

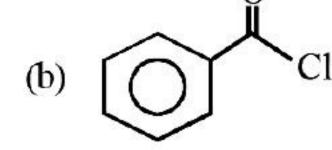
# Aldehydes, Ketones and Carboxylic Acids

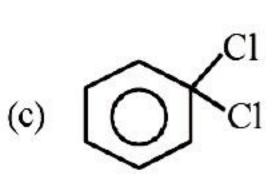
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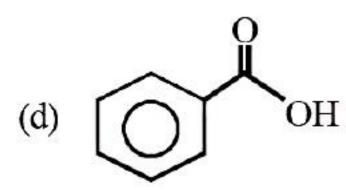
# Multiple Choice Questions (MCQs)

**DIRECTIONS:** This section contains multiple choice questions. Each question has four choices (a), (b), (c) and (d) out of which only one is correct.

- 1. In > C = O group sigma bond is formed by
  - (a)  $sp^2$ -p-overlapping
- (b)  $sp^3$ -p-overlapping
- (c) *sp-p*-overlapping
- (d) *s-p*-overlapping
- 2. Benzaldehyde is obtained from Rosenmund's reduction of







- 3. The conversion  $PhCN \rightarrow PhCOCH_3$ , can be achieved most conveniently by reaction with
  - (a) CH<sub>3</sub>MgBr followed by hydrolysis
  - (b)  $I_2$  NaOH, CH<sub>3</sub>I
  - (c) dil. H<sub>2</sub>SO<sub>4</sub> followed by reaction with CH<sub>2</sub>N<sub>2</sub>
  - (d) LiAlH<sub>4</sub> followed by reaction with CH<sub>3</sub>I
- 4. The reagent which can be used to distinguish acetophenone from benzophenone is
  - (a) 2,4-dinitrophenylhydrazine
  - (b) aqueous solution of NaHSO<sub>3</sub>
  - (c) benedict reagent
  - (d) I<sub>2</sub>and Na<sub>2</sub>CO<sub>3</sub>
- 5. The most appropriate reagent to distinguish between acetaldehyde and formaldehyde is:
  - (a) Fehling's solution
- (b) Tollen's reagent
- (c) Schiff's reagent
- (d) Iodine in presence of base
- 6. Cross aldol condensation occurs between
  - (a) two same aldehydes
    - (b) two same ketones
    - (c) two different aldehydes and ketones
    - (d) None of these

- 7. Imine derivatives of aldehyde and ketone is called as
  - (a) Schiff's reagent
- (b) Fehling's reagent
- (c) Schiff's base
- (d) Schiff's acid
- 8. Identify the product C in the series:

$$CH_3CN \xrightarrow{Na/C_2H_5OH} A \xrightarrow{HNO_2} B \xrightarrow{Cr_2O_7^{2-}/H^+} C$$

- (a) CH<sub>3</sub>COOH
- (b) CH<sub>3</sub>CH<sub>2</sub>NHOH
- (c) CH<sub>3</sub>CONH,
- (d) CH<sub>3</sub>CHO
- 9. Dimerisation of carboxylic acids is due to
  - (a) ionic bond
  - (b) covalent bond
  - (c) coordinate bond
  - (d) intermolecular hydrogen bond
- 10. Which of the following has the maximum acidic strength?
  - (a) o-nitrobenzoic acid
- (b) m-nitrobenzoic acid
- (c) p-nitrobenzoic acid
- (d) p-nitrophenol
- 11. The yield of ester in esterification can be increased by

$$CH_3CH_2OH + CH_3COOH \Longrightarrow$$

- (a) removing water
- (b) taking ethanol in excess
- (c) taking acetic acid in excess
- (d) all the above factors
- 12. Consider the following processes

Process (1) C,H,OH/H<sup>⊕</sup>, LiAlH<sub>4</sub>, PCC

Process (2) NaBH<sub>4</sub>, Cu/300°C

Process (3) PCl<sub>5</sub>, LiAlH<sub>4</sub>, CrO<sub>3</sub>/H<sub>2</sub>SO<sub>4</sub>

Process (4) SOCl, Pd/BaSO<sub>4</sub>/H,

The process by which acetic acid can be converted into acetaldehyde is/are

- (a) 1, 2, 3, 4
- (b) 1.4
- (c) 1,3,4
- (d) 2

3. Propionic acid 
$$\xrightarrow{P/Br_2}$$
 [W]  $\xrightarrow{KCN}$  [X]  $\xrightarrow{H_3O^{\oplus}}$  [Y] is

14.	Among the following acids which has the lowest $pK_a$ value?					(a)	(i), (ii) and (iv) are cor	rect.		
	(a)	CH <sub>3</sub> CH <sub>2</sub> COOH	(b)	$(CH_3)_2CH-COOH$		(b)	(ii), (iii) and (iv) are co	rrect.		
	(c)	HCOOH	(d)	CH <sub>3</sub> COOH		(c)	only(i) is correct.			
15.	OII-					(d) (i), (ii) and (iii) are correct.  What is the correct IUPAC name of the given compound				19
							$CH_3$		· · · · · · · · · · · · · · · · · · ·	
	<ul> <li>the conversion of benzaldehyde to benzyl alcohol?</li> <li>(i) One hydrogen is coming from H<sub>2</sub>O as H<sup>+</sup> and another from C<sub>6</sub>H<sub>5</sub>CHO as H<sup>-</sup>.</li> <li>(ii) One hydrogen is coming from H<sub>2</sub>O as H<sup>-</sup> and another from C<sub>6</sub>H<sub>5</sub>CHO as H<sup>+</sup>.</li> </ul>						***			
							CH <sub>3</sub> —C—C	$CH_2$	CH <sub>3</sub>	
							COC			
						(a)	2, 2-Dimethylbutanoi	e acid	1	
						(b)	2-Carboxyl-2-methylb			
	(111)	(iii) One hydrogen from H <sub>2</sub> O and another from C <sub>6</sub> H <sub>5</sub> CHO,				(c)	2-Ethyl-2-methylprop			
	Gwl	both in the form of H <sup>-</sup> .  (iv) This reaction is an example of disproportionation				(d)	3-Methylbutane carb			
	(1V)	reaction.	ехатр	de of disproportionation			0			
	(a)	(i), (ii) and (iii)	(b)	(i) and (iv)			Ĭ			
	(c)	(ii), (iii) and (iv)	(d)	(iii) and (iv)	21.	Com	pound Ph—O—C-	— Ph	can be prepared by th	16
16.	` '		, ,	d choose the correct option		react	tions of			
	(i)	The carbonyl carbon	atom i	is sp <sup>2</sup> -hybridised		(a)	**************************************		the presence of NaOH	
	(ii)	The carbonyl carbon	is an e	electrophilic (Lewis acid)		(b)	\$ <del>\tau</del>		in the presence of pyridin	
		centre				(c)	N.T		e in the presence of ZnC	4
	(111)	150 15 15 15 15 15 15 15 15 15 15 15 15 15	i is a n	ucleophilic (Lewis base)	22.	(d) Wh	17 T	200	the presence of palladium reacts in different way	
	Gra	Corbonyl compounds	o ora n	on polar in natura	LL.		CH <sub>3</sub> CHO, HCHO and			10
	200	Carbonyl compounds (i), (ii) and (iv) are co		on-poiai in nature.		(a)	Fehling solution	0 3	$C_6H_5NHNH_2$	
	(b)	(i), (ii) and (iii) are con				(c)	Ammonia	255	HCl	
	(c)	(ii), (iii) and (iv) are c			23.		cium acetate, on heatin	g, giv	es:	
		(ii) and (iv) are correct				(a)	acetic anhydride	(b)	acetone	
17.	Addi	tion of water to alkyn	es occu	ers in acidic medium and		(c)	acetaldehyde		ethyl alcohol	
		₹ TW		a catalyst. Which of the	24.	2.0	zaldehyde can be prep		트립트램 - 400 MM	
		Militaria		d on addition of water to		(a)	benzal chloride		benzotrichloride	
	but -	1-yne under these co	nditio	ns?	25	(c)	benzyl chloride	` '	benzonitrile	
		Ö		O	25.	9.3	cium formate on distilla HCOOH	1 TO SERVICE OF THE PARTY OF TH	CH <sub>2</sub> COOH	
	(a)	CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH	(b)	$CH_3CH_2 CCH_3$		(a) (c)	CH <sub>2</sub> CHO	(d)	HCHO	
	(a)	CH3CH2CH2 CH	(0)	C113C112 C C113	26.		3	1	by oxidation of toluene b	7(
		O		o o		(a)	Acidic KMnO <sub>4</sub>		K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> /H <sup>+</sup>	J
	(c)	$CH_3CH_2COH + CO$	(d)	$CH_3 COH + HCH$		(c)	CrO,Cl,	(d)	All of these	
40		(R) (S)	No.	CH3 COH THEH	27.	The	reagent which does no	ot reac	ct with both, acetone an	C
18.		Iodoform test is not given by					zaldehyde?	2 200		
	` '	Ethanol Pentan-2-one	(b) (d)	Ethanal Pentan-3-one			Sodium hydrogen sul	phite		
19.	\ /			and choose the correct			Phenyl hydrazine			
	opti		memo			0.0	Fehling's solution Grignard reagent			
	(i)						tify the product C in th	e seri	es	
	3.5									
	(ii)	Alcohols show inter	rmolec	cular hydrogen bonding		CH <sub>3</sub>	$CN \xrightarrow{Na/C_2 H_5 OH} A$		$\rightarrow B \xrightarrow{Cu/573K} C$	
				ketones do not show		\$ 5	CH <sub>3</sub> COOH	W. C.	CH <sub>3</sub> CH <sub>2</sub> NHOH	
	7000	intermolecular hydro			<b>A</b> 0		CH <sub>3</sub> CONH <sub>2</sub>	20,000,000,000	CH <sub>3</sub> CHO	
	(111)			ehydes and ketones are	29.				oes not form a ketone o	1
		form bydrogen band		proportions, because they		treat	ment with dil H <sub>2</sub> SO <sub>4</sub>	and 1°	% HgSO <sub>4</sub> ?	

(b)  $C_2H_2$ 

(d) All of these

(a)  $C_4H_6$ 

(c)  $C_3H_4$ 

form hydrogen bond with water.

(iv) The solubility of aldehydes and ketones increases

rapidly on increasing the length of alkyl chain

30. Ketones 
$$[R - C - R_1]$$
, where  $R = R_1 =$ alkyl groups $]$ 

can be obtained in one step by

- oxidation of primary alcohols
- hydrolysis of esters
- oxidation of secondary alcohols
- reaction of acid halides with alcohols

31. 
$$OH \longrightarrow OH \longrightarrow H$$

$$(1)NaOH \longrightarrow O$$

$$(2) H^{+} \longrightarrow O$$

Reactant X is:

- CH<sub>2</sub>Cl
- CH,Cl,

32. 
$$CH_3 - C \equiv CH - \frac{40\% H_2 SO_4}{1\% Hg SO_4} \rightarrow A$$

$$\xrightarrow{\text{Isomerisation}} CH_3 - C - CH_3$$

structure of 'A' and type of isomerism in the above reaction are respectively

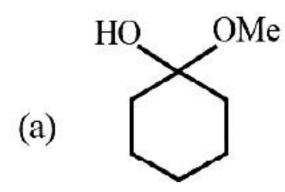
- Prop-1-cn-2-ol, metamerism
- Prop-1-en-1-ol, tautomerism
- Prop-2-en-2-ol, geometrical isomerism
- Prop -1-en-2-ol, tautomerism

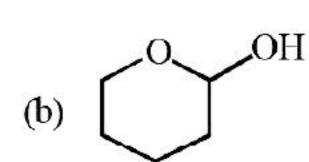
33. 
$$CH_3$$
— $C$ — $CH_2$ —

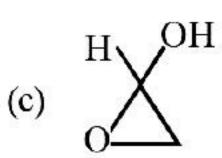
$$CH_3$$
— $(CH_2)_3$ — $CH$ — $CH_2$ 

Above conversion can be achived by:

- Wolff-Kishner reduction
- Clemmensen reduction
- HS CH<sub>2</sub>— CH<sub>2</sub> SH, following by Raney Ni
- None of these
- Which of following compound is hemiacetal?







all of these

- The presence of unsaturation in organic compounds can be tested with:
  - Schiff's reagent
- (b) Tollens' reagent
- Fehling's reagent
- Baeyer's reagent
- Which of the following is correct for the reaction?

$$+ HCN \xrightarrow{pH = 9 \text{ to } 10} (A)$$

- A is cyanohydrin
- Nucleophilic-addition reaction
- The above reaction is not shown by alkenes
- All of these

37. 
$$CH_3CHO \xrightarrow{10\% \text{ NaOH}} \xrightarrow{\Delta} \xrightarrow{H_2} (A);$$

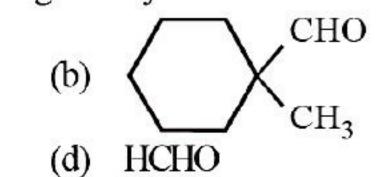
Product (A) of the reaction is:

- (a) propanol
- (b) ethanol
- butanol
- (d) pentanol
- Which is major product formed when acetone is heated with iodine and potassium hydroxide?
  - Iodoacetone
- (b) Acetic acid
- Iodoform
- (d) Acetophenone
- Compounds A and C in following reaction are ........

$$CH_3CHO \xrightarrow{(i) CH_3MgBr} (A) \xrightarrow{H_2SO_4, \Lambda} (B)$$

 $\xrightarrow{\text{Hydroboration oxidation}} (C)$ 

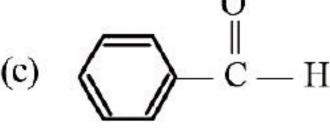
- (a) identical
- (b) positional isomers
- (c) functional isomers
- (d) optical isomers
- Which of the following compounds will give butanone on oxidation with alkaline KMnO<sub>4</sub> solution?
  - Butan-1-ol
- (b) Butan-2-ol
- Both (a) and (b) (d) None of these
- A compound that gives a positive iodoform test is
  - (a) 1-pentanol
- (b) 2-pentanone
- (c) 3-pentanone
- (d) pentanal
- Clemmensen reduction of a ketone is carried out in the presence of:
  - (a) LiAlH<sub>4</sub>
- (b) Zn-Hg with HCl
- Glycol with KOH
- (d) H<sub>2</sub> with Pt as catalyst
- Cannizaro's reaction is not given by:



 $CH_1CHO$ The reaction,

- (a) Cannizaro's reaction (b) Rosenmund reduction
- Wolf-Kishner reduction (d) Clemmenson reduction
- Which of the following compounds is most reactive
- towards nucleophilic addition reactions?

(b) 
$$CH_3 - C - CH_3$$



(d) 
$$\left( \begin{array}{c} C \\ C \end{array} \right) = CH_3$$

- Oxidation of toluene with CrO<sub>3</sub> in the presence of (CH<sub>3</sub>CO)<sub>2</sub>O gives a product (A) which on treatment with aqueous NaOH produces
  - (a)  $C_6H_5CHO$
- (b)  $(C_6H_5CO)_2O$
- (c) C<sub>6</sub>H<sub>5</sub>COONa
- (d) 2, 4-Diacetyltoluene

47. Which carbon atoms are most susceptible to nucleophilic attack?

$$CH_3 - CH = CH - C - CH_3$$

$$A \quad B \quad C \quad D$$

- (a) A and B (b) B and C (c) B and D (d) A and D
- Chlorination of toluene in the presence of light and heat followed by treatment with aqueous NaOH gives
  - o-Cresol
- (b) p-Cresol
- 2, 4-Dihydroxytoluene (d) Benzoic acid
- 49. Cannizzaro's reaction is not given by ......



- **HCHO** (c)
- CH,CHO
- The compound that does **NOT** liberate CO<sub>2</sub>, on treatment with aqueous sodium bicarbonate solution, is
  - Benzoic acid
- (b) Benzenesulphonic acid
- Salicylic acid (c)
- (d) Carbolic acid (Phenol)

51. 
$$\langle \bigcirc \rangle$$
 —  $CH_3 \rightleftharpoons X$   $\langle \bigcirc \rangle$  —  $COOH$ 

In the above sequence of reaction X and Y are respectively

- (a)  $H_2/Pt$ ;  $Br_2$
- (b)  $KMnO_4$ ;  $H_2/Pt$
- $KMnO_4$  (aq); HI/P (d)  $NH_2 NH_2/KOH$ , HI/P
- Hydrolysis of an ester may be achived under acidic as well as basic conditions. Pick up the correct statement regarding this
  - Acidic hydrolysis is faster than alkaline hydrolysis.
  - Alkaline hydrolysis is faster than acidic hydrolysis.
  - Both occur at the same rate.
  - In both, the first step is protonation of the —OH part of the —COOH group.
- Acid amides do not undergo the usual properties of carbonyl, C = O group because
  - it is a weak base
  - it is a weak acid
  - it is amphoteric
  - its carbonyl carbon is not electron deficient
- 54. Which is the most suitable reagent for the following conversion?

$$CH_{3} - CH = CH - CH_{2} - C - CH_{3} \longrightarrow$$

$$CH_{3} - CH = CH - CH_{2} - C - OH$$

- Tollen's reagent (a)
- Benzoyl peroxide (b)
- I, and NaOH solution
- Sn and NaOH solution (d)

- $CH_3CH = CHCOOH + HBr \longrightarrow P$ , here P is
- $CH_3CHCH_2COOH$  (b)  $CH_3CH_2CHCOOH$

Br

- (d) None
- When propionic acid is treated with aqueous sodium bicarbonate, CO, is liberated. The 'C' of CO, comes from
  - (a) methyl group

Both

- (b) carboxylic acid group
- methylene group
- (d) bicarbonate
- Benzoyl chloride is prepared from benzoic acid by
  - (a)  $Cl_2, hv$
- (b) SO<sub>2</sub>Cl<sub>2</sub>
- SOCl,
- (d) Cl<sub>2</sub>, H<sub>2</sub>O
- Among acetic acid, phenol and *n*-hexanol, which of the following compounds will react with NaHCO<sub>3</sub> solution to give sodium salt and carbon dioxide?
  - Acetic acid
- (b) *n*-Hexanol
- Acetic acid and phenol (d) Phenol.
- An ester is boiled with KOH. The product is cooled and acidified with concentrated HCl. A white crystalline acid separates. The ester is
  - methyl acetate
- (b) ethyl acetate
- ethyl formate
- (d) ethyl benzoate
- In the Friedel Craft's acylation reaction, the effective electrophile is
  - (a) RCOCl⊕ (b) AlCl<sub>3</sub>
- (c) RCOCl (d) RCO⊕
- Which one of the following esters cannot undergo Claisen self-condensation?
  - (a)  $CH_3 CH_2 CH_2 CH_2 COOC_2H_5$
  - (b)  $C_6H_5COOC_2H_5$
  - (c) C<sub>6</sub>H<sub>5</sub>CH<sub>5</sub>COOC<sub>5</sub>H<sub>5</sub>
  - (d)  $C_6H_{11}CH_2COOC_2H_5$
- The cyanohydrin of a compound on hydrolysis gives an optically active  $\alpha$ -hydroxy acid. The compound is
  - diethyl ketone
- (b) formaldehyde
- acetaldehyde
- (d) acetone
- $CH_3CH_2COOH \xrightarrow{Cl_2} A \xrightarrow{alc. KOH} B$ . What is B?

- (a) CH<sub>2</sub>CH<sub>2</sub>COCl (b) CH<sub>3</sub>CH<sub>2</sub>CHO (c) CH<sub>2</sub>=CHCOOH (d) ClCH<sub>2</sub>CH<sub>2</sub>COOH.
- When  $CH_2 = CH COOH$  is reduced with  $LiAlH_4$ , the compound obtained will be

  - (a)  $CH_2 = CH CH_2OH$  (b)  $CH_3 CH_2 CH_2OH$
  - (c) CH<sub>3</sub>—CH<sub>2</sub>—CHO (d) CH<sub>3</sub>—CH<sub>2</sub>—COOH
- 65. On mixing ethyl acetate with aqueous sodium chloride, the composition of the resultant solution is
  - (a)  $CH_3COCl+C_2H_5OH+NaOH$
  - (b) CH<sub>3</sub>COONa+C<sub>2</sub>H<sub>5</sub>OH (c) CH<sub>3</sub>COOC<sub>2</sub>H<sub>5</sub>+NaCl

  - (d) CH<sub>3</sub>Cl+C<sub>2</sub>H<sub>5</sub>COONa
- Acetyl bromide reacts with excess of CH<sub>3</sub>MgI followed by treatment with a saturated solution of NH<sub>4</sub>Cl and gives
  - (a) 2-methyl-2-propanol
- (b) acetamide
- acetone
- (d) acetyl iodide

**DIRECTIONS:** Following are the case/passage based questions.

#### Case/Passage-I

Reduction of carboxylic acids and their derivatives plays an important role in organic synthesis, in both laboratory and industrial processes. Traditionally, the reduction is performed using stochiometric amounts of hydride reagents, generating stochiometric amounts of waste. A much more attractive, atomeconomical approach is a catalytic reaction using H<sub>2</sub>; however, hydrogenation of carboxylic acid derivatives under mild conditions is a very challenging task, with amides presenting the highest challenge among all classes of carbonyl compounds. Very few examples of the important hydrogenation of amides to amines, in which the C-O bond is cleaved with the liberation of water (Scheme 1), were reported. C-O cleavage of amides can also be affected with silanes as reducing agents.

Scheme - 1

$$R' \leftarrow \frac{^{2 H_2}}{^{-H_2O}}$$

We have now prepared the new, dearomatized, bipyridine-based pincer complex 3, catalyst 3(Here refered as Cat. 3). Remarkably, it efficiently catalyzes the selective hydrogenation of amides to form amines and alcohols (eq 1). The reaction proceeds under mild pressure and neutral conditions, with no additives being required. Since the reaction proceeds well under anhydrous conditions, hydrolytic cleavage of the amide is not involved in this process.

$$R' + 2 H_2$$

(Balaraman, E., Gnanaprakasam, B., Shimon, L. J., & Milstein, D. (2010). Direct hydrogenation of amides to alcohols and amines under mild conditions. Journal of the American Chemical Society, 132(47), 16756-16758.)

#### [From CBSE Question Bank-2021] In the following questions, a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices on the basis of the above passage.

- Assertion and reason both are correct statements and reason is correct explanation for assertion.
- (b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.
- Assertion is correct statement but reason is wrong (c) statement.
- Assertion is wrong statement but reason is correct (d) statement.
- 67. Assertion: The use of catalyst 3 is an efficient method of preparation of primary amines.

**Reason:** Use of catalyst 3 is a step down reaction.

Assertion: Use of hydride catalyst or hydrogen brings about cleavage of C-O bond in amides.

- **Reason:** Hydride catalyst or hydrogen cause to reduction of amides.
- **Assertion:** N-methyl ethanamide on reaction with catalyst 3 will yield ethanol and methanamine.

**Reason:** Use of Catalyst 3 brings about cleavage of C-N bond of amides

Assertion: Aniline can be prepared from suitable amide using catalyst 3

**Reason:** The use of catalyst 3 is limited to aliphatic amides only.

#### Case/Passage-II

Ketones may be prepared by the oxidation of secondary alcohols in the presence of acid dichromate. The reaction probably proceeds via the rate limiting formation of a chromate ester followed by decomposition of the ester to give the products. In case of primary alcohol, this process may lead to oxidation of the alcohol to carboxylic acid. This problem is overcome by attenuating the oxidizing power of chromic acid based reagents to generate a milder oxidizing agent. The two most popular oxidizing agents for oxidizing primary alcohols to aldehydes are pyridinium chlorochromate (PCC) and pyridinium dichromate (PDC). These reagents are prepared by the treatment of chromic acid with pyridine. The ketones may also be generated from the secondary alcohols using aluminium t-butoxide in the presence of acetone. The secondary alcohol is refluxed with the reagent in an excess of acetone. This reagent is particularly helpful in oxidation of unsaturated alcohol as only the alcoholic OH is oxidized.

- Which of the following statement(s) is/are true regarding preparation of aldehydes and ketones?
  - Both can be prepared by the oxidation of the concerned alcohol with copper at about 250°C.
  - Both can be prepared by the oxidation of the concerned alcohol by Oppenauer oxidation.
  - (iii) Both can be prepared by the oxidation of respective alcohol with acidic dichromate.
  - (i) only
- (b) (ii) and (iii)
- (i) and (iii)
- (d) All the three
- Consider the following statements: Acetophenone can be prepared by
  - oxidation of 1-phenyl ethanol
  - reaction of benzaldehyde with methyl magnesium bromide
  - Friedel's-Crafts reaction of benzene with acetyl chloride
  - distillation of calcium benzoate
  - II and III
- (b) I and IV
- I and III
- (d) III and IV
- 2-Pentanone and 3-pentanone can be distinguished by:
- Cannizzaro's reaction (b) Aldol condensation
  - Iodoform reaction
- Clemmensen's reduction
- Phenylmethyl ketone can be converted into ethylbenzene in one step by which of the following reagents?
  - LiAlH<sub>4</sub>
- (b) Zn-Hg/HCl
- NaBH<sub>4</sub>
- (d) CH<sub>3</sub>MgI
- Which of the following compounds will give butanone on oxidation with alkaline KMnO<sub>4</sub> solution?
  - Butan-1-ol
- Butan-2-ol
- Both of these
- None of these

## >> Assertion & Reason

**DIRECTIONS**: Each of these questions contains an assertion followed by reason. Read them carefully and answer the question on the basis of following options. You have to select the one that best describes the two statements.

- If both Assertion and Reason are correct and the Reason is a correct explanation of the Assertion.
- (b) If both Assertion and Reason are correct but Reason is not a correct explanation of the Assertion.
- If the Assertion is correct but Reason is incorrect.
- If the Assertion is incorrect but the Reason is correct.
- Assertion: Carbonyl compounds when react with NaHSO<sub>3</sub> form white crystalline solid.

**Reason:** Aldehydes are used in the blending of perfumes and flavouring agents.

- 77. **Assertion:** Benzaldehyde gives silver mirror test. **Reason:** Benzaldehyde gives Fehling's solution test.
- **Assertion:** Aldehydes are more reactive than ketones in nucleophilic addition reaction.

Reason: This is due to steric and electronic reasons.

**Assertion:** *p*-Chlorobenzoic acid is a stronger acid than *m*-chlorobenzoic acid.

**Reason:** In *meta* position, only inductive effect is considered.

**80.** Assertion: Carboxylic acids when reacts with NaHCO<sub>3</sub>, the CO<sub>2</sub> evolved comes from carboxylic acid and not from NaHCO<sub>3</sub>.

**Reason:** RCOOH is a stronger acid than H<sub>2</sub>CO<sub>3</sub>.

**Assertion:** The carboxylate ion is more stabilised than phenoxide ion.

**Reason:** This is because the negative charge is delocalised over two electronegative oxygen atoms in carboxylate ion, whereas it is less effectively delocalised over one atom in phenoxide ion.

**82.** Assertion: Compounds containing –CHO group are easily oxidised to corresponding carboxylic acids.

Reason: Carboxylic acids can be reduced to alcohols by treatment with LiAlH<sub>4</sub>.

83. Assertion: The  $\alpha$ -hydrogen atom in carbonyl compounds is less acidic.

**Reason**: The anion formed after the loss of  $\alpha$ -hydrogen atom is resonance stabilised.

**84.** Assertion: Aldehydes and ketones, both react with Tollen's reagent to form silver mirror.

Reason: Both, aldehydes and ketones contain a carbonyl group.

85. Assertion: Acetylene on treatment with alkaline KMnO<sub>4</sub> produces acetaldehyde.

**Reason:** Alkaline  $KMnO_4$  is an oxidising agent.

**86.** Assertion: 2, 2-dimethylpropanal undergoes Cannizzaro reaction with conc. NaOH.

**Reason:** Cannizzaro reaction is a disproportionation reaction.

87. Assertion: Isobutanal does not give iodoform test. **Reason:** It does not have  $\alpha$ -hydrogen.

# >> Match the Following

**DIRECTIONS**: Each question contains statements given in two columns which have to be matched. Statements (A, B, C, D) in column-I have to be matched with statements (p, q, r, s) in column-II.

#### Match the columns

#### Column-I (Common names)

#### Column-II (IUPAC names)

- Cinnamaldehyde
- (p) Pentanal
- Acetophenone
- Prop-2-enal
- Valeraldehyde
- 4-Methylpent-3-en-2-one 3-Phenylprop-2-enal
- Acrolein
- 1-Phenylethanone
- Mesityl oxide
  - A-(s), B-(t), C-(p), D-(q), E-(r)
- (b) A-(p), B-(q), C-(s), D-(t), E-(r)
- (c) A-(t), B-(s), C-(p), D-(r), E-(q)
- (d) A-(q), B-(t), C-(r), D-(s), E-(p)
- Match the columns

#### Column-I

# Column-II

- (A)  $R CO CH_3 \xrightarrow{Zn-Hg/HC1}$  (p) Friedel-Craft's  $R - CH_2 - CH_3$ 
  - reaction
- (B)  $2C_6H_5CHO \xrightarrow{NaOH}$  $C_6H_5COONa + C_6H_5CH_2OH$
- (q) Kolbe's reaction
- (C)  $C_6H_6 + CH_3COCl \xrightarrow{Anhyd.}$  (r) C<sub>6</sub>H<sub>5</sub>COCH<sub>3</sub>
  - reaction

Clemmensen's

- (D)  $C_6H_5OH + CO_2 + NaOH \rightarrow$ HOC<sub>6</sub>H<sub>4</sub>COONa
  - (s) Cannizzaro's reaction
- (a) A-(p), B-(q), C-(r), D-(s)
- (b) A-(r), B-(p), C-(q), D-(s)
- (c) A-(r), B-(s), C-(p), D-(q)
- (d) A-(s), B-(r), C-(p), D-(q)
- Match the columns 90.

## Column-I (Reactions)

## Column-II (Reagents)

- (A) Benzophenone  $\rightarrow$
- (p) LiAlH<sub>4</sub>
- (B) Benzaldehyde  $\rightarrow$
- Diphenylmethane (q) DIBAL-H
- (C) Cyclohexanone  $\rightarrow$
- 1-Phenylethanol
- (r) Zn(Hg)/conc. HCl Cyclohexanol
- (D) Phenyl benzoate  $\rightarrow$
- (s) CH<sub>3</sub>MgBr Benzaldehyde
- (a) A-(p), B-(s), C-(r), D-(q)
- (b) A-(q), B-(s), C-(p), D-(r)
- (c) A-(s), B-(r), C-(q), D-(p)
- (d) A-(r), B-(s), C-(p), D-(q)

>> Fill in the Blanks

**DIRECTIONS**: Complete the following statements with an appropriate word / term to be filled in the blank space(s).

- 91. Tollen's reagent and Fehling solutions are used to distinguish between \_\_\_\_\_.
- **92.** Monocarboxylic acids are functional isomers of .
- 93. Rancidity of butter is due to the formation of \_\_\_\_\_.
- 94. Formic acid is obtained when calcium acetate is heated with .
- 95. The catalyst used in Rosenmund's reduction is \_\_\_\_\_

>>> True / False

**DIRECTIONS**: Read the following statements and write your answer as true or false.

- **96.** The preparation of ethyl acetoacetate involves Cannizzaro's reaction.
- 97. Phthalic acid reacts with resorcinol in the presence of concentrated H<sub>2</sub>SO<sub>4</sub> to give Coumarin.

98. COOH is used as antihistamine:

- 99. The enolic form of acetone contains 9 sigma bonds, 1 pi-bond and 2 lone pairs.
- 100. Formaldehyde can be distinguished from acetaldehyde by the use of I<sub>2</sub>/Alkali.

# ANSWER KEY & SOLUTIONS

#### 1. (a)

2. (b) Rosenmund's reduction:
Acylchloride is hydrogenated over catalyst, Pd-BaSO<sub>4</sub>.

Acylchloride is hydrogenated over catalyst, Pd-BaSO<sub>4</sub> This reaction is called Rosenmund reduction.

3. (a) Alkanenitriles (other than methanenitrile) benzonitrile give ketones with Grignard reagents.

4. (d) I<sub>2</sub> and Na<sub>2</sub>CO<sub>3</sub> react with acetophenone (C<sub>6</sub>H<sub>5</sub>COCH<sub>3</sub>) to give yellow ppt. of CHI<sub>3</sub> but benzophenone (C<sub>6</sub>H<sub>5</sub>COC<sub>6</sub>H<sub>5</sub>) does not and hence can be used to distinguish between them.

5. (d) HCHO does not undergo iodoform test, while acetaldehyde undergoes iodoform test (I<sub>2</sub> in presence of base) to form yellow precipitate of iodoform.

6. (c) In cross aldol condensation aromatic aldehydes or ketones (with or without α-hydrogen) react with aldehydes, ketones or esters having α-hydrogen atoms in the presence of dilute alkali to form a β-unsaturated carbonyl compound.

7. **(c)** 

8. (a) 
$$CH_3CN \xrightarrow{Na} CH_3CH_2NH_2 \xrightarrow{HNO_2} CH_3CH_2OH$$
(B)
$$\downarrow Cr_2O_7^{2-}/H^+$$

$$CH_3COOH$$
(C)

9. **(d)** 

10. (a) o-Nitrobenzoic acid is more acidic due to ortho effect and higher inductive effect compared to -m and -p derivatives.

11. (d) The yield of product in a reversible reaction can be increased by (i) removing one of the products, (ii) taking either of the reactant in excess.

12. (b)

13. (d) 
$$COOH \xrightarrow{P+Br_2} COOH \xrightarrow{W} COOH \xrightarrow{KCN} COOH \xrightarrow{heat} COOH \xrightarrow{heat} COOH$$

14. (c)  $pK_a = -\log K_a$ ; HCOOH is the strongest acid and hence it has the highest  $K_a$  or lowest  $pK_a$  value.

15. (b) The hydrogen atom that is added to the carbonyl carbon of the aldehyde in the reduction is derived directly from the other aldehyde molecule as a hydride ion. The second hydrogen that is added to the negatively charged oxygen is coming from the solvent (consult mechanism of Cannizzaro reaction). Oxidation of one molecule of the compound at the expense of other molecule of the same compound is known as disproportionation.

16. (b) Carbonyl compounds have substantial dipole moments and are polar in nature. The high polarity of the carbonyl group is due to resonance.

17. (b) But-1-yne on reaction with water in presence of Hg<sup>2+</sup> ions as a catalyst produces butan-2-one.

18. (d) The iodoform test is a test for the presence of carbonyl compounds with the structure RCOCH<sub>3</sub> and alcohols with the structure R - CH(OH)CH<sub>3</sub>.

19. (d) The solubility of aldehydes and ketones decreases on increasing the length of alkyl chain.

20. (a) 2,2-Dimethylbutanoic acid

$$CH_{3}$$
 $H_{3}C$ 
 $= {}^{2}C - {}^{3}CH_{2} - {}^{4}CH_{3}$ 
 $= {}^{1}COOH$ 

21. (b) Compound Ph — COO — Ph can be prepared by the reaction of phenol and benzoyl chloride in the presence of pyridine. This is an example of Schotten-Baumann reaction.

(c) With ammonia, HCHO forms hexamethylenetetramine, CH<sub>3</sub>CHO gives acetaldehydeammonia addition product, while C<sub>6</sub>H<sub>5</sub>CHO gives hydrobenzamide.

23. **(b)** 
$$(CH_3COO)_2Ca \xrightarrow{\Delta} CH_3 - C - CH_3 + CaCO_3$$
Calcium acetate Acetone

24. (a) 
$$C_6H_5CHCl_2 \xrightarrow{H^+/H_2O} C_6H_5CHO + H_2O$$

25. (d) 
$$(HCOO)_2 Ca \xrightarrow{\Delta} H - C - H + CaCO_3$$

26. (c)

$$\begin{array}{c} \begin{array}{c} \begin{array}{c} \text{COOH} \\ \end{array} \\ \begin{array}{c} \begin{array}{c} \text{CH}_3 \\ \end{array} \\ \text{or KMnO}_4 \end{array} \end{array} \\ \begin{array}{c} \begin{array}{c} \text{CH}_3 \\ \end{array} \\ \end{array} \\ \begin{array}{c} \text{CrO}_2\text{Cl}_2 \\ \end{array} \\ \end{array} \\ \begin{array}{c} \text{CH}_3 \\ \end{array} \\ \begin{array}{c} \text{CH}_3$$

Calcium formate formaldehyde

Acetone and benzaldehyde both do not react with Fehling's solution.

28. (d) 
$$CH_3CN \xrightarrow{Na/C_2H_5OH} CH_3CH_2NH_2 \xrightarrow{HNO_2} CH_3CH_2OH \xrightarrow{Cu/573K} CH_3CHO$$
[B] [C]

29. **(b)** 
$$C_2H_2 + H_2O \xrightarrow{H_2SO_4} Ethanal$$

 $C_4H_6$  (CH<sub>3</sub>C = CCH<sub>3</sub>) and  $C_3H_4$  (CH<sub>3</sub>C = CH) give ketones.

- (c) By oxidation of secondary alcohols, ketones are formed.
- 31. 32. (d)
- 33. Wolff-Kishner reduction is carried out in basic condition whereas clemmensen reduction carried out in acidic condition. Epoxides are unstable in both acidic and basic medium.
- (d) Hemiacetal → presence of alcohol and ether on same 34. carbon.
- Cold dil. alk KMnO<sub>4</sub>, is Baeyer's reagent.
  - Schiff's reagent is used to test presence of Aldehydes.
  - Tollen's reagent is used to test presence of aldehydes.
  - Fehling's reagent is used to test presence of aldehyde.
  - Baeyer's reagent is cold dil. alk. KMnO<sub>4</sub> is used to test presence of unsaturation.

No partial positive charge developed on carbon in alkene. Nucleophile  $(-C \equiv N)$  is added to carbonyl carbon only.

37. (c) 
$$CH_3CHO \xrightarrow{NaOH} CH_3-CH \equiv CH-CHO$$

$$\xrightarrow{H_2/Ni} CH_3-CH_2-CH_2-CH_2-OH$$

38. (c) 
$$CH_3COCH_3 + 3I_2 + 4KOH \longrightarrow$$
  
 $CHI_3 + CH_3COOK + 3HI + 3H_2O$ 

Chemical reaction can be shown as

$$\begin{array}{c} \text{CH}_3\text{CHO} \xrightarrow{\text{(i) CH}_3\text{MgBr}} \text{CH}_3 \xrightarrow{\text{CH}} \text{CH} \xrightarrow{\text{OH}} \xrightarrow{\text{H}_2\text{SO}_4, \, \Delta} \xrightarrow{\text{Dehydration}} \\ \text{CH}_3 \\ \text{Propan-2-ol} \\ \text{(A)} \end{array}$$

$$\begin{array}{c} \operatorname{CH_3} - \operatorname{C} = \operatorname{CH_2} \xrightarrow{\operatorname{Hydroboration}} \operatorname{CH_3} - \operatorname{CH_2} - \operatorname{CH_2OH} \\ \downarrow & \text{propan -1-ol} \\ \operatorname{Propene} \\ \text{(B)} \end{array}$$

2° alcohols on oxidation with alkaline KMnO<sub>4</sub> solution produce ketones.

Compounds having  $-C - CH_3$  groups show

positive iodoform test.

Hence, 
$$CH_3 - CH_2 - CH_2 - CH_3$$
 (pentanone-2)

gives this test.

42. **(b)** 
$$C = O \xrightarrow{Zn-Hg/HCl} CH_2$$

Ketone Alkane

- 43. (c) Only those aldehydes which do not have  $\alpha$ -H atom undergo Cannizaro's reaction. Hence CH<sub>3</sub>CHO will not undergo Cannizaro's reaction as it has 3 α H atoms.
- The reaction given is a Clemmenson reduction. 44.
- The carbonyl group in ketones being influenced by 45. two alkyl group is less reactive than in aldehydes where the carbonyl group is under the influence of one alkyl group only. As the number of alkyl group increases both the +I effect and the steric hinderance get increases preventing the attack of nucleophile.

Now among benzaldehyde and acetaldehyde former is less electrophilic than carbon atom of carbonyl group present 56. (d)  $C_2H_5COOH + NaHCO_3 \longrightarrow$ in ethanal. The polarity of carbonyl group is reduced in benzaldehyde due to resonance hence it is less reactive than ethanal.

**46.** (a) 
$$C_6H_5CH_3 \xrightarrow{CrO_3} [C_6H_5CHO] \xrightarrow{(CH_3CO)_2O}$$

$$C_6H_5CH(OCOCH_3) \xrightarrow{OH^-} C_6H_5CHO + 2CH_3COONa$$

47. (c) 
$$CH_3 - CH = CH^2 C - CH_3 \longleftrightarrow$$

$$O^-$$

$$\oplus$$

$$CH_3 - CH - CH = C - CH$$

48. (d) 
$$C_6H_5CH_3 + Cl_2(excess) \xrightarrow{\text{light, heat}} C_6H_5CCl_3$$
  
 $\xrightarrow{\text{aq. NaOH}} C_6H_5C(OH)_3 \xrightarrow{-H_2O} C_6H_5COOH$ 

- 49. (d) Cannizzaro reaction is given by those aldehydes which have absence of α-hydrogen atom. So, CH<sub>3</sub>CHO will not give Cannizzaro reaction.
- (d) Carbolic acid (Phenol) is weaker acid than carbonic acid and hence does not liberate CO<sub>2</sub> on treatment with aq. NaHCO<sub>3</sub> solution.
- 51. (c) KMnO<sub>4</sub> converts CH<sub>3</sub> group of toluene into COOH while HI reduces – COOH group into – CH<sub>3</sub> group.
- (b) Alkaline hydrolysis is irreversible because here RCOO is isolated; moreover the product RCOO stabilizes itself due to resonance.
- (d) Due to possibility of resonance, carbonyl carbon is no more electron deficient.

(Electron deficient carbon)

(Carbonyl carbon is not electron deficient)

54. (c) 
$$CH_3 - CH = CH - CH_2 - C - CH_3 - \frac{I_2 / \text{NaOH solution}}{\text{(Iodoform reaction)}}$$

$$Hex-4-en-2-one$$

$$CH_3$$
 —  $CH = CH$  —  $CH_2$  —  $C$  —  $OH$ 

Pent-3-en-1-oic acid

Addition of an unsymmetrical reagent to an  $\alpha$ ,  $\beta$ unsaturated carbonyl compound takes place in such a way that hydrogen becomes attached to the  $\alpha$ -carbon and the negative group becomes attached to the  $\beta$ -carbon.

56. (d) 
$$C_2H_5COOH + NaHCO_3 \longrightarrow$$

$$C_2H_5COONa + H_2O + CO_2$$

57. (c)  $C_6H_5COOH + SOCl_2 \longrightarrow C_6H_5COCl + SO_2 + HCl$ 

58. Among acetic acid, phenol and n-hexanol, only CH<sub>3</sub>COOH reacts with NaHCO3 to evolve CO2 gas.

 $CH_3COOH + NaHCO_3 \longrightarrow CH_3COONa + CO_2 + H_2O$ 

59. (d) Methyl acetate and ethyl acetate on hydrolysis give CH<sub>3</sub>COOH which is a liquid. Similarly, ethyl formate on hydrolysis will give formic acid which is also a liquid. Only ethyl benzoate on hydrolysis will give benzoic acid which is a solid.

60. (d) 
$$R \text{ COCl} + \text{AlCl}_3 \longrightarrow R - C^{\oplus} + \text{AlCl}_4^{-}$$
Electrophile

61. (b) Esters having  $\alpha$ -hydrogen atom show Claisen condensation reaction. We know that ethyl benzoate  $(C_6H_5COOC_2H_5)$  does not contain  $\alpha$ -hydrogen. Therefore, C<sub>6</sub>H<sub>5</sub>COOC<sub>2</sub>H<sub>5</sub> does not undergo Claisen self condensation.

2-Hydroxy propanoic acid

(As it has a chiral C-atom thus it is optically active)

63. (c) 
$$CH_3CH_2COOH \xrightarrow{Cl_2} CH_3CHCICOOH$$

$$\frac{\text{alc.KOH}}{-\text{HCl}} \rightarrow \text{CH}_2 = \text{CHCOOH}$$
Acrylic acid

(a) LiAlH<sub>4</sub> can reduce COOH group, and not the double bond.

$$CH_2 = CH - COOH \xrightarrow{LiAlH_4} CH_2 = CH - CH_2OH$$

- There is no reaction hence the resultant mixture **65.** contains CH<sub>3</sub> COOC<sub>2</sub>H<sub>5</sub> + NaCl.

$$CH_{3} - C - Br \xrightarrow{(i)CH_{3}MgI} CH_{3} - CH_{3} - CH_{3} - CH_{3}$$

$$CH_{3} - C - OH$$

$$CH_{3} - C - O$$

- 67. (b) 68. (b)
- 69. (a) Use of catalyst 3 brings about cleavage of C − N bond of amides and will yield amines and alcohols.

O
$$\parallel$$
 $H_3C-C-NH-CH_3 \xrightarrow{\text{catalyst 3}} \longrightarrow N$ 
N-methylethanamide (C-N bond cleavage)

- 70. (c)
- 71. (a) Primary alcohols on oxidation give carboxylic acids as the final product, of course through aldehydes. Oppenauer oxidation involves oxidation of 2° alcohols to ketones, and not for the oxidation of 1° alcohols.
- 72. (c)
- 73. (c) Iodoform test is given by compounds which have CH<sub>3</sub>CO group.

$$O \\ CH_3 - CH_2 - CH_2 - C - CH_3$$

2-Pentanone

3-Pentanone

∵ 2-pentanone has CH<sub>3</sub>CO group, so it gives iodoform test, while 3-pentanone does not have CH<sub>3</sub>CO group, so it does not give iodoform test.

74. (b) 
$$C-CH_3$$
  $CH_2-CH_3$   $CH_2-CH_3$  Phenyl methyl Ethyl benzene

This reaction is known as Clemmensen's reduction.

- 75. (b) Secondary alcohol on oxidation with alkaline KMnO<sub>4</sub> produces corresponding Ketone.
- 76. (b) Carbonyl compounds shows nucleophilic addition reaction with NaHSO<sub>3</sub> and form white crystalline solid which is on hydrolysis with dil. HCl gives original carbonyl compounds and thus, this property is used for the purification of carbonyl compounds as well as for their separation.
- 77. (c) Benedict's solution (Cu<sup>+2</sup> + citric acid + base) and Fehling's solution (Cu<sup>+2</sup> + tartaric acid + base) are weaker oxidising agent than Tollen's reagent, therefore, they oxidise only aliphatic aldehydes, but not aromatic aldehydes.

$$C_6H_5$$
 — CHO+ $[Ag(NH_3)_2]^+$  —  $C_6H_5COO^-$  +  $Ag(s)$   
Tollen's reagent

- 78. (a)
- 79. (d) In p-chlorobenzoic acid, Cl shows +R and -I effect and due to +R effect, the acidic character of p-chlorobenzoic acid decreases. At meta-position, only inductive effect is considered, therefore, m-chlorobenzoic acid will be more acidic due to -I effect.
- 80. (d) RCOOH+NaHCO<sub>3</sub> → RCOONa+CO<sub>2</sub>+H<sub>2</sub>O
  In this reaction, CO<sub>2</sub> is evolved from NaHCO<sub>3</sub>. Carboxylic acid is a stronger acid than carbonic acid.
- 81. (a) In carboxylate ion, electron cloud is equally delocalised between two oxygen atoms, therefore, each oxygen atom has least charge density, hence carboxylate ion becomes more stable. Phenoxide ion has only one oxygen atom which has higher electron density, hence the ion is less stable.
- 82. (b) Compounds containing —CHO group are easily oxidised to corresponding carboxylic acids. Correct reason is due to electron withdrawing nature of C = O group, C H bond in aldehydes is weak and easily oxidised to the corresponding carboxylic acids even with mild oxidising agent like Fehling's solution and Tollen's reagents.
- 83. (d) The α-hydrogen atom in carbonyl compounds is acidic in nature due to presence of electron withdrawing carbonyl group.
- 84. (d) Aldehydes but not ketones react with Tollen's reagent to form silver mirror.
- 85. (d) KMnO<sub>4</sub> is an oxidising agent and acetylene on treatment with alkaline KMnO<sub>4</sub> produces oxalic acid.

$$\begin{array}{c}
\text{CH} \\
\parallel \\
\text{CH}
\end{array} + 4[O] \xrightarrow{\text{Alk. KMnO}_4} \begin{array}{c}
\text{COOH} \\
\text{COOH}
\end{array}$$

- 86. (b) Aldehydes which do not contain α-hydrogen undergo cannizzaro reaction.
- 87. (c) Isobutanal does not give iodoform test as it does not have -COCH, group.
- 88. (a) (A) Cinnamaldehyde:

3-Phenylprop-2-enal

(B) Acetophenoneone

1-Phenylethan one

(C) Valeraldehyde

$$^{5}$$
  $^{4}$   $^{3}$   $^{2}$   $^{1}$   $^{1}$   $^{2}$   $^{1}$   $^{2}$   $^{2}$   $^{1}$   $^{2}$ 

Pentanal

(D) Acrolein

$$CH_2 = CH - CHO$$

Prop-2-enal

(E) Mesityl oxide

$$H_3^{5} - C = CHCOCH_3$$
 $CH_3$ 

4-Methyl pent-3-en-2-one

89. (c)

90. (d)

91. Ketones and aldehyde.

Note:- Benzaldehyde do not give reaction with Fehling solution.

- 92. Esters
- 93. Butanoic acid; Under moist and warm conditions, glycerides present in butter are hydrolysed to give butanoic acid (along with other volatile acids like caproic and caprylic acids) which produces disagreable smell; this reaction is called rancidity.
- **94.** Conc. H<sub>2</sub>SO<sub>4</sub>.
- **95.** Pd/BaSO<sub>4</sub>
- 96. False; Claisen condensation: Intermolecular condensation of esters containing  $\alpha$ -hydrogen atom in presence of strong base to form  $\beta$ -keto ester

C<sub>2</sub>H<sub>5</sub>ONa 
$$\downarrow$$
 CH<sub>3</sub>C. CH<sub>2</sub>COOC<sub>2</sub>H<sub>5</sub>+ C<sub>2</sub>H<sub>5</sub>OH Ethyl acetoacetate ( $\beta$ -Keto ester)

$$\begin{array}{c} & & & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ & \\ & & \\ & \\ & & \\$$

98. (a) False; Aspirin (acetylsalicylic acid) is a non-narcotic analgesic.

99. True; 
$$CH_3 - C - CH_3 \Longrightarrow CH_3 - C = CH_2$$

keto form

enolic form

No. of  $\sigma$  bonds in enolic form: 3+1+1+1+1+2=9

No. of  $\pi$  bonds in enolic form : 1

No. of lone pairs of electrons in enolic form = 2

**100.** True