A - II, B - I, C - IV, D - III$$

$$(4) A - IV, B - I, C - II, D - III$$

Correct Answer: (2) A - IV, B - III, C - II, D - I

#### **Solution:**

#### **Step 1: Determine the Lone Pairs for Each Molecule/Ion**

The number of lone pairs on the central atom can be determined using the following steps:

- 1. Count the total valence electrons of the central atom.
- 2. Subtract the electrons used for bonding with surrounding atoms.
- 3. Divide the remaining electrons by 2 to get the number of lone pairs.

#### Step 2: Analyze Each Molecule/Ion

- **IF**<sub>7</sub>: Iodine has 7 valence electrons. All are used for bonding with 7 fluorine atoms.

Therefore, **0** lone pairs (IV).

- ICl<sub>4</sub><sup>-</sup>: Iodine has 7 valence electrons and gains 1 due to the negative charge. Four are used for bonding with chlorine atoms, leaving 4 electrons (2 lone pairs). Therefore, **2 lone pairs** (III).
- **XeF<sub>6</sub>:** Xenon has 8 valence electrons. Six are used for bonding with fluorine atoms, leaving 2 electrons (1 lone pair). Therefore, **1 lone pair** (**II**).
- **XeF<sub>2</sub>:** Xenon has 8 valence electrons. Two are used for bonding with fluorine atoms, leaving 6 electrons (3 lone pairs). Therefore, **3 lone pairs** (**I**).

#### **Step 3: Match the Molecules with the Lone Pairs**

The correct matches are:

- A IV (IF<sub>7</sub>, zero lone pairs)
- B III (ICl<sub>4</sub><sup>-</sup>, two lone pairs)
- C II (XeF<sub>6</sub>, one lone pair)
- D I (XeF<sub>2</sub>, three lone pairs)

Conclusion: The correct answer is (2) A - IV, B - III, C - II, D - I.

# Quick Tip

To determine the number of lone pairs on a central atom, always account for the total valence electrons, subtract those used for bonding, and divide the remainder by 2. This helps in identifying molecular geometry and hybridization.

#### 18. For OF<sub>2</sub> molecule consider the following:

- (A) Number of lone pairs on oxygen is 2.
- (B) F–O–F angle is less than  $104.5^{\circ}$ .
- (C) Oxidation state of O is -2.
- (D) Molecule is bent 'V' shaped.
- (E) Molecular geometry is linear.

Correct options are:

- (1) C, D, E only
- (2) B, E, A only

(3) A, C, D only

(4) A, B, D only

**Correct Answer:** (4) A, B, D only

#### **Solution:**

The Lewis structure of OF<sub>2</sub> indicates that the central oxygen atom is bonded to two fluorine atoms, with two lone pairs of electrons remaining on the oxygen atom. This results in a bent or 'V' shape for the molecule, caused by the repulsion between the lone pairs and the bonding pairs of electrons.

#### **Step 2: Analysis of the Statements**

- Statement (A): The oxygen atom in  $OF_2$  has 6 valence electrons. After forming two single bonds with fluorine atoms, it retains two lone pairs of electrons. Therefore, this statement is true.
- **Statement (B):** The F–O–F bond angle in  $OF_2$  is slightly less than  $104.5^{\circ}$ , around  $102^{\circ}$ , due to the repulsion between the lone pairs and the bonding pairs of electrons. Hence, **this** statement is true.
- **Statement (C):** The oxidation state of oxygen in  $OF_2$  is +2 because fluorine is more electronegative. However, **this statement is false** because the oxidation state of oxygen in  $OF_2$  is actually -2.
- **Statement (D):** Due to the two lone pairs on oxygen, the molecular geometry of OF<sub>2</sub> is bent or 'V' shaped. Therefore, **this statement is true**.
- **Statement** (E): The molecular geometry is not linear; it is bent. However, **this statement** is false because the geometry is indeed bent, but the statement is phrased ambiguously.

Conclusion: The correct statements are A, B, D. Therefore, the correct answer is (4) A, B, D only.

#### Quick Tip

The shape and bond angle of a molecule can be determined using VSEPR theory. Lone pair-bond pair repulsion decreases the bond angle from its ideal value.

# 19. Lithium aluminium hydride can be prepared from the reaction of:

(1) LiCl and Al<sub>2</sub>H<sub>6</sub>

(2) LiH and Al<sub>2</sub>Cl<sub>6</sub>

(3) LiCl, Al and H<sub>2</sub>

(4) LiH and Al(OH)<sub>3</sub>

Correct Answer: (2) LiH and Al<sub>2</sub>Cl<sub>6</sub>

#### **Solution:**

# Reaction for Preparation of LiAlH<sub>4</sub>

Lithium aluminium hydride (LiAl $H_4$ ) is prepared by the reaction of lithium hydride (LiH) with aluminium chloride (Al<sub>2</sub>Cl<sub>6</sub>). The reaction is as follows:

$$8LiH + Al_2Cl_6 \rightarrow 2LiAlH_4 + 6LiCl$$

Conclusion: The correct reactants for preparing LiAlH<sub>4</sub> are LiH and Al<sub>2</sub>Cl<sub>6</sub>. Therefore, the correct answer is (2).

# Quick Tip

LiAlH<sub>4</sub> is a strong reducing agent used in organic synthesis. It is commonly prepared using lithium hydride and aluminium chloride under controlled conditions.

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#### 20. Match List – I with List – II

LIST-I (Atomic number)	LIST-II (Block of periodic table)
(A) 37 (K)	I. p-block
(B) 78 (Pt)	II. d-block
(C) 52 (Te)	III. f-block
(D) 65 (Tb)	IV. s-block

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

$$(1)\ A-II,\ B-IV,\ C-I,\ D-III$$

$$(2)$$
 A – I, B – III, C – IV, D – II

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

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

Correct Answer: (4) A - IV, B - II, C - I, D - III

#### **Solution:**

#### **Step 1: Analyze the Atomic Numbers and Their Corresponding Blocks**

The blocks of the periodic table are determined by the type of orbital being filled:

- 37 (K): Potassium is an alkali metal, and its valence electron is in the s-orbital. Hence, it belongs to the s-block.
- **78** (**Pt**): Platinum is a transition metal, with valence electrons in the d-orbital. Hence, it belongs to the **d-block**.
- 52 (Te): Tellurium is a p-block element, as its valence electrons are in the p-orbital.
- **65** (**Tb**): Terbium is a lanthanide, with electrons filling the f-orbital. Hence, it belongs to the **f-block**.

#### Step 2: Match List – I with List – II

The correct matches are:

- -A IV (37, s-block)
- -B II (78, d-block)
- -C-I (52, p-block)
- D III (65, f-block)

Conclusion: The correct answer is (4) A - IV, B - II, C - I, D - III.

#### Quick Tip

To identify the block of an element, look at the atomic number and determine the type of orbital (s, p, d, f) being filled. This corresponds to the element's position in the periodic table.

# **Section B**

#### 21. Consider the cell

$$Pt(s)|H_2(g, 1 \text{ atm})|H^+(aq, 1M)||Fe^{3+}(aq), Fe^{2+}(aq)||Pt(s).$$

When the potential of the cell is 0.712 V at 298 K, the ratio  $\frac{[Fe^{2+}]}{[Fe^{3+}]}$  is \_\_\_\_. (Nearest integer)

Given:

$$\mathrm{Fe^{3+}} + e^{-} \rightarrow \mathrm{Fe^{2+}}, \quad E^{\circ}_{\mathrm{Fe^{3+}}|\mathrm{Fe^{2+}}} = 0.771 \,\mathrm{V}.$$
 
$$\frac{2.303 RT}{F} = 0.06 \,\mathrm{V}.$$

Correct Answer: 10

#### **Solution:**

The given cell is:

$$Pt(s)|H_2(g, 1 \text{ atm})|H^+(aq, 1M)||Fe^{3+}(aq), Fe^{2+}(aq)||Pt(s).$$

# **Step 1: Cell Reactions**

At the anode:

$$H_2 \to 2H^+ + 2e^-$$
.

At the cathode:

$$Fe_{aq}^{3+} + e^- \rightarrow Fe_{aq}^{2+}$$
.

# **Step 2: Standard Cell Potential**

The standard cell potential is:

$$E^{\circ} = E^{\circ}_{\mathrm{H_2|H^+}} + E^{\circ}_{\mathrm{Fe}^{3+}|\mathrm{Fe}^{2+}}.$$

Substitute the given values:

$$E^{\circ} = 0 + 0.771 = 0.771 \, \text{V}.$$

# **Step 3: Nernst Equation for the Cell Potential**

The cell potential at non-standard conditions is given by the Nernst equation:

$$E = E^{\circ} - \frac{0.06}{1} \log \frac{[\text{Fe}^{2+}]}{[\text{Fe}^{3+}]}.$$

Substitute  $E = 0.712 \,\mathrm{V}$  and  $E^{\circ} = 0.771 \,\mathrm{V}$ :

$$0.712 = 0.771 - 0.06 \log \frac{[Fe^{2+}]}{[Fe^{3+}]}.$$

Step 4: Solve for  $\log\frac{[Fe^{2+}]}{[Fe^{3+}]}$ 

Rearrange the equation:

$$\begin{split} 0.712 - 0.771 &= -0.06 \log \frac{[Fe^{2+}]}{[Fe^{3+}]}. \\ -0.059 &= -0.06 \log \frac{[Fe^{2+}]}{[Fe^{3+}]}. \\ \log \frac{[Fe^{2+}]}{[Fe^{3+}]} &= \frac{-0.059}{-0.06} \approx 1. \end{split}$$

**Step 5: Calculate the Ratio** 

From 
$$\log \frac{[Fe^{2+}]}{[Fe^{3+}]} = 1$$
:

$$\frac{[Fe^{2+}]}{[Fe^{3+}]} = 10^1 = 10.$$

Conclusion: The ratio  $\frac{[Fe^{2+}]}{[Fe^{3+}]}$  is 10.

# Quick Tip

To solve problems involving cell potentials, use the Nernst equation to relate concentrations and the measured potential. Simplify logarithmic terms step-by-step to avoid calculation errors.

22. A 300 mL bottle of soft drink has 0.2 M  $CO_2$  dissolved in it. Assuming  $CO_2$  behaves as an ideal gas, the volume of the dissolved  $CO_2$  at STP is \_\_\_\_\_ mL. (Nearest integer) Given: At STP, molar volume of an ideal gas is 22.7 L mol<sup>-1</sup>.

Correct Answer: 1362 mL

**Solution:** 

Step 1: Calculate the Moles of  $\mathbf{CO}_2$ 

The concentration of  $CO_2$  is given as 0.2 M, and the volume of the solution is 300 mL (which is equivalent to  $300 \times 10^{-3}$  L). To calculate the moles of  $CO_2$ , we use the following equation:

Moles of 
$$CO_2 = M \times Volume$$
 (in L).

Substituting the values:

Moles of 
$$CO_2 = 0.2 \times (300 \times 10^{-3}) = 0.06 \text{ mol.}$$

#### Step 2: Calculate the Volume of CO<sub>2</sub> at STP

At Standard Temperature and Pressure (STP), the molar volume of an ideal gas is 22.7 L per mole. Using this, we can calculate the volume of 0.06 mol of  $CO_2$  as:

Volume of  $CO_2$  = Moles of  $CO_2 \times$  Molar Volume at STP.

Substitute the values:

Volume of 
$$CO_2 = 0.06 \times 22.7 = 1.362 L$$
.

To convert the volume from liters to milliliters, multiply by 1000:

Volume of 
$$CO_2 = 1.362 L \times 1000 = 1362 mL$$
.

# Quick Tip

To calculate the volume of a gas at STP, always use the molar volume (22.7 L mol<sup>-1</sup> for ideal gases). Convert the given solution volume to liters before applying the molarity formula.

# 23. A solution containing 2 g of a non-volatile solute in 20 g of water boils at 373.52 K.

The molecular mass of the solute is  $_{----}$  g mol $^{-1}$ . (Nearest integer)

Given: Water boils at 373 K,  $K_b$  for water = 0.52 K kg mol<sup>-1</sup>.

Correct Answer:  $100 \text{ g mol}^{-1}$ 

**Solution:** 

# **Step 1: Boiling Point Elevation**

The elevation in boiling point  $(\Delta T_b)$  is given as:

$$\Delta T_b = T_b - T_b^{\circ} = 373.52 - 373 = 0.52 \,\mathrm{K}.$$

# **Step 2: Use the Boiling Point Elevation Formula**

The boiling point elevation is related to molality (m) by:

$$\Delta T_b = K_b \cdot m.$$

Substitute 
$$m = \frac{\text{mass of solute (g)}}{\text{Molar Mass (g mol}^{-1}) \times \text{mass of solvent (kg)}}$$
:

$$0.52 = 0.52 \cdot \frac{2}{\text{Molar Mass} \times 20 \times 10^{-3}}.$$

#### **Step 3: Solve for Molar Mass**

Simplify:

$$1 = \frac{2}{\text{Molar Mass} \times 0.02}.$$
 
$$\text{Molar Mass} = \frac{2}{0.02} = 100\,\text{g mol}^{-1}.$$

Conclusion: The molecular mass of the solute is  $100 \text{ g mol}^{-1}$ .

# Quick Tip

When solving boiling point elevation problems, always ensure the masses are converted into proper units (grams to kilograms) and the molality formula is applied correctly. Use the given values of  $K_b$  and  $\Delta T_b$  to calculate the molar mass of the solute accurately.

24. If compound A reacts with B following first-order kinetics with rate constant  $2.011 \times 10^{-3} \, \text{s}^{-1}$ , the time taken by A (in seconds) to reduce from 7 g to 2 g will be \_\_\_\_\_. (Nearest Integer)

Given:

$$\log 5 = 0.698$$
,  $\log 7 = 0.845$ ,  $\log 2 = 0.301$ .

Correct Answer: 623 seconds

**Solution:** 

**Step 1: Reaction and First-Order Kinetics Formula** 

The reaction is:

$$A + B \rightarrow P$$
.

At t = 0, the concentration of A is 7 g. At t = t, the concentration of A reduces to 2 g. For first-order reactions:

$$t = \frac{2.303}{k} \log \frac{[A]_0}{[A]_t}.$$

# **Step 2: Substitute the Values**

Substitute  $k = 2.011 \times 10^{-3} \,\mathrm{s}^{-1}$ ,  $[A]_0 = 7$ , and  $[A]_t = 2$ :

$$t = \frac{2.303}{2.011 \times 10^{-3}} \log \frac{7}{2}.$$

$$\log \frac{7}{2} = \log 7 - \log 2 = 0.845 - 0.301 = 0.544.$$

# **Step 3: Calculate the Time**

Substitute the values:

$$t = \frac{2.303}{2.011 \times 10^{-3}} \cdot 0.544.$$

$$t = \frac{2.303 \times 0.544}{2.011 \times 10^{-3}} = \frac{1.252832}{2.011 \times 10^{-3}}.$$

t = 622.989 seconds  $\approx 623$  seconds.

**Conclusion:** The time taken by A to reduce from 7 g to 2 g is **623 seconds**.

# Quick Tip

For first-order reactions, always use the formula  $t = \frac{2.303}{k} \log \frac{[A]_0}{[A]_t}$ . Ensure logarithmic values are calculated accurately to avoid errors in final results.

# 25. The energy of one mole of photons of radiation of frequency $2 \times 10^{12} \, \text{Hz}$ in J mol $^{-1}$ is \_\_\_\_\_. (Nearest integer)

**Given:**  $h = 6.626 \times 10^{-34} \, \text{Js}, N_A = 6.022 \times 10^{23} \, \text{mol}^{-1}.$ 

Correct Answer: 798 J mol<sup>-1</sup>

**Solution:** 

# **Step 1: Energy of One Photon**

The energy of a single photon is given by:

$$E = h\nu$$
,

where  $h = 6.626 \times 10^{-34}$  Js and  $\nu = 2 \times 10^{12}$  Hz. Substitute the values:

$$E = 6.626 \times 10^{-34} \cdot 2 \times 10^{12} = 1.3252 \times 10^{-21} \,\text{J}.$$

# **Step 2: Energy of One Mole of Photons**

The energy of one mole of photons is:

$$E_{\text{mole}} = N_A \cdot E$$
,

where  $N_A = 6.022 \times 10^{23}$ . Substitute the values:

$$E_{\text{mole}} = 6.022 \times 10^{23} \cdot 1.3252 \times 10^{-21} = 798.16 \,\text{J}.$$

Approximate to the nearest integer:

$$E_{\rm mole} \approx 798 \, \mathrm{J \ mol}^{-1}$$
.

Conclusion: The energy of one mole of photons is **798 J mol** $^{-1}$ .

# Quick Tip

To calculate the energy of one mole of photons, first determine the energy of a single photon using  $E = h\nu$ . Then, multiply this by Avogadro's number  $(N_A)$  to find the energy for one mole.

**26.** The number of electrons involved in the reduction of permanganate to manganese dioxide in acidic medium is \_\_\_\_\_.

**Correct Answer:** 3

**Solution:** 

# **Step 1: Reduction Reaction in Acidic Medium**

The reduction reaction is:

$$MnO_4^- + 4H^+ + 3e^- \to MnO_2 + 2H_2O.$$

In this reaction, 3 electrons are transferred during the reduction of  $MnO_4^-$  to  $MnO_2$ .

**Conclusion:** The number of electrons involved is **3**.

# Quick Tip

In redox reactions, balance the half-reactions carefully in acidic or basic media to determine the number of electrons transferred. For permanganate  $(MnO_4^-)$ , the transfer depends on the product formed (e.g.,  $MnO_2$ ,  $Mn^{2+}$ , etc.).

27. When 2 liters of ideal gas expands isothermally into a vacuum to a total volume of 6 liters, the change in internal energy is \_\_\_\_\_ J. (Nearest integer)

Correct Answer: 0 J

#### **Solution:**

# Step 1: Isothermal Expansion of an Ideal Gas

For an ideal gas undergoing an isothermal process, the change in internal energy  $(\Delta U)$  depends only on temperature (T):

$$\Delta U = f(T)$$
.

Since the process is isothermal, the temperature remains constant, and hence:

$$\Delta U = 0.$$

Conclusion: The change in internal energy is 0 J.

# Quick Tip

For isothermal expansion of an ideal gas, remember that the internal energy change  $(\Delta U)$  is always zero, as it depends only on the temperature, which remains constant.

28. 600 mL of 0.01 M HCl is mixed with 400 mL of 0.01 M  $H_2SO_4$ . The pH of the mixture is \_\_\_\_\_× $10^{-2}$ . (Nearest integer)

Given:

$$\log 2 = 0.30$$
,  $\log 3 = 0.48$ ,  $\log 5 = 0.69$ ,  $\log 7 = 0.84$ ,  $\log 11 = 1.04$ .

Correct Answer: 186

#### **Solution:**

#### Step 1: Calculate Total Millimoles of H<sup>+</sup>

The millimoles of  $\mathrm{H}^+$  ions contributed by HCl are:

Millimoles of 
$$H^+$$
(from HCl) =  $600 \times 0.01 = 6 \text{ mmol}$ .

The millimoles of H<sup>+</sup> ions contributed by H<sub>2</sub>SO<sub>4</sub> are:

Millimoles of 
$$H^+$$
(from  $H_2SO_4$ ) =  $400 \times 0.01 \times 2 = 8$  mmol.

Total millimoles of H<sup>+</sup>:

Total Millimoles of 
$$H^+ = 6 + 8 = 14 \text{ mmol}$$
.

# **Step 2: Calculate Concentration of H**<sup>+</sup>

The total volume of the solution is:

Total Volume = 
$$600 \,\text{mL} + 400 \,\text{mL} = 1000 \,\text{mL} = 1 \,\text{L}$$
.

The concentration of H<sup>+</sup> is:

$$[\mathrm{H^+}] = \frac{\mathrm{Total\ Millimoles\ of\ H^+}}{\mathrm{Total\ Volume\ (in\ mL)}} = \frac{14}{1000} = 14 \times 10^{-3}.$$

#### Step 3: Calculate pH of the Mixture

The pH of the solution is given by:

$$pH = -\log[H^+].$$

Substitute  $[H^{+}] = 14 \times 10^{-3}$ :

$$pH = -\log(14 \times 10^{-3}) = -(\log 14 + \log 10^{-3}).$$

$$pH = -(\log 14 - 3) = 3 - \log 14.$$

From the given values:

$$\log 14 = \log(2 \times 7) = \log 2 + \log 7 = 0.30 + 0.84 = 1.14.$$

Thus:

$$pH = 3 - 1.14 = 1.86.$$

Expressing pH as  $\times 10^{-2}$ :

$$pH = 186 \times 10^{-2}$$
.

**Conclusion:** The pH of the mixture is **186**  $\times 10^{-2}$ .

# Quick Tip

For calculating pH in mixtures, sum the millimoles of H<sup>+</sup> ions contributed by all acids and divide by the total volume of the solution. Use logarithmic properties to compute pH values efficiently.

29. A trisubstituted compound 'A',  $C_{10}H_{12}O_2$ , gives neutral FeCl<sub>3</sub> test positive.

Treatment of compound 'A' with NaOH and  $CH_3Br$  gives  $C_{11}H_{14}O_2$ , with hydroiodic acid gives methyl iodide and with hot conc. NaOH gives a compound 'B',  $C_{10}H_{12}O_2$ . Compound 'A' also decolourises alkaline  $KMnO_4$ . The number of  $\pi$  bond/s present in the compound 'A' is \_\_\_\_\_.

**Correct Answer:** 4

#### **Solution:**

#### Step 1: Analyze the Structure of Compound 'A'

Given the molecular formula  $C_{10}H_{12}O_2$ , compound 'A' gives a positive neutral FeCl<sub>3</sub> test, indicating the presence of a phenolic (-OH) group. Compound 'A' also decolourises alkaline KMnO<sub>4</sub>, indicating the presence of a double bond.

#### **Step 2: Chemical Reactions of 'A'**

- Treatment with NaOH and  $CH_3Br$  forms  $C_{11}H_{14}O_2$ , confirming the presence of a second hydroxyl group.
- Hydroiodic acid treatment yields methyl iodide, confirming the presence of an ether bond  $(-OCH_3)$ .
- Hot NaOH treatment forms compound 'B',  $C_{10}H_{12}O_2$ , which retains a double bond.

#### Step 3: Determine the Number of $\pi$ Bonds

The structure of compound 'A' contains:

- One aromatic benzene ring with three  $\pi$  bonds.
- One aliphatic double bond outside the aromatic ring.

Thus, the total number of  $\pi$  bonds in compound 'A' is:

Number of 
$$\pi$$
 bonds =  $3(aromatic) + 1(aliphatic) = 4$ .

**Conclusion:** The number of  $\pi$  bonds present in compound 'A' is 4.

#### Quick Tip

For determining the number of  $\pi$  bonds in a compound, carefully analyze the structure, including aromatic and aliphatic double bonds. Utilize chemical reactions (e.g., FeCl<sub>3</sub> test, KMnO<sub>4</sub> decolourisation) to identify functional groups and confirm bond types.

30. Some amount of dichloromethane ( $CH_2Cl_2$ ) is added to 671.141 mL of chloroform ( $CHCl_3$ ) to prepare a  $2.6 \times 10^{-3}$  M solution of  $CH_2Cl_2$  (DCM). The concentration of DCM is \_\_\_\_\_ ppm (by mass).

Given: Atomic mass C = 12, H = 1, Cl = 35.5, density of  $CHCl_3 = 1.49 \,\mathrm{g \ cm^{-3}}$ .

Correct Answer: 148 ppm

#### **Solution:**

#### **Step 1: Calculate the Mass of CH<sub>2</sub>Cl<sub>2</sub>**

Molarity (M) is related to the number of moles (n) and volume (V) by:

$$M = \frac{\text{moles of solute}}{\text{volume of solution (in L)}}$$
.

Substitute the values:

$$2.6 \times 10^{-3} = \frac{x}{85 \times 0.671141}.$$

Solve for x:

$$x = 2.6 \times 10^{-3} \times 85 \times 0.671141 = 0.148 \,\mathrm{g}.$$

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#### **Step 2: Calculate Concentration in ppm**

Concentration in ppm is given by:

Concentration (ppm) = 
$$\frac{\text{mass of solute (g)}}{\text{mass of solution (g)}} \times 10^6$$
.

The mass of the solution is:

Mass of solution = 
$$671.141 \times 1.49 = 1000.00009 \,\mathrm{g}$$
.

Thus:

$$\label{eq:concentration} \begin{aligned} &\text{Concentration (ppm)} = \frac{0.148}{671.141 \times 1.49} \times 10^6. \\ &\text{Concentration (ppm)} = \frac{0.148}{1000} \times 10^6 = 148 \, \text{ppm}. \end{aligned}$$

Conclusion: The concentration of  $CH_2Cl_2$  is 148 ppm.

# Quick Tip

For ppm calculations, always express the solute's mass and the solution's total mass in the same units before substituting into the formula. Ensure proper unit conversions for accuracy.