



CLASS 12 · CHEMISTRY

ORGANIC CHEMISTRY

# Reaction Problems

Complete Study Guide



**RX**  
NAMED  
REACTIONS

**A → B**  
REACTION  
CHAINS

**→ C**  
PRODUCT  
IDENTIFICATION

★  
EXPLANATIONS  
INCLUDED

**NH<sub>2</sub>**  
AMINES &  
DIAZONIUM

**OH**  
PHENOLS &  
ALCOHOLS

**CHO**  
ALDEHYDES  
& KETONES

**COOH**  
CARBOXYLIC  
ACIDS

**70**  
QUESTIONS

**28**  
NAMED REACTIONS

**2**  
QUESTION SETS

**100%**  
EXAM READY



#### STEP-BY-STEP REACTIONS

Every reaction written with reagents, conditions and products



#### IUPAC & COMMON NAMES

Every compound labelled with formula and both names



#### WHY IT WORKS

Explanations for each mechanism — not just the answer



#### QUICK REFERENCE TABLE

Master named reactions chart at the end for fast revision



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## CLASS 12 — CHEMISTRY ORGANIC CHEMISTRY

*Complete Reaction Problems — With Reactions, Explanations & Named Reactions*

Compiled by

**Embrace Coaching Academy**

**70 Questions | Complete Reactions | Explanations | Answer Tables | Named Reactions**

## HOW TO USE THESE NOTES

### Each question is presented in 4 parts:

1. Question — the original problem statement
2. Named Reaction — the key reaction concept involved
3. Step-by-step Reactions — the actual chemistry
4. Answer Table — labelled compounds with formulas and names

*Explanation: Orange boxes contain key explanations. Always read them — they explain WHY the reaction works, not just WHAT the product is. Named reactions are in purple. Answers are in green tables.*

## SECTION 1 — SET A: 20 QUESTIONS

### Q1. C<sub>6</sub>H<sub>6</sub>O (A) — Phenol with NH<sub>3</sub> and Zn

#### Named Reaction: Reduction & Ammonolysis of Phenol

#### Reactions:

Step	Reaction
A→B	$C_6H_5OH + NH_3 \rightarrow C_6H_5NH_2$ (Aniline)
A→C	$C_6H_5OH + Zn \rightarrow C_6H_6 + ZnO$ (Benzene)

*Explanation: Phenol (C<sub>6</sub>H<sub>5</sub>OH) reacts with ammonia (NH<sub>3</sub>) under catalyst to give Aniline (primary amine). With zinc, the -OH group is removed and benzene is produced (reduction of phenol).*

#### Answer Summary:

Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> OH	Phenol
B	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>	Aniline
C	C <sub>6</sub> H <sub>6</sub>	Benzene

### Q2. C<sub>7</sub>H<sub>6</sub>O (A) — Benzaldehyde via Cannizzaro reaction

#### Named Reaction: Cannizzaro Reaction + Decarboxylation (Sodalime)

#### Reactions:

Step	Reaction
A	$C_7H_6O = C_6H_5CHO$ (Benzaldehyde)
A→B,C	$2C_6H_5CHO + 50\%NaOH \rightarrow C_6H_5CH_2OH$ (B) + $C_6H_5COONa$ (C) [Cannizzaro]
C→D	$C_6H_5COONa + HCl \rightarrow C_6H_5COOH$ (D) + NaCl
D→E	$C_6H_5COOH + NaOH + CaO/heat \rightarrow C_6H_6$ (E) + CaCO <sub>3</sub> [Sodalime]

*Explanation: Benzaldehyde (no alpha-H) undergoes disproportionation with NaOH (Cannizzaro). One molecule is oxidised to sodium benzoate (C), the other is reduced to benzyl alcohol (B). Acid gives benzoic acid (D); dry distillation with sodalime gives benzene (E).*

**Answer Summary:**

Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> CHO	Benzaldehyde
B	C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> OH	Benzyl Alcohol
C	C <sub>6</sub> H <sub>5</sub> COONa	Sodium Benzoate
D	C <sub>6</sub> H <sub>5</sub> COOH	Benzoic Acid
E	C <sub>6</sub> H <sub>6</sub>	Benzene

**Q3. C<sub>2</sub>H<sub>6</sub>O (A) — Ethanol dehydration, hydroxylation, etherification****Named Reaction: Dehydration + Baeyer's Hydroxylation + Etherification****Reactions:**

Step	Reaction
A→B	C <sub>2</sub> H <sub>5</sub> OH + conc.H <sub>2</sub> SO <sub>4</sub> /443K → CH <sub>2</sub> =CH <sub>2</sub> (Dehydration)
B→C	CH <sub>2</sub> =CH <sub>2</sub> + Baeyer's reagent (alk.KMnO <sub>4</sub> ) → HOCH <sub>2</sub> CH <sub>2</sub> OH (Ethane-1,2-diol)
C→D	HOCH <sub>2</sub> CH <sub>2</sub> OH + anhydrous ZnCl <sub>2</sub> → CH <sub>3</sub> CHO (Ethanal)
A→E	C <sub>2</sub> H <sub>5</sub> OH + conc.H <sub>2</sub> SO <sub>4</sub> /413K → C <sub>2</sub> H <sub>5</sub> OC <sub>2</sub> H <sub>5</sub> (Ethoxyethane)

*Explanation: At 443K, ethanol loses water to form ethylene. Baeyer's reagent (cold alkaline KMnO<sub>4</sub>) hydroxylates the double bond to ethane-1,2-diol. At 413K, intermolecular dehydration gives diethyl ether. Temperature controls which product forms.*

**Answer Summary:**

Label	Formula	Name
A	C <sub>2</sub> H <sub>5</sub> OH	Ethanol
B	CH <sub>2</sub> =CH <sub>2</sub>	Ethylene (Ethene)
C	HOCH <sub>2</sub> CH <sub>2</sub> OH	Ethane-1,2-diol (Glycol)

<b>D</b>	CH <sub>3</sub> CHO	<b>Ethanal</b>
<b>E</b>	C <sub>2</sub> H <sub>5</sub> OC <sub>2</sub> H <sub>5</sub>	<b>Ethoxyethane (Diethyl Ether)</b>

#### Q4. CH<sub>4</sub>O (A) — Methanol to Ethanol via Grignard

##### Named Reaction: Grignard Reaction (Chain Extension by 1 Carbon)

##### Reactions:

Step	Reaction
<b>A→B</b>	CH <sub>3</sub> OH + Tollens' reagent → HCHO (Methanal)
<b>B→C</b>	HCHO + CH <sub>3</sub> MgBr → [CH <sub>3</sub> CH <sub>2</sub> OMgBr] + H <sub>2</sub> O/H <sup>+</sup> → CH <sub>3</sub> CH <sub>2</sub> OH (Ethanol)

*Explanation: Methanol oxidised by Tollens' reagent gives methanal (formaldehyde). Methanal reacts with CH<sub>3</sub>MgBr (Grignard reagent) — the carbanion adds to the carbonyl. Hydrolysis gives ethanol. This is a classic one-carbon chain extension.*

##### Answer Summary:

Label	Formula	Name
<b>A</b>	CH <sub>3</sub> OH	<b>Methanol</b>
<b>B</b>	HCHO	<b>Methanal (Formaldehyde)</b>
<b>C</b>	CH <sub>3</sub> CH <sub>2</sub> OH	<b>Ethanol</b>

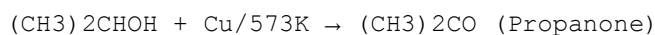
#### Q5. C<sub>2</sub>H<sub>6</sub>O (A) — Ethanol to Propanone via Grignard

##### Named Reaction: Grignard Reaction + Oxidative Dehydrogenation (Cu/573K)

##### Reactions:

Step	Reaction
<b>A→B</b>	C <sub>2</sub> H <sub>5</sub> OH + Cu/573K → CH <sub>3</sub> CHO (Ethanal) [Dehydrogenation]
<b>B→C</b>	CH <sub>3</sub> CHO + CH <sub>3</sub> MgBr + H <sub>2</sub> O/H <sup>+</sup> → (CH <sub>3</sub> ) <sub>2</sub> CHOH (Propan-2-ol)

C→D



*Explanation: Ethanol dehydrogenates at Cu/573K to ethanal. The Grignard reagent CH<sub>3</sub>MgBr adds to ethanal; hydrolysis gives propan-2-ol (secondary alcohol). Cu/573K again oxidises the secondary alcohol to propanone (a ketone).*

**Answer Summary:**

Label	Formula	Name
A	C <sub>2</sub> H <sub>5</sub> OH	Ethanol
B	CH <sub>3</sub> CHO	Ethanal
C	(CH <sub>3</sub> ) <sub>2</sub> CHOH	Propan-2-ol
D	(CH <sub>3</sub> ) <sub>2</sub> CO	Propanone (Acetone)

**Q6. C<sub>6</sub>H<sub>5</sub>Cl (A) — Chlorobenzene to Aniline via Dow's Process****Named Reaction: Dow's Process + Ammonolysis of Phenol****Reactions:**

Step	Reaction
A→B	C <sub>6</sub> H <sub>5</sub> Cl + NaOH (high T, P) → C <sub>6</sub> H <sub>5</sub> OH (Phenol) [Dow's Process]
B→C	C <sub>6</sub> H <sub>5</sub> OH + NH <sub>3</sub> + anhydrous ZnCl <sub>2</sub> /heat → C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub> (Aniline)

*Explanation: Chlorobenzene heated with NaOH under high temperature and pressure (Dow's process) hydrolyses to phenol. Phenol then reacts with ammonia in presence of anhydrous ZnCl<sub>2</sub> to form aniline.*

**Answer Summary:**

Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> Cl	Chlorobenzene
B	C <sub>6</sub> H <sub>5</sub> OH	Phenol
C	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>	Aniline

**Q7. C<sub>6</sub>H<sub>5</sub>Cl (A) — Chlorobenzene to Salicylic Acid (Kolbe's Reaction)****Named Reaction: Dow's Process + Kolbe's Reaction****Reactions:**

Step	Reaction
<b>A→B</b>	$C_6H_5Cl + NaOH \rightarrow C_6H_5OH$ (Phenol) [Dow's Process]
<b>B→C</b>	$C_6H_5OH + NaOH \rightarrow C_6H_5ONa$ (Sodium Phenoxide)
<b>C→D</b>	$C_6H_5ONa + CO_2 / 400K / 4-7 \text{ bar} \rightarrow \text{Sodium Salicylate} \rightarrow H^+/H_2O \rightarrow C_6H_4(OH)COOH$ [Kolbe's Reaction]

*Explanation: Phenol forms sodium phenoxide with NaOH. Treating with CO<sub>2</sub> under pressure at 400K introduces a carboxyl group at the ortho position (Kolbe's reaction). Acidification gives salicylic acid — the parent compound of Aspirin.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	C <sub>6</sub> H <sub>5</sub> Cl	<b>Chlorobenzene</b>
<b>B</b>	C <sub>6</sub> H <sub>5</sub> OH	<b>Phenol</b>
<b>C</b>	C <sub>6</sub> H <sub>5</sub> ONa	<b>Sodium Phenoxide</b>
<b>D</b>	C <sub>6</sub> H <sub>4</sub> (OH)COOH	<b>Salicylic Acid (2-Hydroxybenzoic acid)</b>

**Q8. C<sub>6</sub>H<sub>5</sub>N<sub>2</sub>Cl (A) — Diazonium salt to Toluene****Named Reaction: Hydrolysis + Friedel-Crafts Alkylation****Reactions:**

Step	Reaction
<b>A→B</b>	$C_6H_5N_2Cl + H_2O \rightarrow C_6H_5OH$ (Phenol) + N <sub>2</sub> + HCl
<b>B→C</b>	$C_6H_5OH + Zn \rightarrow C_6H_6$ (Benzene)
<b>C→D</b>	$C_6H_6 + CH_3Cl + \text{anhydrous } AlCl_3 \rightarrow C_6H_5CH_3$ (Toluene) [Friedel-Crafts]

*Explanation: Diazonium chloride hydrolyses to phenol. Zinc reduces phenol to benzene (simplest aromatic hydrocarbon). Benzene undergoes Friedel-Crafts alkylation with  $\text{CH}_3\text{Cl}/\text{AlCl}_3$  to give toluene (methylbenzene).*

**Answer Summary:**

Label	Formula	Name
A	$\text{C}_6\text{H}_5\text{N}_2\text{Cl}$	<b>Benzene Diazonium Chloride</b>
B	$\text{C}_6\text{H}_5\text{OH}$	<b>Phenol</b>
C	$\text{C}_6\text{H}_6$	<b>Benzene</b>
D	$\text{C}_6\text{H}_5\text{CH}_3$	<b>Toluene (Methylbenzene)</b>

**Q9.  $\text{C}_6\text{H}_6$  (A) — Benzene to Phenol via Cumene Process****Named Reaction: Cumene Process (Industrial Phenol Synthesis)****Reactions:**

Step	Reaction
A→B	$\text{C}_6\text{H}_6 + \text{CH}_3\text{CH}=\text{CH}_2 + \text{H}_3\text{PO}_4/523\text{K} \rightarrow \text{C}_6\text{H}_5\text{CH}(\text{CH}_3)_2$ (Cumene)
B→C	$\text{C}_6\text{H}_5\text{CH}(\text{CH}_3)_2 + \text{Air}/\text{O}_2 \rightarrow \text{C}_6\text{H}_5\text{C}(\text{CH}_3)_2\text{OOH}$ (Cumene Hydroperoxide)
C→D	$\text{C}_6\text{H}_5\text{C}(\text{CH}_3)_2\text{OOH} + \text{H}_2\text{SO}_4 \rightarrow \text{C}_6\text{H}_5\text{OH}$ (Phenol) + $\text{CH}_3\text{COCH}_3$ (Acetone)

*Explanation: Benzene alkylated with propylene (Friedel-Crafts,  $\text{H}_3\text{PO}_4$  catalyst) gives cumene. Air oxidation gives cumene hydroperoxide. Acid cleavage simultaneously gives phenol and acetone — an important industrial synthesis.*

**Answer Summary:**

Label	Formula	Name
A	$\text{C}_6\text{H}_6$	<b>Benzene</b>
B	$\text{C}_6\text{H}_5\text{CH}(\text{CH}_3)_2$	<b>Cumene (Isopropylbenzene)</b>
C	$\text{C}_6\text{H}_5\text{C}(\text{CH}_3)_2\text{OOH}$	<b>Cumene Hydroperoxide</b>
D	$\text{C}_6\text{H}_5\text{OH}$	<b>Phenol</b>

**Q10. C<sub>3</sub>H<sub>8</sub>O (A) — Propan-2-ol to Nitroso compound****Named Reaction: Halogenation, Nitration & Nitrosation****Reactions:**

Step	Reaction
<b>A→B</b>	$\text{CH}_3\text{CH}(\text{OH})\text{CH}_3 + \text{P/I}_2 \rightarrow \text{CH}_3\text{CHICH}_3$ (2-Iodopropane)
<b>B→C</b>	$\text{CH}_3\text{CHICH}_3 + \text{AgNO}_2 \rightarrow \text{CH}_3\text{CH}(\text{NO}_2)\text{CH}_3$ (2-Nitropropane)
<b>C→D</b>	$\text{CH}_3\text{CH}(\text{NO}_2)\text{CH}_3 + \text{HNO}_2 \rightarrow \text{CH}_3\text{C}(\text{NO}_2)(\text{NO})\text{CH}_3$ [Blue colour – 2-Nitro-2-nitrosopropane]

*Explanation: Propan-2-ol reacts with P/I<sub>2</sub> to give 2-iodopropane. AgNO<sub>2</sub> (silver nitrite) displaces iodide via S<sub>N</sub>2 to give 2-nitropropane. Treatment with nitrous acid adds a nitroso group, giving a blue-coloured compound.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	CH <sub>3</sub> CH(OH)CH <sub>3</sub>	<b>Propan-2-ol</b>
<b>B</b>	CH <sub>3</sub> CHICH <sub>3</sub>	<b>2-Iodopropane</b>
<b>C</b>	CH <sub>3</sub> CH(NO <sub>2</sub> )CH <sub>3</sub>	<b>2-Nitropropane</b>
<b>D</b>	CH <sub>3</sub> C(NO <sub>2</sub> )(NO)CH <sub>3</sub>	<b>2-Nitro-2-nitrosopropane (Blue colour)</b>

**Q11. C<sub>2</sub>H<sub>6</sub>O (A) — Ethanol to Ethane-1,2-diol****Named Reaction: Dehydration + Baeyer's Hydroxylation****Reactions:**

Step	Reaction
<b>A→B</b>	$\text{CH}_3\text{CH}_2\text{OH} + \text{Al}_2\text{O}_3/620\text{K} \rightarrow \text{CH}_2=\text{CH}_2$ (Ethene)
<b>B→C</b>	$\text{CH}_2=\text{CH}_2 + \text{Baeyer's reagent} \rightarrow \text{CH}_2(\text{OH})\text{CH}_2(\text{OH})$ (Ethane-1,2-diol)

*Explanation: Ethanol dehydrates over alumina at 620K to give ethylene. Baeyer's reagent (cold alkaline KMnO<sub>4</sub> or OsO<sub>4</sub>) oxidises the double bond (syn addition) to give ethane-1,2-diol (ethylene glycol — used as antifreeze).*

**Answer Summary:**

Label	Formula	Name
A	CH <sub>3</sub> CH <sub>2</sub> OH	Ethanol
B	CH <sub>2</sub> =CH <sub>2</sub>	Ethene (Ethylene)
C	HOCH <sub>2</sub> CH <sub>2</sub> OH	Ethane-1,2-diol (Ethylene glycol)

### Q12. C<sub>6</sub>H<sub>6</sub>O (A) — Phenol via Reimer-Tiemann Reaction

#### Named Reaction: Reimer-Tiemann Reaction + Ammonolysis

#### Reactions:

Step	Reaction
A→B	C <sub>6</sub> H <sub>5</sub> OH + CHCl <sub>3</sub> /NaOH → 2-HO-C <sub>6</sub> H <sub>4</sub> -CHO (Salicylaldehyde) [Reimer-Tiemann]
A→C	C <sub>6</sub> H <sub>5</sub> OH + NH <sub>3</sub> /anhydrous ZnCl <sub>2</sub> /heat → C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub> (Aniline)

*Explanation: Phenol treated with chloroform and NaOH — the intermediate :CCl<sub>2</sub> (dichlorocarbene) electrophile attacks the ring at ortho position. After hydrolysis, salicylaldehyde forms (Reimer-Tiemann). Separately, phenol with NH<sub>3</sub>/ZnCl<sub>2</sub> gives aniline.*

#### Answer Summary:

Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> OH	Phenol
B	2-HO-C <sub>6</sub> H <sub>4</sub> -CHO	Salicylaldehyde (2-Hydroxybenzaldehyde)
C	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>	Aniline

### Q13. C<sub>6</sub>H<sub>5</sub>NO<sub>2</sub> (A) — Nitrobenzene to N-Methyl Aniline

#### Named Reaction: Reduction + Carbylamine Reaction + Catalytic Reduction

#### Reactions:

Step	Reaction
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<b>A→B</b>	$C_6H_5NO_2 + Sn/HCl \rightarrow C_6H_5NH_2$ (Aniline)
<b>B→C</b>	$C_6H_5NH_2 + CHCl_3/3KOH \rightarrow C_6H_5NC$ (Phenyl Isocyanide) [Carbylamine test]
<b>C→D</b>	$C_6H_5NC + Na/C_2H_5OH$ or $Ni/H_2 \rightarrow C_6H_5NHCH_3$ (N-Methyl Aniline)

*Explanation: Nitrobenzene reduces to aniline (Sn/HCl). Aniline undergoes carbylamine reaction (isocyanide reaction — foul smell, used to detect primary amines) giving phenyl isocyanide. Reduction of isocyanide gives N-methyl aniline (secondary amine).*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$C_6H_5NO_2$	<b>Nitrobenzene</b>
<b>B</b>	$C_6H_5NH_2$	<b>Aniline</b>
<b>C</b>	$C_6H_5NC$	<b>Phenyl Carbylamine (Isocyanide)</b>
<b>D</b>	$C_6H_5NHCH_3$	<b>N-Methyl Aniline</b>

**Q14.  $C_3H_4$  (A) — Prop-1-yne to multiple products**

**Named Reaction: Hydration (Markovnikov) + Iodoform + Wolff-Kishner + Crossed Cannizzaro**

**Reactions:**

Step	Reaction
<b>A→B</b>	$CH_3C\equiv CH + Hg^{2+}/H_2SO_4/H_2O \rightarrow CH_3COCH_3$ (Propanone) [Markovnikov's rule]
<b>B→(test)</b>	$CH_3COCH_3 + 3I_2/NaOH \rightarrow CHI_3 + CH_3COONa$ [Positive Iodoform Test]
<b>B→C</b>	$CH_3COCH_3 + N_2H_4/C_2H_5ONa \rightarrow CH_3CH_2CH_3$ (Propane) [Wolff-Kishner reduction]
<b>B→D</b>	$CH_3COCH_3 + HCHO + dil.NaOH \rightarrow CH_2(OH)CH_2COCH_3$ [Crossed Cannizzaro]

*Explanation: Prop-1-yne hydrated following Markovnikov's rule gives propanone (methyl ketone). Propanone gives positive iodoform test (yellow  $CHI_3$  precipitate). Wolff-Kishner ( $N_2H_4$  + base) removes  $C=O$  to give propane. Crossed Cannizzaro with  $HCHO$  gives 4-hydroxybutan-2-one.*

**Answer Summary:**

Label	Formula	Name
A	$\text{CH}_3\text{C}\equiv\text{CH}$	Prop-1-yne
B	$\text{CH}_3\text{COCH}_3$	Propanone (Acetone)
C	$\text{CH}_3\text{CH}_2\text{CH}_3$	Propane (via Wolff-Kishner)
D	$\text{CH}_2(\text{OH})\text{CH}_2\text{COCH}_3$	4-Hydroxybutan-2-one

### Q15. $\text{C}_3\text{H}_8\text{O}_3$ (A) — Glycerol via Fenton's Reagent

#### Named Reaction: Fenton's Reagent Oxidation of Glycerol

#### Reactions:

Step	Reaction
A	A = $\text{CH}_2\text{OH}-\text{CHOH}-\text{CH}_2\text{OH}$ (Glycerol / Propan-1,2,3-triol)
A→B,C	Glycerol + $\text{FeSO}_4 + \text{H}_2\text{O}_2 \rightarrow \text{CHO}-\text{CHOH}-\text{CH}_2\text{OH}$ (B) + $\text{CH}_2\text{OH}-\text{CO}-\text{CH}_2\text{OH}$ (C)
Note	Mixture of B and C = Glycerose

*Explanation: Glycerol is a trihydric alcohol. Fenton's reagent ( $\text{FeSO}_4 + \text{H}_2\text{O}_2$ ) oxidises either the central or terminal  $-\text{CH}(\text{OH})-$  to give glyceraldehyde (B = 2,3-dihydroxypropanal) or dihydroxyacetone (C = 1,3-dihydroxypropan-2-one). The mixture is called glycerose.*

#### Answer Summary:

Label	Formula	Name
A	$\text{HOCH}_2\text{CH}(\text{OH})\text{CH}_2\text{OH}$	Propan-1,2,3-triol (Glycerol)
B	$\text{CHO}-\text{CHOH}-\text{CH}_2\text{OH}$	2,3-Dihydroxy Propanal (Glyceraldehyde)
C	$\text{CH}_2\text{OH}-\text{CO}-\text{CH}_2\text{OH}$	1,3-Dihydroxypropan-2-one (Dihydroxyacetone)

### Q16. $\text{C}_7\text{H}_7\text{NO}$ (A) — Benzamide to Azo Dye

#### Named Reaction: Hoffmann Bromamide Reaction + Diazotisation + Azo Coupling

#### Reactions:

Step	Reaction
A→B	$C_6H_5CONH_2 + Br_2/KOH \rightarrow C_6H_5NH_2$ (Aniline) [Hoffmann Bromamide - 1 carbon less]
B→C	$C_6H_5NH_2 + NaNO_2/HCl/273-278K \rightarrow C_6H_5N_2Cl$ (Benzene Diazonium Chloride) [Diazotisation]
C→D	$C_6H_5N_2Cl + p\text{-cresol}/pH\ 9-10 \rightarrow 2\text{-Phenylazo-4-methylphenol}$ (Azo dye) [Coupling]

**Explanation:** Benzamide loses one carbon via Hoffmann bromamide rearrangement to give aniline (C7→C6). Aniline diazotises with NaNO<sub>2</sub>/HCl at 273-278K to give diazonium salt. Coupling with p-cresol at pH 9-10 gives an orange-red azo dye.

**Answer Summary:**

Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> CONH <sub>2</sub>	Benzamide
B	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>	Aniline
C	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl	Benzene Diazonium Chloride
D	CH <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -N=N-C <sub>6</sub> H <sub>5</sub> (OH)	2-Phenylazo-4-methylphenol (Azo Dye)

**Q17. C<sub>2</sub>H<sub>4</sub>O (A) — Methanal reactions****Named Reaction: Acetal Formation + Tischenko Reaction****Reactions:**

Step	Reaction
A	$C_2H_4O = HCHO$ (Methanal / Formaldehyde)
A→B	$HCHO + Methanol/HCl \rightarrow C_4H_{10}O_2$ (Dimethoxymethane - Acetal)
A→C	$HCHO + dil.NaOH \rightarrow HCOOCH_3$ (Methyl Formate) [Tischenko Reaction]

**Explanation:** Methanal reacts with methanol in HCl to form an acetal (dimethoxymethane). With dilute NaOH (Tischenko reaction), methanal disproportionates to methyl formate — one HCHO is oxidised, the other reduced.

**Answer Summary:**

Label	Formula	Name
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<b>A</b>	HCHO	<b>Methanal (Formaldehyde)</b>
<b>B</b>	CH <sub>3</sub> OCH <sub>2</sub> OCH <sub>3</sub>	<b>Dimethoxymethane (Acetal)</b>
<b>C</b>	HCOOCH <sub>3</sub>	<b>Methyl Formate (Tischenko product)</b>

### Q18. C<sub>2</sub>H<sub>5</sub>Br (A) — Bromoethane to Propanoic Acid via Grignard

#### Named Reaction: Grignard Reaction + Carboxylation (CO<sub>2</sub>)

#### Reactions:

Step	Reaction
<b>A→B</b>	C <sub>2</sub> H <sub>5</sub> Br + Mg/dry ether → C <sub>2</sub> H <sub>5</sub> MgBr (Ethyl Magnesium Bromide)
<b>B→C</b>	C <sub>2</sub> H <sub>5</sub> MgBr + CO <sub>2</sub> /dry ether → C <sub>2</sub> H <sub>5</sub> COOMgBr + H <sub>2</sub> O/H <sup>+</sup> → C <sub>2</sub> H <sub>5</sub> COOH

*Explanation: Bromoethane reacts with Mg in dry ether to form a Grignard reagent (strong nucleophile). Reaction with dry CO<sub>2</sub> followed by hydrolysis gives propanoic acid — chain is extended by one carbon (C<sub>2</sub> → C<sub>3</sub> acid).*

#### Answer Summary:

Label	Formula	Name
<b>A</b>	C <sub>2</sub> H <sub>5</sub> Br	<b>Bromoethane</b>
<b>B</b>	C <sub>2</sub> H <sub>5</sub> MgBr	<b>Ethyl Magnesium Bromide (Grignard Reagent)</b>
<b>C</b>	C <sub>2</sub> H <sub>5</sub> COOH	<b>Propanoic Acid</b>

### Q19. (A) Nitrobenzene → N-Methyl Aniline, N-Phenyl Benzamide, Diazonium Salt

#### Named Reaction: Reduction + Schotten-Baumann + Diazotisation

#### Reactions:

Step	Reaction
<b>A→B</b>	C <sub>6</sub> H <sub>5</sub> NO <sub>2</sub> + Sn/HCl → C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub> (Aniline)
<b>B→C</b>	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub> + C <sub>6</sub> H <sub>5</sub> COCl/Pyridine → C <sub>6</sub> H <sub>5</sub> NHCOC <sub>6</sub> H <sub>5</sub> (N-Phenyl Benzamide) [Schotten-Baumann]

<b>B→D</b>	$C_6H_5NH_2 + CH_3Br \rightarrow C_6H_5NHCH_3$ (N-Methyl Aniline)
<b>B→E</b>	$C_6H_5NH_2 + NaNO_2/HCl \rightarrow C_6H_5N_2Cl$ (Benzene Diazonium Chloride) [Diazotisation]

**Explanation:** Nitrobenzene reduces to aniline. Aniline reacts with benzoyl chloride in pyridine (Schotten-Baumann) to give N-phenyl benzamide. Methylation gives N-methyl aniline. Diazotisation at 273-278K gives diazonium salt.

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$C_6H_5NO_2$	<b>Nitrobenzene</b>
<b>B</b>	$C_6H_5NH_2$	<b>Aniline</b>
<b>C</b>	$C_6H_5NHCOC_6H_5$	<b>N-Phenyl Benzamide</b>
<b>D</b>	$C_6H_5NHCH_3$	<b>N-Methyl Aniline</b>
<b>E</b>	$C_6H_5N_2Cl$	<b>Benzene Diazonium Chloride</b>

**Q20.  $C_2H_6O$  (A) — Ethanol complete reaction chain****Named Reaction: Temperature-Controlled Dehydration + Baeyer's + Etherification****Reactions:**

Step	Reaction
<b>A→B</b>	$C_2H_5OH + \text{conc. } H_2SO_4/443K \rightarrow CH_2=CH_2$ (Ethylene)
<b>B→C</b>	$CH_2=CH_2 + \text{Baeyer's reagent} \rightarrow HOCH_2CH_2OH$ (Ethane-1,2-diol)
<b>A→E</b>	$C_2H_5OH + \text{conc. } H_2SO_4/413K \rightarrow C_2H_5OC_2H_5$ (Diethyl Ether)

**Explanation:** Same as Q3. Key point: Temperature determines product. At 443K → alkene (intramolecular dehydration). At 413K → ether (intermolecular dehydration). Baeyer's reagent converts alkene to glycol.

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$C_2H_5OH$	<b>Ethanol</b>
<b>B</b>	$CH_2=CH_2$	<b>Ethylene</b>

<b>C</b>	HOCH <sub>2</sub> CH <sub>2</sub> OH	<b>Ethane-1,2-diol</b>
<b>E</b>	C <sub>2</sub> H <sub>5</sub> OC <sub>2</sub> H <sub>5</sub>	<b>Ethoxyethane (Diethyl Ether)</b>

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## SECTION 2 — SET B: 50 QUESTIONS (SS Prithvi Set)

**Q1.**  $\text{CH}_3\text{-CHO} + \text{NH}_2\text{OH} \rightarrow \text{A}$ ;  $\text{A} + \text{P}_2\text{O}_5 \rightarrow \text{B}$

**Named Reaction: Oxime Formation + Dehydration to Nitrile**

Reactions:

Step	Reaction
A	$\text{CH}_3\text{-CHO} + \text{NH}_2\text{OH} \rightarrow \text{CH}_3\text{-CH=N-OH}$ (Acetaldoxime)
B	$\text{CH}_3\text{-CH=N-OH} + \text{P}_2\text{O}_5 \rightarrow \text{CH}_3\text{-CN}$ (Methylcyanide / Acetonitrile)

*Explanation: Acetaldehyde reacts with hydroxylamine (NH<sub>2</sub>OH) via condensation to form acetaldoxime (an oxime). P<sub>2</sub>O<sub>5</sub> is a dehydrating agent that removes water from the oxime to give methyl cyanide (nitrile).*

Answer Summary:

Label	Formula	Name
A	$\text{CH}_3\text{-CH=N-OH}$	Acetaldoxime
B	$\text{CH}_3\text{-CN}$	Methylcyanide (Acetonitrile)

**Q2.**  $\text{CH}_3\text{-CH=CH}_2$  with  $\text{H}_2\text{O/H}^+$  and  $\text{B}_2\text{H}_6/\text{H}_2\text{O}_2/\text{OH}^-$

**Named Reaction: Markovnikov Addition vs Anti-Markovnikov (Hydroboration-Oxidation)**

Reactions:

Step	Reaction
A (acid-catalysed)	$\text{CH}_3\text{-CH=CH}_2 + \text{H}_2\text{O/H}^+ \rightarrow \text{CH}_3\text{-CH(OH)-CH}_3$ (Isopropyl alcohol)
B (hydroboration)	$\text{CH}_3\text{-CH=CH}_2 + \text{B}_2\text{H}_6$ , then $\text{H}_2\text{O}_2/\text{OH}^- \rightarrow \text{CH}_3\text{-CH}_2\text{-CH}_2\text{-OH}$ (n-Propyl alcohol)

*Explanation: H<sub>2</sub>O/H<sup>+</sup> follows Markovnikov's rule — H adds to less substituted carbon, OH to more substituted — giving isopropyl alcohol (2°). Hydroboration-oxidation (B<sub>2</sub>H<sub>6</sub> then H<sub>2</sub>O<sub>2</sub>/OH<sup>-</sup>) is anti-Markovnikov giving n-propyl alcohol (1°). This is a key comparison!*

**Answer Summary:**

Label	Formula	Name
A	CH <sub>3</sub> CH(OH)CH <sub>3</sub>	Isopropyl Alcohol (Propan-2-ol) [Markovnikov]
B	CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> OH	n-Propyl Alcohol (Propan-1-ol) [Anti-Markovnikov]

**Q3.** C<sub>6</sub>H<sub>5</sub>-OH with Zn and with NaOH + C<sub>6</sub>H<sub>5</sub>COCl**Named Reaction: Reduction of Phenol + Ester Formation (Schotten-Baumann)****Reactions:**

Step	Reaction
A	C <sub>6</sub> H <sub>5</sub> -OH + Zn → C <sub>6</sub> H <sub>6</sub> + ZnO (Benzene)
B	C <sub>6</sub> H <sub>5</sub> -OH + C <sub>6</sub> H <sub>5</sub> -COCl + NaOH → C <sub>6</sub> H <sub>5</sub> -CO-O-C <sub>6</sub> H <sub>5</sub> + NaCl + H <sub>2</sub> O (Phenyl Benzoate)

*Explanation: Zinc reduces phenol to benzene by removing the -OH group. With benzoyl chloride and NaOH, phenol forms phenyl benzoate — an ester. This is the Schotten-Baumann condition for ester formation from phenol.*

**Answer Summary:**

Label	Formula	Name
A	C <sub>6</sub> H <sub>6</sub>	Benzene
B	C <sub>6</sub> H <sub>5</sub> COOC <sub>6</sub> H <sub>5</sub>	Phenyl Benzoate (Ester)

**Q4.** C<sub>2</sub>H<sub>6</sub>O (A) → C<sub>2</sub>H<sub>4</sub>O<sub>2</sub> (B) → C<sub>4</sub>H<sub>6</sub>O<sub>4</sub>Ca (C) → C<sub>3</sub>H<sub>6</sub>O (D)**Named Reaction: Oxidation + Calcium Salt Formation + Dry Distillation (Ketone synthesis)****Reactions:**

Step	Reaction
A	CH <sub>3</sub> CHO (Acetaldehyde)
A→B	CH <sub>3</sub> CHO + K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> → CH <sub>3</sub> COOH (Acetic acid)

<b>B→C</b>	$2\text{CH}_3\text{COOH} + \text{Ca}(\text{OH})_2 \rightarrow (\text{CH}_3\text{COO})_2\text{Ca}$ (Calcium acetate)
<b>C→D</b>	$(\text{CH}_3\text{COO})_2\text{Ca} + \text{heat/distillation} \rightarrow \text{CH}_3\text{COCH}_3 + \text{CaCO}_3$ (Acetone)

**Explanation:** Acetaldehyde oxidised by  $\text{K}_2\text{Cr}_2\text{O}_7$  gives acetic acid. Neutralisation with  $\text{Ca}(\text{OH})_2$  gives calcium acetate. Dry distillation of calcium acetate gives acetone (ketone) — a classical method of ketone synthesis from carboxylic acid calcium salts.

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{CHO}$	<b>Acetaldehyde</b>
<b>B</b>	$\text{CH}_3\text{COOH}$	<b>Acetic Acid</b>
<b>C</b>	$(\text{CH}_3\text{COO})_2\text{Ca}$	<b>Calcium Acetate</b>
<b>D</b>	$\text{CH}_3\text{COCH}_3$	<b>Acetone</b>

**Q5.  $\text{C}_6\text{H}_5\text{-NO}_2$  with three different reagents****Named Reaction: Selective Reduction of Nitrobenzene****Reactions:**

Step	Reaction
<b>A</b>	$\text{C}_6\text{H}_5\text{-NO}_2 + \text{Sn}/\text{HCl} \rightarrow \text{C}_6\text{H}_5\text{-NH}_2$ (Aniline) [Complete reduction]
<b>B</b>	$\text{C}_6\text{H}_5\text{-NO}_2 + \text{Zn}/\text{NH}_4\text{Cl} \rightarrow \text{C}_6\text{H}_5\text{-NH-OH}$ (Phenyl hydroxylamine) [Partial reduction]
<b>C</b>	$\text{C}_6\text{H}_5\text{-NO}_2 + \text{Zn}/\text{NaOH} \rightarrow \text{C}_6\text{H}_5\text{-NH-NH-C}_6\text{H}_5$ (Hydrazobenzene) [Alkaline reduction]

**Explanation:** Nitrobenzene can be selectively reduced based on the reagent.  $\text{Sn}/\text{HCl}$  gives complete reduction to aniline.  $\text{Zn}/\text{NH}_4\text{Cl}$  gives partial reduction to phenyl hydroxylamine.  $\text{Zn}/\text{NaOH}$  in alkaline medium gives hydrazobenzene (bimolecular product). These are called controlled reductions.

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{C}_6\text{H}_5\text{-NH}_2$	<b>Aniline (Sn/HCl)</b>
<b>B</b>	$\text{C}_6\text{H}_5\text{-NH-OH}$	<b>Phenyl Hydroxylamine (Zn/NH<sub>4</sub>Cl)</b>

<b>C</b>	$C_6H_5-NH-NH-C_6H_5$	<b>Hydrazobenzene (Zn/NaOH)</b>
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### Q6. Benzene to o-Cyano toluene via 5-step synthesis

**Named Reaction: Friedel-Crafts → Nitration → Reduction → Diazotisation → Sandmeyer**

#### Reactions:

Step	Reaction
<b>A</b>	$C_6H_6 + CH_3Cl/AlCl_3 \rightarrow C_6H_5CH_3$ (Toluene) [Friedel-Crafts Alkylation]
<b>B</b>	$C_6H_5CH_3 + \text{conc.}HNO_3/H_2SO_4 \rightarrow o\text{-}CH_3\text{-}C_6H_4\text{-}NO_2$ (o-Nitrotoluene)
<b>C</b>	$o\text{-}CH_3\text{-}C_6H_4\text{-}NO_2 + Sn/HCl \rightarrow o\text{-}CH_3\text{-}C_6H_4\text{-}NH_2$ (o-Aminotoluene)
<b>D</b>	$o\text{-}CH_3\text{-}C_6H_4\text{-}NH_2 + NaNO_2/HCl \rightarrow o\text{-}CH_3\text{-}C_6H_4\text{-}N_2Cl$ (o-Methyl benzene diazonium chloride)
<b>E</b>	$o\text{-}CH_3\text{-}C_6H_4\text{-}N_2Cl + CuCN \rightarrow o\text{-}CH_3\text{-}C_6H_4\text{-}CN$ (o-Cyano toluene) [Sandmeyer]

*Explanation: Benzene is alkylated to toluene. Electrophilic nitration gives o-nitrotoluene (ortho director). Sn/HCl reduces to o-aminotoluene. Diazotisation converts -NH<sub>2</sub> to diazonium salt. Sandmeyer reaction with CuCN replaces -N<sub>2</sub><sup>+</sup> with -CN giving o-cyano toluene.*

#### Answer Summary:

Label	Formula	Name
<b>A</b>	$C_6H_5CH_3$	<b>Toluene</b>
<b>B</b>	$o\text{-}CH_3\text{-}C_6H_4\text{-}NO_2$	<b>o-Nitrotoluene</b>
<b>C</b>	$o\text{-}CH_3\text{-}C_6H_4\text{-}NH_2$	<b>o-Aminotoluene</b>
<b>D</b>	$o\text{-}CH_3\text{-}C_6H_4\text{-}N_2Cl$	<b>o-Methylbenzene Diazonium Chloride</b>
<b>E</b>	$o\text{-}CH_3\text{-}C_6H_4\text{-}CN$	<b>o-Cyano Toluene</b>

### Q7. $C_6H_5\text{-CHO}$ with Zn-Hg/HCl, OH<sup>-</sup>/KMnO<sub>4</sub> and NaOH/CaO

**Named Reaction: Clemmensen Reduction + Oxidation + Decarboxylation**

**Reactions:**

Step	Reaction
<b>A</b>	$C_6H_5-CHO + Zn-Hg/HCl \rightarrow C_6H_5-CH_3$ (Toluene) [Clemmensen Reduction]
<b>B</b>	$C_6H_5-CH_3 + OH^-/KMnO_4 \rightarrow C_6H_5-COOH$ (Benzoic acid) [Oxidation]
<b>C</b>	$C_6H_5-COOH + NaOH/CaO/heat \rightarrow C_6H_6$ (Benzene) [Decarboxylation / Sodalime]

*Explanation: Benzaldehyde is reduced by Clemmensen (Zn-Hg/HCl) to toluene (converts C=O to CH<sub>2</sub>). KMnO<sub>4</sub>/OH<sup>-</sup> oxidises the methyl group to carboxyl giving benzoic acid. Heating with NaOH/CaO (sodalime) removes the COOH group to give benzene.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	C <sub>6</sub> H <sub>5</sub> CH <sub>3</sub>	<b>Toluene</b>
<b>B</b>	C <sub>6</sub> H <sub>5</sub> COOH	<b>Benzoic Acid</b>
<b>C</b>	C <sub>6</sub> H <sub>6</sub>	<b>Benzene</b>

**Q8.** Ethanoic acid → A (SOCl<sub>2</sub>) → B (Pd/BaSO<sub>4</sub>) → C (NaOH)

**Named Reaction: Rosenmund Reduction + Aldol Condensation****Reactions:**

Step	Reaction
<b>A</b>	$CH_3COOH + SOCl_2 \rightarrow CH_3COCl$ (Acetyl chloride)
<b>B</b>	$CH_3COCl + Pd/BaSO_4 \rightarrow CH_3CHO$ (Acetaldehyde) [Rosenmund Reduction]
<b>C</b>	$2CH_3CHO + dil. NaOH \rightarrow CH_3CH(OH)CH_2CHO$ (Aldol = 3-Hydroxybutanal) [Aldol Condensation]

*Explanation: Acetic acid reacts with SOCl<sub>2</sub> to form acetyl chloride (acid chloride). Pd/BaSO<sub>4</sub> (Rosenmund catalyst) selectively reduces acid chloride to aldehyde (acetaldehyde). Two molecules of acetaldehyde undergo aldol condensation with dilute NaOH to form aldol.*

**Answer Summary:**

Label	Formula	Name
A	CH <sub>3</sub> COCl	Acetyl Chloride
B	CH <sub>3</sub> CHO	Acetaldehyde
C	CH <sub>3</sub> CH(OH)CH <sub>2</sub> CHO	Aldol (3-Hydroxybutanal)

### Q9. C<sub>6</sub>H<sub>5</sub>-NO<sub>2</sub> → A (Sn/HCl) → B (Br<sub>2</sub>/H<sub>2</sub>O)

#### Named Reaction: Reduction + Electrophilic Bromination of Aniline

#### Reactions:

Step	Reaction
A	C <sub>6</sub> H <sub>5</sub> -NO <sub>2</sub> + Sn/HCl → C <sub>6</sub> H <sub>5</sub> -NH <sub>2</sub> (Aniline)
B	C <sub>6</sub> H <sub>5</sub> -NH <sub>2</sub> + Br <sub>2</sub> /H <sub>2</sub> O → 2,4,6-Tribromoaniline (white precipitate)

*Explanation: Nitrobenzene reduces to aniline. The -NH<sub>2</sub> group is a powerful ortho/para director. With Br<sub>2</sub>/H<sub>2</sub>O (no catalyst needed), all three available positions (2,4,6) are brominated immediately to give 2,4,6-tribromoaniline (white precipitate — used as a test for aniline).*

#### Answer Summary:

Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>	Aniline
B	2,4,6-Br <sub>3</sub> -C <sub>6</sub> H <sub>2</sub> -NH <sub>2</sub>	2,4,6-Tribromoaniline

### Q10. C<sub>6</sub>H<sub>5</sub>N<sub>2</sub>Cl → A (CuCN) → B (H<sub>2</sub>O/H<sup>+</sup>)

#### Named Reaction: Sandmeyer Reaction + Hydrolysis of Nitrile

#### Reactions:

Step	Reaction
A	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl + CuCN → C <sub>6</sub> H <sub>5</sub> -CN (Phenyl cyanide / Benzonitrile) [Sandmeyer]

<b>B</b>	$C_6H_5-CN + H_2O/H^+ \rightarrow C_6H_5-COOH$ (Benzoic acid) [Hydrolysis]
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*Explanation: Sandmeyer reaction replaces  $-N_2^+$  with  $-CN$  using cuprous cyanide ( $CuCN$ ), giving benzonitrile. Acid hydrolysis of the nitrile converts  $-CN$  to  $-COOH$ , giving benzoic acid. This is a useful 2-step route from amine to carboxylic acid.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$C_6H_5-CN$	<b>Phenyl Cyanide (Benzonitrile)</b>
<b>B</b>	$C_6H_5-COOH$	<b>Benzoic Acid</b>

**Q11.**  $C_6H_5-COOH \rightarrow A (PCl_5) \rightarrow B (C_6H_6/AlCl_3)$ **Named Reaction: Acid Chloride Formation + Friedel-Crafts Acylation****Reactions:**

Step	Reaction
<b>A</b>	$C_6H_5-COOH + PCl_5 \rightarrow C_6H_5-COCl$ (Benzoyl chloride)
<b>B</b>	$C_6H_5-COCl + C_6H_6/\text{anhydrous } AlCl_3 \rightarrow C_6H_5-CO-C_6H_5$ (Benzophenone) [Friedel-Crafts Acylation]

*Explanation: Benzoic acid reacts with  $PCl_5$  (or  $SOCl_2$ ) to convert  $-COOH$  to  $-COCl$  (acid chloride). Benzoyl chloride undergoes Friedel-Crafts acylation with benzene and anhydrous  $AlCl_3$  to give benzophenone (diphenyl ketone).*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$C_6H_5COCl$	<b>Benzoyl Chloride</b>
<b>B</b>	$C_6H_5CO-C_6H_5$	<b>Benzophenone (Diphenyl Ketone)</b>

**Q12.**  $C_6H_5-CONH_2 \rightarrow A (Br_2/NaOH) \rightarrow B (NaNO_2/HCl)$ **Named Reaction: Hoffmann Bromamide Reaction + Diazotisation**

## Reactions:

Step	Reaction
A	$C_6H_5-CONH_2 + Br_2/NaOH \rightarrow C_6H_5-NH_2$ (Aniline) [Hoffmann – 1 C less]
B	$C_6H_5-NH_2 + NaNO_2/HCl/273-278K \rightarrow C_6H_5-N \equiv N-Cl$ (Benzene Diazonium Chloride)

**Explanation:** Benzamide (C7) undergoes Hoffmann bromamide rearrangement with  $Br_2/NaOH$  — the  $-CO-$  is lost, giving aniline (C6). Note: this is one of the few reactions that reduce carbon count. Aniline then diazotises to form the diazonium salt at 273-278K.

## Answer Summary:

Label	Formula	Name
A	$C_6H_5NH_2$	Aniline
B	$C_6H_5N_2Cl$	Benzene Diazonium Chloride

**Q13.**  $C_6H_5-NO_2 \rightarrow A (Fe/HCl) \rightarrow B (HNO_2) \rightarrow C (C_6H_5OH)$

## Named Reaction: Reduction + Diazotisation + Azo Coupling

## Reactions:

Step	Reaction
A	$C_6H_5-NO_2 + Fe/HCl \rightarrow C_6H_5-NH_2$ (Aniline)
B	$C_6H_5-NH_2 + HNO_2 \rightarrow C_6H_5-N=N-Cl$ (Benzene Diazonium Chloride)
C	$C_6H_5-N_2Cl + C_6H_5OH \rightarrow C_6H_5-N=N-C_6H_4-OH$ (p-Hydroxy azobenzene)

**Explanation:**  $Fe/HCl$  reduces nitrobenzene to aniline. Aniline diazotised with  $HNO_2$  (=  $NaNO_2 + HCl$ ). The diazonium salt couples with phenol (at para position) at pH 9-10 to give p-hydroxy azobenzene — an orange-yellow azo dye.

## Answer Summary:

Label	Formula	Name
A	$C_6H_5NH_2$	Aniline
B	$C_6H_5N_2Cl$	Benzene Diazonium Chloride

<b>C</b>	$C_6H_5-N=N-C_6H_4-OH$	<b>p-Hydroxy Azobenzene (Azo Dye)</b>
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**Q14.**  $C_6H_5-OH \rightarrow A (Zn) \rightarrow B (CH_3Cl/AlCl_3) \rightarrow C (O/KMnO_4)$

**Named Reaction: Reduction + Friedel-Crafts + Oxidation**

**Reactions:**

Step	Reaction
<b>A</b>	$C_6H_5-OH + Zn \rightarrow C_6H_6$ (Benzene)
<b>B</b>	$C_6H_6 + CH_3Cl/AlCl_3 \rightarrow C_6H_5-CH_3$ (Toluene) [Friedel-Crafts Alkylation]
<b>C</b>	$C_6H_5-CH_3 + (O)/KMnO_4 \rightarrow C_6H_5-COOH$ (Benzoic acid) [Side-chain oxidation]

*Explanation: Phenol reduces to benzene with Zn. Friedel-Crafts alkylation adds a methyl group to give toluene. Hot acidic  $KMnO_4$  oxidises the entire methyl group to  $-COOH$  regardless of chain length, giving benzoic acid.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$C_6H_6$	<b>Benzene</b>
<b>B</b>	$C_6H_5CH_3$	<b>Toluene</b>
<b>C</b>	$C_6H_5COOH$	<b>Benzoic Acid</b>

**Q15.** Cyclohexanone + methylenecyclohexylamine, trace  $H^+$

**Named Reaction: Schiff's Base Formation**

**Reactions:**

Step	Reaction
<b>Product</b>	Cyclohexanone + $H_2N-CH_2$ -[ring] + trace $H^+$ $\rightarrow$ Schiff's base (ring=N- $CH_2$ -[ring]) + $H_2O$

*Explanation: A Schiff's base (imine) is formed by condensation of a carbonyl compound (aldehyde or ketone) with a primary amine. Water is eliminated. Trace acid catalyses the reaction. General form:  $R-CHO + H_2N-R' \rightarrow R-CH=N-R' + H_2O$ .*

**Answer Summary:**

Label	Formula	Name
Product	$R-C=N-CH_2-R'$	Schiff's Base (Aldimine / Ketimine)

**Q16.**  $C_6H_5-NH_2 + C_6H_5-CHO \rightarrow ?$ 

**Named Reaction: Schiff's Base Formation from Aniline + Benzaldehyde**

**Reactions:**

Step	Reaction
Product	$C_6H_5-CHO + H_2N-C_6H_5 + H^+ \rightarrow C_6H_5-CH=N-C_6H_5 + H_2O$ (Benzal aniline)

*Explanation: Benzaldehyde (aromatic aldehyde) condenses with aniline (primary aromatic amine) to form benzal aniline — an aromatic Schiff's base. Acid catalysis helps. Water is lost in this condensation. Schiff's bases are used as ligands and dyes.*

**Answer Summary:**

Label	Formula	Name
Product	$C_6H_5-CH=N-C_6H_5$	Benzal Aniline (Schiff's Base)

**Q17.**  $C_6H_5-NH_2 + C_6H_5-COCl \rightarrow ?$  (Pyridine)

**Named Reaction: Schotten-Baumann Reaction (N-Acylation)**

**Reactions:**

Step	Reaction
Product	$C_6H_5-COCl + H_2N-C_6H_5 + \text{Pyridine} \rightarrow C_6H_5-CO-NH-C_6H_5 + HCl$

*Explanation: Aniline (nucleophile) reacts with benzoyl chloride (electrophile) in the presence of pyridine (base — neutralises HCl) to form N-phenyl benzamide. This is Schotten-Baumann condition. Pyridine prevents the HCl produced from reacting with amine.*

**Answer Summary:**

Label	Formula	Name
Product	C <sub>6</sub> H <sub>5</sub> -CO-NH-C <sub>6</sub> H <sub>5</sub>	N-Phenyl Benzamide

**Q18.**  $\text{CH}_3\text{CH}_2\text{NC} \rightarrow \text{A} (\text{HgO}) \rightarrow \text{B} (\text{H}_2\text{O}) \rightarrow \text{C} (\text{NaNO}_2\text{-HCl/H}_2\text{O})$

**Named Reaction: Isocyanide Hydrolysis + Diazotisation → Alcohol**

**Reactions:**

Step	Reaction
A	$\text{CH}_3\text{CH}_2\text{NC} + \text{HgO} \rightarrow \text{CH}_3\text{CH}_2\text{-N=C=O}$ (Ethyl isocyanate)
B	$\text{CH}_3\text{CH}_2\text{-N=C=O} + \text{H}_2\text{O} \rightarrow \text{CH}_3\text{CH}_2\text{-NH}_2$ (Ethylamine)
C	$\text{CH}_3\text{CH}_2\text{-NH}_2 + \text{NaNO}_2/\text{HCl}/\text{H}_2\text{O} \rightarrow \text{CH}_3\text{CH}_2\text{-OH}$ (Ethyl alcohol) + N <sub>2</sub>

*Explanation: Ethyl isocyanide oxidised by HgO gives ethyl isocyanate. Hydrolysis of isocyanate gives ethylamine. Diazotisation of primary aliphatic amine with NaNO<sub>2</sub>/HCl gives unstable diazonium salt which immediately decomposes to alcohol + N<sub>2</sub>.*

**Answer Summary:**

Label	Formula	Name
A	CH <sub>3</sub> CH <sub>2</sub> NCO	Ethyl Isocyanate
B	CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub>	Ethylamine
C	CH <sub>3</sub> CH <sub>2</sub> OH	Ethyl Alcohol

**Q19.** *Glutaric acid* → A (SOCl<sub>2</sub>) → B (NH<sub>3</sub>) → C (LiAlH<sub>4</sub>)

**Named Reaction: Acid Chloride → Amide → Amine (LiAlH<sub>4</sub> Reduction)**

**Reactions:**

Step	Reaction
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<b>A</b>	Glutaric acid + SOCl <sub>2</sub> → Glutaryl dichloride (Glycocyl chloride)
<b>B</b>	Glutaryl dichloride + NH <sub>3</sub> → Glutaramide (Glutamide)
<b>C</b>	Glutamide + LiAlH <sub>4</sub> → Pentan-1,5-diamine [Reduction of amide to amine]

**Explanation:** Dicarboxylic acid reacts with SOCl<sub>2</sub> to give diacid chloride. Reaction with ammonia gives diamide. LiAlH<sub>4</sub> is a powerful reducing agent that converts amide (-CONH<sub>2</sub>) to amine (-CH<sub>2</sub>NH<sub>2</sub>). Product is a diamine (1,5-pentanediamine).

**Answer Summary:**

Label	Formula	Name
<b>A</b>	C <sub>1</sub> CO (CH <sub>2</sub> ) <sub>3</sub> COCl	<b>Glutaryl Dichloride (Glycocyl Chloride)</b>
<b>B</b>	H <sub>2</sub> NOC (CH <sub>2</sub> ) <sub>3</sub> CONH <sub>2</sub>	<b>Glutaramide (Glutamide)</b>
<b>C</b>	H <sub>2</sub> N (CH <sub>2</sub> ) <sub>5</sub> NH <sub>2</sub>	<b>Pentan-1,5-diamine</b>

**Q20. C<sub>6</sub>H<sub>5</sub>-N<sub>2</sub>Cl coupling with three different compounds****Named Reaction: Azo Coupling Reactions (pH dependent)****Reactions:**

Step	Reaction
<b>A</b>	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl + p-(CH <sub>3</sub> ) <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> → p-(CH <sub>3</sub> ) <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -N=N-C <sub>6</sub> H <sub>5</sub> [p-N,N-dimethylaminoazobenzene]
<b>B</b>	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl + p-CH <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> -OH → p-CH <sub>3</sub> -C <sub>6</sub> H <sub>4</sub> (N=N-C <sub>6</sub> H <sub>5</sub> )-OH [2-Phenylazo-4-methylphenol]
<b>C</b>	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl + p-NH <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> → p-NH <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -N=N-C <sub>6</sub> H <sub>5</sub> [p-Aminoazobenzene]

**Explanation:** Diazonium salt couples with aromatic amines (pH 4-5) and phenols (pH 9-10). N,N-dimethylaniline couples at para to give orange azo dye (A). p-cresol gives 2-phenylazo-4-methylphenol (B). p-aminoaniline gives p-aminoazobenzene (C). Azo dyes are coloured due to -N=N- chromophore.

**Answer Summary:**

Label	Formula	Name
<b>A</b>	p-(CH <sub>3</sub> ) <sub>2</sub> N-C <sub>6</sub> H <sub>4</sub> -N=N-C <sub>6</sub> H <sub>5</sub>	<b>p-N,N-Dimethylaminoazobenzene</b>

<b>B</b>	$\text{CH}_3\text{-C}_6\text{H}_3(\text{OH})\text{-N=N-C}_6\text{H}_5$	<b>2-Phenylazo-4-methylphenol</b>
<b>C</b>	$\text{p-NH}_2\text{-C}_6\text{H}_4\text{-N=N-C}_6\text{H}_5$	<b>p-Aminoazobenzene</b>

**Q21.**  $A \rightarrow \text{CH}_3\text{CH}_2\text{NH}_2$  and  $B \rightarrow \text{CH}_3\text{NHCH}_3$  with  $\text{Na-Hg/C}_2\text{H}_5\text{OH}$

**Named Reaction: Reduction of Cyanide vs Isocyanide**

**Reactions:**

Step	Reaction
<b>A</b>	$\text{CH}_3\text{-CN} + \text{Na-Hg/C}_2\text{H}_5\text{OH} \rightarrow \text{CH}_3\text{-CH}_2\text{-NH}_2$ (Ethylamine – 1° amine)
<b>B</b>	$\text{CH}_3\text{-NC} + \text{Na-Hg/C}_2\text{H}_5\text{OH} \rightarrow \text{CH}_3\text{-NH-CH}_3$ (Dimethylamine – 2° amine)

*Explanation: Key distinction: Cyanide (R-CN) on reduction gives PRIMARY amine (chain extended). Isocyanide (R-NC) on reduction gives SECONDARY amine (chain not extended, N inserted between carbons). This is how you distinguish cyanide from isocyanide.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{-CN}$	<b>Methyl Cyanide (Acetonitrile)</b>
<b>B</b>	$\text{CH}_3\text{-NC}$	<b>Methyl Isocyanide</b>

**Q22.**  $\text{CH}_3\text{CONH}_2 \rightarrow A (\text{Br}_2/\text{NaOH}) \rightarrow B (\text{NaNO}_2/\text{HCl}) \rightarrow C (\text{O then NH}_3)$

**Named Reaction: Hoffmann → Diazotisation → Urotropine Formation**

**Reactions:**

Step	Reaction
<b>A</b>	$\text{CH}_3\text{CONH}_2 + \text{Br}_2/\text{NaOH} \rightarrow \text{CH}_3\text{NH}_2$ (Methylamine) [Hoffmann Bromamide]
<b>B</b>	$\text{CH}_3\text{NH}_2 + \text{NaNO}_2/\text{HCl} \rightarrow \text{CH}_3\text{OH}$ (Methyl alcohol) + $\text{N}_2$
<b>C</b>	$\text{HCHO} + (\text{O}) \rightarrow \text{HCHO}$ ; $6\text{HCHO} + 4\text{NH}_3 \rightarrow (\text{CH}_2)_6\text{N}_4 + 6\text{H}_2\text{O}$ (Urotropine)

*Explanation: Acetamide loses CO via Hoffmann bromamide to give methylamine (C2→C1). Methylamine with NaNO<sub>2</sub>/HCl gives methyl alcohol. Oxidation of methanol gives formaldehyde. 6HCHO + 4NH<sub>3</sub> condense to hexamethylenetetramine (Urotropine — used as antiseptic).*

**Answer Summary:**

Label	Formula	Name
A	CH <sub>3</sub> NH <sub>2</sub>	Methylamine
B	CH <sub>3</sub> OH	Methyl Alcohol
C	(CH <sub>2</sub> ) <sub>6</sub> N <sub>4</sub>	Urotropine (Hexamethylenetetramine)

**Q23.** Anisole + *t*-butyl chloride/AlCl<sub>3</sub> → A; A + Cl<sub>2</sub>/FeCl<sub>3</sub> → B

**Named Reaction: Friedel-Crafts Alkylation + Electrophilic Chlorination**

**Reactions:**

Step	Reaction
A	Anisole + (CH <sub>3</sub> ) <sub>3</sub> C-Cl/AlCl <sub>3</sub> → 4- <i>t</i> -butyl-1-methoxybenzene [Friedel-Crafts]
B	4- <i>t</i> -butyl-1-methoxybenzene + Cl <sub>2</sub> /FeCl <sub>3</sub> → 2-Chloro-4- <i>t</i> -butyl-1-methoxybenzene

*Explanation: Anisole (-OCH<sub>3</sub> group) is an ortho/para director. *t*-Butyl group adds at para position (Friedel-Crafts). Second electrophilic substitution by Cl<sub>2</sub>/FeCl<sub>3</sub> adds Cl at ortho to -OCH<sub>3</sub> (position 2), since para is already occupied by *t*-butyl.*

**Answer Summary:**

Label	Formula	Name
A	(CH <sub>3</sub> ) <sub>3</sub> C-C <sub>6</sub> H <sub>4</sub> -OCH <sub>3</sub>	4- <i>t</i> -Butyl-1-methoxybenzene
B	Cl-(CH <sub>3</sub> ) <sub>3</sub> C-C <sub>6</sub> H <sub>3</sub> -OCH <sub>3</sub>	2-Chloro-4- <i>t</i> -butyl-1-methoxybenzene

**Q24.** CH<sub>3</sub>-CHO + HCN → A; A + H<sub>2</sub>O/H<sup>+</sup> → B; A + LiAlH<sub>4</sub> → C

**Named Reaction: Cyanohydrin Formation + Hydrolysis + LiAlH<sub>4</sub> Reduction**

## Reactions:

Step	Reaction
A	$\text{CH}_3\text{-CHO} + \text{HCN} \rightarrow \text{CH}_3\text{-CH(OH)-CN}$ (Acetaldehyde cyanohydrin)
B	$\text{CH}_3\text{-CH(OH)-CN} + \text{H}_2\text{O/H}^+ \rightarrow \text{CH}_3\text{-CH(OH)-COOH}$ (Lactic acid) [Hydrolysis]
C	$\text{CH}_3\text{-CH(OH)-CN} + \text{LiAlH}_4 \rightarrow \text{CH}_3\text{-CH(OH)-CH}_2\text{-NH}_2$ (1-Amino-2-propanol)

*Explanation: HCN (nucleophile) adds to acetaldehyde's carbonyl carbon to form cyanohydrin. Acid hydrolysis converts -CN to -COOH giving lactic acid (found in sour milk). LiAlH<sub>4</sub> reduces -CN to -CH<sub>2</sub>NH<sub>2</sub> without touching -OH, giving an amino alcohol.*

## Answer Summary:

Label	Formula	Name
A	$\text{CH}_3\text{CH(OH)CN}$	Acetaldehyde Cyanohydrin
B	$\text{CH}_3\text{CH(OH)COOH}$	Lactic Acid (2-Hydroxy Propanoic Acid)
C	$\text{CH}_3\text{CH(OH)CH}_2\text{NH}_2$	1-Amino-2-propanol

**Q25.**  $\text{C}_6\text{H}_5\text{-OH}$  with i)NaOH ii)CO<sub>2</sub> iii)H<sub>2</sub>O/H<sup>+</sup> → X; and CHCl<sub>3</sub>/NaOH → Y

## Named Reaction: Kolbe's Reaction + Reimer-Tiemann Reaction

## Reactions:

Step	Reaction
X	$\text{C}_6\text{H}_5\text{OH} + \text{NaOH} \rightarrow \text{C}_6\text{H}_5\text{ONa}$ ; + CO <sub>2</sub> /pressure → sodium salicylate; + H <sup>+</sup> /H <sub>2</sub> O → Salicylic acid (X)
Y	$\text{C}_6\text{H}_5\text{OH} + \text{CHCl}_3/\text{NaOH} \rightarrow 2\text{-HO-C}_6\text{H}_4\text{-CHO}$ (Salicylaldehyde Y) [Reimer-Tiemann]

*Explanation: Both Kolbe's and Reimer-Tiemann introduce substituents ortho to -OH in phenol. Kolbe's uses CO<sub>2</sub> under pressure → -COOH group. Reimer-Tiemann uses CHCl<sub>3</sub>/NaOH (dichlorocarbene intermediate) → -CHO group.*

## Answer Summary:

Label	Formula	Name
X	$2\text{-HO-C}_6\text{H}_4\text{-COOH}$	Salicylic Acid (Kolbe's Reaction)

Y	2-HO-C <sub>6</sub> H <sub>4</sub> -CHO	Salicylaldehyde (Reimer-Tiemann Reaction)
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**Q26.** Benzene → A (HNO<sub>3</sub>/H<sub>2</sub>SO<sub>4</sub>) → B (Sn/HCl) → C (NaNO<sub>2</sub>/HCl)

**Named Reaction: Nitration + Reduction + Diazotisation**

Reactions:

Step	Reaction
A	C <sub>6</sub> H <sub>6</sub> + conc.HNO <sub>3</sub> /conc.H <sub>2</sub> SO <sub>4</sub> → C <sub>6</sub> H <sub>5</sub> -NO <sub>2</sub> (Nitrobenzene)
B	C <sub>6</sub> H <sub>5</sub> -NO <sub>2</sub> + Sn/HCl → C <sub>6</sub> H <sub>5</sub> -NH <sub>2</sub> (Aniline)
C	C <sub>6</sub> H <sub>5</sub> -NH <sub>2</sub> + NaNO <sub>2</sub> /HCl → C <sub>6</sub> H <sub>5</sub> -N <sub>2</sub> Cl (Benzene Diazonium Chloride)

*Explanation: Benzene undergoes electrophilic aromatic nitration with mixed acid (HNO<sub>3</sub>+H<sub>2</sub>SO<sub>4</sub>) giving nitrobenzene. Reduction by Sn/HCl gives aniline. Diazotisation at 273-278K gives benzene diazonium chloride — a key intermediate in synthesis.*

Answer Summary:

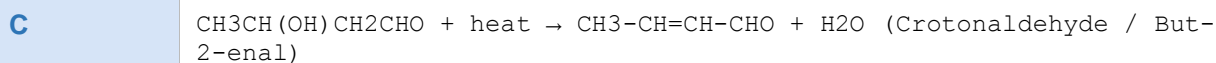
Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> NO <sub>2</sub>	Nitrobenzene
B	C <sub>6</sub> H <sub>5</sub> NH <sub>2</sub>	Aniline
C	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl	Benzene Diazonium Chloride

**Q27.** CH<sub>3</sub>-COCl → A (Pd/BaSO<sub>4</sub>) → B (NaOH) → C (heat)

**Named Reaction: Rosenmund Reduction + Aldol Condensation + Dehydration**

Reactions:

Step	Reaction
A	CH <sub>3</sub> -COCl + Pd/BaSO <sub>4</sub> /H <sub>2</sub> → CH <sub>3</sub> -CHO (Acetaldehyde) [Rosenmund]
B	2CH <sub>3</sub> -CHO + dil.NaOH → CH <sub>3</sub> CH(OH)CH <sub>2</sub> CHO (Aldol / 3-Hydroxybutanal)



*Explanation: Rosenmund reduces acid chloride to aldehyde (Pd/BaSO<sub>4</sub> prevents over-reduction). Aldol condensation gives 3-hydroxybutanal (Aldol). On heating, aldol dehydrates to give crotonaldehyde (conjugated aldehyde — alpha,beta-unsaturated).*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{CHO}$	<b>Acetaldehyde</b>
<b>B</b>	$\text{CH}_3\text{CH}(\text{OH})\text{CH}_2\text{CHO}$	<b>Aldol (3-Hydroxybutanal)</b>
<b>C</b>	$\text{CH}_3\text{CH=CHCHO}$	<b>Crotonaldehyde (But-2-enal)</b>

**Q28. NH<sub>3</sub> with HCHO, CH<sub>3</sub>-CHO, and C<sub>6</sub>H<sub>5</sub>-CHO****Named Reaction: Amine Condensation with Aldehydes****Reactions:**

Step	Reaction
<b>A</b>	$6\text{HCHO} + 4\text{NH}_3 \rightarrow (\text{CH}_2)_6\text{N}_4 + 6\text{H}_2\text{O}$ (Hexamethylenetetramine / Urotropine)
<b>B</b>	$\text{CH}_3\text{-CHO} + \text{NH}_3 \rightarrow \text{CH}_3\text{-CH=NH}$ (Aldimine / Ethylideneamine)
<b>C</b>	$3\text{C}_6\text{H}_5\text{-CHO} + 2\text{NH}_3 \rightarrow \text{C}_6\text{H}_5\text{-CH=N-CH(C}_6\text{H}_5\text{)-N=CH-C}_6\text{H}_5$ (Hydrobenzamide)

*Explanation: Different aldehydes react differently with NH<sub>3</sub>. HCHO (excess) with NH<sub>3</sub> forms urotropine (cage compound). Acetaldehyde with NH<sub>3</sub> forms simple aldimine. Benzaldehyde (3:2 ratio) forms hydrobenzamide (a trimer-like Schiff's base).*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$(\text{CH}_2)_6\text{N}_4$	<b>Hexamethylenetetramine (Urotropine)</b>
<b>B</b>	$\text{CH}_3\text{-CH=NH}$	<b>Aldimine (Ethylideneamine)</b>
<b>C</b>	$(\text{C}_6\text{H}_5\text{CH=N})_2\text{CHCH}_6\text{H}_5$	<b>Hydrobenzamide</b>

**Q29.**  $C_2H_6O$  (A)  $\rightarrow$  B (Cu)  $\rightarrow$  C ( $N_2H_4/C_2H_5ONa$ )**Named Reaction: Oxidative Dehydrogenation + Wolff-Kishner Reduction****Reactions:**

Step	Reaction
A	$CH_3CH_2OH$ (Ethyl Alcohol)
B	$CH_3CH_2OH + Cu/heat \rightarrow CH_3CHO$ (Acetaldehyde) [Dehydrogenation]
C	$CH_3CHO + N_2H_4/C_2H_5ONa \rightarrow CH_3CH_3$ (Ethane) [Wolff-Kishner Reduction]

*Explanation: Ethanol dehydrogenates over copper to give acetaldehyde. Wolff-Kishner reduction (hydrazine + base) converts the aldehyde carbonyl to  $-CH_2-$  giving ethane. This converts oxygenated compound to hydrocarbon.*

**Answer Summary:**

Label	Formula	Name
A	$CH_3CH_2OH$	Ethyl Alcohol
B	$CH_3CHO$	Acetaldehyde
C	$CH_3CH_3$	Ethane

**Q30.**  $CH_3COCH_3$  with Zn/Hg and Mg/Hg**Named Reaction: Clemmensen Reduction vs Pinacol Coupling****Reactions:**

Step	Reaction
A	$CH_3COCH_3 + Zn-Hg/HCl \rightarrow CH_3CH_2CH_3$ (Propane) [Clemmensen - $C=O \rightarrow CH_2$ ]
B	$2 CH_3COCH_3 + Mg/Hg \rightarrow (CH_3)_2C(OH)-C(OH)(CH_3)_2$ [Pinacol - 2,3-Dimethylbutane-2,3-diol]

*Explanation: Clemmensen (Zn-Hg/HCl) removes the carbonyl oxygen giving propane (full reduction). Mg/Hg is used for pinacol coupling — two ketone molecules couple reductively to give a 1,2-diol called pinacol. Two different reducing methods, two different products!*

**Answer Summary:**

Label	Formula	Name
A	CH <sub>3</sub> CH <sub>2</sub> CH <sub>3</sub>	Propane (Clemmensen Reduction)
B	(CH <sub>3</sub> ) <sub>2</sub> C(OH)C(OH)(CH <sub>3</sub> ) <sub>2</sub>	Pinacol (2,3-Dimethylbutane-2,3-diol)

**Q31. C<sub>6</sub>H<sub>5</sub>-NO<sub>2</sub> with HNO<sub>3</sub>/H<sub>2</sub>SO<sub>4</sub> at 373K and 473K****Named Reaction: Temperature-Controlled Polynitration****Reactions:**

Step	Reaction
A	C <sub>6</sub> H <sub>5</sub> NO <sub>2</sub> + HNO <sub>3</sub> /H <sub>2</sub> SO <sub>4</sub> at 373K → 1,3-Dinitrobenzene (m-dinitrobenzene)
B	1,3-C <sub>6</sub> H <sub>4</sub> (NO <sub>2</sub> ) <sub>2</sub> + HNO <sub>3</sub> /H <sub>2</sub> SO <sub>4</sub> at 473K → 1,3,5-Trinitrobenzene (TNB)

*Explanation: Nitrobenzene's -NO<sub>2</sub> group deactivates the ring (meta director). A second nitro group enters at meta position at 373K. At higher temperature (473K) a third nitro group is introduced. 1,3,5-Trinitrobenzene (TNB) is an explosive.*

**Answer Summary:**

Label	Formula	Name
A	1,3-C <sub>6</sub> H <sub>4</sub> (NO <sub>2</sub> ) <sub>2</sub>	m-Dinitrobenzene (1,3-Dinitrobenzene)
B	1,3,5-C <sub>6</sub> H <sub>3</sub> (NO <sub>2</sub> ) <sub>3</sub>	1,3,5-Trinitrobenzene (TNB)

**Q32. Glycol with anhydrous ZnCl<sub>2</sub> and conc.H<sub>2</sub>SO<sub>4</sub>****Named Reaction: Dehydration of Glycol****Reactions:**

Step	Reaction
A	$\text{CH}_2(\text{OH})-\text{CH}_2(\text{OH}) + \text{anhydrous ZnCl}_2 \rightarrow \text{CH}_3-\text{CHO}$ (Acetaldehyde) [Intramolecular dehydration]
B	$2 \text{CH}_2(\text{OH})-\text{CH}_2(\text{OH}) + \text{conc. H}_2\text{SO}_4 \rightarrow 1,4\text{-Dioxane}$ (cyclic ether) [Intermolecular]

**Explanation:** Glycol (ethane-1,2-diol) with  $\text{ZnCl}_2$  undergoes intramolecular dehydration rearranging to acetaldehyde. With conc.  $\text{H}_2\text{SO}_4$ , two molecules undergo intermolecular dehydration to form 1,4-dioxane (cyclic diether — 6-membered ring with 2 oxygens).

**Answer Summary:**

Label	Formula	Name
A	$\text{CH}_3\text{CHO}$	Acetaldehyde ( $\text{ZnCl}_2$ , intramolecular)
B	1,4-dioxane	1,4-Dioxane ( $\text{H}_2\text{SO}_4$ , intermolecular)

**Q33. Glycol with  $\text{HIO}_4$  and  $\text{PI}_3$** **Named Reaction: Glycol Cleavage ( $\text{HIO}_4$ ) and Deoxygenation ( $\text{PI}_3$ )****Reactions:**

Step	Reaction
A	$\text{HOCH}_2-\text{CH}_2\text{OH} + \text{HIO}_4 \rightarrow 2 \text{HCHO}$ (Formaldehyde) [Glycol cleavage]
B	$\text{HOCH}_2-\text{CH}_2\text{OH} + \text{PI}_3 \rightarrow \text{CH}_2=\text{CH}_2$ (Ethylene) [Dehydration / deoxygenation]

**Explanation:**  $\text{HIO}_4$  (periodic acid) cleaves the C-C bond of vicinal diol, giving 2 moles of formaldehyde (one per carbon).  $\text{PI}_3$  converts the two -OH groups to iodide and eliminates to give ethylene. This is a key diagnostic test for 1,2-diol (vicinal diol).

**Answer Summary:**

Label	Formula	Name
A	2 HCHO	Formaldehyde (x2) [ $\text{HIO}_4$ cleavage]
B	$\text{CH}_2=\text{CH}_2$	Ethylene [ $\text{PI}_3$ elimination]

**Q34.**  $C_2H_5-OH$  with  $H_2SO_4$  at 413K and 443K**Named Reaction: Temperature-Dependent Dehydration of Ethanol****Reactions:**

Step	Reaction
A	$C_2H_5OH + H_2SO_4$ at 413K $\rightarrow$ $C_2H_5-O-C_2H_5$ (Diethyl Ether) [Intermolecular]
B	$C_2H_5OH + H_2SO_4$ at 443K $\rightarrow$ $CH_2=CH_2$ (Ethylene) [Intramolecular]

*Explanation: Temperature determines the product: 413K  $\rightarrow$  bimolecular (ether formation). 443K  $\rightarrow$  unimolecular (elimination to alkene). At lower temp., two alcohol molecules react. At higher temp., single molecule loses water. This is a classic temperature-controlled reaction.*

**Answer Summary:**

Label	Formula	Name
A	$C_2H_5OC_2H_5$	Diethyl Ether (Ethoxyethane) [413K]
B	$CH_2=CH_2$	Ethylene (Ethene) [443K]

**Q35.** Diethyl ether with  $PCl_5$  and dil.  $H_2SO_4$ **Named Reaction: Ether Cleavage Reactions****Reactions:**

Step	Reaction
A	$C_2H_5-O-C_2H_5 + PCl_5 \rightarrow 2 C_2H_5Cl$ (Ethyl chloride) + $POCl_3$
B	$C_2H_5-O-C_2H_5 + \text{dil. } H_2SO_4 \rightarrow 2 C_2H_5OH$ (Ethyl alcohol)

*Explanation: Diethyl ether is cleaved by  $PCl_5$  — the oxygen is replaced by two Cl atoms giving ethyl chloride. Dilute  $H_2SO_4$  hydrolyses (cleaves) the ether's C-O bond with water to give ethanol. Ethers are generally resistant to acids but HI and  $PCl_5$  can cleave them.*

**Answer Summary:**

Label	Formula	Name
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<b>A</b>	C <sub>2</sub> H <sub>5</sub> Cl	<b>Ethyl Chloride (PCl<sub>5</sub> cleavage)</b>
<b>B</b>	C <sub>2</sub> H <sub>5</sub> OH	<b>Ethyl Alcohol (H<sub>2</sub>SO<sub>4</sub> hydrolysis)</b>

**Q36.** C<sub>6</sub>H<sub>5</sub>-CHO with alc.KCN and NaOH**Named Reaction: Benzoin Condensation + Cannizzaro Reaction****Reactions:**

Step	Reaction
<b>A</b>	2 C <sub>6</sub> H <sub>5</sub> -CHO + alc.KCN → C <sub>6</sub> H <sub>5</sub> -CH(OH)-CO-C <sub>6</sub> H <sub>5</sub> (Benzoin) [Benzoin Condensation]
<b>B,C</b>	2 C <sub>6</sub> H <sub>5</sub> -CHO + NaOH → C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> OH (Benzyl alcohol B) + C <sub>6</sub> H <sub>5</sub> COONa (Sodium benzoate C) [Cannizzaro]

*Explanation: Two different reactions of benzaldehyde. KCN catalyses benzoin condensation — one molecule acts as nucleophile (after CN addition) and attacks the other; product is benzoin (alpha-hydroxy ketone). NaOH causes Cannizzaro disproportionation (no alpha-H).*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	C <sub>6</sub> H <sub>5</sub> CH(OH)COC <sub>6</sub> H <sub>5</sub>	<b>Benzoin</b>
<b>B</b>	C <sub>6</sub> H <sub>5</sub> CH <sub>2</sub> OH	<b>Benzyl Alcohol (Cannizzaro)</b>
<b>C</b>	C <sub>6</sub> H <sub>5</sub> COONa	<b>Sodium Benzoate (Cannizzaro)</b>

**Q37.** CH<sub>3</sub>-COOH with HI/red P and Cl<sub>2</sub>/red P**Named Reaction: Hell-Volhard-Zelinsky (HVZ) Reaction + HI Reduction****Reactions:**

Step	Reaction
<b>A</b>	CH <sub>3</sub> -COOH + HI/red P → CH <sub>3</sub> -CH <sub>3</sub> (Ethane) [Reduction of acid]

<b>B</b>	$\text{CH}_3\text{-COOH} + \text{Cl}_2/\text{red P} \rightarrow \text{Cl-CH}_2\text{-COOH}$ (Monochloroacetic acid) [HVZ Reaction]
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*Explanation: HI reduces acetic acid all the way to ethane ( $\text{COOH} \rightarrow \text{CH}_3$ ). Red P +  $\text{Cl}_2$  halogenates at alpha-carbon (Hell-Volhard-Zelinsky reaction) — red P converts  $\text{Cl}_2$  to  $\text{PCl}_3$  which activates the acid. Alpha-H is replaced by Cl to give chloroacetic acid.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{CH}_3$	<b>Ethane (HI reduction)</b>
<b>B</b>	$\text{ClCH}_2\text{COOH}$	<b>Monochloroacetic Acid (HVZ Reaction)</b>

**Q38.** A and B are isomers: A with  $\text{H}_2\text{O}/\text{H}^+$  gives  $\text{CH}_3\text{COOH}$ ; B gives  $\text{CH}_3\text{CH}_2\text{OH}$

**Named Reaction: Hydrolysis of Nitroalkane vs Alkyl Nitrite****Reactions:**

Step	Reaction
<b>A</b>	$\text{CH}_3\text{CH}_2\text{NO}_2 + \text{H}_2\text{O}/\text{H}^+ \rightarrow \text{CH}_3\text{COOH}$ (Acetic acid) [Nitroethane — C-N bond]
<b>B</b>	$\text{CH}_3\text{CH}_2\text{-O-N=O} + \text{H}_2\text{O}/\text{H}^+ \rightarrow \text{CH}_3\text{CH}_2\text{OH}$ (Ethanol) [Ethyl nitrite — O-N bond]

*Explanation: Nitroethane ( $\text{R-NO}_2$  — C bonded to N) on hydrolysis gives acetic acid. Ethyl nitrite ( $\text{R-O-N=O}$  — O bonded to N) on hydrolysis gives ethanol. Both have formula  $\text{C}_2\text{H}_5\text{NO}_2$  — same molecular formula, structural isomers. Key: C-N vs O-N bond determines product.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{CH}_2\text{NO}_2$	<b>Nitroethane</b>
<b>B</b>	$\text{CH}_3\text{CH}_2\text{ONO}$	<b>Ethyl Nitrite</b>

**Q39.** A and B are isomers: A +  $\text{Sn}/\text{HCl} \rightarrow \text{CH}_3\text{CH}_2\text{NH}_2$ ; B  $\rightarrow \text{CH}_3\text{CH}_2\text{OH}$

**Named Reaction: Reduction of Nitroalkane vs Alkyl Nitrite****Reactions:**

Step	Reaction
<b>A</b>	$\text{CH}_3\text{CH}_2\text{NO}_2 + \text{Sn}/\text{HCl} \rightarrow \text{CH}_3\text{CH}_2\text{NH}_2$ (Ethylamine) [Nitroethane – $\text{NO}_2$ reduces to $\text{NH}_2$ ]
<b>B</b>	$\text{CH}_3\text{CH}_2\text{-O-N=O} + \text{Sn}/\text{HCl} \rightarrow \text{CH}_3\text{CH}_2\text{OH}$ (Ethanol) [Ethyl nitrite – O-N bond cleaves]

*Explanation: Same structural isomers as Q38 but with Sn/HCl. Nitroethane's -NO<sub>2</sub> reduces to -NH<sub>2</sub> giving ethylamine. Ethyl nitrite's O-N bond cleaves to give ethanol. The difference in bonding (C-N vs O-N) leads to different products under the same conditions.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{CH}_2\text{NO}_2$	<b>Nitroethane → Ethylamine</b>
<b>B</b>	$\text{CH}_3\text{CH}_2\text{ONO}$	<b>Ethyl Nitrite → Ethanol</b>

**Q40.**  $\text{CH}_3\text{-Br} + \text{NaN}_3 \rightarrow \text{A}$ ;  $\text{A} + \text{LiAlH}_4 \rightarrow \text{B} + \text{N}_2$ **Named Reaction: Azide Formation + Reduction to Amine****Reactions:**

Step	Reaction
<b>A</b>	$\text{CH}_3\text{-Br} + \text{NaN}_3 \rightarrow \text{CH}_3\text{-N}_3$ (Methyl azide) [ $\text{S}_{\text{N}}2$ substitution]
<b>B</b>	$\text{CH}_3\text{-N}_3 + \text{LiAlH}_4 \rightarrow \text{CH}_3\text{-NH}_2 + \text{N}_2$ (Methylamine) [Reduction]

*Explanation: Bromomethane undergoes  $\text{S}_{\text{N}}2$  with  $\text{NaN}_3$  (sodium azide) to give methyl azide (organic azide  $\text{R-N}_3$ ).  $\text{LiAlH}_4$  reduces the azide — the  $\text{N}=\text{N}=\text{N}$  group loses  $\text{N}_2$  and the remaining N bonds to H, giving methylamine (primary amine).*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{N}_3$	<b>Methyl Azide</b>
<b>B</b>	$\text{CH}_3\text{NH}_2$	<b>Methylamine</b>

**Q41.**  $\text{CH}_3\text{CH}_2\text{Br}$  with  $\text{KCN}$  and  $\text{AgCN}/\text{C}_2\text{H}_5\text{OH}$ **Named Reaction: KCN vs AgCN — Cyanide vs Isocyanide formation****Reactions:**

Step	Reaction
<b>A</b>	$\text{CH}_3\text{CH}_2\text{Br} + \text{KCN} \rightarrow \text{CH}_3\text{CH}_2\text{CN}$ (Ethyl cyanide) [C-attack, $\text{S}_{\text{N}}2$ ]
<b>B</b>	$\text{CH}_3\text{CH}_2\text{Br} + \text{AgCN}/\text{C}_2\text{H}_5\text{OH} \rightarrow \text{CH}_3\text{CH}_2\text{NC}$ (Ethyl isocyanide) [N-attack]

*Explanation: KCN is ionic —  $\text{CN}^-$  is a good nucleophile, attacks via carbon to give cyanide (R-CN). AgCN is covalent — nitrogen attacks alkyl halide to give isocyanide (R-NC). Same reagent (CN), different bonding: K gives C-alkylation; Ag gives N-alkylation.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	$\text{CH}_3\text{CH}_2\text{CN}$	<b>Ethyl Cyanide (Propanenitrile) [KCN]</b>
<b>B</b>	$\text{CH}_3\text{CH}_2\text{NC}$	<b>Ethyl Isocyanide [AgCN]</b>

**Q42.**  $\text{CH}_3\text{CH}_2\text{NH}_2$  with  $\text{NaNO}_2/\text{HCl}$  and  $\text{CHCl}_3/\text{KOH}$ **Named Reaction: Diazotisation of Aliphatic Amine + Carbylamine Test****Reactions:**

Step	Reaction
<b>A</b>	$\text{CH}_3\text{CH}_2\text{NH}_2 + \text{NaNO}_2/\text{HCl} \rightarrow \text{CH}_3\text{CH}_2\text{OH}$ (Ethanol) + $\text{N}_2$ [Aliphatic diazonium — unstable]
<b>B</b>	$\text{CH}_3\text{CH}_2\text{NH}_2 + \text{CHCl}_3/\text{KOH} \rightarrow \text{CH}_3\text{CH}_2\text{NC}$ (Ethyl isocyanide — foul smell) [Carbylamine test]

*Explanation: Aliphatic primary amines with  $\text{NaNO}_2/\text{HCl}$  form unstable diazonium salts that immediately decompose to alcohol +  $\text{N}_2$  (unlike aromatic amines).  $\text{CHCl}_3/\text{KOH}$  is the carbylamine test — characteristic foul smell of isocyanide confirms primary amine.*

**Answer Summary:**

Label	Formula	Name
A	CH <sub>3</sub> CH <sub>2</sub> OH	Ethanol (unstable aliphatic diazonium decomposition)
B	CH <sub>3</sub> CH <sub>2</sub> NC	Ethyl Isocyanide (carbylamine test — foul smell)

### Q43. C<sub>6</sub>H<sub>5</sub>N<sub>2</sub>Cl with C<sub>6</sub>H<sub>5</sub>OH and C<sub>6</sub>H<sub>6</sub>/NaOH

#### Named Reaction: Azo Coupling + Gomberg-Bachmann Reaction

#### Reactions:

Step	Reaction
A	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl + C <sub>6</sub> H <sub>5</sub> -OH → C <sub>6</sub> H <sub>5</sub> -N=N-C <sub>6</sub> H <sub>4</sub> -OH (p-Hydroxy azobenzene) [Azo coupling]
B	C <sub>6</sub> H <sub>5</sub> N <sub>2</sub> Cl + C <sub>6</sub> H <sub>6</sub> /KOH → C <sub>6</sub> H <sub>5</sub> -C <sub>6</sub> H <sub>5</sub> (Biphenyl) + HCl + N <sub>2</sub> [Gomberg-Bachmann]

*Explanation: Diazonium salt can undergo two reactions. With phenol at pH 9-10, electrophilic coupling gives azo dye. With benzene/KOH (Gomberg-Bachmann reaction), the diazonium forms phenyl radical which attacks benzene ring to give biphenyl.*

#### Answer Summary:

Label	Formula	Name
A	C <sub>6</sub> H <sub>5</sub> -N=N-C <sub>6</sub> H <sub>4</sub> -OH	p-Hydroxy Azobenzene (Azo Coupling)
B	C <sub>6</sub> H <sub>5</sub> -C <sub>6</sub> H <sub>5</sub>	Biphenyl (Gomberg-Bachmann Reaction)

### Q44. C<sub>6</sub>H<sub>5</sub>-COOH with HNO<sub>3</sub>/H<sub>2</sub>SO<sub>4</sub> and Fuming H<sub>2</sub>SO<sub>4</sub>

#### Named Reaction: Electrophilic Nitration + Sulphonation of Benzoic Acid

#### Reactions:

Step	Reaction
A	C <sub>6</sub> H <sub>5</sub> -COOH + conc.HNO <sub>3</sub> /conc.H <sub>2</sub> SO <sub>4</sub> → m-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -COOH (m-Nitrobenzoic acid)

**B**  $C_6H_5-COOH + \text{fuming } H_2SO_4 \rightarrow m-SO_3H-C_6H_4-COOH$  (m-Sulphobenzoic acid)

*Explanation: Benzoic acid's -COOH group is an electron-withdrawing meta director. Both nitration and sulphonation give meta products. Fuming H<sub>2</sub>SO<sub>4</sub> (oleum) introduces -SO<sub>3</sub>H. Both reactions follow EAS (electrophilic aromatic substitution) with meta selectivity.*

### Answer Summary:

Label	Formula	Name
<b>A</b>	m-NO <sub>2</sub> -C <sub>6</sub> H <sub>4</sub> -COOH	<b>m-Nitrobenzoic Acid</b>
<b>B</b>	m-SO <sub>3</sub> H-C <sub>6</sub> H <sub>4</sub> -COOH	<b>m-Sulphobenzoic Acid</b>

**Q45.** Acetyl chloride + CH<sub>3</sub>MgBr → X; X + acidic K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> → A

**Named Reaction: Grignard Addition to Acid Chloride + Oxidation**

### Reactions:

Step	Reaction
<b>X</b>	$CH_3COCl + CH_3MgBr \rightarrow CH_3COCH_3$ (Acetone) + MgClBr [Grignard adds to acyl chloride]
<b>A</b>	$CH_3COCH_3 + \text{acidic } K_2Cr_2O_7 \rightarrow CH_3COOH$ (Acetic acid) [Oxidation of ketone]

*Explanation: Grignard reagent adds to acetyl chloride — the CH<sub>3</sub><sup>-</sup> (carbanion) attacks the carbonyl carbon, and after elimination of MgClBr, acetone forms. Acidic K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> oxidises acetone to acetic acid (oxidative cleavage of ketone at lower carbon).*

### Answer Summary:

Label	Formula	Name
<b>X</b>	CH <sub>3</sub> COCH <sub>3</sub>	<b>Acetone (Propanone)</b>
<b>A</b>	CH <sub>3</sub> COOH	<b>Acetic Acid (Ethanoic Acid)</b>

**Q46.** 2-Methylbut-2-ene + i)O<sub>3</sub> ii)Zn/H<sub>2</sub>O; and CH<sub>3</sub>COCH<sub>3</sub> + Mg/Hg

**Named Reaction: Ozonolysis + Pinacol Coupling**

## Reactions:

Step	Reaction
x	$\text{CH}_3\text{-CH}=\text{C}(\text{CH}_3)\text{-CH}_3 + \text{O}_3 \rightarrow [\text{ozonide}] + \text{Zn}/\text{H}_2\text{O} \rightarrow \text{CH}_3\text{CHO} + \text{CH}_3\text{COCH}_3$ (Ethanal + Propanone)
y	$2 \text{CH}_3\text{COCH}_3 + \text{Mg}/\text{Hg} \rightarrow (\text{CH}_3)_2\text{C}(\text{OH})\text{-C}(\text{OH})(\text{CH}_3)_2$ (Pinacol)

**Explanation:** Ozonolysis cleaves the C=C bond. Each carbon of the double bond gets an oxygen — left carbon (CH<sub>3</sub>-CH=) gives ethanal, right carbon (=C(CH<sub>3</sub>)<sub>2</sub>) gives propanone. Pinacol coupling: Mg/Hg reduces two ketone carbonyls and they couple to give diol.

## Answer Summary:

Label	Formula	Name
x-products	$\text{CH}_3\text{CHO} + \text{CH}_3\text{COCH}_3$	Ethanal + Propanone (Ozonolysis)
y	$(\text{CH}_3)_2\text{C}(\text{OH})\text{C}(\text{OH})(\text{CH}_3)_2$	Pinacol (2,3-Dimethylbutane-2,3-diol)

Q47. C<sub>3</sub>H<sub>8</sub>O<sub>3</sub> (A) oxidised by Fenton's reagent → B and C

## Named Reaction: Fenton's Reagent Oxidation of Glycerol (Paragraph type)

## Reactions:

Step	Reaction
A	A = CH <sub>2</sub> OH-CHOH-CH <sub>2</sub> OH (Glycerol) — sweetening agent, trihydric alcohol
A→B	Glycerol + [O] → CHO-CHOH-CH <sub>2</sub> OH (Glyceraldehyde / 2,3-Dihydroxypropanal)
A→C	Glycerol + [O] → CH <sub>2</sub> OH-CO-CH <sub>2</sub> OH (Dihydroxyacetone / 1,3-Dihydroxypropan-2-one)

**Explanation:** Glycerol (C<sub>3</sub>H<sub>8</sub>O<sub>3</sub>) is a sweet-tasting trihydric alcohol used as sweetening agent and in cosmetics. Fenton's reagent (FeSO<sub>4</sub> + H<sub>2</sub>O<sub>2</sub>) oxidises either the terminal or central carbon. Oxidation of end carbon gives glyceraldehyde (aldose type). Oxidation of central carbon gives dihydroxyacetone (ketose type).

## Answer Summary:

Label	Formula	Name
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<b>A</b>	HOCH <sub>2</sub> CH(OH)CH <sub>2</sub> OH	<b>Glycerol (Propane-1,2,3-triol)</b>
<b>B</b>	CHO-CHOH-CH <sub>2</sub> OH	<b>Glyceraldehyde (2,3-Dihydroxypropanal)</b>
<b>C</b>	CH <sub>2</sub> OH-CO-CH <sub>2</sub> OH	<b>Dihydroxyacetone (1,3-Dihydroxypropan-2-one)</b>

**Q48.** C<sub>2</sub>H<sub>3</sub>N (A) → C<sub>2</sub>H<sub>7</sub>N (B) → C<sub>2</sub>H<sub>6</sub>O (C) — paragraph type

**Named Reaction: Cyanide Reduction + Carbylamine Test + Diazotisation**

**Reactions:**

Step	Reaction
<b>A</b>	A = CH <sub>3</sub> -CN (Methyl cyanide, C <sub>2</sub> H <sub>3</sub> N) – nitrile
<b>A→B</b>	CH <sub>3</sub> -CN + Na-Hg/C <sub>2</sub> H <sub>5</sub> OH → CH <sub>3</sub> -CH <sub>2</sub> -NH <sub>2</sub> (Ethylamine, C <sub>2</sub> H <sub>7</sub> N)
<b>B test</b>	CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub> + CHCl <sub>3</sub> /KOH → CH <sub>3</sub> CH <sub>2</sub> NC (Isocyanide – foul smell, carbylamine test positive)
<b>B→C</b>	CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub> + HNO <sub>2</sub> → CH <sub>3</sub> CH <sub>2</sub> OH (Ethyl alcohol, C <sub>2</sub> H <sub>6</sub> O) + N <sub>2</sub>

*Explanation: Methyl cyanide (A) reduces to ethylamine (B). Ethylamine gives positive carbylamine test (foul smelling isocyanide) confirming it's a primary amine. Ethylamine with HNO<sub>2</sub> (= NaNO<sub>2</sub>+HCl, aliphatic diazotisation) gives unstable diazonium → ethyl alcohol + N<sub>2</sub>.*

**Answer Summary:**

Label	Formula	Name
<b>A</b>	CH <sub>3</sub> -CN	<b>Methyl Cyanide (C<sub>2</sub>H<sub>3</sub>N)</b>
<b>B</b>	CH <sub>3</sub> CH <sub>2</sub> NH <sub>2</sub>	<b>Ethylamine (C<sub>2</sub>H<sub>7</sub>N)</b>
<b>C</b>	CH <sub>3</sub> CH <sub>2</sub> OH	<b>Ethyl Alcohol (C<sub>2</sub>H<sub>6</sub>O)</b>

**Q49.** C<sub>6</sub>H<sub>6</sub> (A) + propylene → B → C (C<sub>9</sub>H<sub>12</sub>O<sub>2</sub>) → D — paragraph type

**Named Reaction: Cumene Process (Industrial Phenol Production)**

**Reactions:**

Step	Reaction
<b>A</b>	A = C <sub>6</sub> H <sub>6</sub> (Benzene)
<b>A→B</b>	C <sub>6</sub> H <sub>6</sub> + CH <sub>3</sub> CH=CH <sub>2</sub> + H <sub>3</sub> PO <sub>4</sub> /523K → C <sub>6</sub> H <sub>5</sub> CH(CH <sub>3</sub> ) <sub>2</sub> (Cumene)
<b>B→C</b>	Cumene + Air/O <sub>2</sub> + 5% aq. Na <sub>2</sub> CO <sub>3</sub> → C <sub>6</sub> H <sub>5</sub> C(CH <sub>3</sub> ) <sub>2</sub> OOH (Cumene Hydroperoxide, C <sub>9</sub> H <sub>12</sub> O <sub>2</sub> )
<b>C→D</b>	C <sub>6</sub> H <sub>5</sub> C(CH <sub>3</sub> ) <sub>2</sub> OOH + H <sub>2</sub> SO <sub>4</sub> → C <sub>6</sub> H <sub>5</sub> OH (Phenol) + CH <sub>3</sub> COCH <sub>3</sub> (Acetone)

*Explanation: Cumene process is the industrial method for phenol production. Benzene alkylated with propylene gives cumene. Air oxidation in alkaline medium gives cumene hydroperoxide. Acid cleavage simultaneously gives phenol AND acetone (both are commercially valuable). This is an atom-efficient process.*

### Answer Summary:

Label	Formula	Name
<b>A</b>	C <sub>6</sub> H <sub>6</sub>	<b>Benzene</b>
<b>B</b>	C <sub>6</sub> H <sub>5</sub> CH(CH <sub>3</sub> ) <sub>2</sub>	<b>Cumene (Isopropylbenzene)</b>
<b>C</b>	C <sub>6</sub> H <sub>5</sub> C(CH <sub>3</sub> ) <sub>2</sub> OOH	<b>Cumene Hydroperoxide</b>
<b>D</b>	C <sub>6</sub> H <sub>5</sub> OH	<b>Phenol (+ Acetone as by-product)</b>

**Q50.** C<sub>2</sub>H<sub>6</sub>O (A) → C<sub>2</sub>H<sub>4</sub> (B) → C<sub>2</sub>H<sub>6</sub>O<sub>2</sub> (C) → C<sub>4</sub>H<sub>8</sub>O (D); A at 413K → C<sub>4</sub>H<sub>10</sub>O (E) — paragraph type

### Named Reaction: Ethanol — Complete Multi-Step Reaction Chain

#### Reactions:

Step	Reaction
<b>A→B</b>	CH <sub>3</sub> CH <sub>2</sub> OH + conc.H <sub>2</sub> SO <sub>4</sub> /443K → CH <sub>2</sub> =CH <sub>2</sub> (Ethylene B) [Dehydration]
<b>B→C</b>	CH <sub>2</sub> =CH <sub>2</sub> + alkaline KMnO <sub>4</sub> → HOCH <sub>2</sub> CH <sub>2</sub> OH (Glycol C) [Baeyer's hydroxylation, antifreeze]
<b>C→D</b>	2 HOCH <sub>2</sub> CH <sub>2</sub> OH + conc.H <sub>2</sub> SO <sub>4</sub> → 1,4-Dioxane (D, cyclic ether C <sub>4</sub> H <sub>8</sub> O)
<b>A→E</b>	CH <sub>3</sub> CH <sub>2</sub> OH + conc.H <sub>2</sub> SO <sub>4</sub> /413K → C <sub>2</sub> H <sub>5</sub> OC <sub>2</sub> H <sub>5</sub> (Diethyl Ether E) [Etherification]

*Explanation: Ethanol (A) is the starting material for many products depending on conditions: 443K gives ethylene; Baeyer's reagent gives glycol (used as antifreeze in car radiators); glycol + H<sub>2</sub>SO<sub>4</sub> gives 1,4-dioxane (cyclic compound). At 413K, ethanol gives diethyl ether. This illustrates the versatility of ethanol.*

**Answer Summary:**

Label	Formula	Name
A	C <sub>2</sub> H <sub>5</sub> OH	Ethyl Alcohol (Ethanol)
B	CH <sub>2</sub> =CH <sub>2</sub>	Ethylene (Ethene)
C	HOCH <sub>2</sub> CH <sub>2</sub> OH	Ethylene Glycol (antifreeze, C <sub>2</sub> H <sub>6</sub> O <sub>2</sub> )
D	1,4-dioxane	1,4-Dioxane (cyclic ether, C <sub>4</sub> H <sub>8</sub> O)
E	C <sub>2</sub> H <sub>5</sub> OC <sub>2</sub> H <sub>5</sub>	Diethyl Ether (C <sub>4</sub> H <sub>10</sub> O)

## MASTER NAMED REACTIONS QUICK REFERENCE

Named Reaction	Reagent / Condition	Product / What Happens
<b>Cannizzaro Reaction</b>	Aldehyde without $\alpha$ -H + 50% NaOH	Disproportionation — one oxidised, one reduced
<b>Aldol Condensation</b>	2 Aldehydes/ketones + dil. NaOH	Beta-hydroxy aldehyde (Aldol); heating $\rightarrow$ $\alpha,\beta$ -unsaturated
<b>Grignard Reaction</b>	RMgX (Grignard reagent) + carbonyl + H <sub>2</sub> O	Nucleophilic addition — chain extension
<b>Dow's Process</b>	ArCl + NaOH at high T,P	Industrial phenol from chlorobenzene
<b>Kolbe's Reaction</b>	ArONa + CO <sub>2</sub> / 400K / 4-7 bar	Carboxylation of phenol $\rightarrow$ Salicylic acid
<b>Reimer-Tiemann Reaction</b>	Phenol + CHCl <sub>3</sub> /NaOH	Formylation of phenol $\rightarrow$ Salicylaldehyde
<b>Friedel-Crafts Alkylation</b>	ArH + RCl + anhydrous AlCl <sub>3</sub>	Alkyl group added to ring
<b>Friedel-Crafts Acylation</b>	ArH + RCOCl + anhydrous AlCl <sub>3</sub>	Acyl group added to ring — gives ketone
<b>Cumene Process</b>	Benzene + propylene $\rightarrow$ Cumene $\rightarrow$ Phenol	Industrial phenol + acetone co-production
<b>Hoffmann Bromamide</b>	Primary amide + Br <sub>2</sub> /NaOH	Amine with 1 less C (decarboxylation)
<b>Diazotisation</b>	Primary ArNH <sub>2</sub> + NaNO <sub>2</sub> /HCl at 273-278K	Diazonium salt formation
<b>Sandmeyer Reaction</b>	ArN <sub>2</sub> Cl + CuX	Replaces -N <sub>2</sub> <sup>+</sup> with -CN, -Cl, -Br
<b>Gomberg-Bachmann</b>	ArN <sub>2</sub> Cl + ArH/KOH	Coupling of two arenes via radical
<b>Azo Coupling</b>	ArN <sub>2</sub> Cl + phenol/amine at pH 9-10	Coloured azo dye (-N=N-)
<b>Carbylamine Test</b>	Primary amine + CHCl <sub>3</sub> /KOH	Foul-smelling isocyanide — test for 1° amine
<b>Clemmensen Reduction</b>	C=O + Zn-Hg/HCl	Reduces C=O to CH <sub>2</sub> (acidic medium)
<b>Wolff-Kishner Reduction</b>	C=O + N <sub>2</sub> H <sub>4</sub> /base	Reduces C=O to CH <sub>2</sub> (basic medium)
<b>Rosenmund Reduction</b>	Acid chloride + Pd/BaSO <sub>4</sub> /H <sub>2</sub>	Reduces -COCl to -CHO (selective)
<b>HVZ Reaction</b>	Carboxylic acid + Cl <sub>2</sub> /red P	Alpha-halogenation of carboxylic acid
<b>Pinacol Coupling</b>	2 Ketones + Mg/Hg	Reductive coupling $\rightarrow$ 1,2-diol (pinacol)
<b>Benzoin Condensation</b>	2 ArCHO + alc.KCN	Alpha-hydroxy ketone (Benzoin)

<b>Markovnikov Addition</b>	Alkene + HX or H <sub>2</sub> O/H <sup>+</sup>	H adds to less substituted C
<b>Anti-Markovnikov</b>	Alkene + B <sub>2</sub> H <sub>6</sub> /H <sub>2</sub> O <sub>2</sub> /OH <sup>-</sup>	H adds to more substituted C (hydroboration)
<b>Schiff's Base Formation</b>	Aldehyde/ketone + 1° amine + trace H <sup>+</sup>	Imine (C=N) formed with loss of water
<b>Schotten-Baumann</b>	Acid chloride + amine in pyridine	N-acylation to form amide
<b>Baeyer's Hydroxylation</b>	Alkene + cold alk. KMnO <sub>4</sub> or OsO <sub>4</sub>	Syn dihydroxylation → vicinal diol
<b>HIO<sub>4</sub> Cleavage</b>	Vicinal diol (1,2-diol) + HIO <sub>4</sub>	Cleaves C-C bond of diol
<b>Ozonolysis</b>	Alkene + O <sub>3</sub> then Zn/H <sub>2</sub> O	Cleaves C=C to give aldehydes/ketones

## ALL THE BEST — SCORE CENTUM!

*Compiled by Embrace Coaching Academy | Source: Kaviyarasu Set + SS Prithvi Set*