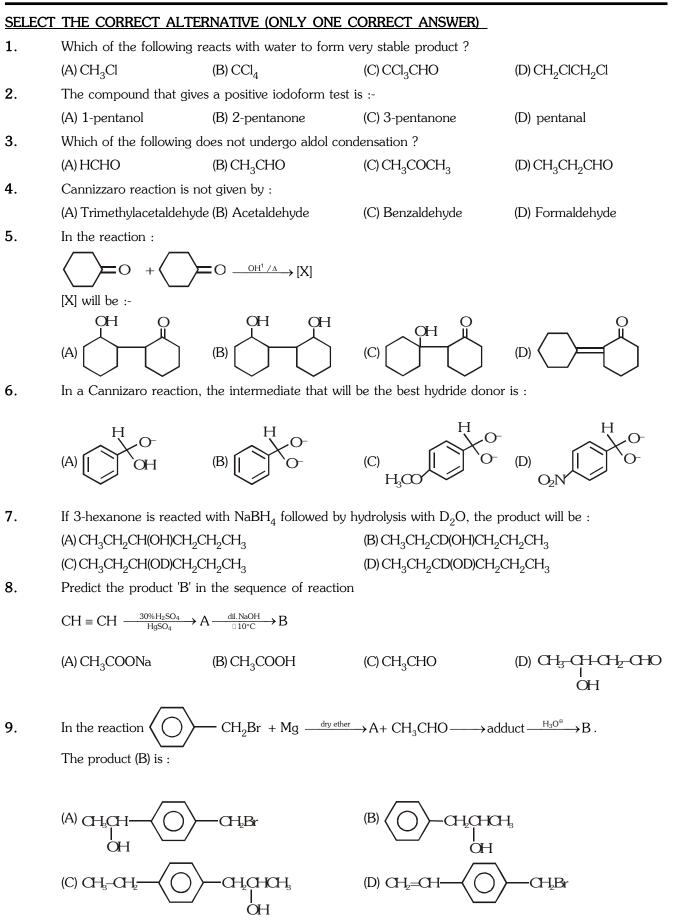
EXERCISE-01

CHECK YOUR GRASP



10. Which one of the following reagents is suitable for the conversion of 2-cyclohexenone into 3-methylcyclohexanone ?

(A) CH_3MgI (B) $(CH_3)_2CuLi$ (C) CH_3

C)
$$CH_3I_2$$
 and Zn (D) CH_3AlCl_2

11. Acetaldehyde reacts with NaOH to form :-

$$(A) \ CH_{3}-CH_{2}-C$$

12. Benzaldehyde reacts with formaldehyde in the presence of alkali to form :-

- (A) Methyl alcohol and sodium benzoate
- (B) Benzyl alcohol and sodium formate

(B) RCHO, RCH(OH)CN, RCH(OH)COOH

- (C) Benzoic acid and ethanol (D) Formic acid and benzyl alcohol
- **13.** The compounds A, B and C in the reaction sequence

$$\xrightarrow{\text{CH}_3} C = O \xrightarrow{I_2} A_{\text{lkali}} \xrightarrow{A_g} B \xrightarrow{\text{dil} H_2 SO_4} Hg^{+2} C$$

are given by the set :-

- (A) Iodoform, ethylene, ethyl alcohol (B) Iodoform, acetylene, acetaldehyde
- (C) Iodoform, propyne, acetone (D) Iodoform, 2-propanol, propanone

14. In the reaction sequence

$$\text{RCOCl} + \text{H}_2 \xrightarrow{\text{Pd}+\text{BaSO}_4} A \xrightarrow{\text{HCN}} B \xrightarrow{\text{H}_3\text{O}^{\oplus}} C$$

A,B and C are given by the set :-

(A) RCHO, RCH(OH)CN, RCH(OH)CH₂NH₂

- 15.
 Oxidation of 2-methyl propane–1,2-diol with periodic acid gives :

 (A) Propionic acid and formaldehyde
 (B) Acetone and formaldehyde

 (C) Acetone and acetic acid
 (D) Acetone and propionic acid
- 16. A carbonyl compound gives a positive iodoform test but does not reduce Tollen's reagent or Fehling's solution. It forms a cyanohydrin with HCN, which on hydrolysis gives a hydroxy acid with a methyl side chain. The compound is :-
 - (A) Acetaldehyde (B) Propionaldehyde (C) Acetone (D) Crotonaldehyde
- 17. A carbonyl compound 'A' reacts with hydrogen cyanide to form a cyanohydrin 'B' which on hydrolysis gives an optically active alpha hydroxy acid 'C'. 'A' gives a positive iodoform test 'A', 'B' and 'C' are given by the set :-

(A) HCHO;
$$\stackrel{H}{H} \subset \stackrel{OH}{ON}$$
; $\stackrel{H}{H} \subset \stackrel{OH}{COOH}$
(B) CH₃CHO; $\stackrel{OH}{H} \subset \stackrel{OH}{ON}$; $\stackrel{OH}{H} \subset \stackrel{OH}{COOH}$

(C)
$$CH_3CH_2CHO$$
; C_2H_5 C_2OH ; C_2H_5 C_2OH ; H C_2OH

$$(D) \xrightarrow{CH_3} C = O; \xrightarrow{CH_3} C \xleftarrow{OH} ; \xrightarrow{CH_3} C \xleftarrow{OH} C \xrightarrow{OH} C \xrightarrow{O} C \xrightarrow{OH} C \xrightarrow{OH} C \xrightarrow{O} C \xrightarrow{OH} C \xrightarrow{OH} C \xrightarrow{OH} C$$

18. In which of the following reactions aldehydes and ketones are distinguished :

- (A) Reaction with phenyl hydrazine (B) Reaction with hydroxylamine
- (C) Reaction with semicarbazide (D) Reaction with silver nirate mixed with ammonia

19. Cyanohydrin of the following compound on hydrolysis gives optically active product :

(A) HCHO (B) CH_3CHO (C) CH_3COCH_3 (D) All the above

20. The major organic product formed from the following reaction is :-

$$\underbrace{\begin{array}{c} O \\ \hline (i) CH_3NH_2 \end{array}}_{(ii) LiAlH_4 (iii)H_2O} \xrightarrow{(i) CH_3NH_2}$$

 $(A) \xrightarrow{\text{NHCH}_3} (B) \xrightarrow{\text{NHCH}_3} (C) \xrightarrow{\text{NHCH}_3} (D) \xrightarrow{\text{ONHCH}_3} (D) \xrightarrow{\text{ONHCH$

- **21.** Which one of the following on treatment with 50% aq. NaOH yields the corresponding alcohol and acid (A) C_6H_5CHO (B) $CH_3CH_2CH_2CHO$ (C) CH_3COCH_3 (D) $C_6H_5CH_2CHO$
- **22.** Which compound gives reaction with 2,4-dinitro phenyl hydrazine?
 - (A) $\begin{array}{c} CH_{3} \\ CH_{3} \end{array} C = O$ (B) $\begin{array}{c} CH_{3} \\ H \end{array} C = O$ (C) $\begin{array}{c} H \\ H \end{array} C = O$ (D) All of these
- 23. Which of the following reaction leads to the formation of secondary alcohol ?

24. Which compound is unable to react with $NaHSO_3$?

$$\begin{array}{cccc} \text{(A) } \text{CH}_3\text{CHO} & \text{(B) } \text{CH}_3\text{-}\text{CH}\text{-}\text{OCH}_3 & \text{(C) } \text{CH}_3\text{-}\text{C=O} \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ \end{array}$$

(D) HCHO

(D) Methyl iodide

25.Which of the following compounds gives a ketone with grignard reagent(A) formaldehyde(B) Ethyl alcohol(C) Methyl cyanide

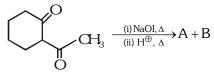
CHECK YOUR GRASP					A	ANSWER KEY			ΞΥ	EXERCISE -1										
Que.	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20
Ans.	С	В	А	В	D	С	С	D	В	В	В	В	В	В	В	С	В	D	В	В
Que.	21	22	23	24	25				_		-									
Ans.	А	D	В	В	С															

EXERCISE-02

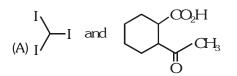
BRAIN TEASERS

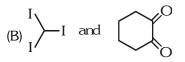
SELECT THE CORRECT ALTERNATIVES (ONE OR MORE THEN ONE CORRECT ANSWERS)

1. End products of the following sequence of reaction is



A & B are :







2. Which of the following compound on treatment with LiAlH_4 will give a product that will give positive iodoform test?

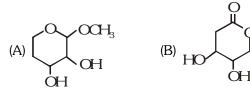
(A) CH_3CH_2CHO (B) $CH_3CH_2CO_2CH_3$

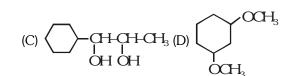
Among the given compounds, the most susceptible to nucleophilic attack at the carbonyl group is
(A) CH₃COCI
(B) CH₃CHO
(C) CH₃COOCH₃
(D) CH₃COOCOCH₃

4. Which of the following does not give iodoform reaction ?

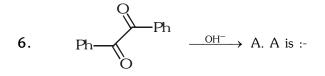
(A) $CH_3CH_2CH_2OH$ (B) CH_3OH

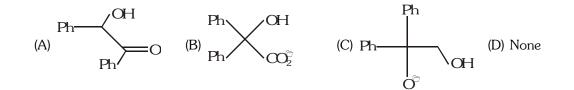
- (C) CH₃CHO
- (D) PhCOCH₃
- 5. Which of the following compounds undergo periodic oxidation





(C) CH₃CH₂OCH₂CH₃ (D) CH₃COCH₃



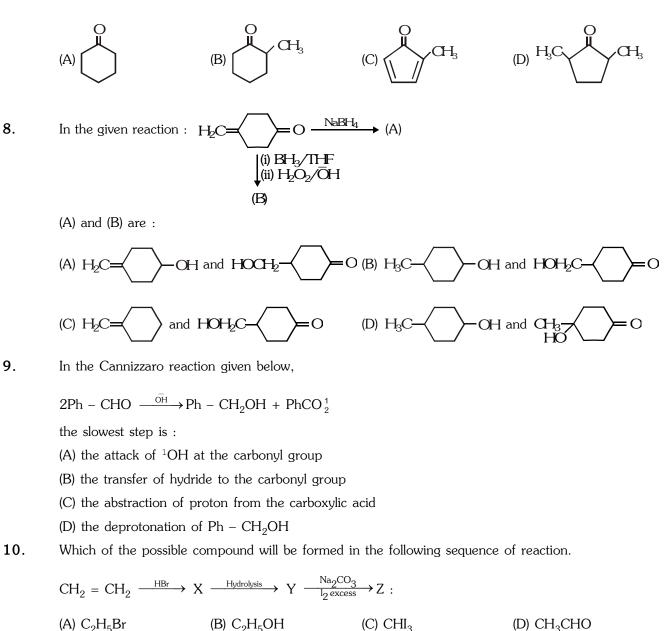


7. In the reaction sequence, [X] is ketone :

$$[X] \xrightarrow{KMnO_4/OH^-/\Delta} HOOQ(CH_2)_3 - CH - COOH$$

[X] will be :-

8.



(A) Benzaldehyde reduces Fehling's solution

(B) $C_6H_5CHO + C_6H_5CHO \longrightarrow C_6H_5CH = CHC_6H_5 + O_2$ is a Claisen-Schmidt reaction. (C) pK_a (formic acid) is less than pK_a (acetic acid)

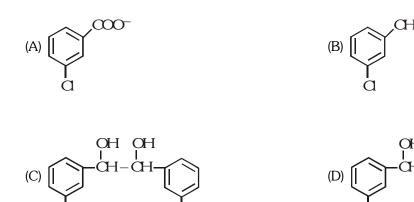
(D)
$$CH_3CCH_3 + CH_3CHO \xrightarrow{NaOH} CH_3 - CH_2CHO$$
 is an example of aldol condensation.

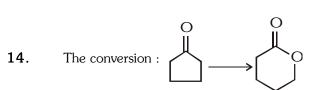
12. $CH_3 - CHO \xrightarrow{\overline{O}H} CH_3CH(OH)CH_2CHO$

In the aldol condensation of acetaldehyde represented above, which of the following intermediate species are obtained ?

OH

13. When m-Chlorobenzaldehyde is treated with 50% KOH solution, the products obtained is (are)

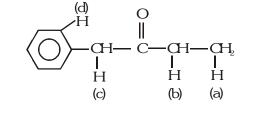




can be effected by using the reagent

(A)
$$H_2O$$
, H_2SO_4 (B) O_2 (C) $C_2H_3 - COOH$ (D) CrO_3 , H_2SO_4

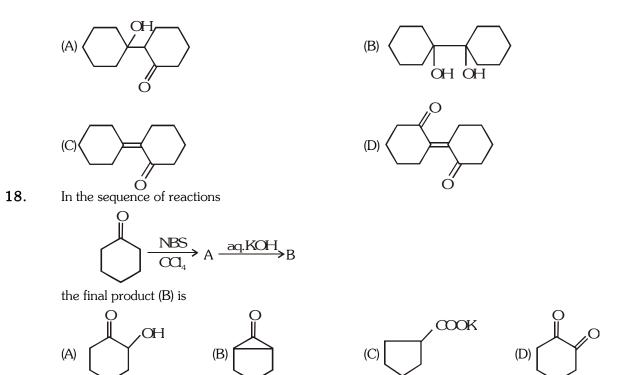
15. Which of the following hydrogens will be the most acidic?



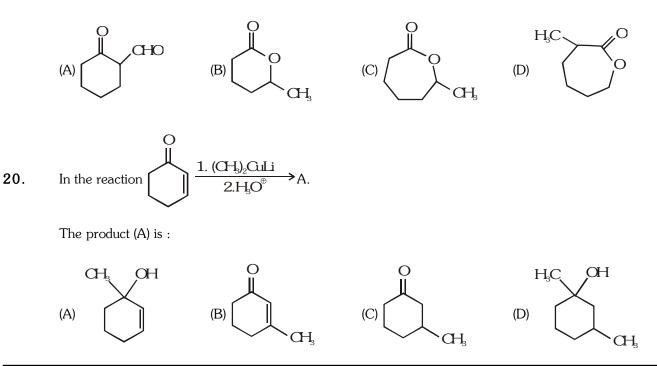
(A) a (B) b (C) c (D) d

16. An organic compound with the molecular formula C_3H_6O does not respond positively to the silver mirror test with Tollens reagent but produces an oxime. The compound is

(A) $CH_2 = CHCH_2OH$ (B) CH_3CH_2CHO (C) $CH_2 = CHOCH_3$ (D) CH_3COCH_3



19. 2- Methylcyclohexanone is allowed to react with metachloroperoxobenzoic acid. The major product formed in the reaction is



BRAIN TEASERS							ANSWER KEY				EXERCISE -2				
Que.	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
Ans.	D	D	А	A,B	A,B,C	В	В	А	В	B,C,D	C,D	A,B,C	A,B	С	С
Que.	16	17	18	19	20										
Ans.	D	С	С	С	С										

EXERCISE-03

TRUE OR FALSE :

- 1. The reaction of methyl magnesium bromide with acetone followed by hydrolysis gives secondary alcohol.
- 2. Aldehydes are more reactive than ketones.
- 3. The yield of ketone, when a secondary alcohol is oxidised, is more than the yield of aldehyde when primary alcohol is oxidised by $K_2Cr_2O_7/H^{\oplus}$
- 4. Both aldehydes and ketones reduce Tollen's reagent.
- 5. Aldol condensation is given by all carbonyl compounds.
- 6. Acetaldehyde and acetone can be distinguished by iodoform test.
- 7. LiAlH_4 converts ketones into secondary alcohols.
- 8. Methanol can be distinguished from ethanol by haloform reaction.
- 9. Propanone does not show tautomerism.
- 10. Ketones restore pink colour of Schiff's reagent.

FILL IN THE BLANKS :

- 1. To prepare ethanol, CH_3MgI is treated with the other reagent
- 2. Urotropine is formed by the action of with
- 3. The conversion of acid chlorides into aldehydes by reduction is termed
- 4. Aldehyde show reducing properties due to their ready conversion into
- 5. Hydrazone of an aldehyde when heated with sodium ethoxide forms This is known as reaction.
- 6. Cannizzaro's reaction is followed by those aldehydes which $\dots \alpha$ -hydrogen atom.
- 7. Two separate solutions, Fehling's solution A(.....) and Fehling's solution B(NaOH +.....) are at first mixed up together and is then heated with the aldehyde. A precipitate is formed.
- 8. Tollen's reagent gives with acetaldehyde.
- 9. Aldehydes have boiling points lower than those of and higher than those of of comparable molecular masses.
- 10. When calcium acetate is distilled alone is formed.

MATCH THE COLUMN

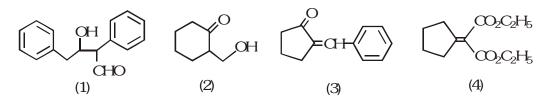
1. Match the compounds/ions in Column I with their properties/reactions is Column. II

\square	Column -I	Colu	mn -II
(A)	C ₆ H ₅ CHO	(p) gives	precipitate with
		2,4di	initrophenylhydrazine
(B)	$CH_3C \equiv CH$	(q) give	preciptitate with AgNO ₃
(C)	CN-	(r) is a r	nucleophile
(D)	I-	(s) is inv	olved in cyanohydrin formation

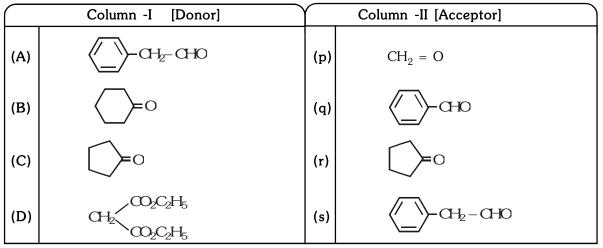
2. Match the column I with column II.

	Column-I	γ	Column-II
(A)	$\begin{array}{c} RMgX + HCHO \longrightarrow \\ n-carbon & H_{3}O^{\oplus} \end{array} $	(p)	Ketone
(B)	$\underset{n-\text{carbon}}{\text{RMgX}} + (CH_2)_2 O \longrightarrow$	(q)	1ºAlcohol (n + 1) carbon
(C)	Adduct $\xrightarrow{H_3O^{\oplus}}$ $\underset{n-carbon}{RMgX} + CO_2 \longrightarrow$	(r)	Acid (n + 1) carbon
(D)	Adduct $\xrightarrow{H_3O^{\oplus}}$ $\underset{n-\text{carbon}}{\text{RMgX}}$ + Ph-C=N \longrightarrow	(s)	1ºAlcohol (n + 2) carbon
	Adduct $\xrightarrow{H_3O^{\oplus}}$		

3. Aldol dondensation proceeds by carbon-carbon bond fromation between an enolate donor and a carbonyl acceptor. For each of the following aldol products (1 through 4)



match the donor and acceptor compound.



ASSERTION & REASON QUESTION :

These questions contains, Statement-I (assertion) and Statement-II (reason).

- (A) Statement-I is True, Statement-II is True ; Statement-II is a correct explanation for Statement-I
- (B) Statement-I is True, Statement-II is True ; Statement-II is NOT a correct explanation for Statement-I
- (C) Statement-I is True, Statement-II is False.
- (D) Statement-I is False, Statement-II is True.
- 1. Statement-I : 2, 2,-dimethyl propanol undergoes Cannizaro reaction with concentrated NaOH. Because

Statement-II : Cannizarro reaction is a disproportionation reaction.

2.	Statement-I : Benzaldehyde does not undergoes aldol condensation.
	Because
	Statement-II : Benzaldehyde does not contains acidic α -Hydrogen.
3.	Statement-I : Acetaldehyde is less reactive than trichloro acetaldehyde.
	Because
	Statement-II : Chlorine atom exhibit –I effect in trichloro acetaldehyde.
4.	Statement-I : Benzaldehyde gives a positive test with Benedict's and Fehling's solution.
	Because
	Statement-II : Benzaldehyde gives silver mirror with Tollen's reagent.
5.	Statement-I : $R-C=O^+$ is more stable than $R-C^+=O$.
	Because
	Statement-II : Resonance in carbonyl compound provides $\supset C^{\oplus}$ and O^{-} .
6.	Statement-I : Rate of addition of HCN on carbonyl compounds increases in presence of NaCN.
	Because
	Statement-II : Reaction involves the addition of CN ⁻ in rate determining step.
7.	Statement-I : Fehling's solution can be used to distinguish benzaldehyde from acetaldehyde.
	Because
	Statement-II : Both benzaldehyde and acetaldehyde reduces tollen's reagent.
8.	Statement-I : Ketones are less reactive than aldehydes.
	Because
	Statement-II : Ketones do not give Schiff's test.
9.	Statement-I : Benzaldehyde is more reactive than ethanol towards nucleophilic attack.
	Because
	Statement-II : The overall effect of $-I$ and $+R$ effect of phenyl group decreases the electron density on
	the carbon atom of $>C = O$ group in benzaldehydes.
10.	Statement-I : In formaldehyde all the four atoms are in same plane.
	Because
	Statement-II : The carbon atom in formaldehyde is sp^2 hybridised.

COMPREHENSION BASED QUESTIONS :

Comprehension # 1

Aldehyde, ketone, acid and acid derivatives contain >C = O group. Aldehyde and ketones gives nucleophilic addition reactions where as acid and acid derivatives gives nucleophilic addition followed by elimination reactions. Nucleophilic addition reactions followed by elimination of acid derivatives is known as acyl substitution reaction. This substitution reaction takes places by formation of tetrahedral intermediate.

1. For the given reaction

$$\bigwedge_{R}^{\ominus} L + \bigwedge_{Nu}^{\ominus} \bigvee_{R}^{O} Nu + \overset{\ominus}{L}$$

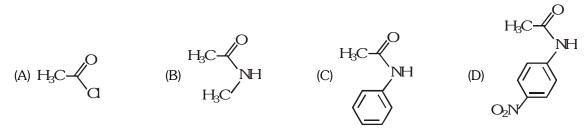
which of these is correct ?

- (A) L must be better leaving group than Nu
- (B) $\mathrm{Nu^{-}}$ must be strong enough nucleophile to attack carbonyl carbon
- (C) Carbonyl carbon must be enough electrophilic to react with Nu-
- (D) All of these

2. Which of the following compounds has very poor leaving group?



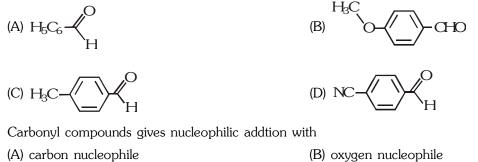
3. Which one of the following is least reactive compound for nucleophilic acyl substitution.



Comprehension # 2

Aldehydes and ketones are amphoteric. Thus they can act both as acids and bases. Under acidic conditions, the carbon of the protonated carbonyl group is much more electrophilic, reacting even with weak nucleophilie. Carbonyl compound gives nucleophilic addition reaction. In this reaction the nucleophilic attack proceedes the electrophilic attack.

- 1. Which of these statements are correct ?
 - (A) Carbonyl compound is amphoteric in character
 - (B) Acid catalyst makes the carbonyl carbon more electrophilic
 - (C) basic catalyst makes the nucleophilic attack more faster.
 - (D) All of these
- 2. Which of the statements are/is correct?
 - (A) The rate determining step of addition reaction is the addition of nucleophile
 - (B) The rate-determining step is addition of electrophile
 - (C) The reaction intermeidate of the reaction is alkoxide ion
 - (D) both (A) and (C)
- 3. Which one of the carbonyl compounds is more reactive towards NaCN/H⁺?



4.

- - (C) Nitrogen nucleophile

(D) All of these

Comprehension # 3

Aldehyde and ketones are specially susceptible to nucleophile addition because carbonyl group $\sum -O$ is polar (due to electronegativity different between carbon and oxygen).

>°+ °-C=O

Positive charge on carbon makes it reactive towards the nucleophile. This addition is catalysed by acid.

Reactivity of carbonyl compound towards nucleophilic addition increases with increase in the electron deficiency at carbonyl carbon. Thus, (-I.E.) group increase while (+I.E.) groups decrease the reactivity of carbonyl compound.

(D) all of these

(D) H C=O

- 1. Which of the following is most reactive to give nucleophilic addition?
 - (A) FCH_2CHO (B) $CICH_2CHO$ (C) $BrCH_2CHO$ (D) ICH_2CHO Carbonyl compounds show nucleophilic addition with :
- Carbonyl compounds show nucleophilic addition with :
 (A) HCN
 (B) NaHSO₃
 (C) (CH₃OH + HCl)
- **3**. Which among the following carbonyl compounds is most polar?

$$(A) \xrightarrow{C_2H_3} C=O \qquad (B) \xrightarrow{CH_3} C=O \qquad (C) \xrightarrow{CH_3} C=O$$

4. Select the least reactive carbonyl compound for nucleophilic addition:

5. Which among the following isomeric compound is most reactive?

(A)
$$CH_3 - CH_2 - CH_2 - CH_2 - CH_3$$

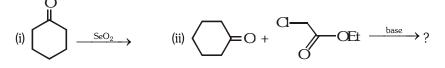
(B) $CH_3 - CH_2 - CH_2 - CH_3$
(C) $CH_3 - CH_2 - CH_2 - CH_3$
(D) All are equally reactive

MIS	SCELLANEOU	s type que	STION	AN	SWER KEY	<u>ľ</u>		EXERCISE -3
•	<u>True / Fa</u>	<u>lse</u>						
	1 . F	2. T	3. T	4. F	5 . F			
	6. F	7. T	8. T	9. F	10. F			
•	<u>Fill in the</u>	e Blanks						
	 HCHO acids CuSO₄, n acetone 	roschell salt, e	Cu ₂ O	5 . alkaı	HO, NH ₃ ne, wolf-kishne r mirror	r 6. do :	senmund reduction not have bhol, alkane	n
•	<u>Match the</u>	<u>e Column</u>						
	1. (A) \rightarrow p,s 3. (A) \rightarrow s ;	,	,	,	2. (A) →	$q; B \rightarrow s$	$; (C) \rightarrow r ; (D) \rightarrow j$	р
•	Assertion	- Reason	Question	<u>ns</u>				
	1 . D	2.	А	3 . A	4. D	5 . B		
	6 . A	7.	В	8. B	9 . A	10 . A		
•	<u>Comprehe</u>	ension Ba	sed Que	stions				
		nsion #2 :	1 . (D)	2. (A) 2. (D) 2. (D)		4. (D) 4. (A)	5. (A)	

EXERCISE-04 [A]

CONCEPTUAL SUBJECTIVE EXERCISE

- 1. Which of the carbonyl groups in p-MeOC₆H₄COMe and p-NO₂C₆H₄COMe protonates more readily in acid solution and why ?
- **2**. Identify (A) and (B) in the given sequence of reaction PhCH₂CHO $\xrightarrow{\text{SeO}_2}$ (A) $\xrightarrow{(i)CD}$ (B)
- **3.** (a) Distinguish between PhCOEt and p-MeC₆H₄COMe by a chemical method.
 - (b) Arrage the following in the order of increasing reactivity towards nucleophilic addition reactions.
 (i) CH₃CHO, C₆H₅COC₆H₅, CH₃COC₆H₅, CH₃COCH₃
 (ii) CH₃CHO, CF₃CHO, CH₂ = CHCHO
- 4. Compound X with molecular formula $C_9H_{10}O$ form a semicarbazone and give negative Tollen's and Iodoform tests. Upon reduction it gives n-propyl benzene. Deduce the structure of X.
- 5. (a) The double bond in aldehydes and ketones is reactive towards nucleophilic reagents like CN⁻ whereas that in an alkene is not.
 - (b) Alkenes undergo electrophilic addition whereas aldehydes and ketones undergo nucleophilic addition.
- 6. Complete the following equations giving the structures of the major organic product.



7. Complete the following equation :

(a) H₃C
$$\xrightarrow{O}$$
 +HC \equiv CH $\xrightarrow{(i)NaNH_2}$?

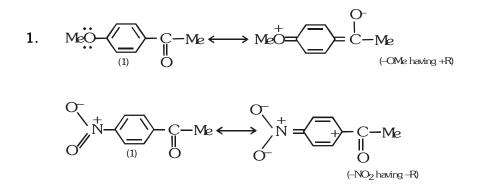
(b)
$$H_3C \xrightarrow{O} + \xrightarrow{O} \xrightarrow{O} \xrightarrow{NaNH_2} \xrightarrow{NaNH_2}$$

- **8**. Two organic compounds (A) and (B) have same empirical formula CH₂O. Vapour density of (B) is twice the vapour density of (A). (A) reduces Fehling solution but does not react with NaHCO₃. Compound (B) neither reacts with NaHCO₃ nor reduces Fehling solution. What are (A) and (B) ? Also report an isomer of (B) if it reacts with NaHCO₃.
- **9.** Complete the following equation :

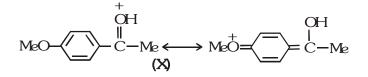
$$HC = CH \xrightarrow{NaNH_2} ? \xrightarrow{CH_3Br} ? \xrightarrow{Hg^{2\oplus}/H^{\oplus}} ? \xrightarrow{H^{\oplus}} H_3C \xrightarrow{OH} H_3C \xrightarrow{I_2} NaOH$$

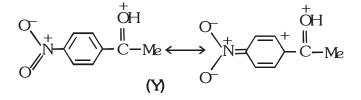
10. A comopound has two isomers (A) and (B) of formula $C_5H_{10}O$. Isomer (A) on treating with NaOH (aq.) give 2, 2-demethylpropan-1-ol and 2, 2-dimethylpropanoic acid salt. The isomer (B) on treating with NaOH (aq.) gives 3-hydroxy-2-propylheptanal. What are A and B ?

CONCEPTUAL SUBJECTIVE EXERCISE



I will be more readily protonated than (II). Alternative protoned (I) is more stabilised by resonance than protonated (II).





In (X) there is extended conjugation and only one charge is involved. In (Y) there is not this extended conjugation and the relative close proximity of two positive charges is a destabilising factor. Hence (X) is more stable than (Y).

2. (A)
$$\therefore$$
 C₈H₆O₂ = C₆H₅ - C-CHO ; (B) C₈H₈O₃ = C₆H₅-CHOHCO₂H

3. (a) I does not undergo haloform reaction whereas II does.

(b)
$$A = C_6H_5COC_6H_5 < CH_3COC_6H_5 < CH_3COCH_3 < CH_3CHO$$

 $B = CH_2=CHCHO < CH_3CHO < CF_3CHO$

- 4. (i) X from semicarbazone and thus possesses carbonyl group. $>C=O + H_2N.NHCONH_2 \longrightarrow >C=N.NHCONH_2$
 - (ii) It does not given Tollen's reagent test and thus it is ketone.
 - (iii) It does not given iodiform test and thus it is not methyl ketone.
 - (iv) Keepting in view of the above facts and molecular formula ; X is

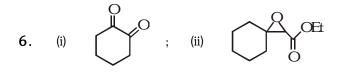
(v) This on reduction will given n-propyl benzene.

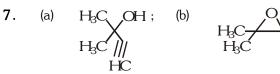
5. (a) The carbonyl group in aldehydes and ketones add on CN⁻ resulting in the fromation of an anion where the negative charge resides on oxygen. However if a nucleophile adds on to an alkene the negative charge resides on carbon. Since carbon is much less strongly electron attracting than oxygen, this species is less stable and hence not readily formed.



(b) In alkenes the double bond joins two carbon atoms and there is no resultant polarity. In carbonyl compounds, the carbonyl group is highly polar and the high partial positive charge on the C atom makes it subsceptible to nucleophilic attack.

 C_2H_5





CHO

8.

- A = HCHO (Formaldehyde) $B = HCOOCH_3$ (Methyl methanoate)
- 9. $HC = C^{-}Na^{+}$; $H_{3}C C = CH$; $H_{3}C C = CH_{3}$; $H_{3}C CH_{3}$; $H_{$

EXERCISE-04 [B]

BRAIN STORMING SUBJECTIVE EXERCISE

1. Identify (A) to (D) as reactant, reagent, product as name of the reaction in following :

(i) $3(CH_3)_2C=O \xrightarrow{HCl} (A)$ Aldol condensation

(ii) $CH_3COCI + H_2 \xrightarrow{(B)} (C)$ Rosenmund's reaction

(iii) (D) $\xrightarrow{(i) \operatorname{NH}_2\operatorname{NH}_2}{(ii) \operatorname{C}_2\operatorname{H}_5\operatorname{ONa}} \rightarrow \operatorname{CH}_3\operatorname{CH}_2\operatorname{CH}_3$

(iv) (D) $\xrightarrow{1}{OBr} CHBr_3 + CH_3COO^1$

- **2.** Two different Grignard reagents, (X) and (Y) produce $C_6H_5CH_2C(CH_3)_2OH$ on reaction with (P) and (Q) respectively. Give structures of (X), (Y) and (Q).
- 3. (a) 2-methyl-1,3-cyclohexanedione is more acidic than cyclohexanone explain with reason
 - (b) Explain why HCN will add to the double bond in CH2=CHCOOH but not in RCH=CHR
- 4. A compound $C_5H_{10}O$ does not reduce Fehling's solution, form a phenyl hydrazone, shows the haloform reaction, and can be converted into n-pentane by Zn—Hg and conc. HCl. What is this comopound ?
- 5. (a) Convert

PhCHO into PhCH = CHCOPh

(b) Identify A, B, C and D in the following reaction.

$$\begin{array}{cccc} C_7H_6O & \xrightarrow{Alc.} & C_{14}H_{12}O_2 \\ [A] & [B] \\ & & CH_3COONa \\ & & Acetic anhydride \end{array} \xrightarrow{C_9H_8O_2} \xrightarrow{Br_2} C_9H_8O_2Br_2 \\ [C] & [D] \end{array}$$

- **6**. A ketone (A) which undergoes haloform reaction gives compound B on reduction. B on heating with sulphuric acid gives compound C, which form mono ozonide D. D on hydrolysis in presence of Zn dust gives only acetaldehyde. Identify A, B and C. Write down the reactions involved.
- **7.** Complete the following reactions :

Write the structures of A to D and give the IUPAC name of (D).

(ii)
$$H_3C \xrightarrow{O} CH_3 \xrightarrow{NaOH} (A)$$

8. The sodium salt of a carboxylic acid, A, was produced by passing a gas, B, into an aqueous solution of caustic alkali at an elevated temperature and pressure. A on heating in presence of sodium hydroxide followed by treatment with sulphuric acid gave a dibasic acid C. A sameple of 0.4 g of C on combustion gave 0.08 g of water and 0.39 g of carbondioxide. The silver salt of acid weighin 1g on ignition yielded 0.71 g of silver as residue. Identify A, B and C.

9. Give reasons for the following :

BRAIN STORMING SUBJECTIVE EXERCISE

- (a) Ketones are less electrophilic than aldehydes.
- (b) Aldehydes are reducing agents and ketones are not.
- (c) HBr fails to give addition products with carbonyl compounds.
- **10.** Identify A, B and C?

$$(A) \xrightarrow{\text{NaBH}_4} (B) \xrightarrow{\text{dil.OH}^1} (C)$$

1. (i)
$$3 \xrightarrow{H_3C} 0 \xrightarrow{dryHCl} H_3C \xrightarrow{H_3C} 0 \xrightarrow{dryHCl} H_3C \xrightarrow{H_3C} 0 \xrightarrow{dryHCl} H_3C \xrightarrow{H_3C} 0 \xrightarrow{H_3C} 0$$

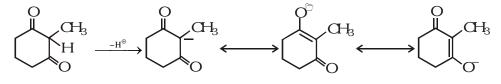
ANSWER

KEY

EXERCISE -4(B)

2. If X is Ph—CH₂MgX, then P is $CH_3 - C - CH_3$ and if Y is CH₃-MgX, then Q is Ph \mathcal{H}_3 CH_3

3. (a) 2-methyl-1, 3-cyclohexanedione is more acidic because its enolate ion is stabilized by an additional resanance structure.



Resonance stabilized conjugate base.

(b) In the case of the acid the double bond is activated due to the presence of a -COOH group

$$H_{2}C = CH - C - OH \leftrightarrow H_{2}C - CH = C - OH \xrightarrow{HCN} NC - CH_{2} - HC = C - OH$$

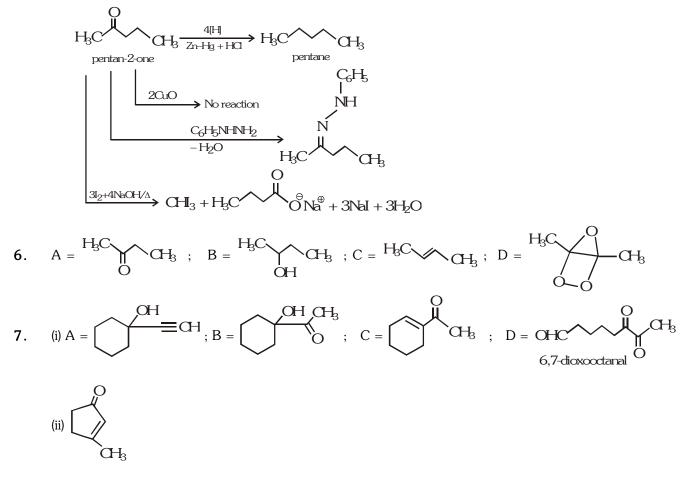
- **4.** (i) Since $C_5H_{10}O$ forms a phenyl hydrazone, hence it is an aldehyde or ketone.
 - (ii) As it does not reduce Fehling's solution, hence it is not aldehyde but a ketone.
 - (iii) Since it undergoes haloform reaction, therefore it is $CH_3 C R$ type of ketone. The value of R can be derived as follows : $R = -C_3H_7$

Hence this ketone is
$$H_3C$$
 H_3C H_3C

(iv) Ketone on Clemmensen reduction yields n-pentane; hence R is n-propyl and not isopropyl.

5. (a)
$$CH_3MgI, H_3O^+, H^+/K_2Cr_2O_7, PhCHO/OH, \Delta$$

(b) $A = C_6H_5CHO$; B = PhCH(OH)COPh; $C = C_6H_5CH=CH$ —COOH; $D = C_6H_5CHBr$ —CHBr—COOH



8.
$$A = HCOOH$$
 ; $B = CO$; $C = COOH$

- **9.** (a) The positive inductive effect of the second alkyl radical reinforces that of the first one decreasing still further the partial positive charge on the carbonyl carbon atom. This reduces the attraction of the atom for nucleophilic reagents. Hence ketones are less electrophilic.
 - (b) The >C=O group in aldehydes activates the H atom attached to the carbonyl group. This is due to the relaying of the –I effect of the oxygen atom to the C—H bond so that partial positive charge is created on the H atom. The result of this activation is that the H atom of the –CHO group can be oxidised readily to a (OH) group. Thus aldehydes are reducers.
 - (c) HBr is strongly polar and is hence readily added to the polarized >C=O group. The addition product

 H_3C Br is however unstable and decomposes to give the original carbonyl compound and HBr.

10. (A) :
$$(B)$$
 : (B) : (B) : (C) : (C) : (C)

H₃C_OH

EXERCISE-05

PREVIOUS YEARS QUESTIONS

1.	The formation of cyanohydrin from a ketone is a	n example of	[IIT-90]				
	(A) Electrophilic addition	(B) Nucleophilic addition					
	(C) Nucleophilic substitution	(D) Electrophilic substitution					
2.	The enolic form of acetone contains -		[IIT-90]				
	(A) 9 sigma bonds, 1 pi bond and 2 lone pairs	(B) 8 sigma bonds, 2 pi bond and 2 lone	e pairs				
	(C) 10 sigma bonds, 1 pi bond and 1 lone pairs	(D) 9 sigma bonds, 2 pi bond and 1 lone pairs					
3.	m—chlorobenzaldehyde on reaction with conc. K	OH at room temperature gives [IIT-91]					
	(A) Potassium m-chlorobenzoate and m-hydroxyb	enzaldehyde					
	(B) m-hydroxy benzaldehyde and m-chlorobenzyl	alcohol					
	(C) m-chlorobenzyl alcohol and m-hydroxybenzyl	alcohol					
	(D) Potassium m-chlorobenzoate and m-chlorobe	nzyl alcohol					
4.	Hydrogenation of benzoyl chloride in the presence	ce of Pd on BaSO ₄ gives	[IIT-92]				
	(A) Benzyl alcohol	(B) Benzaldehyde					
	(C) Benzoic acid	(D) Phenol					
5.	An organic compound C_3H_6O does not give a pr	ecipitate with 2,4-Dinitrophenyl hydrazine re	eagent-				
			[IIT-93]				
	(A) CH ₃ CH ₂ CHO	(B) CH ₃ COCH ₃					
	(C) CH ₂ =CH-CH ₂ OH	(B) CH ₃ COCH ₃ (D) -OH					
6.	Under Wolff Kishner reduction conditions, the co	nversion which may be through about is -	[IIT-95]				
	(A) Benzaldehyde into Benzyl alcohol	(B) Cyclohexanol into Cyclohexane					
	(C) Cyclohexanol into Cyclohexanol	(D) Benzophenone into Diphenylmethar	ne				
7.	In the reaction, P is,		[IIT-95]				
	$\begin{array}{c} CH_3 \\ CH_3 \end{array} \xrightarrow{SeO_2} P + Se + H_2O \end{array}$						
	(A) CH_3COCHO (B) CH_3COOCH_3	(C) CH ₃ COCH ₂ OH (D) None					
8.	In the Cannizzaro reaction given below 2 Ph–CH	$IO \xrightarrow{OH^2} Ph-CH_0OH + PhCO_0^1$ the slow	est step is-				
	(A) the attack of OH^1 at the carbonyl group		[IIT-96]				
	(B) the transfer of hydride to the carbonyl group		[50]				
	(C) the abstraction of proton from the carboxylic	acid					
	(D) the deprotonation of Ph–CH $_2$ OH						
9.	Which of the following will give yellow precipitate	with I _o /NaOH-	[IIT-97]				
	(A) ICH ₂ COCH ₂ CH ₂	$(B) CH_3COOCOCH_3$					
	(C) CH_3CONH_2	(D) $CH_3CH(OH)CH_2CH_3$					
10.	Among the given compounds, the most succepti	0 2 0	up is -				
		. , , , , , , , , , , , , , , , , , , ,	-				
	(A) MeCOCl	(B) MeCHO	[IIT-97]				

	H (A) C,H,-C,-O ^つ	(B) С ₆ H ₅ -С-О [¬]	(C) O	
12.		O^2 $\rightarrow CH_3 - CH = N - OH.$ The		[IIT-97]
12.	(A) $pH = 1$	(B) pH <u>~</u> 4.5	(C) Any value of pH	
13.				give a product containing [IIT-98]
	(A) $C_6H_5NH_2$	(B) (CH ₃) ₃ N	(C) C ₆ H ₅ NHC ₆ H ₅	(D) C ₆ H ₅ NHNH ₂
14.	The product obtained v	via oxymercuration (HgSO4	+H ₂ SO ₄) of 1-butyne wou	ld be - [IIT-98]
	Ö			
	(A) CH_3CH_2 -C-CH ₃		(B) CH ₃ CH ₂ CH ₂ CHO	
	(C) CH ₃ CH ₂ CHO + H	СНО	(D) CH ₃ CH ₂ COOH+H	СООН
15.	Which of the following	will not undergo aldol con	densation -	[IIT-98]
	(A) Acetaldehyde		(B) Propanaldehyde	
	(C) Benzaldehyde		(D) Trideutero acetaldeh	nyde
16.		will react with water to for		[IIT-98]
	(A) CHCl ₃	(B) Cl ₃ CCHO	(C) CCl ₄	(D) $CICH_2CH_2CI$
17.	The enol form of aceto	one, after treatment with D	₂ O gives -	[IIT-99]
	QН	OH	OH	QD
	(A) CH2-C-CH2D	OH (B) CD ₂ =C-CD ₃	(C) CH2=C-CH2D	(D) $CD_2 = C - CD_3$
18.	Which of the following	has the most acidic hydrog	gen-	[IIT-2000]
	(A) 3-hexanone	(B) 2, 4-hexanedione	(C) 2, 5-hexanedione	(D) 2, 3 hexandione
19.	The appropriate reage	nt for the following transfo	rmation -	[IIT-2000]
		OH OH	OH CHICH	3
	(A) Zn (Hg), HCl	(B) NH ₂ NH ₂ ,OH⁻	(C) H ₂ /Ni	(D) NaBH ₄
20.	Read the following stat	ement and explanation and	l answer as per the option	given below : [IIT-01]
	Statment-I : Dimethyls carbonyl compounds.	ulphide is commonly used	for the reduction of an oz	onide of an alkene to get the
	Statment-II : It reduces	the ozonide giving water s	soluble dimethyl sulphoxide	e and excess of it evaporates.
	(A) If both assertion an	d reason are correct and r	eason is the correct evolan	ation of the assertion

- (A) If both assertion and reason are correct, and reason is the correct explanation of the assertion
- (B) If both assertion and reason are correct, but reason is not correct explanation of the assertion
- (C) If assertion is correct but reason is incorrect

(D) If assertion is incorrect but reason is correct

 21.
 A mixture of benzaldehyde and formaldehyde on heating with aqueous NaOH solution gives [IIT-01]

 (A) benzyl alcohol and sodium formate
 (B) sodium benzoate and methyl alcohol

(C) sodium benzoate and sodium formate

(D) benzyl alcohol and methyl alcohol

22. Compound A (molecular formula C_3H_8O) is treated with acidified potassium dichromate to from a product B (molecular formula C_3H_6O). B forms a shining silver mirror on warming with ammonical silver nitrate. B when treated with an aqueous solution of $H_2NCONHNH_2$. HCl and sodium acetate gives a product C. Identify the structure of C-

(A)
$$CH_3CH_2CH=NNHCONH_2$$

$$(B)CH_3-C=NNHCONH_2 \\ | \\ CH_3$$

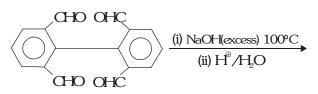
(C)CH3-C=NCONHNH2 | CH3 (D) $CH_3CH_2CH=NCONHNH_2$

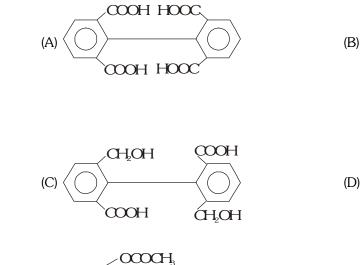
CHOH

CH₂OH

|| O

23. Any one of the product formed is :





Ρ

Q

[IIT-03]

Product formed by P & Q can be differentietod by :

OCOCH.

Acidic

	(A) 2, 4-DNP		(B) Lucas reagent (ZnCl	$_2$ + conc. HCl)
	(C) NaHSO ₃		(D) Fehlings solution	
25.	Which of the reagent i	s used to convert 2-Butanc	one into propanoic acid -	[IIT-05]
	(A) NaOH, I ₂ /H [⊕]	(B) Tollen's reagent	(C) Fehling solution	(D) NaOH, NaI/H⊕

[IIT-02]

CH₂OH

CH_OH

26. CH₃-C-CH₃+ next homologue

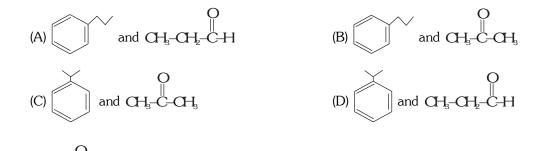
Following statements is/are correct about mixture :

(A) mixture is 3-types of oximes

(C) all are optically active

(B) mixture is 2-types of oximes(D) one is optically active

27.
$$\underbrace{CH_3-CH_2-CH_2-C}_{A|C_3} P \xrightarrow{(i) O_2/\Delta} Q + Phenol, P \& Q \text{ are } -$$
 [IIT-06]

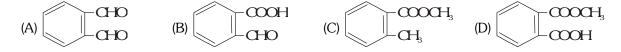


28.

O The above reagent can be synthesise from the following compounds in alkaline medium

following by acidification -

[IIT-06]



- 29. An unknown compound of carbon, hydrogen and oxygen contains 69.77%C and 11.63% H, and has a molecular weight of 86. It does not reduce Fehling solution but forms a bisulphate addition compound and gives a positive iodiform test. What are the possible structures ? [IIT-87]
- 30. Iodoform is obtained by the reaction of acetone with hypoiodite but not with iodide. Why ? [IIT-91]
- 31. An organic compound 'A' on treatment with ethyl alcohol gives a carboxylic acid 'B' and compound 'C'. Hydrolysis of 'C' under acidic conditions gives 'B' and 'D'. Oxidation of 'D' with KMnO₄ also gives 'B'. 'B' on heating with Ca(OH)₂ gives 'E' (C₃H₆O). E does not give Tollen's test and does not reduce Fehling's solution but form a 2,4-dinitrophenyl hydrazone. Identify (A), (B), (C), (D) and (E). [IIT-92]
- **32**. Complete the following :

(i)
$$H_3^{CO} \longrightarrow CHO+ HCHO \longrightarrow (A) + (B)$$
 [IIT-92]

33. Complete the following :

$$C_{6}H_{5} - CHO + CH_{3} - COOC_{2}H_{5} \xrightarrow{NaOC_{2}H_{5} \text{ in absolute}}{C_{2}H_{5}OH \text{ and heat}} (A)$$
[IIT-95]

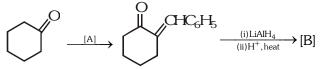
34. Complete the following reaction with appropriate structure :



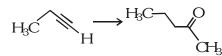
35. Suggest appropriate structures for the missing compounds. (The number of carbon atoms remains the same throughout the reactions). [IIT-96]

$$(A) \xrightarrow{HIO_4} (B) \xrightarrow{OH^-} (C)$$

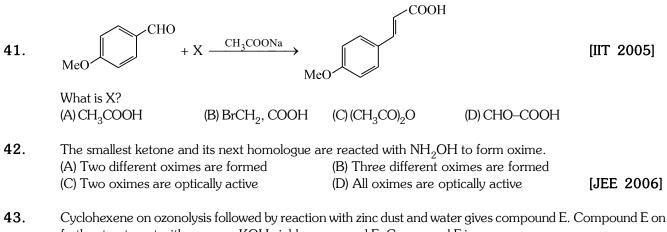
36. Complete the following :



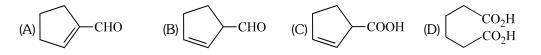
- **37.** Compound 'A' (C_8H_8O) on treatment with NH₂OH.HCl given 'B' and 'C'. 'B' and 'C' rearrange to give 'D' and 'E', respectively, on treatment with acid. B, C, D and E are all isomers of molecular formula (C_8H_9NO). When 'D' is boiled with alcoholic KOH an oil 'F' (C_6H_7N) separates out. 'F' reacts rapidly with CH₃COCl to give back 'D'. On the other hand, H' on boiling with alkali followed by acidification gives a white solid 'G' ($C_7H_6O_2$). Identify 'A' to 'G'.
- **38**. Carry out the following transformation in not more than three steps.



- An organic compound (A), C₆H₁₀O, on reaction with CH₃MgBr followed by acid treatment gives compound (B). The compound (B) on ozonolysis gives compound (C), which in presence of a base give 1-acetyl cyclopentene (D). The compound (B) on reaction with HBr gives compound (E). Write the strucures of (A), (B), (C) and (E). Show how (D) is formed from (C).
- **40.** An organic compound (A) reacts with H_2 to give (B) and (C) successively. On ozonolysis of (A), two aldehydes (D) C_2H_4O and (E) $C_2H_2O_2$ and formed. On ozonolysis of (B) only propanal is formed. What are (A) to (E) ? [IIT-01]



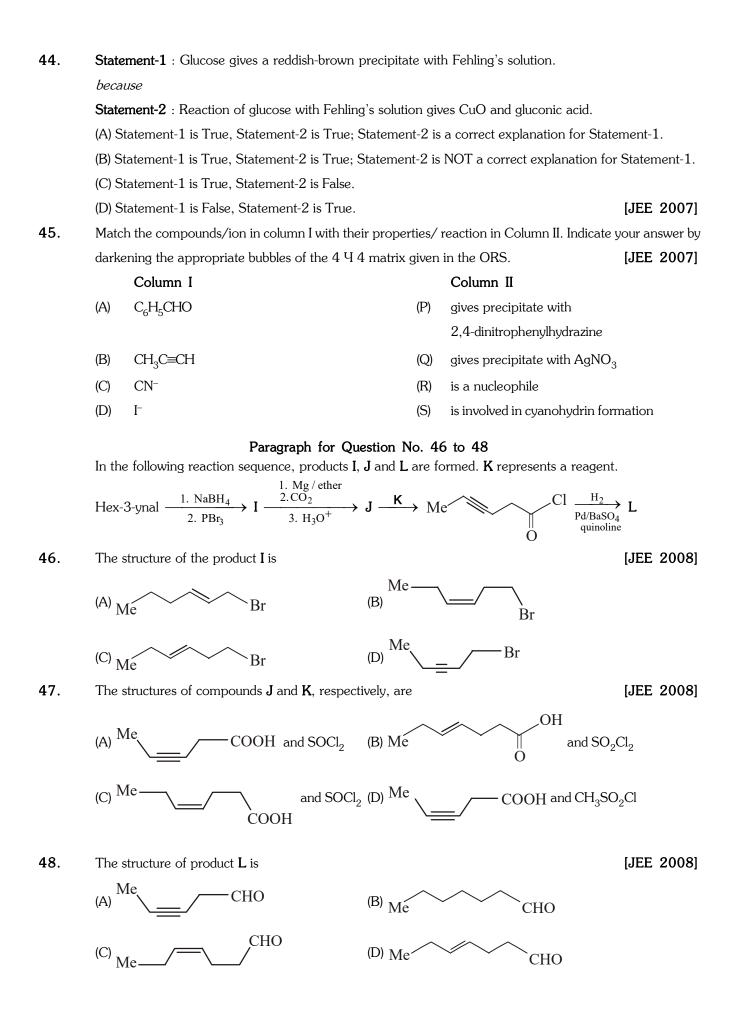
further treatment with aqueous KOH yields compound F. Compound F is
[JEE 2007]



[IIT-96]

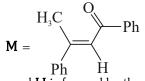
[IIT-98]

[IIT-99]



Paragraph for Question No. 49 to 51

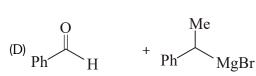
A tertiary alcohol H upon acid catalysed dehydration gives a product I. Ozonolysis of I leads to compounds J and K. Compound J upon reaction with KOH gives benzyl alcohol and a compound L, whereas K on reaction with KOH gives only M.



49. Compound **H** is formed by the reaction of



(C) H + PhCH₂MgBr



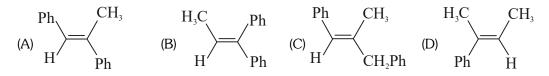
(B) Ph CH_3 + PhCH₂MgBr

50. The structure of compound **I** is



[JEE 2008]

[JEE 2008]



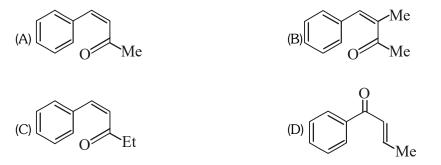
51. The structures of compounds J, K and L, respectively, are (A) PhCOCH₃, PhCH₂COCH₃ and PhCH₂COO□K⁺
(B) PhCHO, PhCH₂CHO and PhCOO□K⁺
(C) PhCOCH₃, PhCH₂CHO and CH₃COO□K⁺
(D) PhCHO, PhCOCH₃ and PhCOO□K⁺

Paragraph for Question Nos. 52 to 54

A carbonyl compound P, which gives positive iodoform test, undergoes reaction with MeMgBr followed by dehydration to give an olefin Q. Ozonolysis of Q leads to a dicarbonyl compound R, which undergoes intramolecular addol reaction to give predominantly S.

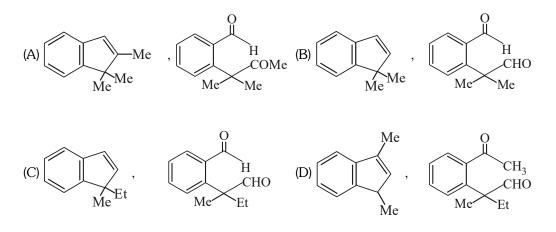
$$\mathbf{P} \xrightarrow[3,H^+,H_2O]{2.H^+,H_2O} \mathbf{Q} \xrightarrow[3,H_2SO_4,\Delta]{1.O_3} \mathbf{R} \xrightarrow[2.\Delta]{1.OH^-} \mathbf{S}$$

52. The structure of the carbonyl compound **P** is



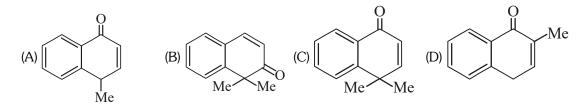
[JEE 2009]

53. The structure of the products Q and R, respectively, are



54. The structure of the product **S** is

[JEE 2009]



Paragraph for Questions Nos. 55 to 56

An acyclic hydrocarbon P, having molecular formula C_6H_{10} , gave acetone as the only organic product through the following sequence of reactions, in the which Q is an intermediate organic compound.

$$\begin{array}{c}
\mathbf{P} \\
(C_{6}H_{10}) & \xrightarrow{(i) \text{ dil } H_{2}SO_{4}/HgSO_{4}} \\
(ii) \text{ NaBH}_{4}/\text{ethanol} \\
(iii) \text{ dil.acid}
\end{array} \qquad \mathbf{Q} \qquad \begin{array}{c}
(i) \text{ conc.}H_{2}SO_{4} \\
(Catalytic amount) \\
(-H_{2}O) \\
(ii) O_{3} \\
(iii) Zn/H_{2}O
\end{array} \qquad \mathbf{2} \qquad \begin{array}{c}
\mathbf{O} \\
\mathbf{U} \\$$

(A)
$$CH_{3}CH_{2}CH_{2}CH_{2}-C\equiv C-H$$

(B) $H_{3}CH_{2}C\equiv C-CH_{2}CH_{3}$
(C) $H_{-}C-C\equiv C-CH_{3}$
(D) $H_{3}C-C-C\equiv C-H$
 $H_{3}C$

56. The structure of the compound Q is -

(C)
$$\begin{array}{c} H_3C & OH \\ H_-C - CH_2CHCH_3 \\ H_3C \end{array}$$
 (D) $CH_3CH_2CH_2CHCH_2CH_3$

57. The number of aldol reaction(s) that occurs in the given transformation is

$$CH_{3}CHO + 4HCHO \xrightarrow{\text{conc. aq. NaOH}} HO \xrightarrow{\text{OH}} OH HO \xrightarrow{\text{OH}} OH HO \xrightarrow{\text{OH}} OH$$

(A) 1 (B) 2 (C) 3 (D) 4 58. Among P, Q, R and S, the aromatic compound(s) is / are : AICl₃ Ρ NaH Q $\xrightarrow{(\mathrm{NH}_4)_2 \mathrm{CO}_3}_{100-115^{\circ}\mathrm{C}} \rightarrow \mathrm{R}$

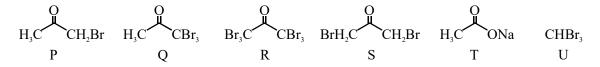
59. After completion of the reactions (I and II), the organic compound(s) in the reaction mixtures is(are)

Reaction I:
$$H_3C$$
 $CH_3 \xrightarrow{Br_2(1.0 \text{ mol})}{(1.0 \text{ mol})}$ [JEE 2013]

HCl

S

Reaction II :
$$H_3C$$
 $CH_3 \xrightarrow{Br_2(1.0 \text{ mol})} (1.0 \text{ mol})$



(A) Reaction I : P and Reaction II : P

(B) Reaction I : U, acetone and Reaction II : Q acetone

(C) Reaction I : T, U, acetone and Reaction II : P

(D) Reaction I : R, acetone and Reaction II : S acetone

[JEE 2012]

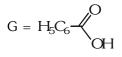
[JEE 2013]

PREVIOUS	S YEARS QU	ESTIONS		ANSWER	KEY		EΣ	KERCISE -5			
1 . (B)	2 . (A)	3. (D)	4. (B)	5. (C, D)	6. (D)	7 .(A)	8. (B)	9 . (A)			
10. (A)	11. (D)	12. (B)	13. (A)	14. (A)	15. (C)	16. (B)	17. (C)	18. (B)			
19 . (B)	20. (A)	21. (A)	22. (A)	23. (C)	24. (D)	25. (A)	26. (A)	27 . (C)			
28. (A) 29. (i)	For empir Eleme		%	Relativ	e no. of at	oms	Simplest	ratio			
	C H O		69.77 11.63 19.20		5.76 11.63 1.2		5 1(1)			
	and		ormula wt. =	nd is C ₅ H ₁₀ O 86							
	(ii) Co			ind is C ₅ H ₁₀ O ddition compo		s, has carbony	l group. i.e., a	aldehyde or			
		(iii) It does not reduce Fehling solution and thus, it is not aldehyde but it is ketone.									
	H ₃ C-										
	(v) Ab	ove facts reve	eals that com	pound is							
	Œ	H3.CH2.CH2. Pentan-2-or		H3C H3C methylbutan-2							
30.	The forma	ation of CHI ₃	takes place a	-							
	-	•		$\xrightarrow{OH^1} \to CH_3C$ ne active speci	-	-	and an iodina	ating agent.			
31.	A = H ₃ C H ₃ C	Ko ; B =	= H ₃ C-	H3(; C =	C	D = H ₅ C ₂ - 0	OH ; E = H	₽C – CH₃			
32.	A = HOH	I_C-	}00H₃	B = HCOO	ЭK						
33.	$C_6H_5CH =$	= CHCOOC ₂ H	H ₅								
34.	KCN + H ₂	$_2SO_4 \longrightarrow KH$	HSO ₄ + HCN	I							
	$ \overset{CH_3}{\underset{H}{\succ}} O_+ $		H ² C ₂ OH H ² CN Gyanohydrine	LiAIH₄ →	H-5C2 OH H-N	2					

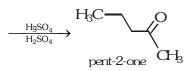
35.
$$A = \begin{pmatrix} CH_3 \\ OH \\ OH \\ OH_3 \end{pmatrix}$$
; $B = \begin{pmatrix} O & O \\ H_3C \end{pmatrix}$; $C = \begin{pmatrix} OH \\ OH \\ OH \\ OH \end{pmatrix}$

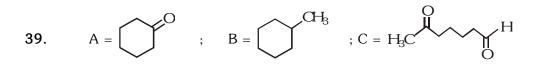
36. A = C₆H₅CHO, base B =
$$C_6H_5$$

$$D = \frac{H_3C - C - NH}{O}; \quad E = \frac{H_5C_6 - C - NH}{O}; \quad F = H_5C_6 - NH_2$$



38.
$$CH_3CH_2C \equiv CH \xrightarrow{\text{Naln}} CH_3CH_2C \equiv \overline{C}Na^{\oplus} \xrightarrow{CH_3X} CH_3CH_2C \equiv C-CH_3$$





$$D = \bigcup_{i=1}^{COCH_3} ; E = \bigcup_{i=1}^{CH_3} Br$$

40.
$$A = H_3C$$
 $C = H_3C$ $C =$