

CBSE Class 12 Chemistry 2026

Questions with Solutions

Set 56/3/3 · Section A · Detailed step-by-step solutions



General Instructions

- (i) This booklet contains 10 questions, each provided with a complete, step-by-step solution.
- (ii) It comprises 10 single-correct multiple-choice questions.
- (iii) Attempt each question on your own before reviewing the given solution.

1. According to Werner's theory, the primary valencies of the central metal atom :

- (A) are satisfied by neutral molecules or negative ions.
- (B) are equal to its coordination number.
- (C) are satisfied by negative ions.
- (D) are non-ionisable.

Correct Answer: (C) are satisfied by negative ions.

SOLUTION:

Approach by matching each property to the right valency. Werner described the metal's **primary** valency as the ionisable charge that equals the metal's oxidation state, and it is neutralised by anions; the **secondary** valency is the fixed, non-ionisable coordination number filled by ligands (neutral or anionic).

Eliminate: 'equal to coordination number' and 'satisfied by neutral molecules' and 'non-ionisable' all describe the secondary valency, so they are ruled out. Only 'satisfied by negative ions' is a property of the primary valency.

Hence the correct option is (C).

Quick Tip: Primary valency = oxidation number, ionisable, met by anions; secondary valency = coordination number.

2. The oxidation number of Pt in $[\text{Pt}(\text{en})_2\text{Cl}_2]^{2+}$ is :

- (A) + 3
- (B) + 4
- (C) + 2
- (D) + 6

Correct Answer: (B) + 4

SOLUTION:

Balance the charges on the complex ion. The two *en* ligands are neutral and contribute 0; the two Cl^- contribute $2 \times (-1) = -2$. Writing the metal oxidation state as *n* and equating to the ionic charge +2:

$$n + (-2) = +2 \Rightarrow n = +4$$

So platinum is in the +4 state. **Correct option: (B) +4.**

Quick Tip: en is neutral (0); each Cl is -1; sum of charges must equal the +2 on the ion.

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3. Which of the following product is formed when salicylic acid is treated with $(\text{CH}_3\text{CO})_2\text{O}$ in the presence of acid ?

- (A) $\text{o-C}_6\text{H}_4(\text{COOCH}_3)(\text{OH})$ — methyl 2-hydroxybenzoate
- (B) $\text{o-C}_6\text{H}_4(\text{OCOCH}_3)(\text{OH})$
- (C) $\text{o-C}_6\text{H}_4(\text{COCH}_3)(\text{COOH})$
- (D) $\text{o-C}_6\text{H}_4(\text{OCOCH}_3)(\text{COOH})$ — aspirin (acetylsalicylic acid)

Correct Answer: (D) $\text{o-C}_6\text{H}_4(\text{OCOCH}_3)(\text{COOH})$ — aspirin (acetylsalicylic acid)

SOLUTION:

Identify the functional groups and the reagent's job. Salicylic acid carries a *phenolic* hydroxyl next to a carboxylic acid. Acetic anhydride is an acetylating agent: under acid it converts a phenol -OH into an acetate ester -OCOCH_3 , leaving the -COOH group unchanged.

The result is acetylsalicylic acid (aspirin), with substituents -OCOCH_3 and -COOH on the ring. Options keeping a free -OH (no acetylation) or esterifying the acid instead are incorrect. **Correct option: (D).**

Quick Tip: Acetic anhydride acetylates the phenolic -OH of salicylic acid $\rightarrow$ aspirin; -COOH stays.

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4. Which reagent is used to distinguish between primary, secondary and tertiary amines ?

- (A) $\text{C}_6\text{H}_5\text{SO}_2\text{Cl}$
- (B) $\text{C}_6\text{H}_5\text{COCl}$
- (C) $\text{CHCl}_3 + \text{ethanolic KOH}$
- (D) $\text{NaOH} + \text{I}_2$

Correct Answer: (A) $\text{C}_6\text{H}_5\text{SO}_2\text{Cl}$

SOLUTION:

Decide which reagent gives three different responses. Carbylamine ($\text{CHCl}_3 + \text{ethanolic KOH}$) and the iodoform test ($\text{NaOH} + \text{I}_2$) are confirmatory tests, not a way to separate all three amine classes; benzoyl chloride acylates 1° and 2° alike. *Hinsberg's* reagent, benzenesulphonyl chloride, is the standard choice: $1^\circ \rightarrow$ alkali-soluble sulphonamide, $2^\circ \rightarrow$ alkali-insoluble sulphonamide, $3^\circ \rightarrow$ no reaction — three distinguishable results. **Correct option: (A).**

Quick Tip: Hinsberg's reagent = benzenesulphonyl chloride; gives 3 different responses for 1/2/3-amines.

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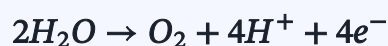
5. On electrolysis of very dilute aqueous solution of NaCl using platinum electrodes :

- (A) H₂ gas is evolved at anode.
- (B) Na is produced at anode.
- (C) O₂ gas is evolved at anode.
- (D) H₂ gas is evolved at cathode.

Correct Answer: (C) O₂ gas is evolved at anode.

SOLUTION:

Compare the anode candidates for dilute brine. Oxidising Cl⁻ to Cl₂ requires a reasonable chloride concentration; in a *very dilute* solution that route is suppressed, so water is oxidised instead, releasing O₂ at the anode:



Sodium is never deposited from aqueous solution and no gas is reduced at the anode, ruling out the other anode options. **The condition-specific answer is (C), O₂ at the anode** (H₂ is simultaneously released at the cathode).

Quick Tip: Dilute brine → water oxidised at anode → O₂ (Cl₂ needs concentrated NaCl); H₂ forms at cathode.

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6. The correct order of decreasing basic strength in aqueous solution of the following is :

- (A) (C₂H₅)₃N > C₂H₅NH₂ > (C₂H₅)₂NH
- (B) (C₂H₅)₂NH > C₂H₅NH₂ > (C₂H₅)₃N
- (C) C₂H₅NH₂ > (C₂H₅)₂NH > (C₂H₅)₃N
- (D) (C₂H₅)₂NH > (C₂H₅)₃N > C₂H₅NH₂

Correct Answer: (D) $(\text{C}_2\text{H}_5)_2\text{NH} > (\text{C}_2\text{H}_5)_3\text{N} > \text{C}_2\text{H}_5\text{NH}_2$

SOLUTION:

Recall the NCERT result for ethyl-substituted amines in aqueous medium. Although in the gas phase basicity rises $1^\circ < 2^\circ < 3^\circ$ (pure inductive effect), in *water* the solvation of the ammonium ion and steric factors rearrange the order. For the ethyl series the measured order is diethylamine $>$ triethylamine $>$ ethylamine.

Matching this to the options, only (D) reads $(\text{C}_2\text{H}_5)_2\text{NH} > (\text{C}_2\text{H}_5)_3\text{N} > \text{C}_2\text{H}_5\text{NH}_2$. **Correct option: (D).**

Quick Tip: Ethyl amines in water: 2-amine $>$ 3-amine $>$ 1-amine (solvation + steric + inductive balance).

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7. The mole fraction of a solute in 2.0 molal aqueous solution is :

- (A) 1.87
- (B) 0.347
- (C) 0.0347
- (D) 0.00347

Correct Answer: (C) 0.0347

SOLUTION:

Use the definition of molality and convert to mole fraction. Molality 2.0 means $n_{\text{solute}} = 2.0$ mol in 1000 g water, and $n_{\text{water}} = 1000/18 \approx 55.6$ mol.

$$x_{\text{solute}} = \frac{2.0}{2.0 + 55.6} \approx \frac{2.0}{57.6} \approx 0.0347$$

The value is close to $m \times M_{\text{water}}/1000 = 2.0 \times 18/1000 = 0.036$, confirming the order of magnitude. **Correct option: (C) 0.0347.**

Quick Tip: 2 molal = 2 mol solute in 1000 g water (55.56 mol); $x = 2/(2+55.56)$.

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8. Which of the following transition metals shows +1 and +2 oxidation states ?

- (A) Mn
- (B) Zn
- (C) Cu
- (D) Sc

Correct Answer: (C) Cu

SOLUTION:

Test each option against its characteristic states. Manganese ranges from +2 up to +7, scandium is fixed at +3, and zinc — with a full d^{10} core — shows only +2. **Copper** is the one element that is stable in both a +1 (cuprous) and a +2 (cupric) state, so it is the metal exhibiting +1 and +2. **Correct option: (C) Cu.**

Quick Tip: Cu = cuprous (+1) and cupric (+2); Zn only +2, Sc only +3, Mn +2 to +7.

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9. On hydrolysis, which of the following carbohydrates gives only β -glucose ?

- (A) Starch
- (B) Sucrose
- (C) Maltose
- (D) Cellulose

Correct Answer: (D) Cellulose

SOLUTION:

Work from the building blocks of each carbohydrate. Sucrose is glucose + fructose, so it cannot give only glucose; starch (amylose/amylopectin) and maltose are built from α -D-glucose units.

Cellulose alone is a polymer of β -D-glucose linked β -1,4, so its complete hydrolysis yields β -glucose exclusively. **Correct option: (D) Cellulose.**

Quick Tip: Cellulose = beta-1,4 polymer of beta-D-glucose; starch/maltose give alpha-glucose; sucrose gives glucose+fructose.

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10. Consider the following compounds : Chlorobenzene (I), 2,4,6-trinitrochlorobenzene (II), 2,4-dinitrochlorobenzene (III), 4-nitrochlorobenzene (IV). The correct order of increasing ease of nucleophilic substitution reactions of these compounds is :

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

Correct Answer: (A) $I < IV < III < II$

SOLUTION:

Rank by the count of ortho/para nitro groups, since each $-\text{NO}_2$ withdraws electron density and stabilises the intermediate formed when the nucleophile adds. Plain chlorobenzene has none and reacts the least; 4-nitro has one, 2,4-dinitro has two, and 2,4,6-trinitro (picryl chloride) has three and reacts the most. So ease increases $I < IV < III < II$. **Correct option: (A).**

Quick Tip: More o/p $-\text{NO}_2$ groups stabilise the intermediate $\rightarrow$ faster NAS: $0 < 1 < 2 < 3$ nitro groups.