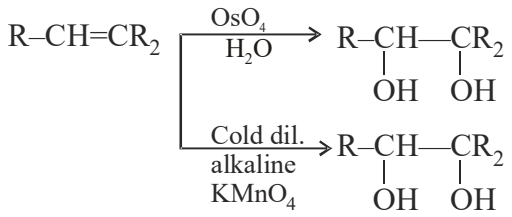
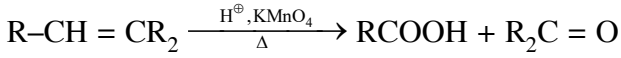
OXIDATION

(I) OXIDATION OF ALKENES



* Cold dil. alkaline $KMnO_4$ is called as Bayer's reagent, use as a test of unsaturation.
* Overall syn addition
* Given by alkenes & alkynes
* Benzene & Cyclopropane can not give this reaction.

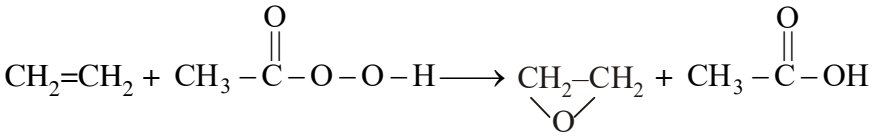
If we use acidic $KMnO_4$ or warm $KMnO_4$ or concentrated $KMnO_4$ the oxidative cleavage of Glycol occurs resulting in mixture of Carboxylic acids & Ketones.



Hot acidic $KMnO_4$, Hot acidic $K_2Cr_2O_7$ & hot acidic $NaIO_4$ gives same result with alkene. The effect is similar to that of oxidative ozonolysis on alkenes.

Preilschaive reaction :

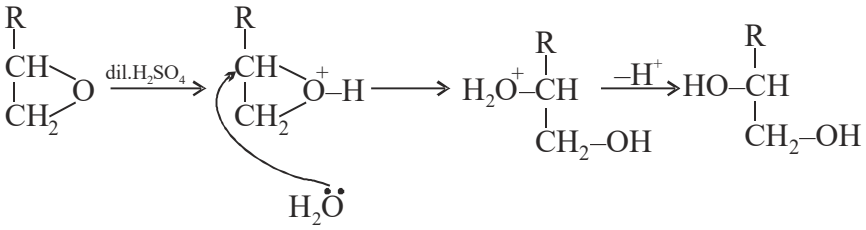
Epoxidation of alkenes is reaction of alkenes with peroxyacids.

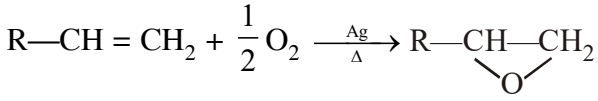


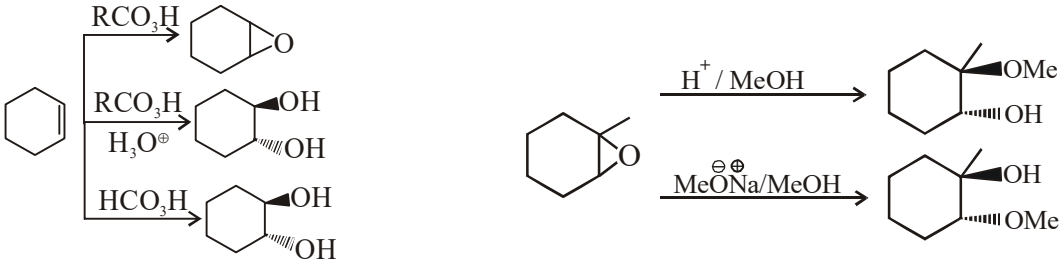
→ Rate α nucleophicity of alkene.

→ With the increase in e^- withdrawing substituents in leaving group, rate increases.

→ Hydrolysis of epoxides form anti diols.



→ 



OXO PROCESS :

Hydroformylation/rolyen synthesis :

$$CH_2 = CH_2 \xrightarrow[\text{Lewis acid high T/P}]{CO+H_2} CH_3–CH_2–CHO$$

Carbonylation / Koch's reaction :

$$CH_2 = CH_2 \xrightarrow[\text{Lewis acid high T/P}]{CO+H_2O} CH_3–CH_2–COOH$$

$$CH_2 = CH_2 \xrightarrow[\text{Lewis acid high T/P}]{CO+ROH} CH_3–CH_2–\overset{\overset{O}{\|}}{C}–OR$$

$$HC \equiv CH \xrightarrow[\text{Lewis acid high T/P}]{CO+H_2O} \underset{\text{Acrylic acid}}{CH_2=CH–COOH}$$

$$HC \equiv CH \xrightarrow[\text{Lewis acid high T/P}]{CO+MeOH} \underset{\text{Methyl acrylate}}{CH_2=CH–\overset{\overset{O}{\|}}{C}–OMe} \xrightarrow{\text{Polymerisation}} \underset{\text{PMA}}{\left(CH_2—\overset{\overset{COOMe}{|}}{CH}\right)_n}$$

(II) OXIDATION OF ALCOHOLS

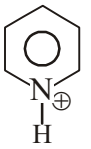
Oxidising agents

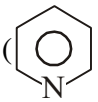
(1) $H^{\oplus}/K_2Cr_2O_7$, Δ (Strong oxidising agent)

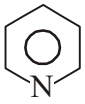
(2) $H^{\oplus}/KMnO_4$, Δ (Strong oxidising agent)

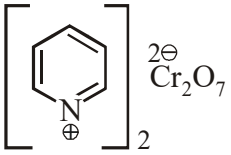
(3) **Jones reagent :** CrO_3 / H_2SO_4 treated with alcohol usually taken in acetone (strong oxidising agent)

(4) Cu / 300°C (or Red hot Cu tube)

(5) PCC (Pyridinium chloro chromate)  $CrO_3Cl^{\ominus}$ or  $CrO_3 + HCl$

(6) Collin's reagent ( (2 mol) + $CrO_3 + CH_2Cl_2$)

(7) Sarett reagent (i.e. PCC in CH_2Cl_2)  + CrO_3 + HCl + CH_2Cl_2

(8) $\underset{\text{(Pyridinium dichromate)}}{\text{PDC}}$  $Cr_2O_7^{2\ominus}$

(9) TsCl + DMSO + $NaHCO_3$

$$RCH_2OH \xrightarrow{Ts–Cl} RCH_2OTs \xrightarrow[NaHCO_3]{DMSO} RCHO$$

$$R_2CHOH \xrightarrow{Ts–Cl} R_2CH–OTs \xrightarrow[NaHCO_3]{DMSO} R_2CO$$

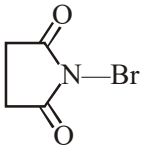
$$R_3COH \xrightarrow{Ts–Cl} R_3C–OTs \xrightarrow[NaHCO_3]{DMSO} \times$$

(10) MnO_2-Oxidises only allylic or benzylic– OH. 1° Allylic or benzylic OH $\xrightarrow{MnO_2}$ Aldehyde

2° Allylic or benzylic OH $\xrightarrow{MnO_2}$ Ketone

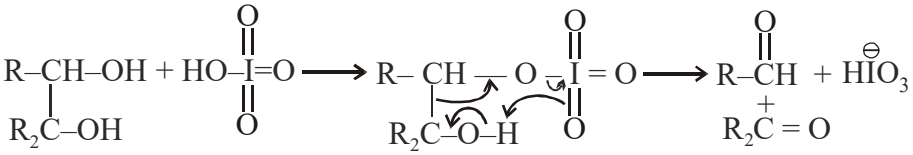
No effect on 3° ROH and on Carbon-Carbon multiple bond.

(11) NBS (N-Bromosuccinamide)



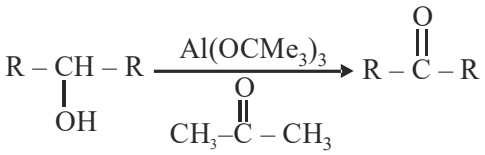
(12) **Periodic cleavage**

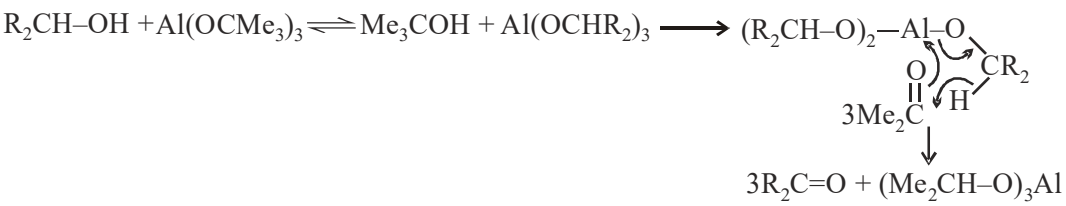
A similar oxidation is obtained in case of HIO_4 known as periodic cleavage.



Reaction is observed for Vic-diols.

(13) **Openaur oxidation**





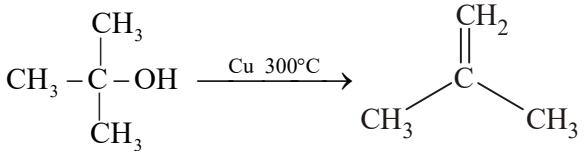
Oxidation of alcohol with aluminium tertiary butoxide is Openaur oxidation.

Ex. Different oxidising agents are used to oxidise alcohols in corresponding carbonyl compounds and carboxylic acids.

e.g. (I) $\underset{1°\text{alcohol}}{R-CH_2-OH} \xrightarrow{1,2,3,} R-\overset{O}{\overset{\|}{C}}-OH$

(II) $\underset{1°\text{alcohol}}{R-CH_2-OH} \xrightarrow{4,5,6,7,8,9,10,11} R-\overset{O}{\overset{\|}{C}}-H$ (Aldehyde)

(III) $\underset{2°\text{ Alcohol}}{R-\overset{OH}{\overset{|}{C}}H-R'} \xrightarrow{1,2,3,4,5,6,7,8,8,9,10,11} R-\overset{O}{\overset{\|}{C}}-R'$ (Ketone)

(IV)  Dehydration takes place.

(V) Double bond or Tripple bond is not affected by 4,5,6,7,8,9,10

(VI) No effect on 3° alcohol by 1,2,3,5,6,7,8,9,10,12,13

(III) OXIDATION OF CARBONYL COMPOUNDS

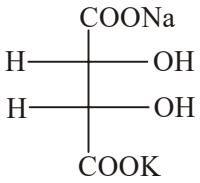
1. Tollen's Test (Silver mirror test)

$RCHO + [Ag(NH_3)_2]OH$ (Tollen's Reagent) $\xrightarrow{\Delta}$ $RCOOH + NH_3 + Ag \downarrow$ (Silver mirror)

Aldehyde acts as reducing agent, they can reduce mild oxidizing agents like Tollen's Reagent.

Tollen's test gentle Heating for 20 to 25 mins.

2. **Fehling's Solutions**



Fehling's A	Fehling's B
aq. $CuSO_4$	Alkaline solution of Roschelye salt (sodium potassium tartrate)

It act's a carrier for Cu^{2+} as it make reversible complex with Cu^{2+}

This test is also used is Blood and Urine test.

$$RCHO + Cu^{2+} \xrightarrow[\Delta]{H_2O} \underset{\downarrow}{RCOOH} + \underset{\downarrow}{Cu^{\oplus}}$$
$$\underset{\downarrow}{RCOO^-} \quad \underset{\downarrow}{Cu_2O\,(\text{red ppt.})}$$

Note : Aromatic aldehyde shows negative test with fehling reagent.

3. **Benedict's solution**

Sodium Citrate + NaOH + $NaHCO_3$ + $CuSO_4$

$$RCHO + Cu^{2+} \xrightarrow[\Delta]{H_2O} \underset{\downarrow}{RCOOH} + \underset{\downarrow}{Cu^{\oplus}}$$
$$\underset{\downarrow}{RCOO^-} \quad \underset{\downarrow}{Cu_2O\,(\text{red ppt.})}$$

It is similar to Fehling test

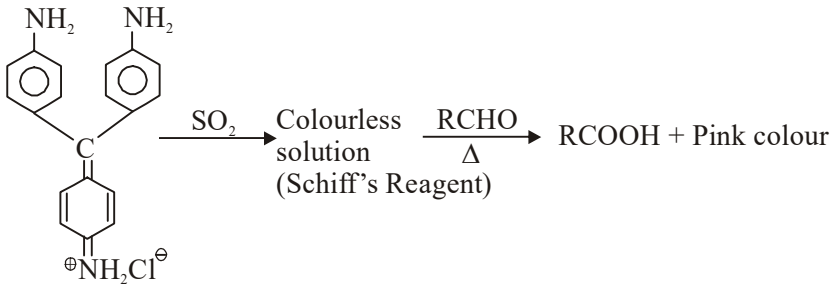
4. $RCHO + HgCl_2 + H_2O \xrightarrow[\Delta]{} RCOOH + 2HCl + Hg_2Cl_2$ (white)

$RCHO + Hg_2Cl_2 + H_2O \xrightarrow[\Delta]{} RCOOH + 2HCl + 2Hg \downarrow$ greyish black

5. **Schiff's Reagent**

Schiff's Reagent is aq. solution of following base decolourised by passing SO_2.

Aldehyde restore pink colour of Schiff's reagent.



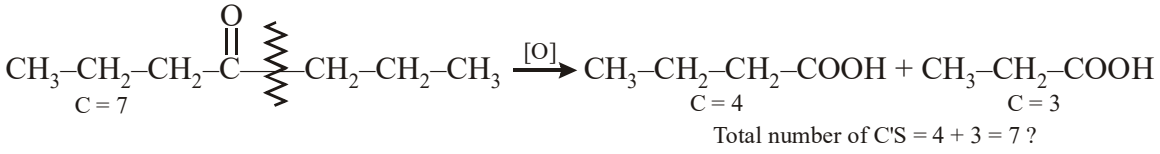
p-Rosaniline Hydrochloride
Magenta colour (Fuschin)

Ketons are not easy to oxidize so they do not give these 5 tests. These five tests can be used to distinguish aldehyde and ketones. Both gives 2,4 DNP test

Oxidation of ketones : Ketones undergo oxidation only in drastic conditions.

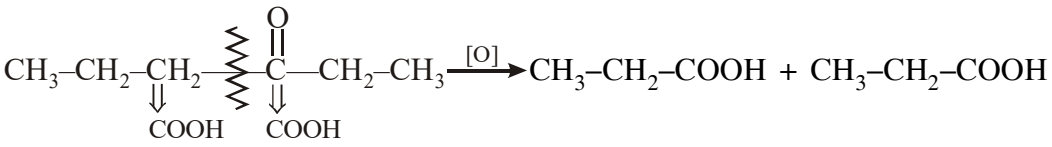
During the oxidation of ketones there is breaking of carbon-carbon bond between α-carbon and carbonyl carbon. In this process both carbons convert into carboxylic groups. This leads to the formation of two moles of monocarboxylic acids.

Case - I : Oxidation of Symmetrical ketones :

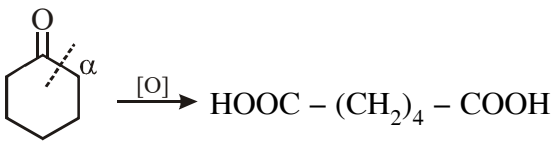


Thus number of carbons in any product is less than the number of carbons in ketone.

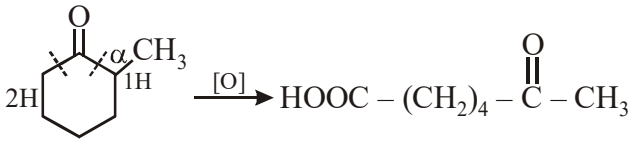
Case - II : Oxidation of unsymmetrical ketones : In case of unsymmetrical ketones, those —C(=O)—R bond break in which alkyl group has more number of carbons. This rule is known as **Poff's rule.**



Case - III : Oxidation of cyclic ketones : Formation of dibasic acid takes place from cyclic ketones. In this case number of carbons in ketone and dibasic carboxylic acid is always same.

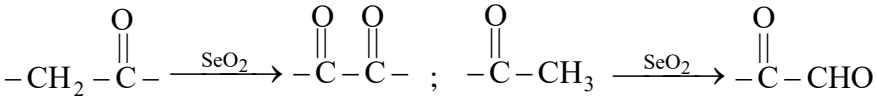


Note : If both α-carbons are not identical then ;

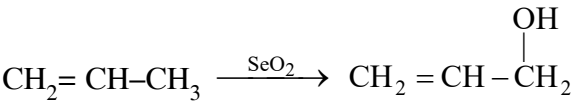


Allylic oxidation

SeO_2 is a selective oxidizing agent with converts $-CH_2-$ group adjacent to carbonyl group into carbonyl group.The reagent, in general, oxidises active methylene and methyl groups to ketonic and aldehydic groups respectively.

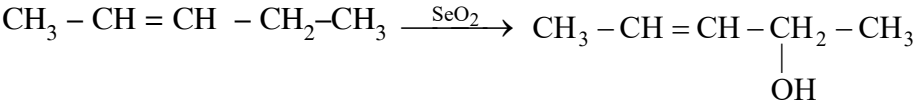


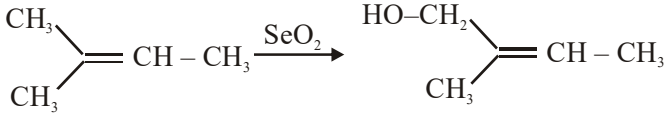
Double bonds, triple bonds and aromatic rings may also activate the methylene group.The methylene or methyl group α to the most highly substitued end of the double bond is hydroxylated according to the order of preference of oxidation $CH_2 > CH_3 > CH$ groups.

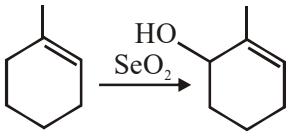


$2° \text{ C} - \text{H} > 1° \text{ C} - \text{H} > 3° \text{ C} - \text{H}$

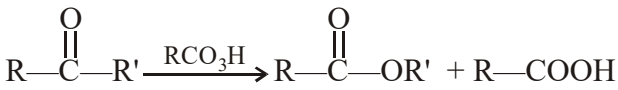
Rate of reactivity order







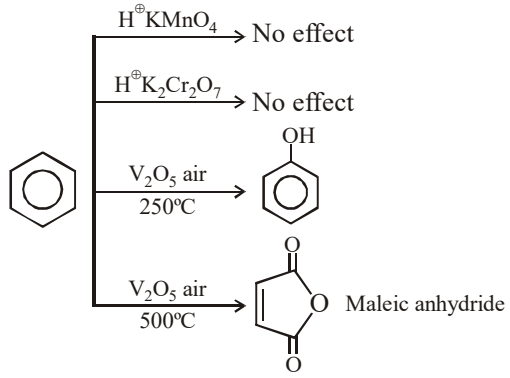
Baeyer villiger oxidation

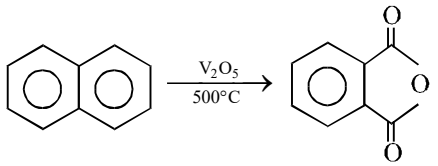


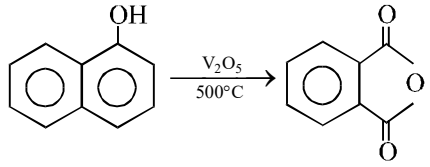
Migratory order for above reaction

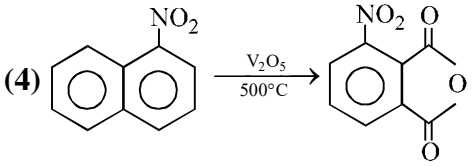
$-H > 3^o > 2^o > -Ph > 1^o > -Me$

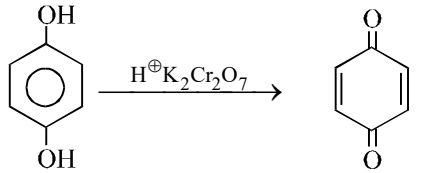
Oxidation of Aromatic Compounds.

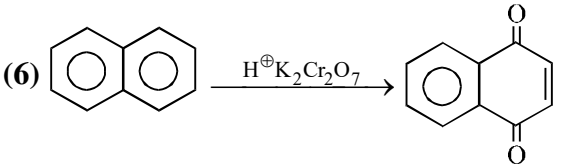
(1) 

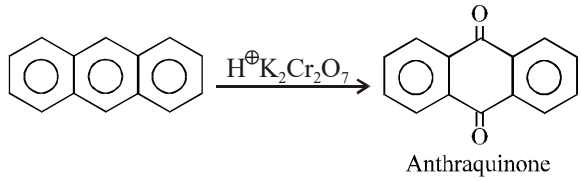
(2) 

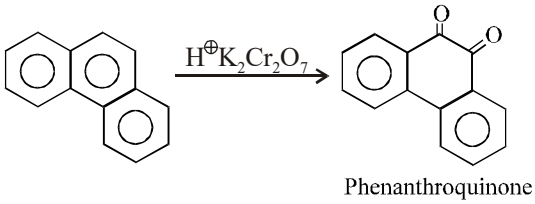
(3) 

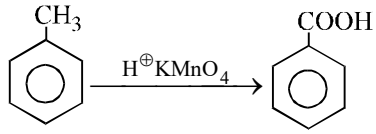
(4) 

(5) 

(6) 

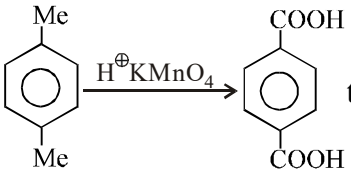
(7) 

(8) 

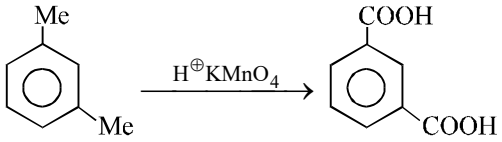
(9) 

(10) $Ph\text{-}(CH_2)_n CH_3$, $PhCHMe_2$, $Ph–CH_2OH$, $Ph–CH_2Br$, $PhCHCl_2$, PhCHO, Ph-  & all the compounds having at least one α H give PhCOOH.

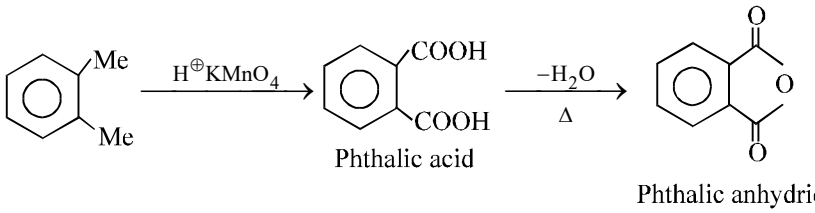
(11) $Ph–CMe_3 \xrightarrow[\Delta]{H^{\oplus}KMnO_4} Me_3C–COOH$

(12) 

terephthalic acid

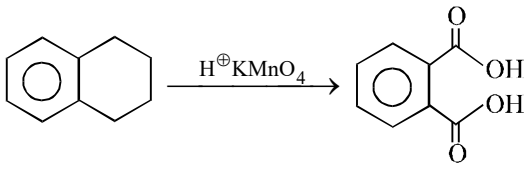
(13) 

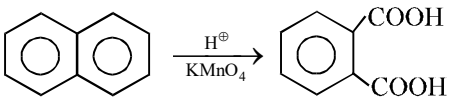
isophthalic acid

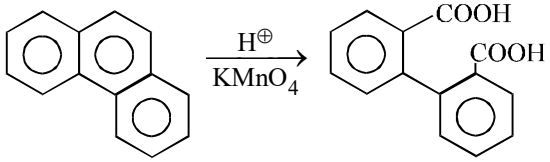
(14) 

Phthalic acid

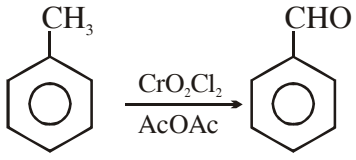
Phthalic anhydride

(15) 

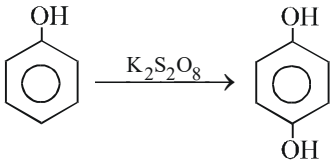
(16) 

(17) 

Etard oxidation

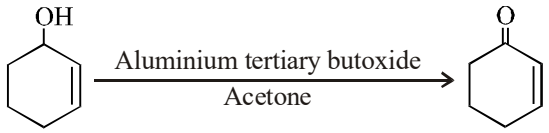
(18) 

Elb's persulphate oxidation:

(19) 

EXERCISE#O-I

1. The given reaction

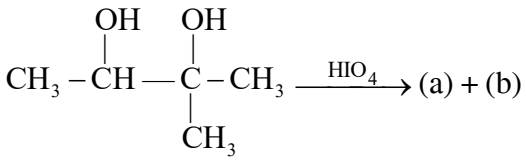


is knows as :

(A) Kolbe reaction (B) Tischenko reaction (C) MPV reaction (D) Oppeneur oxidation

OX0001

2. In the given reaction :

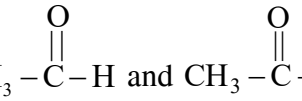


(a) and (b) respectively be :

(A) CH_3CHO and CH_3CHO (B) CH_3COCH_3 and CH_3CHO

(C) CH_3COCH_3 and CH_3COCH_3 (D) CH_3COOH and CH_3COCH_3

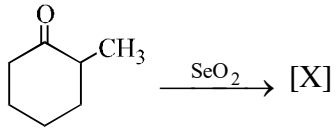
OX0002

3.  and is differentiated by

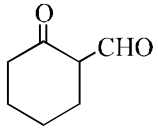
(A) Tollen's reagent (B) Lucas test (C) Iodoform test (D) $NaHSO_3$ test

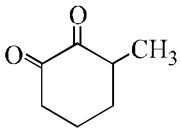
OX0003

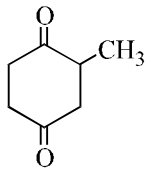
4. In the given reaction

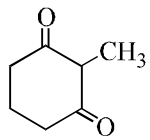


[X] will be:

(A) 

(B) 

(C) 

(D) 

OX0004

5. Which will give silver mirror test with Tollens reagent :

(A) C_6H_5CHO (B) CH_3–CHO (C) HCOOH (D) All of these

OX0005

6. Which one of the following compounds will give dimethyl glyoxal with SeO_2:

(A) Acetone (B) Acetophenone

(C) Ethyl methyl ketone (D) Propanaldehyde

OX0006

7. Which one of the following alcohols can be oxidised by K_2CrO_4?

(A) Ethanol (B) Tert butyl alcohol

(C) Isopropyl alcohol (D) Allyl alcohol

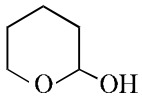
OX0007

8. Fehling solution gives red precipitate with:

(A) Aromatic aldehyde (B) Aliphatic aldehyde (C) Ketones (D) $CH_2(OH)-C(=O)-CH_3$

OX0008

9. Which of the following compound will give positive Tollens test

(A) CH_3CHO (B) 

(C) $CH_3CH_2CH_2CH_2CH(OH)-OCH_3$ (D) $CH_3-CH(OCH_3)_2$

OX0009

10. Schiff's reagent gives pink colour with :

(A) Acetaldehyde (B) Formic acid (C) Acetic acid (D) Methyl acetate

OX0010

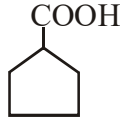
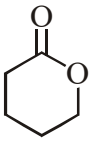
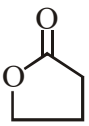
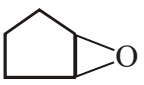
11. Which one of the following alcohols are oxidised by MnO_2?

(A) $C_6H_5-CH_2-CH_2-OH$ (B) $CH_2=CH-CH_2-CH(OH)-CH_3$

(C) $CH_3-CH=CH-CH(OH)-CH_3$ (D) $CH_3-CH_2-CH_2-OH$

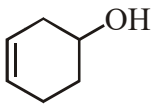
OX0011

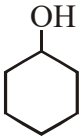
12.  Major product is :

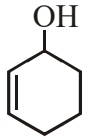
(A)  (B)  (C)  (D) 

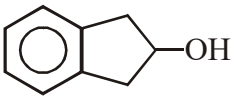
OX0012

13. Which of the following can be oxidised by MnO_2:

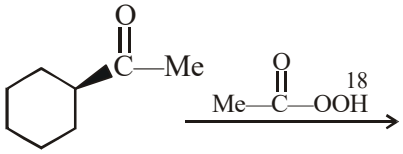
(A) 

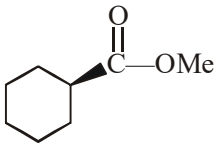
(B) 

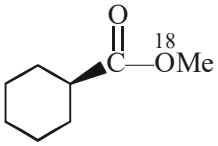
(C) 

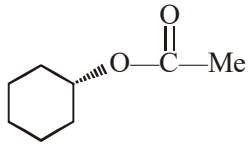
(D) 

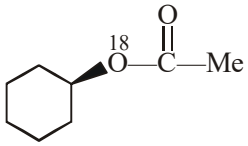
OX0013

14.  Major product is :

(A) 

(B) 

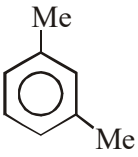
(C) 

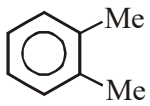
(D) 

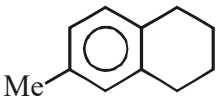
OX0014

15. A $\xrightarrow{H^+ / KMnO_4}$ B $\xrightarrow{\Delta}$ Phthalic Anhydride ; A is :

(A) 

(B) 

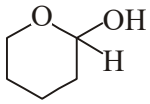
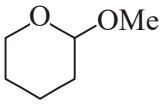
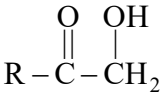
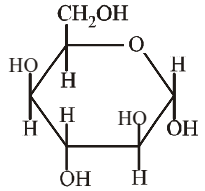
(C) 

(D) 

OX0015

EXERCISE#O-II

1. Which will give the Tollen test.

(A)  (B)  (C)  (D) 

OX0016

2. Which of the following is an example of oxidation reaction :

(A) $CH_3 - CH_2 - OH \xrightarrow{PCl_3}$ (B) $CH_3 - CH = CH_2 \xrightarrow{Ni/H_2}$ (C) $CH_3 - CH = CH_2 \xrightarrow[CCl_4]{Br_2}$

OX0017

3. (i) $CH_2 = CH_2 \xrightarrow{H^\oplus / KMnO_4}$

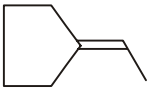
OX0018

(ii) $CH_3–CH= CH_2 \xrightarrow{H^\oplus / KMnO_4}$

OX0019

(iii)  $\xrightarrow{H^\oplus / KMnO_4}$

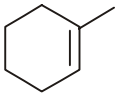
OX0020

(iv)  $\xrightarrow{H^\oplus / KMnO_4}$

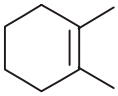
OX0020

(v)  $\xrightarrow{H^\oplus / KMnO_4}$

OX0021

(vi)  $\xrightarrow{H^\oplus / KMnO_4}$

OX0021

(vii)  $\xrightarrow{H^\oplus / KMnO_4}$

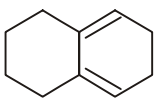
OX0022

(viii)  $\xrightarrow[-CO_2]{H^\oplus / KMnO_4, \Delta}$

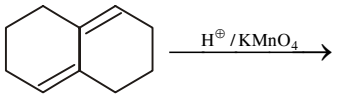
OX0022

(ix)  $\xrightarrow{H^\oplus / KMnO_4}$

OX0023

(x)  $\xrightarrow{H^\oplus / KMnO_4}$

OX0023

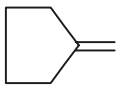
(xi) 

OX0023

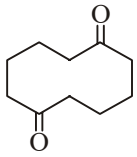
4. A to F alkenes with minimum possible carbon.

(i) A $\xrightarrow[\Delta]{H^{\oplus}/KMnO_4}$ MeCOOH as the only product

OX0024

(ii) B $\xrightarrow[\Delta]{H^{\oplus}/KMnO_4}$  as the only product

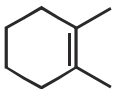
OX0024

(iii) D $\xrightarrow[\Delta]{H^{\oplus}/KMnO_4}$ 

OX0025

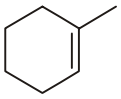
(iv) F $\xrightarrow[\Delta]{H^{\oplus}/KMnO_4}$ Acetone + Ethanoic acid

OX0025

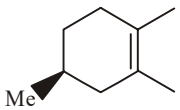
5. (i)  $\xrightarrow[KMnO_4]{1\%\text{ alkaline}}$ A

$\xrightarrow{m\text{-CPBA} / H_3O^+}$ B

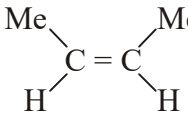
OX0026

(ii)  $\xrightarrow{mCPBA}$

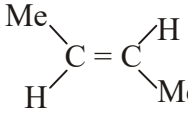
OX0026

(iii)  $\xrightarrow[\text{hydrolysis}]{mCPBA}$

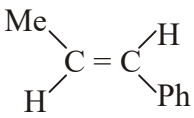
OX0027

(iv)  $\xrightarrow[\text{hydrolysis}]{mCPBA}$

OX0027

(v)  $\xrightarrow[\text{hydrolysis}]{mCPBA}$

OX0028

(vi)  $\xrightarrow[\Delta]{Ag_2O \text{ or } 2Ag + \frac{1}{2}O_2}$

OX0028

6. (i) $CH_3-CH_2-CH_2-OH \xrightarrow{KMnO_4/OH^-,\Delta} ?$

$\xrightarrow[\text{or conc. } HNO_3,\Delta \text{ or } KMnO_4/H^+,\Delta]{K_2Cr_2O_7/H^+,\Delta}$

OX0029

(ii) $CH_3-\underset{\displaystyle OH}{\underset{|}{CH}}-CH_2-CH_3 \xrightarrow[\text{or } K_2Cr_2O_7,H^\oplus]{KMnO_4,H^\oplus} ?$

OX0029

7. (i) $CH_2=CH-(CH_2)_3-CH_2-OH \xrightarrow[\text{in glacial acetic acid}]{\text{Chromic anhydride}}$

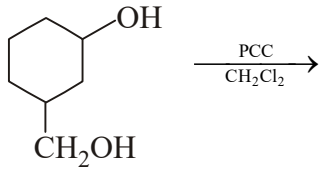
OX0030

(ii) $C_6H_5-CH=CH-CH_2-OH \xrightarrow[\text{in glacial acetic acid}]{\text{Chromic anhydride}}$

OX0030

(iii) $CH_3-\underset{\displaystyle OH}{\underset{|}{CH}}-CH_2-CH_2-CH_2-OH \xrightarrow[CH_2Cl_2]{PCC}$ (A)

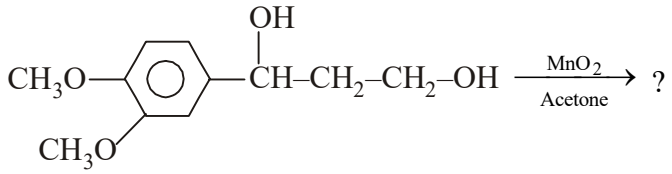
OX0030

(iv) 

OX0031

(v) $CH_2=CH-CH_2-OH \xrightarrow{MnO_2} ?$

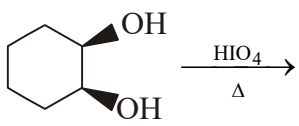
OX0031

(vi) 

OX0031

8. (i) $Me-\underset{\displaystyle OH}{\underset{|}{CH}}-CH_2-OH \xrightarrow[\Delta]{HIO_4}$

OX0032

(ii) 

OX0032

(iii) $HO-CH_2-CH_2-\underset{\underset{OH}{|}}{CH}-CH_2-OH \xrightarrow[\Delta]{HIO_4}$

OX0032

(iv) $\underset{\underset{OH}{|}}{CH_2}-\underset{\underset{OH}{|}}{CH}-\underset{\underset{OH}{|}}{CH}-CH_3 \xrightarrow[\Delta]{HIO_4}$

OX0032

(v) $\underset{\underset{OH}{|}}{CH_2}-\underset{\underset{OH}{|}}{CH}-\underset{\underset{OH}{|}}{CH}-\underset{\underset{OH}{|}}{CH_2} \xrightarrow[\Delta]{HIO_4}$

OX0033

(vi) $Me-\underset{\underset{O}{||}}{C}-\underset{\underset{OH}{|}}{CH}-Me \xrightarrow[\Delta]{HIO_4}$

OX0033

(vii) $Me-\underset{\underset{O}{||}}{C}-\underset{\underset{O}{||}}{C}-Me \xrightarrow[\Delta]{HIO_4}$

OX0033

9. (a) $CH_3-CHO \xrightarrow[\Delta]{SeO_2}$

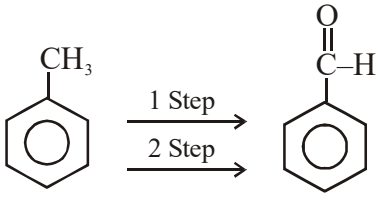
OX0034

(b) $Me_2CO \xrightarrow[\Delta]{SeO_2}$

OX0034

(c) $CH_3-CH=CH_2 \xrightarrow{?}$ Acrolein

OX0034

(d) 

OX0035

(e) $CH_3-\underset{\underset{O}{||}}{C}-H \xrightarrow{SeO_2} P_1 \xrightarrow{\text{Conc. NaOH}} \underset{\underset{COO^-Na^+}{|}}{CH_2OH} \xrightarrow{H^+/\Delta} P_2$

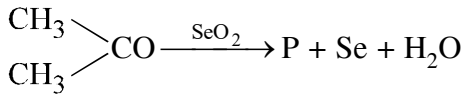
OX0036

10. (a) How will you differentiate HCHO and PhCHO ?
(b) How will you differentiate HCHO and MeCHO ?

OX0037

EXERCISE# (MAIN & ADVANCE)

1. In the reaction, P is **[IIT 1995]**



(A) CH_3COCHO (B) CH_3COOCH_3 (C) CH_3COCH_2OH (D) None

OX0038

2. The best reagent to convert pent–3–en–2–ol into pent –3–en –2–one is - **[AIEEE-2005]**

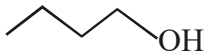
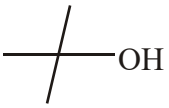
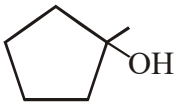
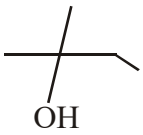
(A) Acidic dichromate (B) Acidic permanganate

(C) Pyridinium chloro-chromate (D) Chromic anhydride in glacial acetic acid

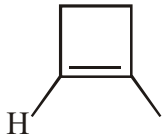
OX0039

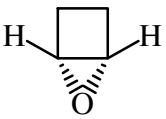
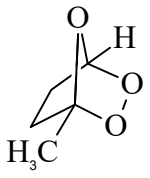
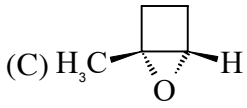
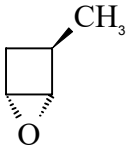
3. Which of the following will change the colour of acidic dichromate solution.

[JEE-MAINS 2013]

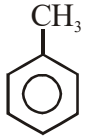
(A)  (B)  (C)  (D) 

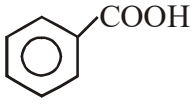
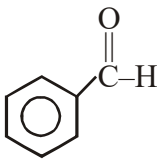
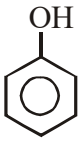
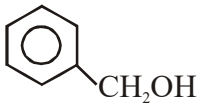
OX0040

4.  Product **[JEE-MAINS 2013]**

(A)  (B)  (C)  (D) 

OX0041

5.  Major product of reaction is? **[JEE-MAINS 2013]**

(A)  (B)  (C)  (D) 

OX0042

6. The most suitable reagent for the conversion of $R–CH_2–OH \rightarrow R–CHO$ is :-

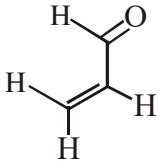
[JEE-MAINS 2014]

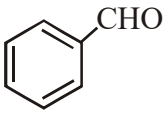
(A) CrO_3 (B) PCC (Pyridinium chlorochromate)

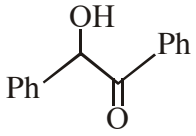
(C) $KMNO_4$ (D) $K_2Cr_2O_7$

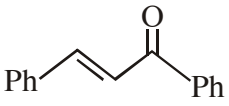
OX0043

7. Positive Tollen's test is observed for **[JEE-Advance 2016]**

(A) 

(B) 

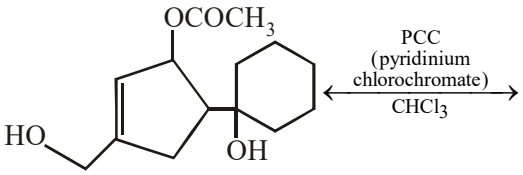
(C) 

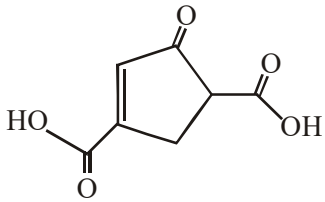
(D) 

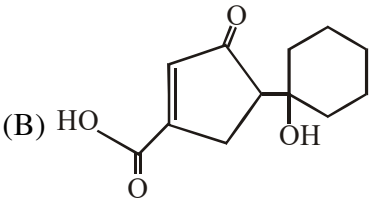
OX0044

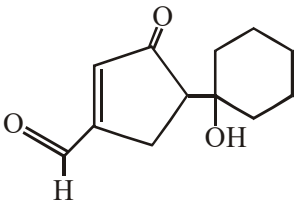
8. The major product formed in the following reaction is :-

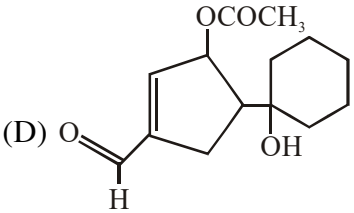
[Jee Main - Onl_line Evening 2018]



(A) 

(B) 

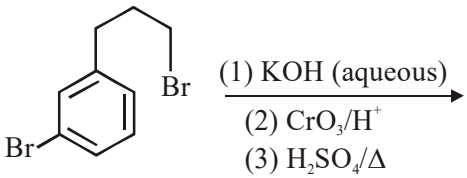
(C) 

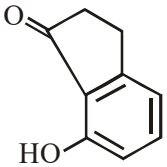
(D) 

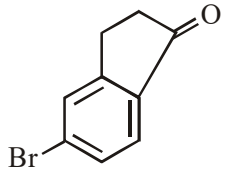
OX0045

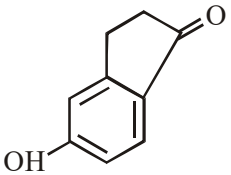
9. The major product of the following reaction is:

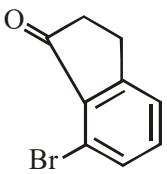
[JEE-MAIN-On-line-(Jan)-2019]



(1) 

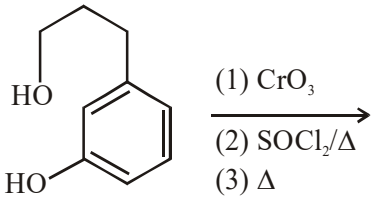
(2) 

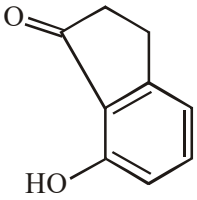
(3) 

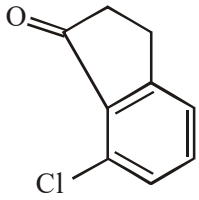
(4) 

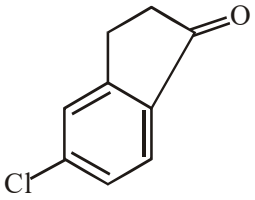
OX0046

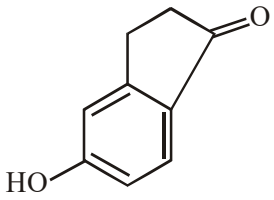
10. The major product of the following reaction **[JEE-MAIN-On-line-(April)-2019]**



(1) 

(2) 

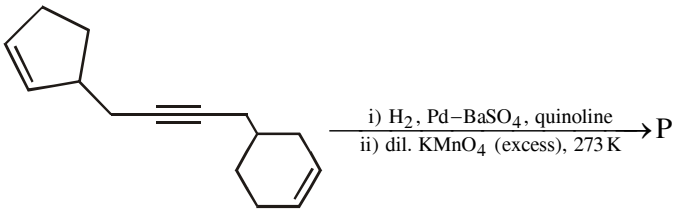
(3) 

(4) 

OX0047

11. Total number of hydroxyl groups present in a molecule of the major product P is ___

[JEE-Advance 2019]



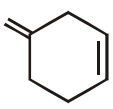
OX0048

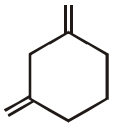
12. An unsaturated hydrocarbon X absorbs two hydrogen molecules on catalytic hydrogenattion, and also gives following reaction : **[JEE-MAIN-On-line-(Jan)-2020]**

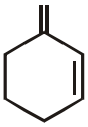
$$X \xrightarrow[Zn/H_2O]{O_3} A \xrightarrow{[Ag(NH_3)_2]^+}$$

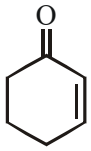
B(3 - oxo - hexanedicarboxylic acid)

X will be :-

(1) 

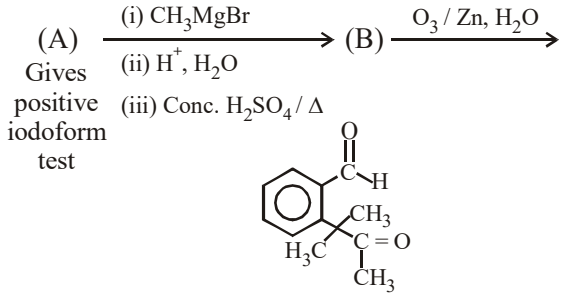
(2) 

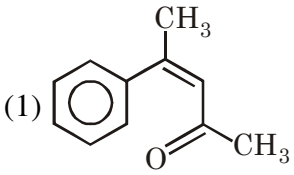
(3) 

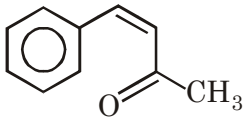
(4) 

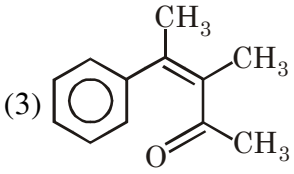
OX0049

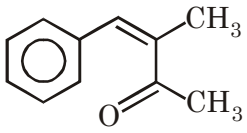
13. Identify (A) in the following reaction sequence : **[JEE-MAIN-On-line-(Jan)-2020]**



(1) 

(2) 

(3) 

(4) 

OX0050

ANSWER-KEY

EXERCISE#O-I

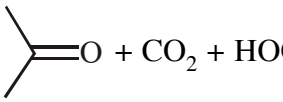
1.	**Ans. (D)**	**2.**	**Ans. (B)**	**3.**	**Ans. (A)**	**4.**	**Ans. (B)**
5.	**Ans. (D)**	**6.**	**Ans. (C)**	**7.**	**Ans. (A,C,D)**	**8.**	**Ans. (B,D)**
9.	**Ans. (A,B,C)**	**10.**	**Ans. (A,B)**	**11.**	**Ans. (C)**	**12.**	**Ans. (B)**
13.	**Ans. (C)**	**14.**	**Ans. (D)**	**15.**	**Ans.** (C)		

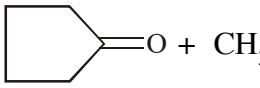
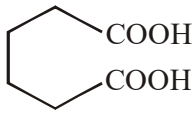
EXERCISE#O-II

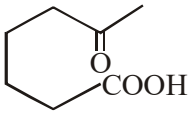
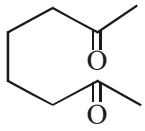
1. **Ans. (A,C,D)** **2.** **Ans. (C)**

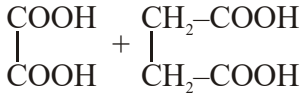
3. **Ans.**

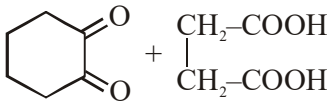
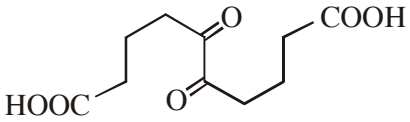
(i) CO_2 (ii) $CH_3–COOH + CO_2$

(iii)  $+ CO_2 + HOOC–C–C–C–COOH$

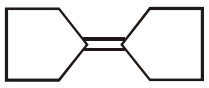
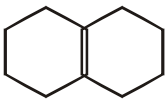
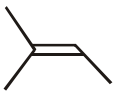
(iv)  $+ CH_3–COOH$ (v) 

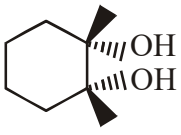
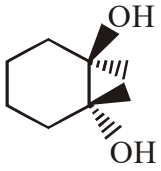
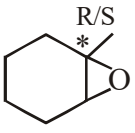
(vi)  (vii) 

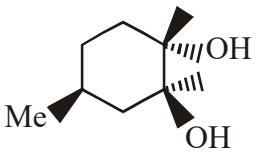
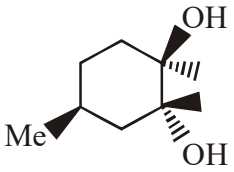
(viii) $2CH_3–COOH$ (ix) 

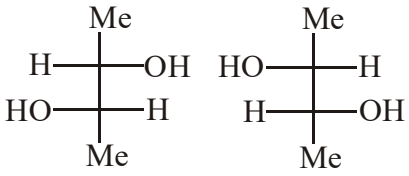
(x)  (xi) 

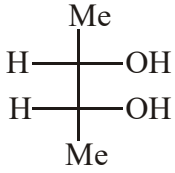
4. **Ans.**

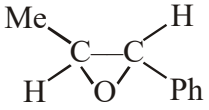
(i) $Me–CH=CH–CH_3$ (ii)  (iii)  (iv) 

5. **Ans.** (i) (A)  (B)  (ii) 

(iii)  

(iv) 

(v) 

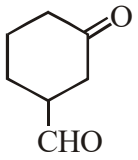
(vi) 

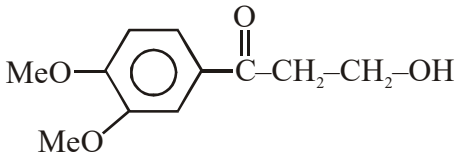
6. Ans.

(i) $CH_3–CH_2–COO^-$, $CH_3–CH_2–COOH$ (ii) $CH_3–\overset{\overset{O}{\|}}{C}–CH_2–CH_3$

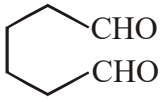
7.

Ans. (i) $CH_2=CH–(CH_2)_3–CHO$ (ii) Ph–CH=CH–CHO

(iii) (A) $H_3C–\overset{\overset{O}{\|}}{C}–CH_2–CH_2–CHO$ (iv) 

(v) $H_2C=CH–CHO$ (vi) 

8. Ans.

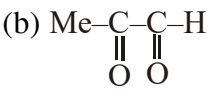
(i) Me–CHO + HCHO (ii) 

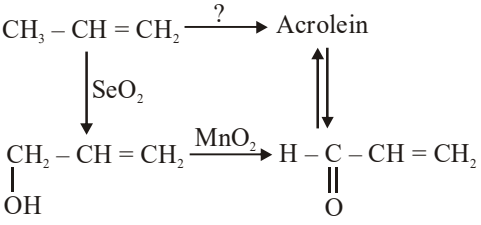
(iii) $HO–CH_2–CH_2–CHO$ + HCHO (iv) HCHO + HCOOH + $CH_3–CHO$

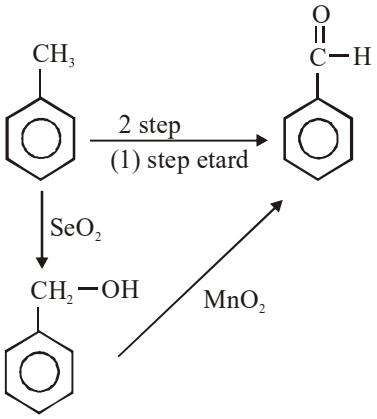
(v) 2HCHO + 2HCOOH (vi) Me–COOH + Me–CHO

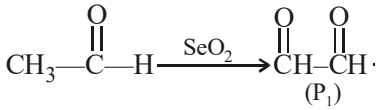
(vii) 2Me–COOH

9. Ans.

(a)  (b) 

(c) 

(d) 

(e) 

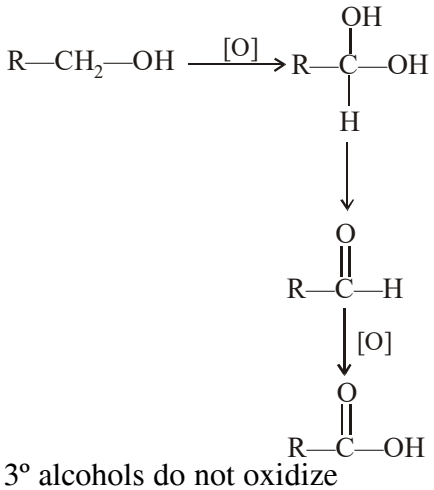
10. (a) Ans. F and B test

(b) Ans. Iodoform test

EXERCISE# (MAIN & ADVANCE)

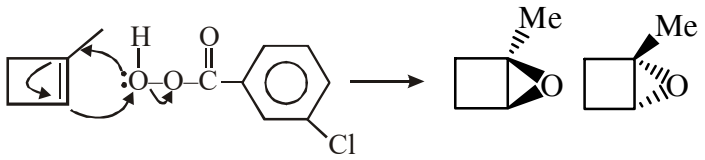
1. **Ans. (A)** **2.** **Ans. (D)**

3. **Ans. (A)**



3º alcohols do not oxidize
they undergo dehydration

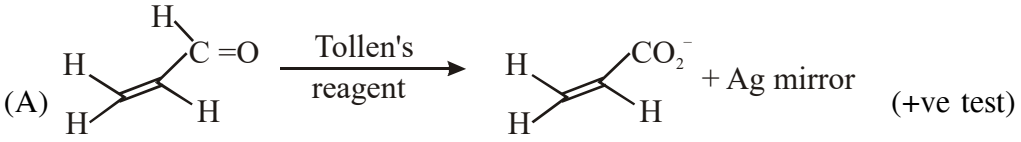
4. **Ans. (C)**



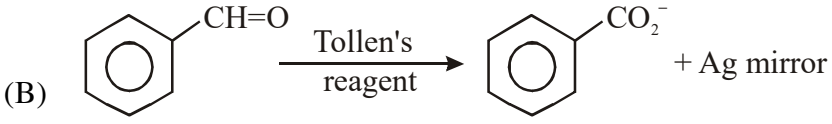
5. **Ans. (A)** **6.** **Ans. (B)**

7. **Ans. (A,B,C)**

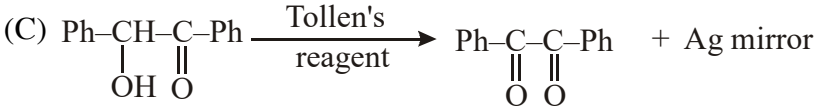
Sol. Tollens's test is given by compounds having aldehyde group. Also α-hydroxy carbonyl gives positive tollen's test.



(A) Acraldehyde + Ag mirror (+ve test)



(B) Benzaldehyde + Ag mirror (+ve test)



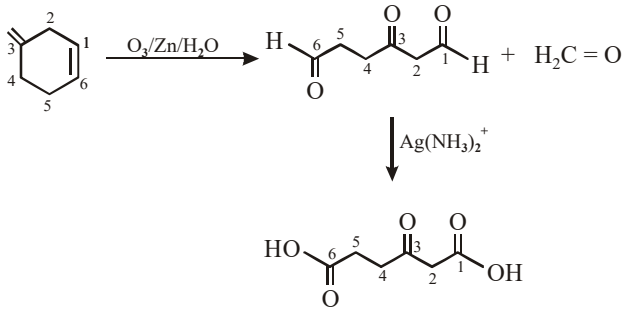
(C) Benzoin + Ag mirror (+ve test)

(D) No reaction (–ve test)

8. **Ans. (D)** **9.** **Ans. (2)** **10.** **Ans. (4)** **11.** **Ans. (6.00)**

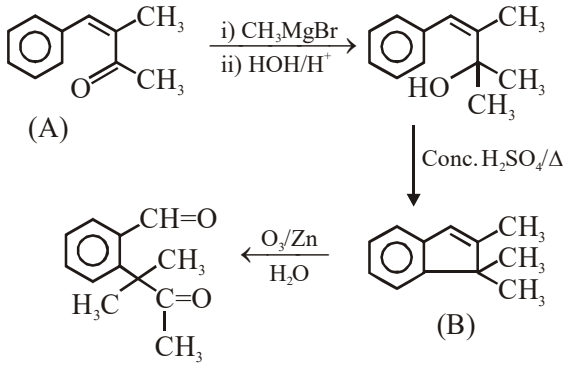
12. Ans. (1)

Sol.



13. Ans. (4)

Sol.



REDUCING AGENTS AND THEIR ROLE

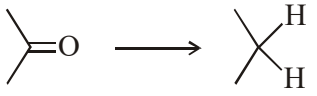
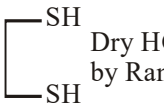
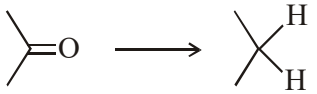
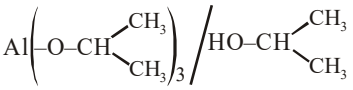
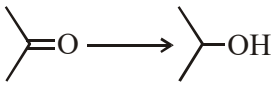
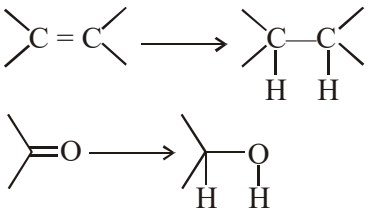
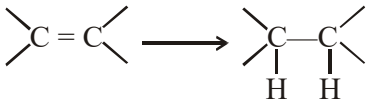
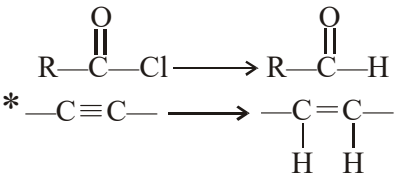
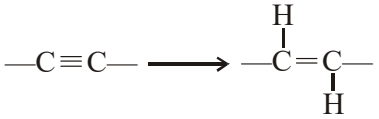
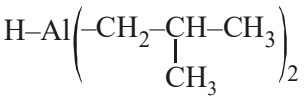
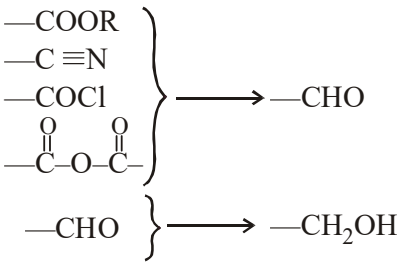
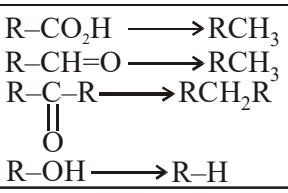
Group	Product	LAH in Ether	$NaBH_4$ in H_2O	B_2H_6 in THF	H_2 / Catalyst Δ
–CHO	$-CH_2OH$	+	+	+	+
>C = O	>CH – OH	+	+	+	+
$-CO_2H$	$-CH_2OH$	+	–	+	–
$-CO_2R$	$-CH_2OH$	+	–	+	+
–COCl	$-CH_2OH$	+	+	–	+
$-CONH_2$	$-CH_2NH_2$	+	–	+	+
$(RCO)_2O$	RCH_2OH	+	–	+	+
–CN	$-CH_2NH_2$	+	–	+	+
>C = NOH	$-CH_2NH_2$	+	–	–	+
>C = C<	>CH – CH<	–	–	+	+
–C ≡ C–	–CH = CH–	–	–	+	+
1° RX	RH	+	–	–	+

Note :

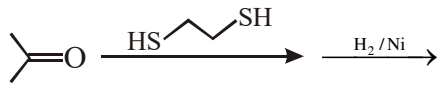
1. B_2H_6 in THF not able to reduce cyclic ester.

2. $NaBH_4$ can also reduce imine group ($\diagdown\!\!\diagup C = NH$).

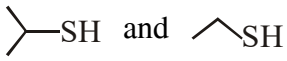
3. Reactivity order towards Ni / H_2 , $\Delta \Rightarrow -C \equiv N > -\overset{\overset{O}{\|}}{C} - O - R$

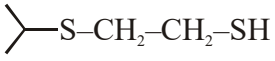
Name	Reagent	Function
Wolf Kishner Reduction	(i) N_2H_4 / (ii) KOH, Δ	
Clemenson Reduction	Zn–Hg / HCl	
Mozingo Reduction	 Dry HCl, followed by Raney Ni	
Stephen's Reduction	$SnCl_2$ / HCl followed by H_3O^+	$R–C\equiv N \longrightarrow R–CH=O$
MPV Reduction		
Hydroboration Reduction	B_2H_6 / AcOH, H_2O	
Bouvoult Blank Reduction	Na / EtOH	$R–COO–R \longrightarrow RCH_2OH + ROH$
Transfer Hydrogenation	N_2H_4 / H_2O_2	
Rosenmund Reduction	H_2, Pd–$BaSO_4$	
Birch Reduction	Na / Liq. NH_3	
DIBAL-H (–78°C)	 followed by $H_3O^{\oplus}$	
Red phosphorus in presence of HI	Red P + HI	

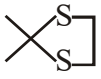
EXERCISE # O-I

1. 

The end products of the reaction are :

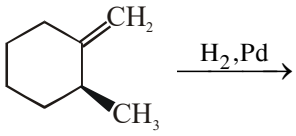
(A)  and

(B) 

(C)  and H_2O

(D) 

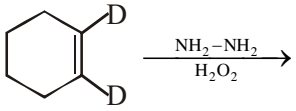
RD0001

2. 

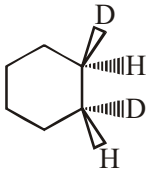
Products of the above reaction will be :

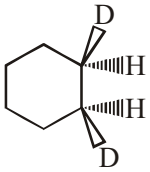
(A) Racemic mixture (B) Diastereomers (C) Meso (D) Structural isomer

RD0002

3.  Product :

(A) 

(B) 

(C) 

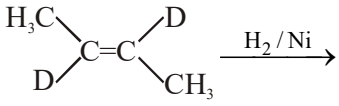
(D) Both (B) and (C)

RD0003

4. On catalytic reduction with H_2/Pt how many alkenes will give n-butane ?

(A) 1 (B) 2 (C) 3 (D) 4

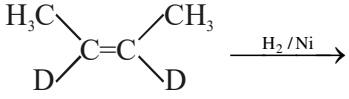
RD0004

5. 

Product of above reaction will be :

(A) Racemic mixture (B) Diastereomers (C) Meso (D) Constitutional isomers

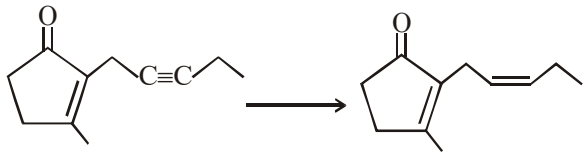
RD0005

6. 

Product of above reaction will be :

(A) Racemic mixture (B) Diastereomers (C) Meso (D) Constitutional isomers

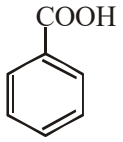
RD0006

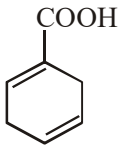
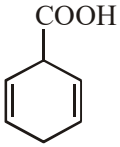
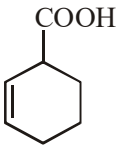
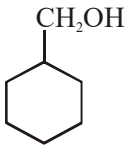
7. 

Which reagent will be used for the above conversion ?

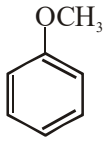
(A) Na/Liq. NH_3 (B) H_2, Pd–$CaCO_3$ (C) Li, Ph–NH_2 (D) H_2, Pt

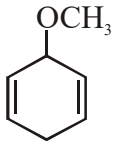
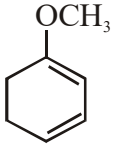
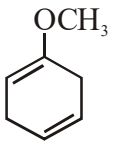
RD0007

8.  $\xrightarrow[\text{Liq.}NH_3]{\text{Li}}$ Major product :

(A)  (B)  (C)  (D) 

RD0008

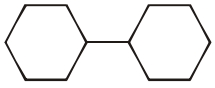
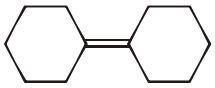
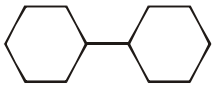
9.  $\xrightarrow{Na,Liq.,NH_3}$ Major product :

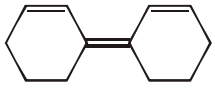
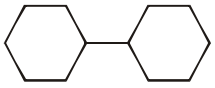
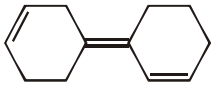
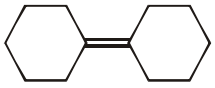
(A)  (B)  (C)  (D) None of these

RD0009

10.  $\xrightarrow[\Delta]{\text{Conc.}H_2SO_4}$ A $\xrightarrow{Pt/H_2}$ B

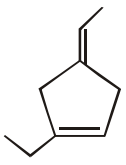
A and B respectively are :

(A) Both  (B)  , 

(C)  ,  (D)  , 

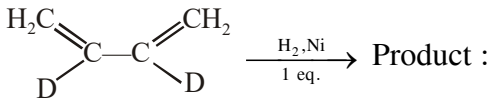
RD0010

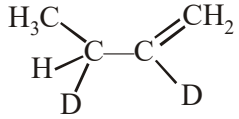
11. If the following compound is treated with Pd in excess of H_2 gas, how many stereoisomers of the product will be obtained ?

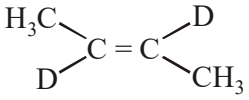


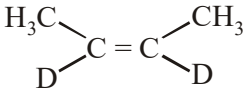
(A) 1 (B) 2 (C) 3 (D) 4

RD0011

12.  Product :

(A) 

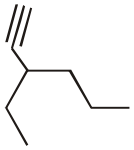
(B) 

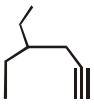
(C) 

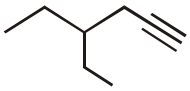
(D) All of these

RD0012

13. Which alkyne gives 3-ethylhexane on catalytic hydrogenation ?

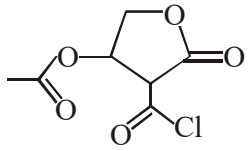
(A) 

(B) 

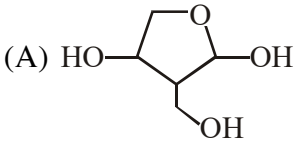
(C) 

(D) All of these

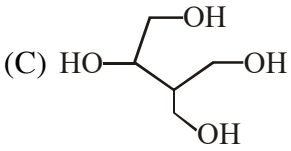
RD0013

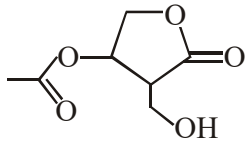
14.  Product

The product is :

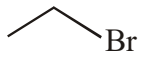
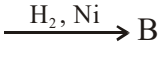
(A) 

(B) $CH_3–CH_2–OH$

(C) 

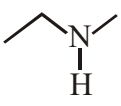
(D) 

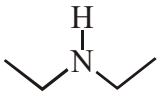
RD0014

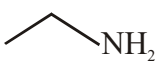
15.  A  B

The final product (B) is :

(A) 

(B) 

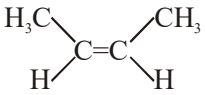
(C) 

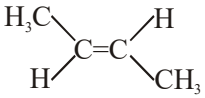
(D) 

RD0015

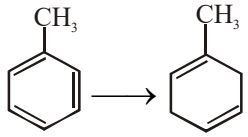
EXERCISE # O-II

1. **Column-I (Conversion)** | **Column-II (Required Reagent)**

(A) $H_3C–C\equiv C–CH_3 \longrightarrow$  (P) H_2 , Pd-$BaSO_4$

(B) $H_3C–C\equiv C–CH_3 \longrightarrow$  (Q) Li , Liq.NH_3

(C) $H_3C–CH=CH–CH_3 \longrightarrow H_3C–CH_2–CH_2–CH_3$ (R) N_2H_4 / H_2O_2 (1 eq.)

(D)  (S) B_2H_6 , CH_3COOH

RD0016

2. $R–\overset{\overset{O}{\|}}{C}–R \xrightarrow{?} R–CH_2–R$

Identify numbers of reagent that can be used for above conversion.

(a) Zn–Hg / HCl (b) $LiAlH_4$ (c) N_2H_4 / $\overset{\ominus}{O}H$ (d) $\begin{matrix}CH_2–SH\\|\\CH_2–SH\end{matrix}$, H_2 / Ni

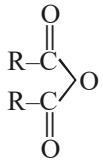
RD0017

3. **Column-I (Conversion)** | **Column-II (Required Reagent)**

(A) $–\overset{\overset{O}{\|}}{C}–OH \longrightarrow –CH_2OH$ (P) $LiAlH_4$

(B) $–\overset{\overset{O}{\|}}{C}–Cl \longrightarrow –CH_2OH$ (Q) $NaBH_4$

(C) $–\overset{\overset{O}{\|}}{C}–OR' \longrightarrow –CH_2OH$ (R) B_2H_6 / THF

(D)  $\longrightarrow R–CH_2OH$ (S) H_2 / Catalyst, Δ

RD0018

4. How many alkene on catalytic reduction gives following product.

(i) (A) $\xrightarrow{H_2/Pt}$ n-butane

(ii) (B) $\xrightarrow{H_2/Pt}$ Iso-pentane

(iii) (C) $\xrightarrow{H_2/Pt}$ Neo-pentane

(iv) (D) $\xrightarrow{H_2/Pt}$ Cyclopentane

(v) (E) $\xrightarrow[Pt]{H_2}$ 

RD0019

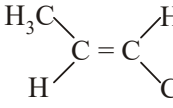
5. Give the expected major product for each reaction, including stereochemistry where applicable.

(a) $CH_3–CH_2–CH=CH_2 \xrightarrow[Pt]{H_2} P_1$

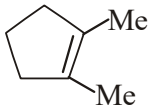
RD0020

(b)  $\xrightarrow[Pt]{D_2} P_2$

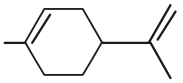
RD0021

(c)  $\xrightarrow[Ni]{D_2} P_3$

RD0022

(d)  $\xrightarrow{Ni/H_2} P_4$

RD0023

(e)  $\xrightarrow[excess]{H_2/Pt} P_5$; $\xrightarrow[1\ eq.]{H_2/Pt} P_6$

RD0024

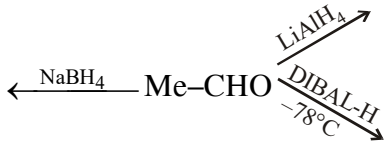
6. **(i)** $CH_2=CH–CH_2–CH=O \xrightarrow[excess]{H_2/Pt} P_1$

RD0025

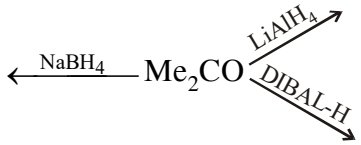
(ii) $CH_2=CH–CH_2–C\equiv N \xrightarrow[excess]{H_2/Pt} P_2$

RD0026

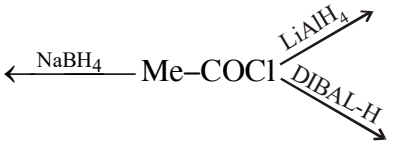
7. **(a) Identify the product ?**

(i) 

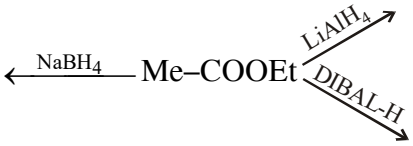
RD0027

(ii) 

RD0028

(iii) 

RD0029

(iv) 

RD0030

(b) Identify the product ?

(v) $\xleftarrow{NaBH_4}$ Me–COOH $\xrightarrow{LiAlH_4}$

RD0031

(vi) $\xleftarrow{NaBH_4}$ Me–COOMe $\xrightarrow{LiAlH_4}$

RD0031

(vii) $\xleftarrow{NaBH_4}$ $Me–CONH_2$ $\xrightarrow{LiAlH_4}$

RD0031

(viii) $\xleftarrow{NaBH_4}$ Me–CONH–Me $\xrightarrow{LiAlH_4}$

RD0032

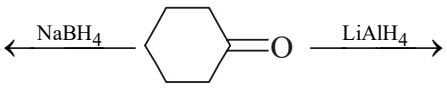
(ix) $\xleftarrow{NaBH_4}$ $Me–CONMe_2$ $\xrightarrow{LiAlH_4}$

RD0032

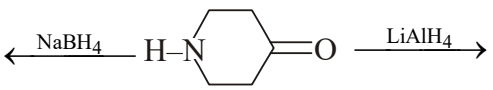
(x) $\xleftarrow{NaBH_4}$ $CH_3–CH = CH_2$ $\xrightarrow{LiAlH_4}$

RD0032

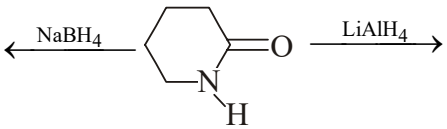
8. Give product in following reactions.

(i) 

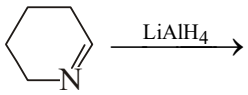
RD0033

(ii) 

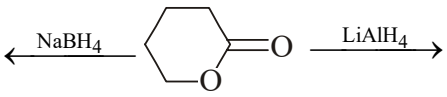
RD0034

(iii) 

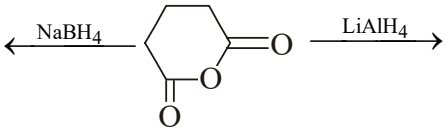
RD0033

(iv) 

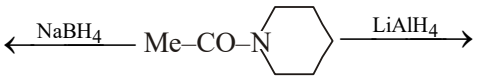
RD0034

(v) 

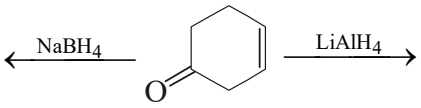
RD0033

(vi) 

RD0034

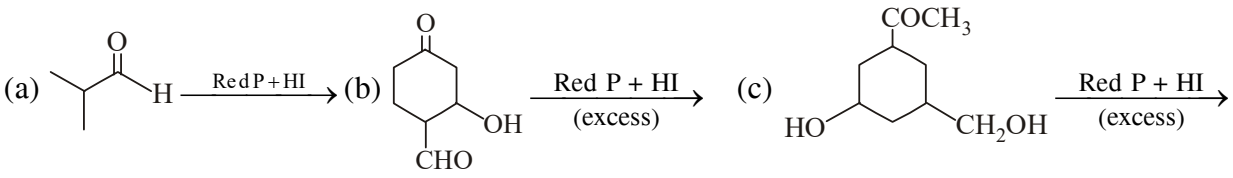
(vii) 

RD0033

(viii) 

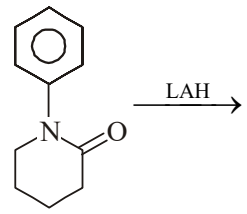
RD0034

9. Give product in following reactions.

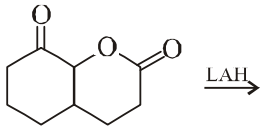


RD0035

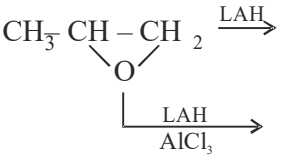
10. Give product in following reactions.

(i) 

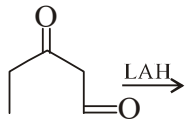
RD0036

(ii) 

RD0037

(iii) 

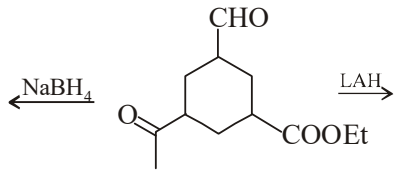
RD0038

(iv) 

RD0039

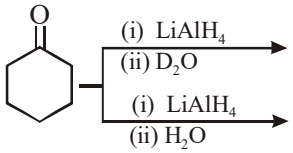
11. Give product in following reactions.

(i)



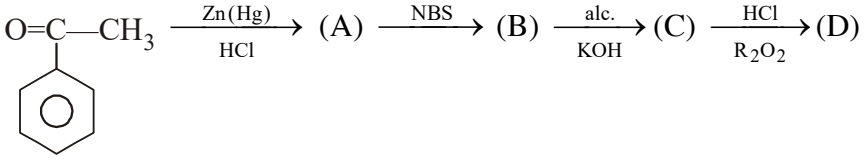
RD0040

(ii)



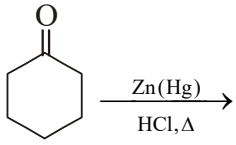
RD0041

12. (a)



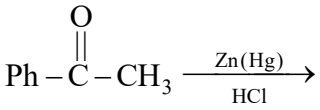
RD0042

13. (a)



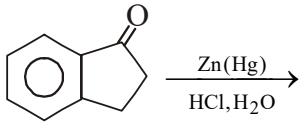
RD0043

(b)



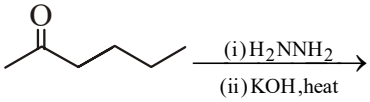
RD0043

(c)



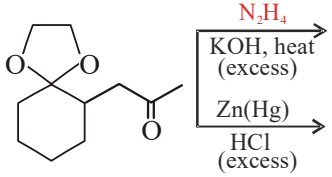
RD0043

(d)

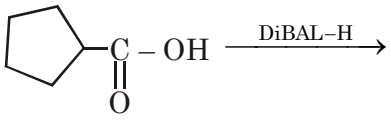


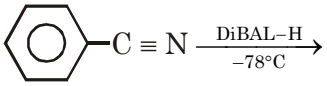
RD0044

(e)



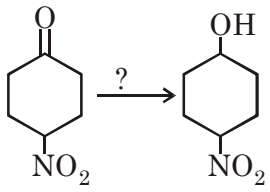
RD0045

14. **(a)** 

(b) 

(c) $CH_3 - \underset{\underset{O}{\|}}{C} - O - CH_2 - CH_3 \xrightarrow[-78°C]{DiBAL-H}$

RD0046

15. 

above conversion can be achieved by

(A) $LiAlH_4$ (B) H_2 / Ni, Δ (C) $NaBH_4$ (D) All

RD0047

EXERCISE # (MAIN & ADVANCE)

1. Hydrogenation of benzoyl chloride in the presence of Pd and $BaSO_4$ gives : **[IIT 1992]**

(A) Benzyl alcohol (B) Benzaldehyde (C) Benzoic acid (D) Phenol

RD0048

2. Draw the stereochemical structure of the product in the following reactions. **[IIT 1994]**

$$R–C\equiv C–R \xrightarrow[\text{Lindlar catalyst}]{H_2}$$

RD0049

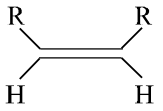
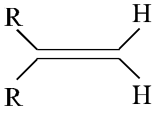
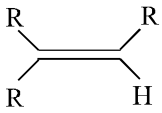
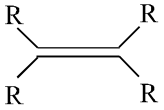
3. Under Wolff Kishner reduction conditions, the conversions which may be brought about is?

(A) Benzaldehyde into Benzyl alcohol (B) Cyclohexanol into Cyclohexane

(C) Cyclohexanone into Cyclohexanol (D) Benzophenone into Diphenylmethane **[IIT 1995]**

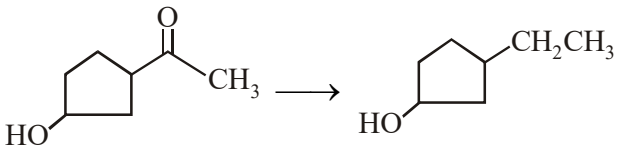
RD0050

4. Which one of the following alkenes will react fastest with H_2 under catalytic hydrogenation condition **[IIT 2000]**

(A)  (B)  (C)  (D) 

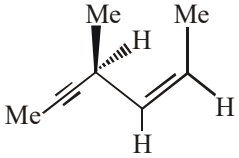
RD0051

5. The appropriate reagent for the following transformation: **[IIT-2000]**



(A) Zn(Hg), HCl (B) NH_2NH_2, OH^- (C) H_2/Ni (D) $NaBH_4$

RD0052

6. 

Hydrogenation of the above compound in the presence of poisoned paladium catalyst gives-

(A) An optically active compound (B) An optically inactive compound **[IIT-2001]**

(C) A racemic mixture (D) A diastereomeric mixture

RD0053

7. When $CH_2=CH–COOH$ is reduced with $LiAlH_4$, the compound obtained will be -**[AIEEE-2003]**

(A) $CH_3–CH_2–CH_2OH$ (B) $CH_3–CH_2–CHO$

(C) $CH_3–CH_2–COOH$ (D) $CH_2=CH–CH_2OH$

RD0054

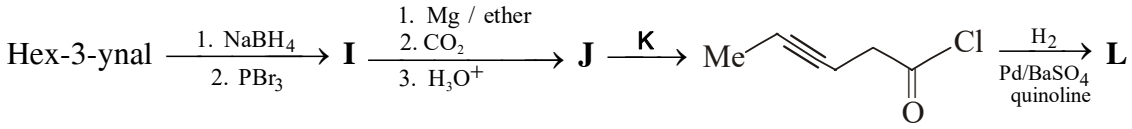
8. 2–hexyne can be converted into trans–2–hexene by the action of : **[IIT-2004]**

(A) H_2–Pd-$BaSO_4$ (B) Li in liq. NH_3 (C) H_2–PtO_2 (D) $NaBH_4$

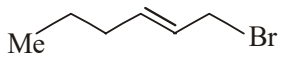
RD0055

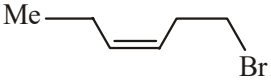
Paragraph for Question No. 09 to 11

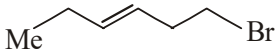
In the following reaction sequence, products **I**, **J** and **L** are formed. **K** represents a reagent.

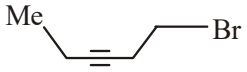


9. The structure of the product **I** is **[JEE-2008]**

(A) 

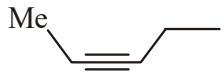
(B) 

(C) 

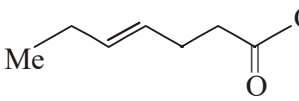
(D) 

RD0056

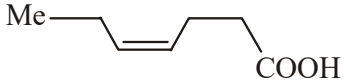
10. The structures of compounds **J** and **K**, respectively, are **[JEE-2008]**

(A) 

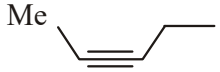
and $SOCl_2$

(B) 

and SO_2Cl_2

(C) 

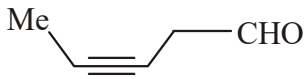
and $SOCl_2$

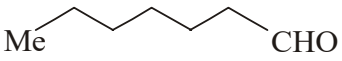
(D) 

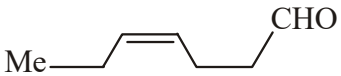
and CH_3SO_2Cl

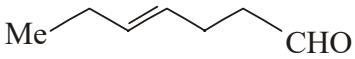
RD0056

11. The structure of product **L** is : **[JEE-2008]**

(A) 

(B) 

(C) 

(D) 

RD0056

12. Consider all possible isomeric ketones including stereoisomers of MW = 100, All these isomers are independently reacted with $NaBH_4$ (**NOTE :** stereoisomers are also reacted separately). The total number of ketones that give a racemic product(s) is/are. **[JEE-2014]**

RD0057

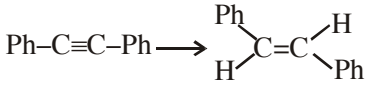
13. In the reaction, **[JEE-2014]**

$CH_3COOH \xrightarrow{LiAlH_4} A \xrightarrow{PCl_5} B \xrightarrow{AlC.KOH} C$, Final product C is.

(A) Ethylene (B) Acetyl chloride (C) Acetaldehyde (D) Acetylene

RD0058

14. The reagent needed for converting is : **[JEE-2014]**

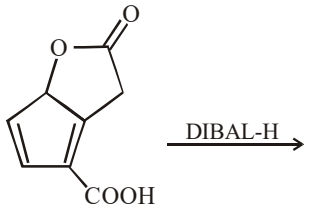


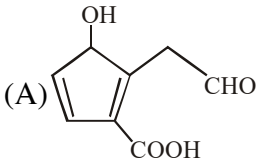
(A) H_2 / Lindlar Cat. (B) Cat. hydrogenation

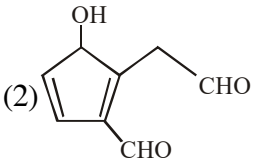
(C) $LiAlH_4$ (D) Li / NH_3

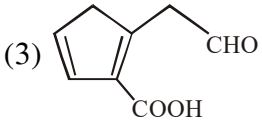
RD0059

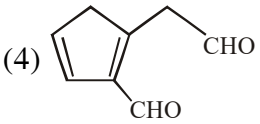
15. The major product obtained in the following reaction is : **[JEE MAIN-2017]**



(A) 

(2) 

(3) 

(4) 

RD0060

16. Glucose on prolonged heating with HI gives : **[Jee Main - 2018]**

(1) 1–Hexene (2) Hexanoic acid (3) 6-iodohexanal (4) n-Hexane

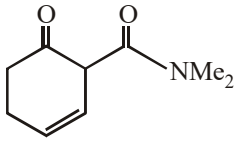
RD0061

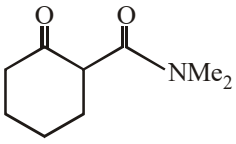
17. The *trans*-alkenes are formed by the reduction of alkynes with : **[Jee Main - 2018]**

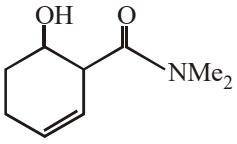
(1) $NaBH_4$ (2) Na/liq.NH_3 (3) Sn–HCl (4) H_2–Pd/C, $BaSO_4$

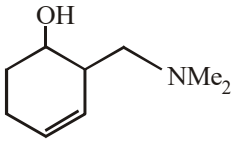
RD0062

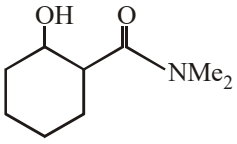
18. The main reduction product of the following compound with $NaBH_4$ in methanol is :- **[Jee Main - Onl_line Morning 2018]**



(1) 

(2) 

(3) 

(4) 

RD0063

19. The major product of following reaction is : **[JEE-MAIN-On-line-(Jan)-2019]**

$R-C\equiv N \xrightarrow[(2)H_2O]{(1)AlH(i-Bu_2)} ?$

(1) RCHO (2) RCOOH (3) RCH_2NH_2 (4) $RCONH_2$

RD0064

20. Wilkinson catalyst is : **[JEE-MAIN-On-line-(Jan)-2019]**

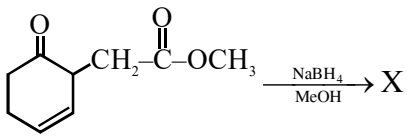
(1) $[(Ph_3P)_3RhCl]$ (Et = C_2H_5) (2) $[Et_3P)_3IrCl]$

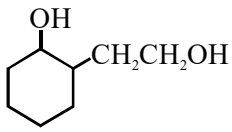
(3) $[Et_3P)_3RhCl]$ (4) $[Ph_3P)_3IrCl]$

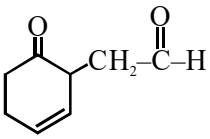
RD0065

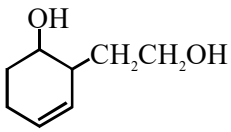
21. The major product 'X' formed in the following reaction is :

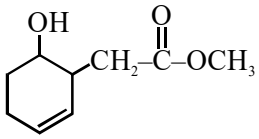
[JEE-MAIN-On-line-(Jan)-2019]



(1) 

(2) 

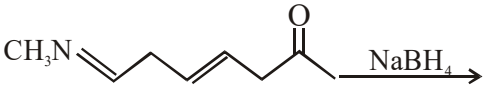
(3) 

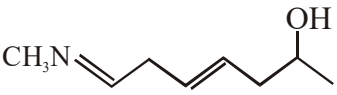
(4) 

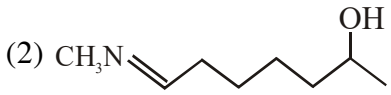
RD0066

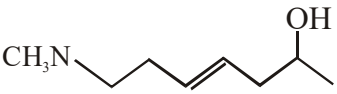
22. The major product of the following recation is:

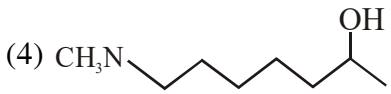
[JEE-MAIN-On-line-(Jan)-2019]



(1) 

(2) 

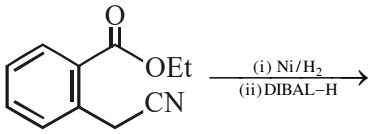
(3) 

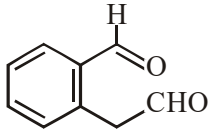
(4) 

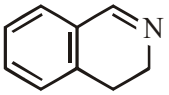
RD0067

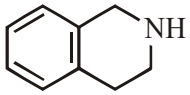
23. The major product of the following reaction is:

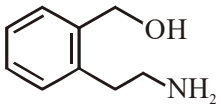
[JEE-MAIN-On-line-(Jan)-2019]



(1) 

(2) 

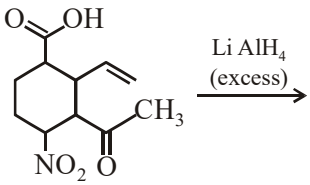
(3) 

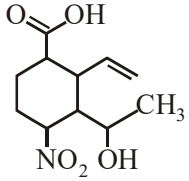
(4) 

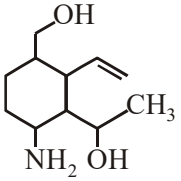
RD0068

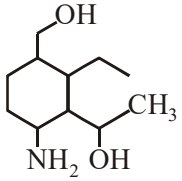
24. The major product obtained in the following reaction is :-

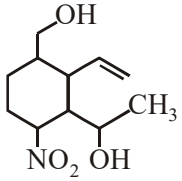
[JEE-MAIN-On-line-(Jan)-2019]



(1) 

(2) 

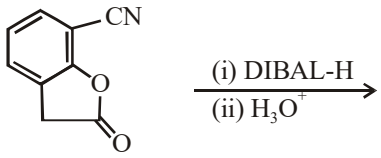
(3) 

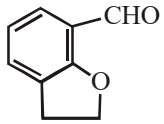
(4) 

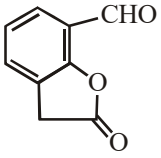
RD0069

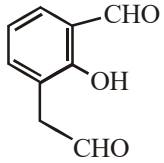
25. The major product of the following reaction is:

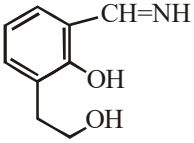
[JEE-MAIN-On-line-(Jan)-2019]



(1) 

(2) 

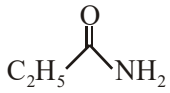
(3) 

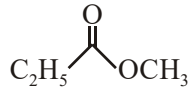
(4) 

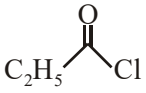
RD0070

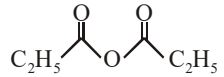
26. The increasing order of the reactivity of the following with $LiAlH_4$ is :

[JEE-MAIN-On-line-(Jan)-2019]

(A) 

(B) 

(C) 

(D) 

(1) (A) < (B) < (D) < (C)

(2) (A) < (B) < (C) < (D)

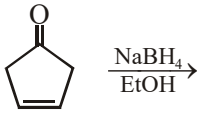
(3) (B) < (A) < (D) < (C)

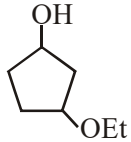
(4) (B) < (A) < (C) < (D)

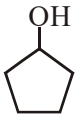
RD0071

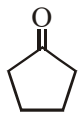
27. The major product of the following reaction is:

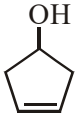
[JEE-MAIN-On-line-(Jan)-2019]



(1) 

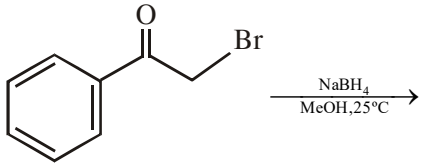
(2) 

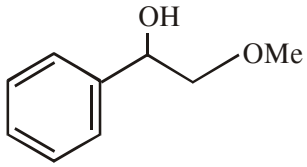
(3) 

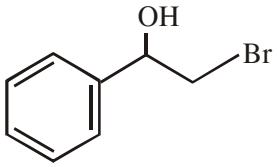
(4) 

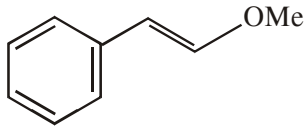
RD0072

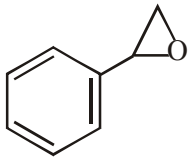
28. The major product of the following reaction is: **[JEE-MAIN-On-line-(April)-2019]**



(1) 

(2) 

(3) 

(4) 

RD0073

29. The major product of the following reaction is : **[JEE-MAIN-On-line-(April)-2019]**

$CH_3CH = CHCO_2CH_3 \xrightarrow{LiAlH_4}$

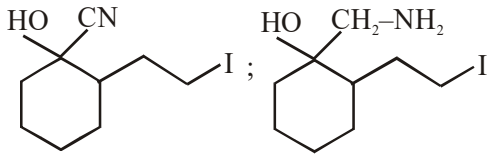
(1) $CH_3CH_2CH_2CHO$ (2) $CH_3CH = CHCH_2OH$

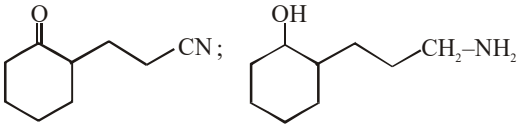
(3) $CH_3CH_2CH_2CO_2CH_3$ (4) $CH_3CH_2CH_2CH_2OH$

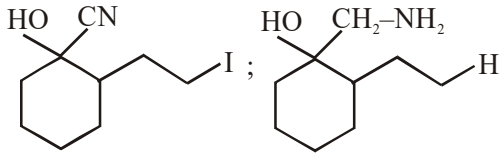
RD0074

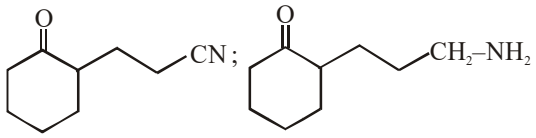
30. The major products A and B for the following reactions are, respectively:

[JEE-MAIN-On-line-(April)-2019]

(1) 

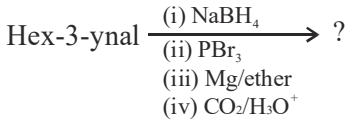
(2) 

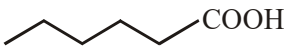
(3) 

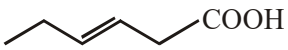
(4) 

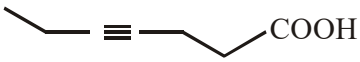
RD0075

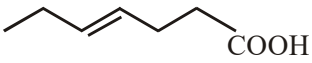
31. What is the product of following reaction ? **[JEE-MAIN-On-line-(Jan)-2020]**



(1) 

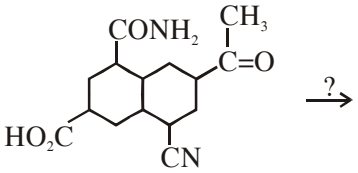
(2) 

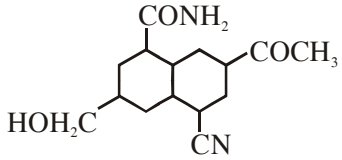
(3) 

(4) 

RD0076

32. The most suitable reagent for the given conversion is : **[JEE-MAIN-On-line-(Jan)-2020]**





(1) $LiAlH_4$ (2) $NaBH_4$ (3) H_2/Pd (4) B_2H_6

RD0077

ANSWER KEY

EXERCISE # O-I

1.	**Ans. (D)**	**2.**	**Ans. (B)**	**3.**	**Ans. (C)**	**4.**	**Ans. (C)**	**5.**	**Ans. (A)**
6.	**Ans. (C)**	**7.**	**Ans. (B)**	**8.**	**Ans. (B)**	**9.**	**Ans. (C)**	**10.**	**Ans. (B)**
11.	**Ans. (C)**	**12.**	**Ans. (C)**	**13.**	**Ans. (D)**	**14.**	**Ans. (D)**	**15.**	**Ans. (B)**

EXERCISE # O-II

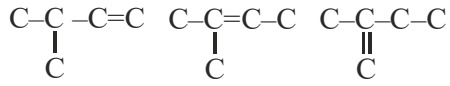
1. **Ans. (A)→P, R, S; (B)→Q; (C)→R, S; (D)→Q**

2. **Ans. (3)**

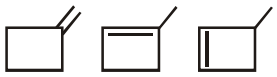
3. **Ans. (A)→P, R, S; (B)→P, Q, S; (C)→P, R, S; (D)→P, R, S**

4. **Ans.**

(i) Cis & trans 2-butene & 1-butene ;

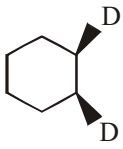
(ii) 

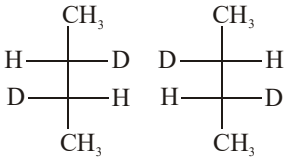
(iii) Zero (Neo-pentane can not be prepared by catalytic hydrogenation of alkene) ; (iv) One

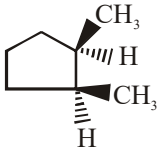
(v) Including optical = 4  , Excluding optical = 3.

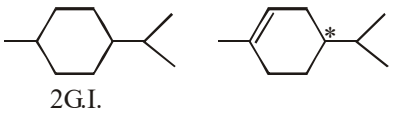
5. **Ans.**

(a) 

(b) 

(c) 

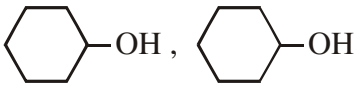
(d) 

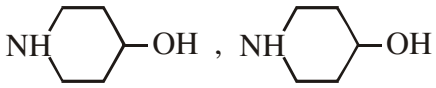
(e) 

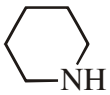
2G.I.

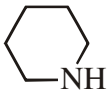
6. **Ans.** (i) $P_1 = CH_3–CH_2–CH_2–CH_2–OH$ (ii) $P_2 = CH_3–CH_2–CH_2–CH_2–NH_2$

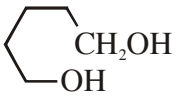
8. **Ans.**

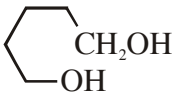
(i) 

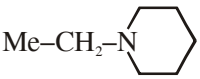
(ii) 

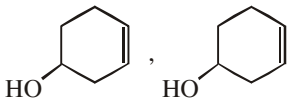
(iii) No reaction, 

(iv) 

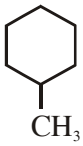
(v) No reaction, 

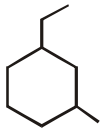
(vi) No reaction, 

(vii) No reaction, 

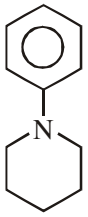
(viii) 

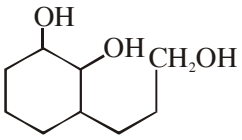
9. **Ans.** (a) 

(b) 

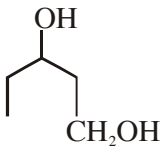
(c) 

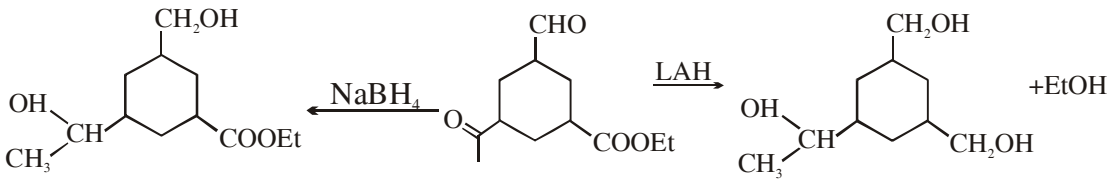
10. Ans.

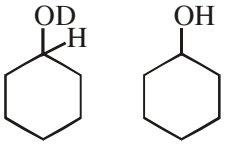
(i) 

(ii) 

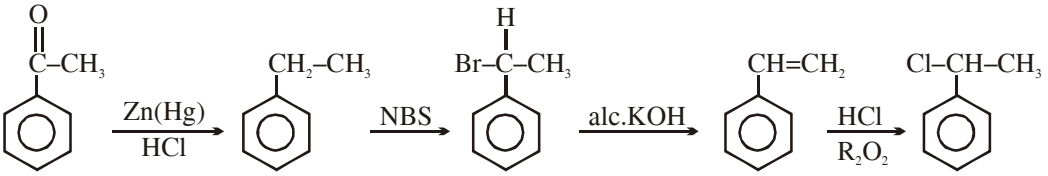
(iii) $CH_3–\underset{\displaystyle OH}{\underset{|}{CH}}–CH_3$, $CH_3–CH_2–CH_2OH$

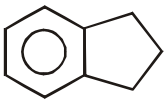
(iv) 

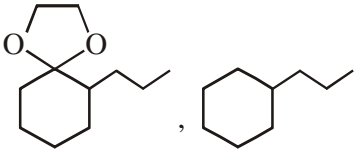
11. Ans. (i) 

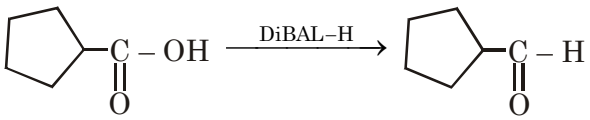
(ii) 

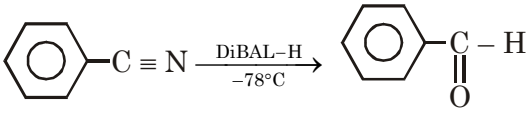
12. Ans.

(a) 

13. (a)  (b) $Ph–CH_2–CH_3$ (c)  (d) 

(e) 

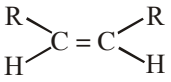
14. (a) 

(b) 

(c) CH_3 – CHO

15. Ans.(C)

EXERCISE # (MAIN & ADVANCE)

1. **Ans. (B)** **2.** **Ans.**  **3.** **Ans. (D)** **4.** **Ans. (A)**

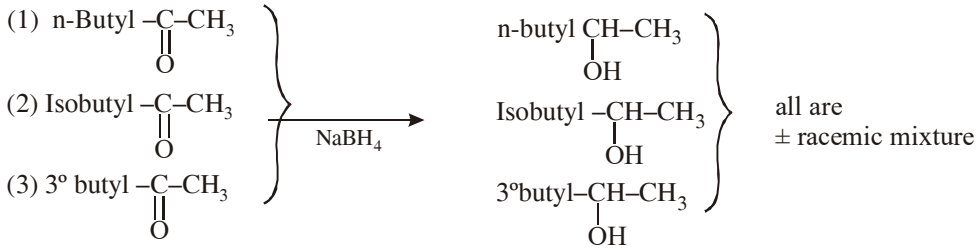
5. **Ans. (B)** **6.** **Ans. (B)** **7.** **Ans. (D)** **8.** **Ans. (B)**

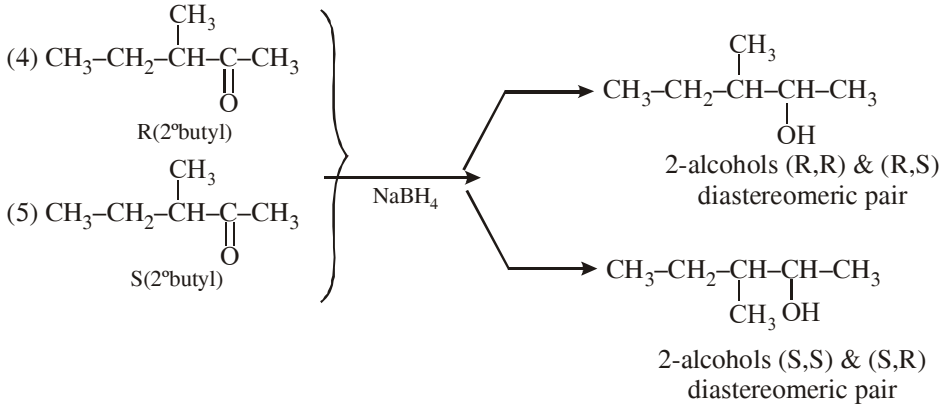
9. **Ans. (D)** **10.** **Ans. (A)** **11.** **Ans. (C)**

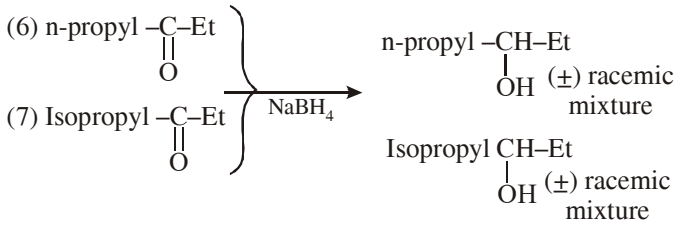
12. **Ans. (5)**

M. wt 100 of ketone

So m. formula = $C_6H_{12}O$

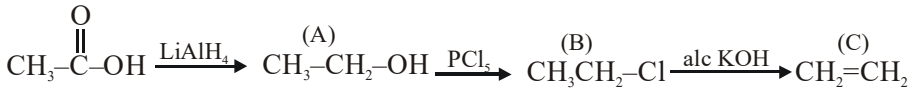






(1 ; 2 ; 3 ; 6 ; 7)

13. **Ans.** (A)



Reduction of CH_3COOH will produce $CH_3–CH_2–OH$(A)

Now reaction of $CH_3–CH_2–OH$(A) with PCl_5 will produce $CH_3–CH_2–Cl$(B).

Now alcoholic KOH with Et–Cl will produce $CH_2=CH_2$(C) Ethylene by E_2 elemination.

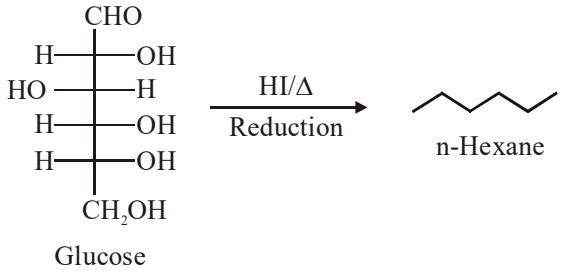
Thus Ans. is (1) Ethylene.

14. **Ans. (D)**

15. **Ans. (1)**

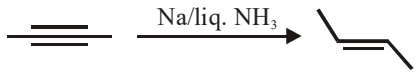
16. Ans. (4)

Sol.



17. Ans. (2)

Sol.



Birch reduction is anti addition. So trans alkene will be produced.

18. Ans. (4) **19. Ans. (1)** **20. Ans. (1)** **21. Ans. (4)**

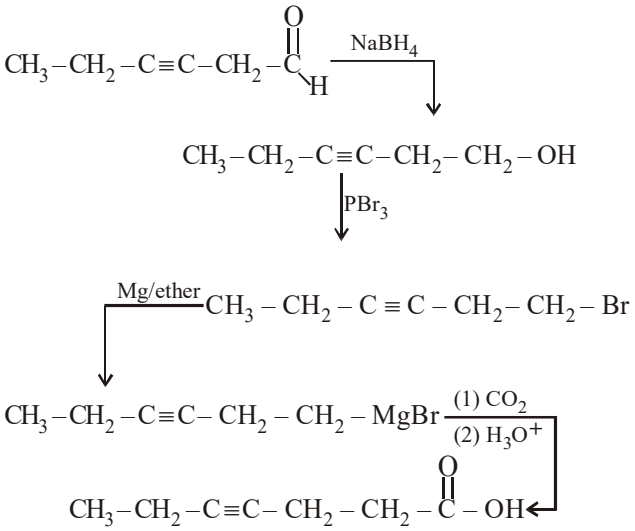
22. Ans.(3) **23. Ans. (2)** **24. Ans. (2)** **25. Ans. (3)**

26. Ans. (1) **27. Ans. (4)** **28. Ans. (4)** **29. Ans. (2)**

30. Ans. (2)

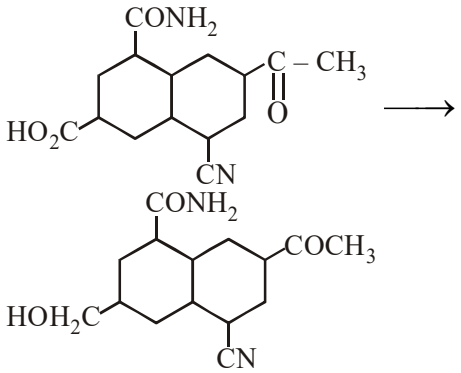
31. Ans. (3)

Sol.



32. Ans. (4)

Sol.



Most suitable reagent for given conversion is B_2H_6 (electrophilic reducing agent)