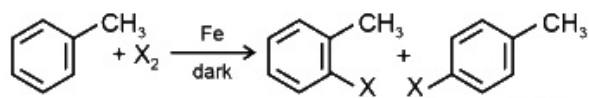


# Haloalkanes and Haloarenes

## Question1

The following reaction method



is not suitable for the preparation of the corresponding haloarene products, due to high reactivity of halogen, when X is :

[NEET 2024 Re]

Options:

A.

F

B.

I

C.

Cl

D.

Br

Answer: A

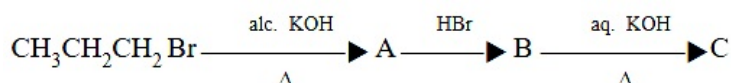
Solution:

- Aryl chlorides and bromides can easily be prepared by electrophilic substitution of arenes (toluene) with  $\text{Cl}_2$  and  $\text{Br}_2$  respectively in the presence of Lewis acid catalyst (  $\text{Fe}$  in dark).
- Reaction with  $\text{I}_2$  is reversible and requires the presence of oxidising agent.
- Corresponding fluoroarene is not prepared by this method due to high reactivity of fluorine.

Hence, 'X' is F .

## Question2

The major product C in the below mentioned reaction is:



## [NEET 2024 Re]

### Options:

A.

Propan-1-ol

B.

Propan-2-ol

C.

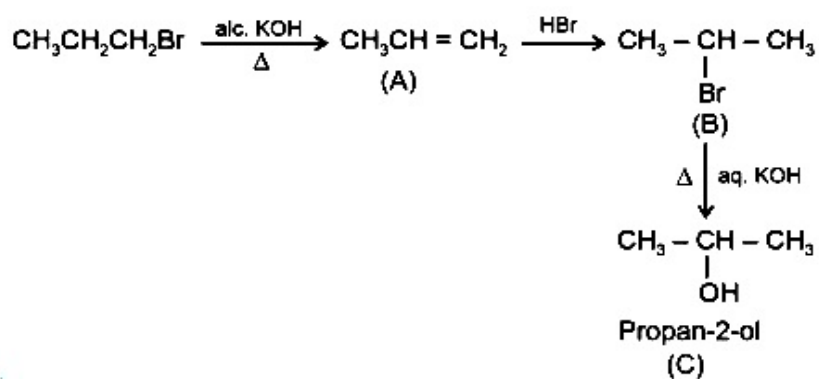
Propane

D.

Propyne

**Answer: B**

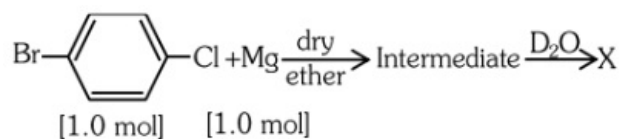
### Solution:



---

## Question3

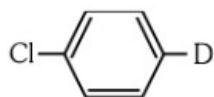
Identify ' X ' in the following reaction.



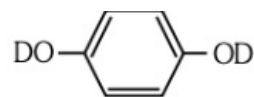
## [NEET 2024]

### Options:

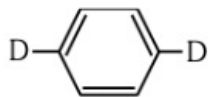
A.



B.



C.

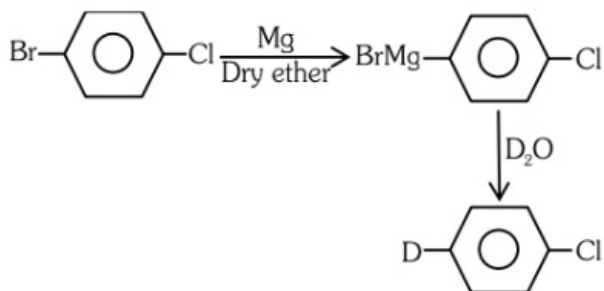


D.



**Answer: A**

**Solution:**



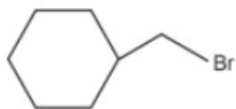
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## Question4

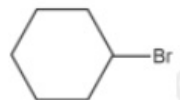
The compound that will undergo  $\text{S}_{\text{N}}1$  reaction with the fastest rate is  
[NEET 2024]

**Options:**

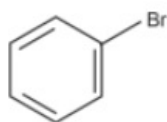
A.



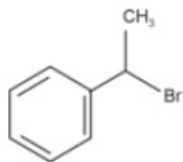
B.



C.



D.

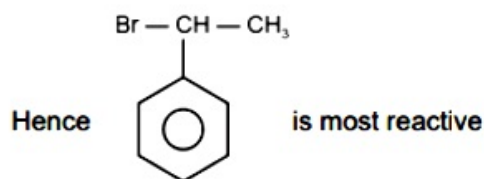
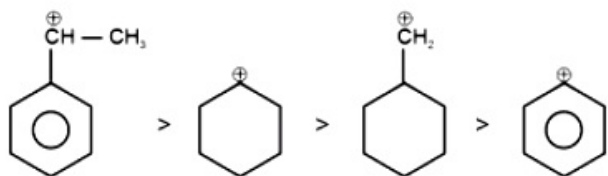


**Answer: D**

### Solution:

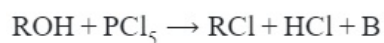
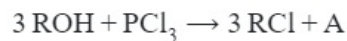
Reactivity towards  $S_N1$  depends upon stability of carbocation.

Order of stability is



## Question5

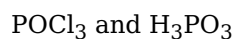
The products A and B obtained in the following reactions, respectively, are



**[NEET 2024]**

**Options:**

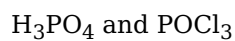
A.



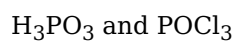
B.



C.



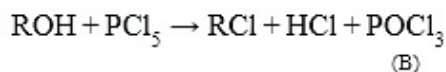
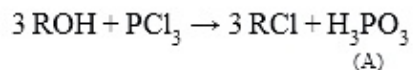
D.



**Answer: D**

### Solution:

These reactions are preparation of haloalkanes from alcohols.

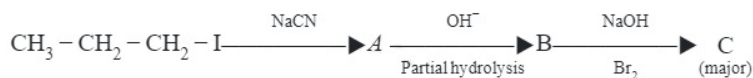


A and B are  $\text{H}_3\text{PO}_3$  and  $\text{POCl}_3$  respectively.

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## Question6

Identify the major product C formed in the following reaction sequence:



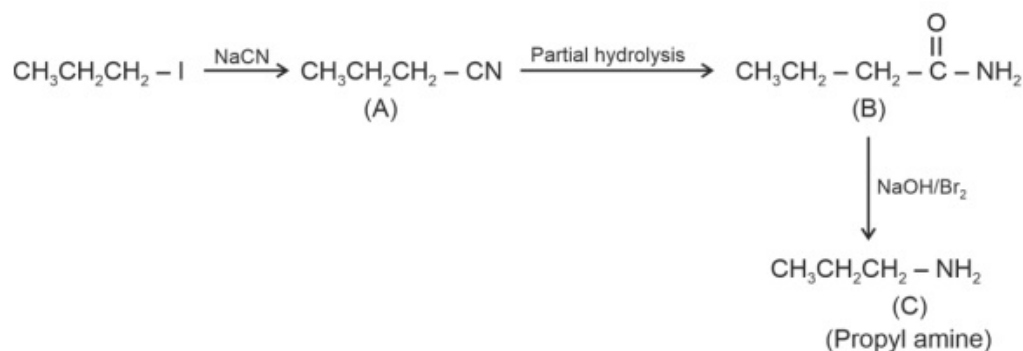
[NEET 2024]

Options:

- A.  
propylamine
- B.  
butylamine
- C.  
butanamide
- D.  
 $\alpha$ -bromobutanoic acid

Answer: A

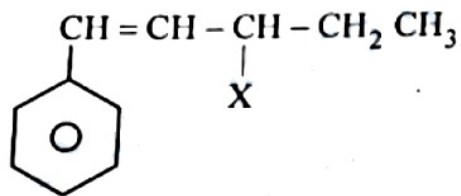
Solution:



- Step-I is  $\text{S}_{\text{N}}^2$  reaction with  $\text{CN}^-$  nucleophile.
  - Step-II will give amide.
  - Step-III is Hoffmann bromamide degradation reaction.
- 

## Question7

The given compound is an example of



[NEET 2023]

Options:

A.

Aryl halide

B.

Allylic halide

C.

Vinyl halide

D.

Benzylic halide

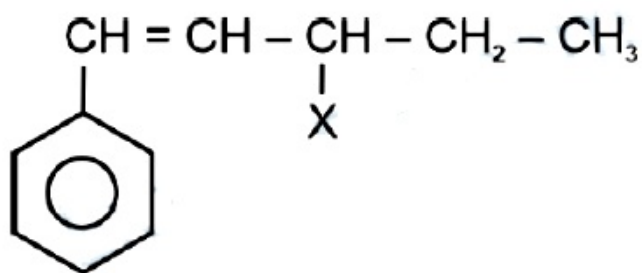
**Answer: B**

**Solution:**

$\alpha$ -carbon is  $sp^3$  carbon which is right next to  $>\text{C}=\text{C}<$

This  $\alpha$ -position is known as allylic position

Hence,

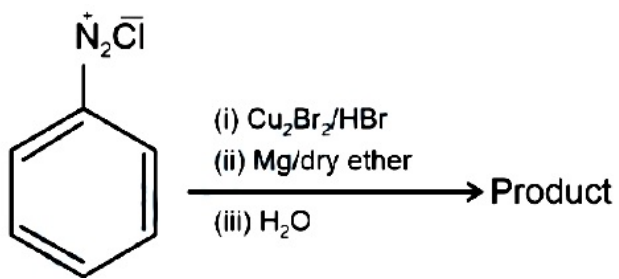


is allylic halide

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## Question8

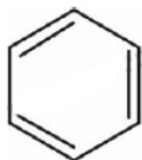
Identify the product in the following reaction:



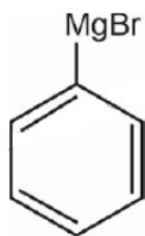
**[NEET 2023]**

**Options:**

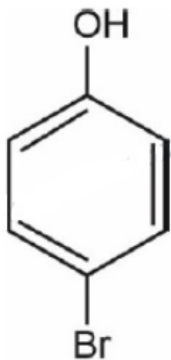
A.



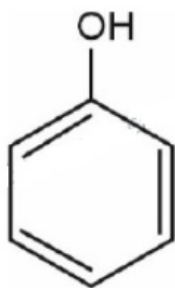
B.



C.

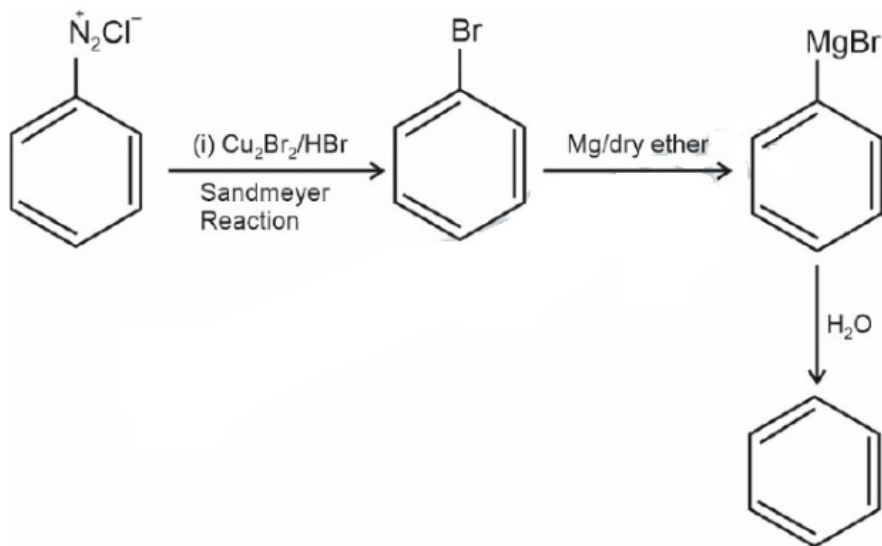


D.



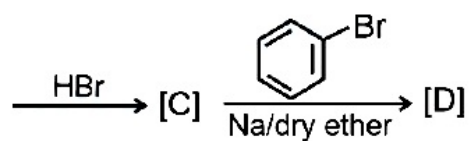
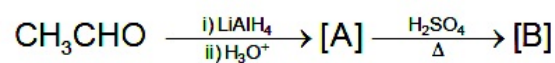
**Answer: A**

**Solution:**



## Question9

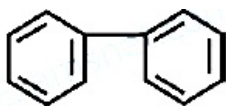
Identify the final product [D] obtained in the following sequence of reactions.



[NEET 2023]

Options:

A.



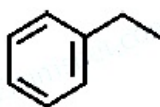
B.



C.



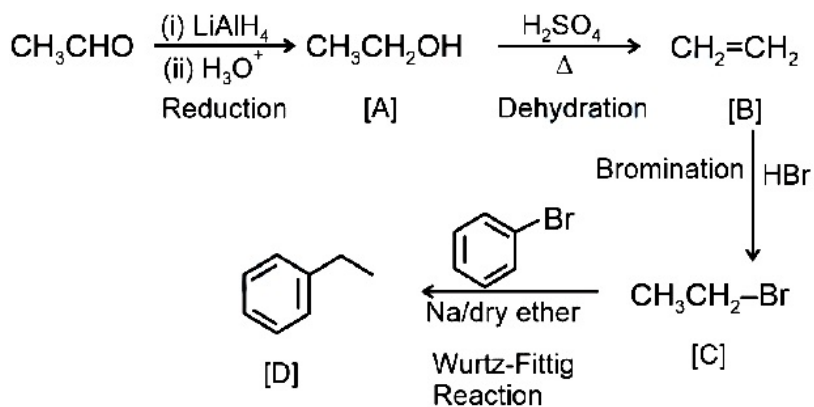
D.



Answer: C

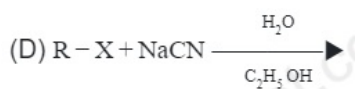
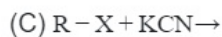
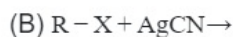
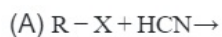
Solution:





## Question10

Which amongst the following reactions of alkyl halides produces isonitrile as a major product?



Choose the most appropriate answer from the options given below :

[NEET 2023 mpr]

Options:

A.

(D) only

B.

(C) and (D) only

C.

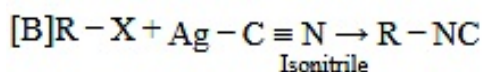
(B) only

D.

(A) and (B) only

**Answer: C**

**Solution:**



## Question11

Choose the correct sequence of reagents in the conversion of 4-nitrotoluene to 2-bromotoluene.

[NEET 2023 mpr]

Options:

A.

$\text{NaNO}_2/\text{HCl}$ ;  $\text{Sn}/\text{HCl}$ ;  $\text{Br}_2$ ;  $\text{H}_2\text{O}/\text{H}_3\text{PO}_2$

B.

$\text{Sn}/\text{HCl}$ ;  $\text{NaNO}_2/\text{HCl}$ ;  $\text{Br}_2$ ;  $\text{H}_2\text{O}/\text{H}_3\text{PO}_2$

C.

$\text{Br}_2$ ;  $\text{Sn}/\text{HCl}$ ;  $\text{NaNO}_2/\text{HCl}$ ;  $\text{H}_2\text{O}/\text{H}_3\text{PO}_2$

D.

$\text{Sn}/\text{HCl}$ ;  $\text{Br}_2$ ;  $\text{NaNO}_2/\text{HCl}$ ;  $\text{H}_2\text{O}/\text{H}_3\text{PO}_2$

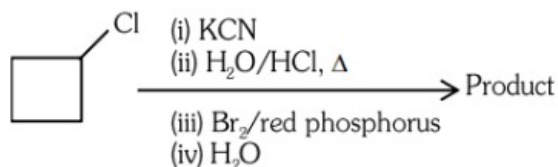
**Answer: C**

**Solution:**



## Question12

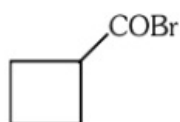
Identify the product in the following reaction



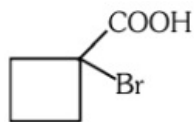
## [NEET 2023 mpr]

Options:

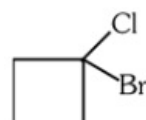
A.



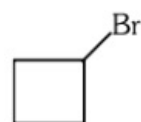
B.



C.

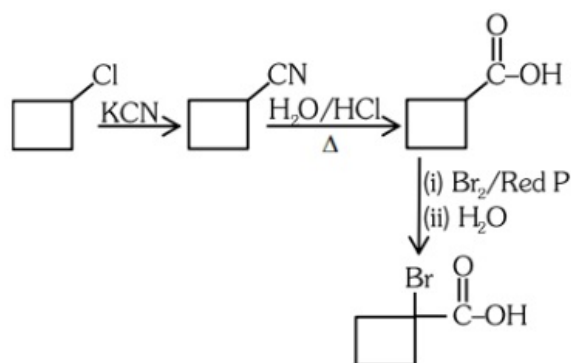


D.



**Answer: B**

**Solution:**



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## Question13

**The incorrect statement regarding chirality is  
[NEET-2022]**

Options:

A.  $\text{S}_{\text{N}}1$  reaction yields 1 : 1 mixture of both enantiomers

B. The product obtained by  $\text{S}_{\text{N}}2$  reaction of haloalkane having chirality at the reactive site shows inversion of configuration

C. Enantiomers are superimposable mirror images on each other

D. A racemic mixture shows zero optical rotation

**Answer: C**

**Solution:**

The stereoisomers related to each other as non-superimposable mirror image are called enantiomers.

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## Question14

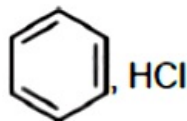
**Which of the following sequence of reactions is suitable to synthesize chlorobenzene?**  
**[NEET-2022]**

**Options:**

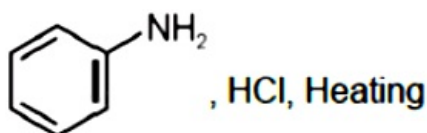
A. Benzene,  $\text{Cl}_2$ , anhydrous  $\text{FeCl}_3$

B. Phenol,  $\text{NaNO}_2$ ,  $\text{HCl}$ ,  $\text{CuCl}$

C.

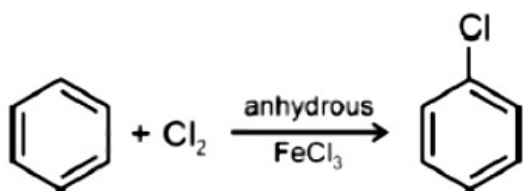


D.



**Answer: A**

**Solution:**



Benzene reacts with chlorine in presence of anhydrous  $\text{FeCl}_3$  to give chlorobenzene

-----

## Question15

**Given below are two statements : one is**

labelled as Assertion (A) and the other is labelled as Reason (R).

**Assertion (A) :**

**Chlorine is an electron withdrawing group but it is ortho, para directing in electrophilic aromatic substitution.**

**Reason (R) :**

**Inductive effect of chlorine destabilises the intermediate carbocation formed during the electrophilic substitution, however due to the more pronounced resonance effect, the halogen stabilises the carbocation at ortho and para positions.**

**In the light of the above statements, choose the most appropriate answer from the options given below :**

**[NEET Re-2022]**

**Options:**

- A. (A) is not correct but (R) is correct.
- B. Both (A) and (R) are correct and (R) is the correct explanation of (A).
- C. Both (A) and (R) are correct but (R) is not the correct explanation of (A).
- D. (A) is correct but (R) is not correct.

**Answer: D**

**Solution:**

Halogen are electron withdrawing groups due to high electronegativity. They have high electron withdrawing inductive effect. In electrophilic substitution reaction it shows both electron withdrawing inductive effect and electron donating resonance effect, but inductive effect overpowers resonance effect so, it deactivates benzene ring and directs the incoming electrophile to ortho and para positions. So, here assertion is correct and Reason is false.

-----

## Question16

**Predict the order of reactivity of the following four isomers towards  $S_N2$  reaction.**

**(I)  $CH_3CH_2CH_2CH_2Cl$**

**(II)  $CH_3CH_2CH(Cl)CH_3$**

**(III)  $(CH_3)_2CHCH_2Cl$**

**(IV)  $(\text{CH}_3)_3\text{CCl}$**   
**[NEET Re-2022]**

**Options:**

- A. (IV) > (II) > (III) > (I)
- B. (IV) > (III) > (II) > (I)
- C. (I) > (II) > (III) > (IV)
- D. (I) > (III) > (II) > (IV)

**Answer: D**

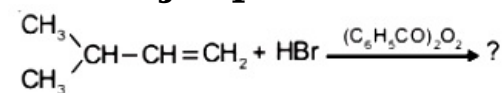
**Solution:**

Greater the steric hindrance, lesser will be the rate of  $S_N2$  reaction. So order of reactivity of  $S_N2$  reaction is- (I) > (III) > (II) > (IV)

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## Question17

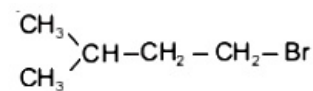
**The major product of the following chemical reaction is :**



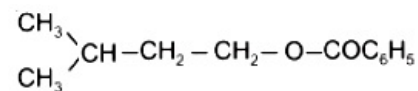
**[NEET 2021]**

**Options:**

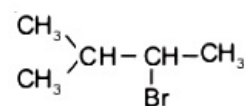
A.



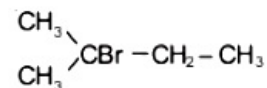
B.



C.

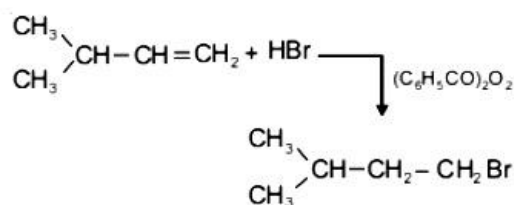


D.

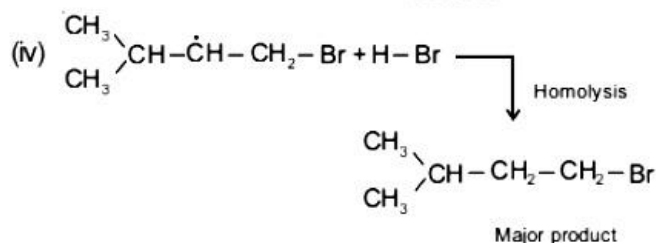
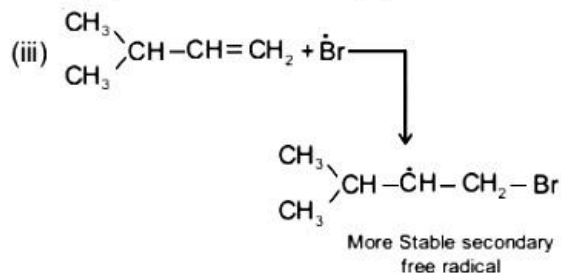
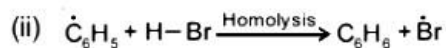
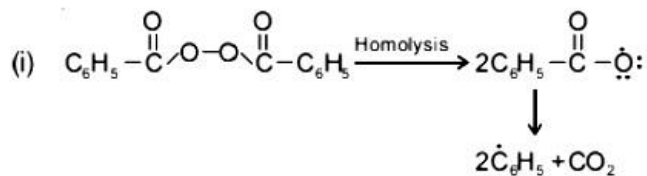


**Answer: A**

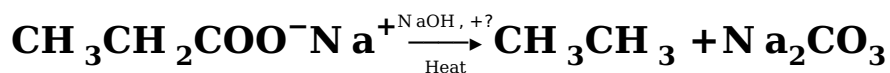
**Solution:**



Mechanism : Peroxide effect proceeds via free radical chain mechanism.



## Question18



Consider the above reaction and identify the missing reagent/chemical.  
[NEET 2021]

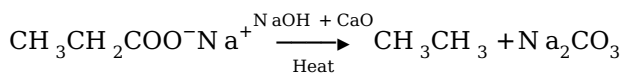
**Options:**

- A.  $\text{B}_2\text{H}_6$
- B. Red Phosphorus
- C. CaO
- D. DI BAL – H

**Answer: C**

**Solution:**

Alkane is produced by heating sodium salt of carboxylic acid with sodalime (NaOH and CaO in the ratio of 3 : 1)



## Question19

**Elimination reaction of 2-Bromo-pentane to form pent-2-ene is:**

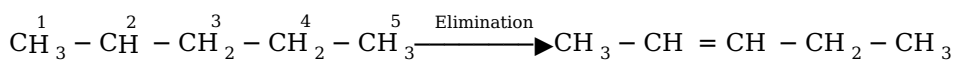
- (A)  $\beta$ -Elimination reaction  
 (B) Follows Zaitsev rule  
 (C) Dehydrohalogenation reaction  
 (D) Dehydration reaction  
 [2020]

**Options:**

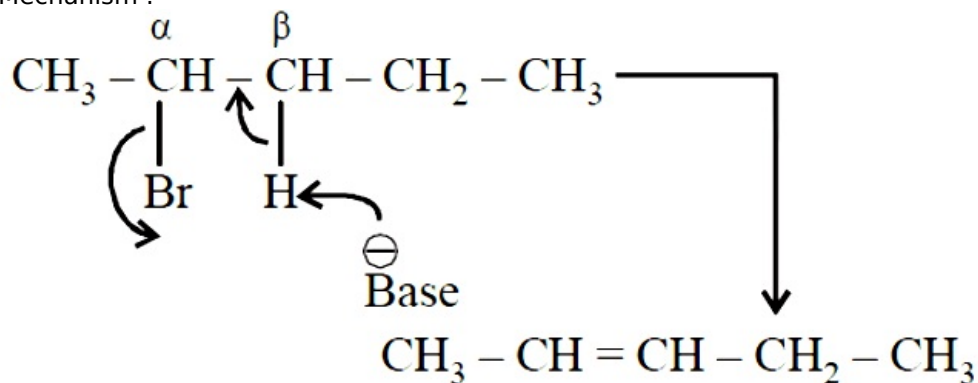
- A. (A), (C), (D)  
 B. (B), (C), (D)  
 C. (A), (B), (D)  
 D. (A), (B), (C)

**Answer: D**

**Solution:**



Mechanism :



It is an  $\beta$  elimination reaction as  $\beta$  hydrogen is abstracted and results more substituted alkene. which follows Zaitsev's rule

Since 'H' and 'Br' are removed, it is dehydrohalogenation.

---

## Question20

**Which of the following alkane cannot be made in good yield by Wurtz reaction?  
 [2020]**

**Options:**



A. 2,3-Dimethylbutane

B. n-Heptane

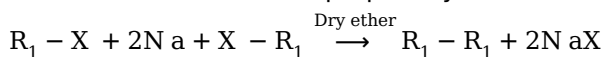
C. n-Butane

D. n-Hexane

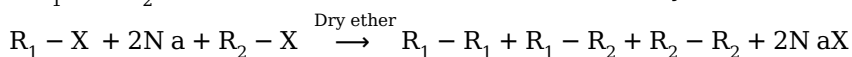
**Answer: B**

**Solution:**

Wurtz reaction is used to prepare symmetrical alkanes like  $R_1 - R_1$ , as



If  $R_1$  and  $R_2$  are different, then mixture of alkanes may be obtained as



Hence, n-heptane cannot be made in good yield using Wurtz reaction, since it is unsymmetrical alkane

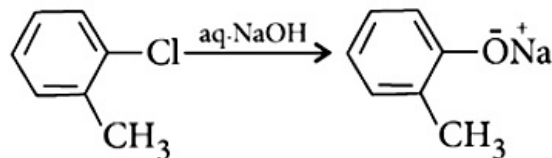
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## Question 21

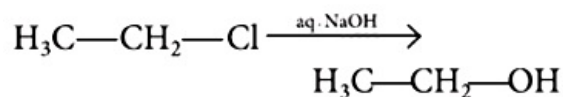
**The hydrolysis reaction that takes place at the slowest rate, among the following is (Odisha NEET 2019)**

**Options:**

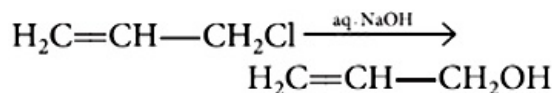
A.



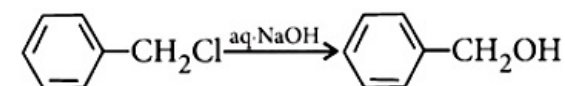
B.



C.



D.



**Answer: A**

## Solution:

Aryl halides are less reactive as compared to alkyl halides as the halogen atom in these compounds is firmly attached and cannot be replaced by nucleophiles such as  $\text{OH}^-$ ,  $\text{NH}_2^-$ , etc. In chlorobenzene, the electron pair of chlorine atom is in conjugation with  $\pi$ -electrons of benzene ring. Thus C – Cl bond acquires double bond character and is difficult to break.

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## Question22

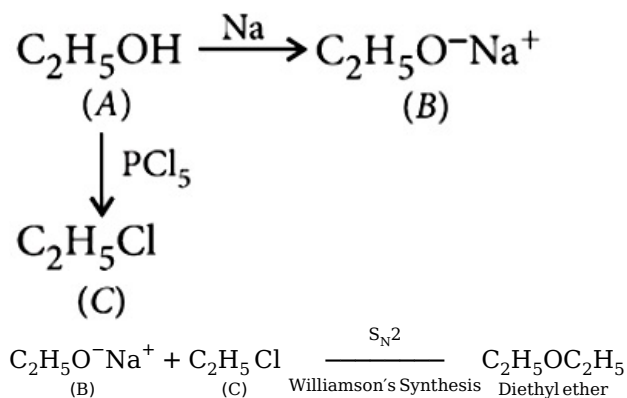
The compound A on treatment with Na gives B, and with  $\text{PCl}_5$  gives C. B and C react together to give diethyl ether. A, B and C are in the order (NEET 2018)

### Options:

- A.  $\text{C}_2\text{H}_5\text{OH}$ ,  $\text{C}_2\text{H}_6$ ,  $\text{C}_2\text{H}_5\text{Cl}$
- B.  $\text{C}_2\text{H}_5\text{OH}$ ,  $\text{C}_2\text{H}_5\text{Cl}$ ,  $\text{C}_2\text{H}_5\text{ONa}$
- C.  $\text{C}_2\text{H}_5\text{Cl}$ ,  $\text{C}_2\text{H}_6$ ,  $\text{C}_2\text{H}_5\text{OH}$
- D.  $\text{C}_2\text{H}_5\text{OH}$ ,  $\text{C}_2\text{H}_5\text{ONa}$ ,  $\text{C}_2\text{H}_5\text{Cl}$

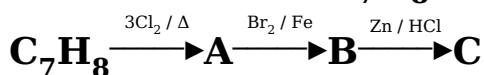
**Answer: D**

### Solution:



## Question23

The compound  $\text{C}_7\text{H}_8$  undergoes the following reactions:



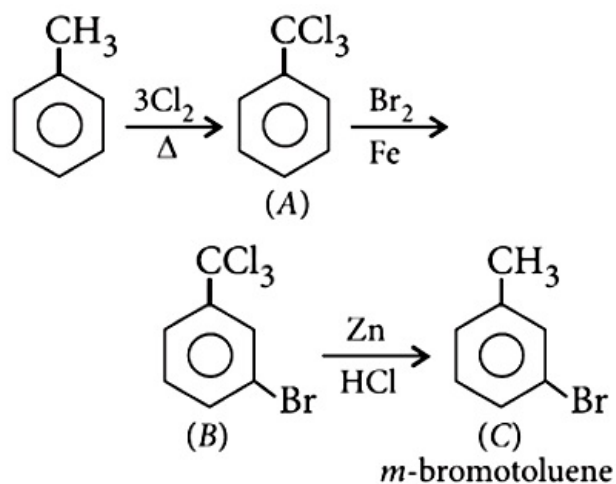
The product C is.  
(NEET 2018)

### Options:

- A. m -bromotoluene
- B. o -bromotoluene
- C. 3 -bromo- 2,4,6 -trichlorotoluene
- D. p -bromotoluene

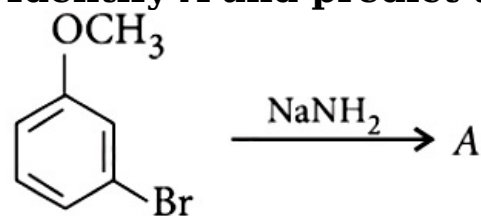
**Answer: A**

**Solution:**



## Question24

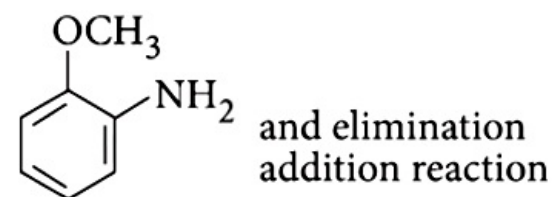
**Identify A and predict the type of reaction.**



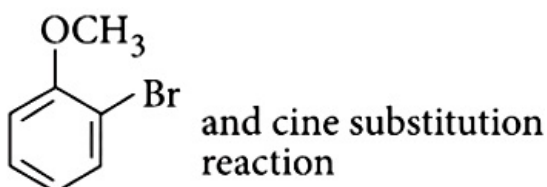
**(NEET 2017)**

**Options:**

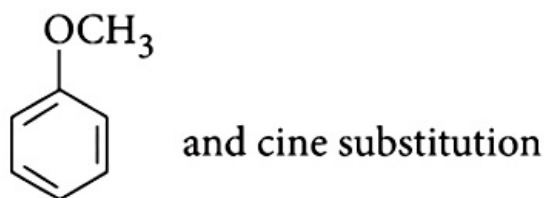
A.



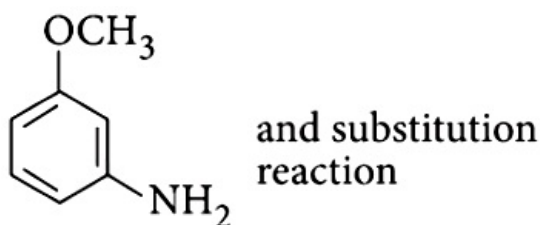
B.



C.

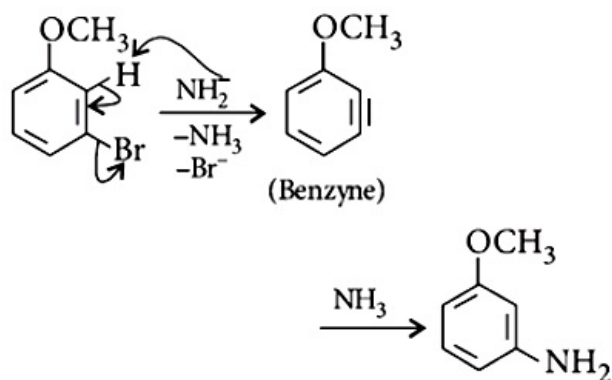


D.



**Answer: D**

**Solution:**



m -Bromoanisole gives only the respective meta substituted aniline. This is a substitution reaction which goes by an elimination-addition pathway.

## Question25

**Consider the reaction,  $\text{CH}_3\text{CH}_2\text{CH}_2\text{Br} + \text{NaCN} \rightarrow \text{CH}_3\text{CH}_2\text{CH}_2\text{CN} + \text{NaBr}$ . This reaction will be the fastest in (NEET-II 2016)**

**Options:**

A. ethanol

B. methanol

C. N , N ' -dimethylformamide (DMF)

D. water. (N E E T – I I 2016)

**Answer: C**

## Solution:

The reaction,  $\text{CH}_3\text{CH}_2\text{CH}_2\text{Br} + \text{NaCN} \rightarrow \text{CH}_3\text{CH}_2\text{CH}_2\text{CN} + \text{NaBr}$  follows  $\text{S}_{\text{N}}2$  mechanism which is favoured by polar aprotic solvent i.e., N, N' dimethylformamide (DMF),  $\text{H} - \overset{\text{O}}{\overset{\parallel}{\text{C}}} - \text{N}(\text{CH}_3)_2$

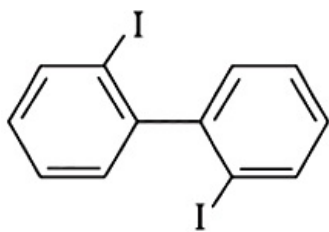
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## Question26

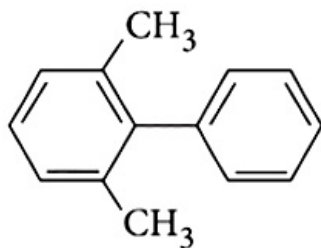
Which of the following biphenyls is optically active?  
(NEET-I 2016)

Options:

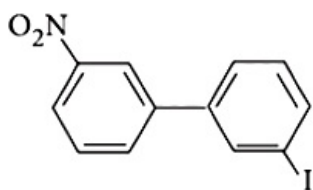
A.



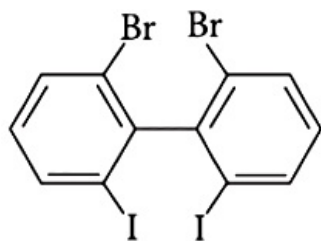
B.



C.



D.



**Answer: D**

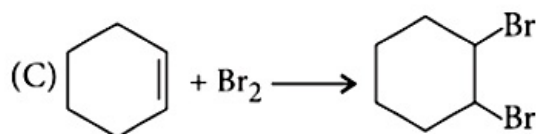
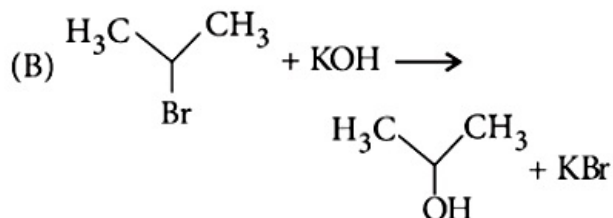
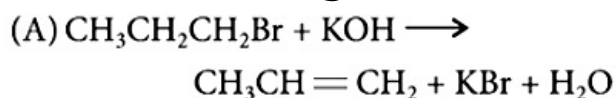
**Solution:**

o- Substituted biphenyls are optically active as both the rings are not in one plane and their mirror images are non-superimposable.

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## Question27

**For the following reactions:**



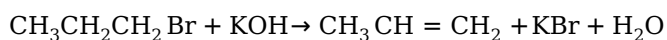
**Which of the following statements is correct?  
(NEET-I 2016)**

**Options:**

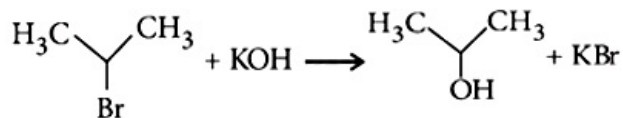
- A. (A) is elimination, (B) and (C) are substitution reactions.
- B. (A) is substitution, (B) and (C) are addition reactions.
- C. (A) and (B) are elimination reactions and (C) is addition reaction.
- D. (A) is elimination, (B) is substitution and (C) is addition reaction.

**Answer: D**

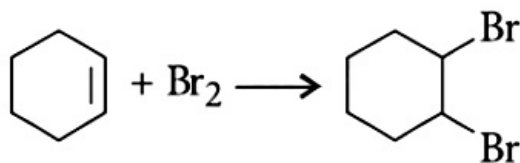
**Solution:**



Saturated compound is converted into unsaturated compound by removal of group of atoms hence, it is an elimination reaction.



- Br group is replaced by - OH group hence, it is a substitution reaction.



Addition of  $\text{Br}_2$  converts an unsaturated compound into a saturated compound hence, it is an addition reaction.

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## Question28

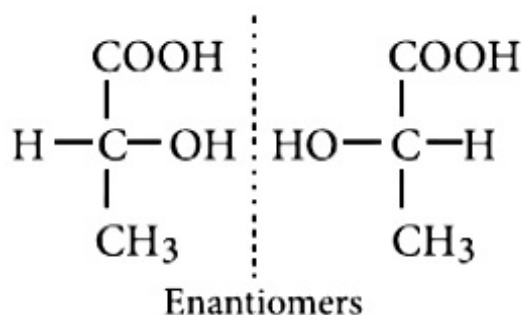
**Two possible stereo-structures of  $\text{CH}_3\text{CH(OH)COOH}$  which are optically active are called (2015)**

**Options:**

- A. atropisomers
- B. enantiomers
- C. mesomers
- D. diastereomers

**Answer: B**

**Solution:**



---

## Question29

**In an  $\text{S}_{\text{N}}1$  reaction on chiral centers,there is (2015)**

**Options:**

- A. inversion more than retention leading to partial racemisation
- B. 100% retention
- C. 100% inversion
- D. 100% racemisation.

**Answer: A**

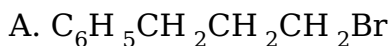
**Solution:**

In case of optically active alkyl halides,  $\text{S}_{\text{N}}1$  reaction is accompanied by racemisation. The carbocation formed in the slow step being  $\text{sp}^2$  hybridised is planar and attack of nucleophile may take place from either side resulting in a mixture of products, one having the same configuration and other having inverted configuration. The isomer corresponding to inversion is present in slight excess because  $\text{S}_{\text{N}}1$  also depends upon the degree of shielding of the front side of the reacting carbon

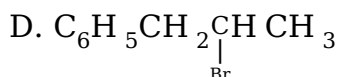
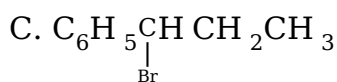
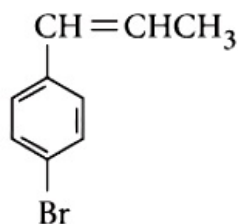
## Question30

The reaction of  $\text{C}_6\text{H}_5\text{CH}=\text{CHCH}_3$  with  $\text{HBr}$  products  
(2015 Cancelled)

Options:

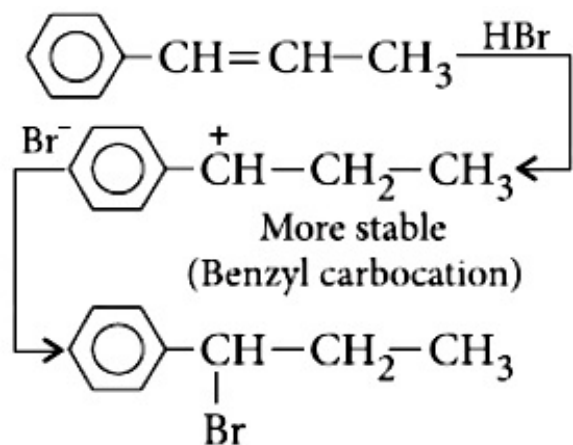


B.



Answer: C

Solution:

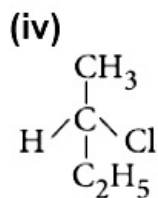
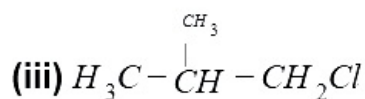
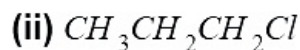
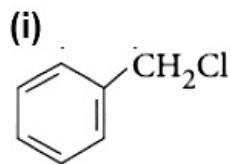


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## Question31

Which of the following compounds will undergo racemisation when solution of  $\text{KOH}$  hydrolyses?





**(2014)**

**Options:**

- A. (i) and (ii)
- B. (ii) and (iv)
- C. (iii) and (iv)
- D. Only (iv)

**Answer: D**

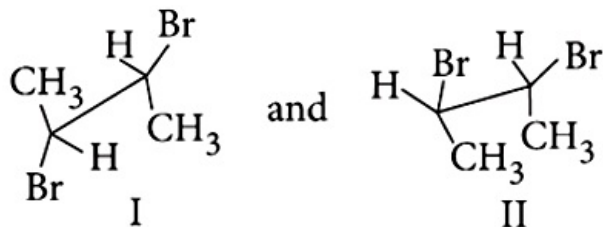
**Solution:**

**Solution:**

Only (iv) option will undergo racemization when the solution of KOH hydrolyzes because it is the only compound which contains a chiral carbon atom

## Question32

**Given:**



**I and II are  
(Karnataka NEET 2013)**

**Options:**

- A. identical
- B. a pair of conformers
- C. a pair of geometrical isomers
- D. a pair of optical isomers.

**Answer: B**

**Solution:**

I and II are staggered and eclipsed conformers.

---

## Question33

**Which of the following acids does not exhibit optical isomerism? (2012)**

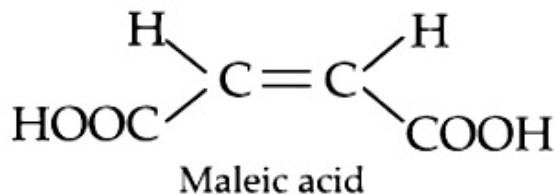
**Options:**

- A. Maleic acid
- B. alpha -amino acids
- C. Lactic acid
- D. Tartaric acid

**Answer: A**

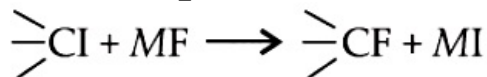
**Solution:**

Maleic acid shows geometrical isomerism and not optical isomerism.



## Question34

**In the replacement reaction**



**The reaction will be most favourable if M happens to be (Mains 2012)**

**Options:**

- A. Na
- B. K
- C. Rb
- D. Li

**Answer: C**

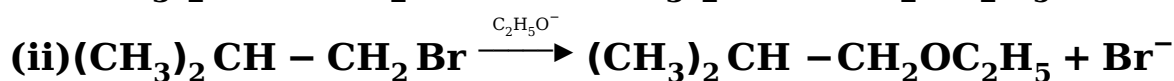
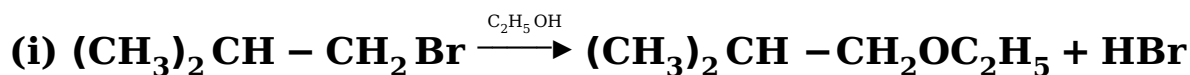
## Solution:

Tertiary halide shows  $S_N1$  mechanism i.e., ionic mechanism. In the given reaction negative ion will attack on carbocation. Thus greater the tendency of ionisation (greater ionic character in  $M - F$  bond more favourable will be reaction. The most ionic bond is  $Rb - F$  in the given examples thus most favourable reaction will be with  $Rb - F$ .

-----

## Question35

Consider the reactions.



The mechanisms of reactions (i) and (ii) are respectively  
(Mains 2011)

Options:

A.  $S_N1$  and  $S_N2$

B.  $S_N1$  and  $S_N1$

C.  $S_N2$  and  $S_N2$

D.  $S_N2$  and  $S_N1$

Answer: C

## Solution:

Positive charge will be on the Carbon with  $1^\circ$  in Both Examples  
as order of Reaction for  $SN_2$  is  $1^\circ > 2^\circ > 3^\circ$

So, Answer is  $S_N2$  and  $S_N2$

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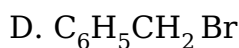
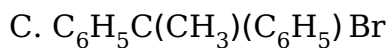
## Question36

Which one is most reactive towards  $S_N1$  reaction?  
(2010)

Options:

A.  $C_6H_5CH(C_6H_5)Br$

B.  $C_6H_5CH(CH_3)Br$



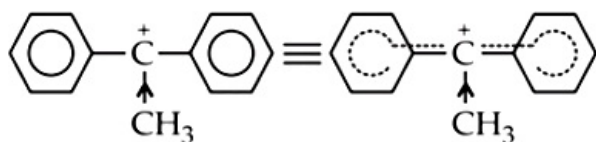
**Answer: C**

### Solution:

$\text{S}_{\text{N}}1$  reactions proceed via the formation of a carbocation intermediate.

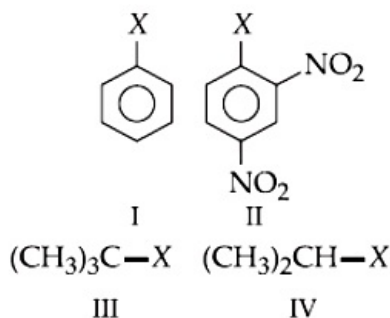
More stable is the carbocation more reactive is the alkyl/aryl halide towards  $\text{S}_{\text{N}}1$ .

In  $\text{C}_6\text{H}_5\text{C}^+(\text{CH}_3)(\text{C}_6\text{H}_5)$  carbocation, the two phenyl rings by their  $-\text{R}$  effect and  $-\text{CH}_3$  by its  $+\text{I}$  effect diminish the positive charge and make it stable.



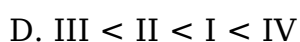
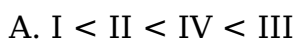
## Question37

The correct order of increasing reactivity of



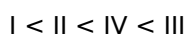
**C—X bond towards nucleophile in the following compounds is (2010)**

**Options:**



**Answer: A**

### Solution:



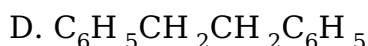
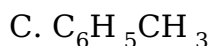
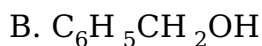
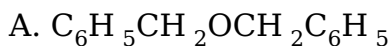
The order of reactivity is dependent on the stability of the intermediate carbocation formed by cleavage of C—X bond. The  $3^\circ$  carbocation (formed from III) will be more stable than its  $2^\circ$  counter part (formed from IV) which in turn will be more stable than the arcnium ion (formed from I). Also, the aryl halide has a double bond character in the C—X bond which makes the cleavage more difficult. However, inspite of all the stated factors, II will be more reactive than I due to the presence of the electron withdrawing  $-\text{NO}_2$  group. C—X bond becomes weak and undergoes nucleophilic substitution reaction

## Question38

In the following reaction

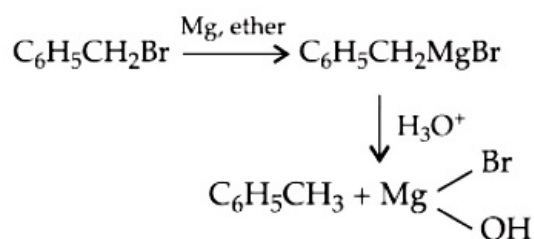
$\text{C}_6\text{H}_5\text{CH}_2\text{Br} \xrightarrow[2.\text{H}_3\text{O}^+]{1.\text{Mg Ether}} \text{X}$ , the product 'X' is  
(2010 Mains)

Options:



Answer: C

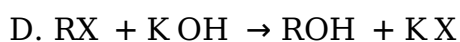
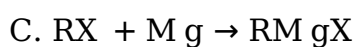
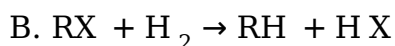
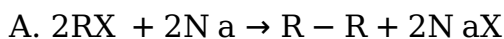
Solution:



## Question39

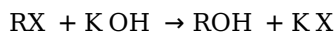
Which of the following reactions is an example of nucleophilic substitution reaction?  
(2009)

Options:

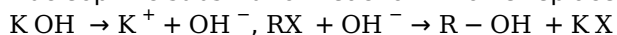


Answer: D

Solution:



Nucleophilic substitution reaction involve replacement of a group or atom by a nucleophilic



Nucleophile are either negative charge or a lone pair of  $e^-$  bearing species,  $\text{OH}^-$  etc.

## Question40

**How many stereoisomers does this molecule have?**



**(2008)**

**Options:**

A. 8

B. 2

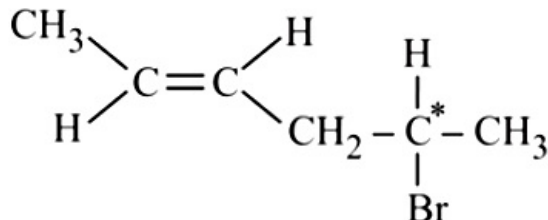
C. 4

D. 6

**Answer: C**

**Solution:**

The given compound may be written as



Both geometrical isomerism (cis-trans form) and optical isomerism is possible in the given compound.

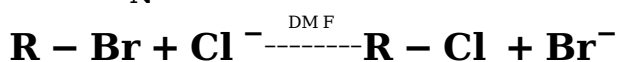
No. of optical isomer =  $2^n = 2^1 = 2$

(where  $n$  = no. of asymmetric carbon)

Hence total no. of stereoisomers =  $2 + 2 = 4$

## Question41

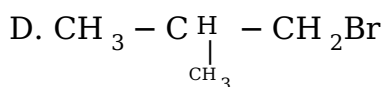
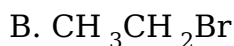
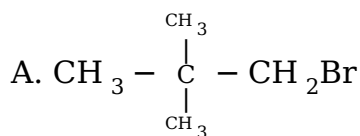
**In a  $\text{S}_\text{N}2$  substitution reaction of the type**



**which one of the following has the highest relative rate?**

**(2008)**

**Options:**



**Answer: B**

**Solution:**

$\text{S}_{\text{N}}2$  mechanism is followed in case of primary and secondary alkyl halides i.e.  $\text{S}_{\text{N}}2$  reaction is favoured by small groups on the carbon atoms attached to halogen so,  
 $\text{CH}_3 - \text{X} > \text{R} - \text{CH}_2 - \text{X} > \text{R}_2\text{CH} - \text{X} > \text{R}_3\text{C} - \text{X}$

Primary is more reactive than secondary and tertiary alkyl halides.  
 $\text{S}_{\text{N}}2$  order :Methyl > Ethyl > Isopropyl > Tertiary butyl > Allyl > Benzyl

## Question42

**If there is no rotation of plane polarised light by a compound in a specific solvent, though to be chiral, it may mean that (2007)**

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**Options:**

- A. the compound is certainly meso
- B. there is no compound in the solvent
- C. the compound may be a racemic mixture
- D. the compound is certainly a chiral.

**Answer: A**

**Solution:**

**Solution:**

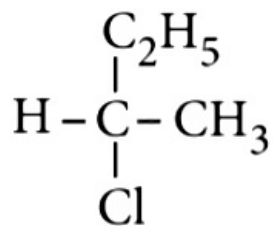
Meso compound does not rotate plane polarised light. Compound which contains tetrahedral atoms with four different groups but the whole molecule is achiral, is known as meso compound. It possesses a plane of symmetry and is optically inactive. One of the asymmetric carbon atoms turns the plane of polarised light to the right and other to the left and to the same extent so that the rotation due to upper half is compensated by the lower half, i.e., internally compensated, and finally there is no rotation of plane polarised light.

## Question43

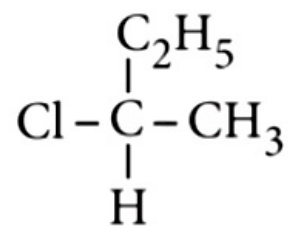
**$\text{CH}_3 - \text{CHCl} - \text{CH}_2 - \text{CH}_3$  has a chiral centre. Which one of the following represents its R -configuration? (2007)**

**Options:**

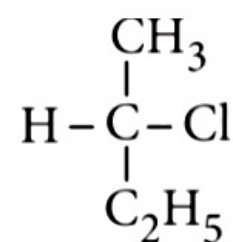
A.



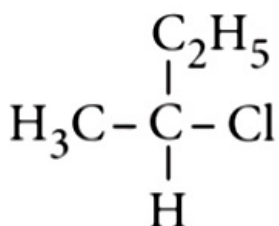
B.



C.

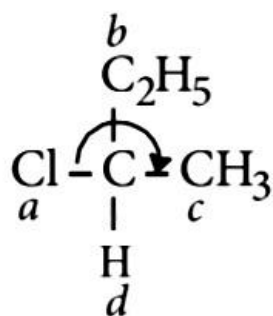


D.



**Answer: B**

**Solution:**



R-configuration



## Question44

Which of the following is not chiral?  
(2006)

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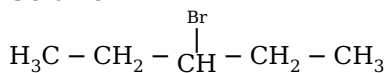
**Options:**

- A. 2 -Hydroxypropanoic acid
- B. 2 -Butanol
- C. 2,3 -Dibromopentane
- D. 3 -Bromopentane

**Answer: D**

**Solution:**

**Solution:**



Due to absence of asymmetric carbon atom.

-----

## Question45

Which of the following undergoes nucleophilic substitution exclusively by S<sub>N</sub>1 mechanism?  
(2005)

©

**Options:**

- A. Ethyl chloride
- B. Isopropyl chloride
- C. Chlorobenzene
- D. Benzyl chloride

**Answer: D**

**Solution:**

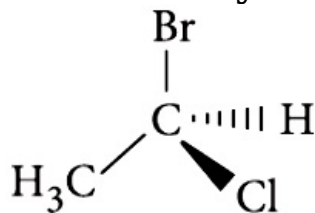
**Solution:**

S<sub>N</sub>1 reaction is favoured by heavy (bulky) groups on the carbon atom attached to halogens and nature of carbonium ion in substrate is

Benzyl > Allyl > Tertiary > Secondary > Primary > Methyl halides.

## Question46

The chirality of the compound



is  
(2005)

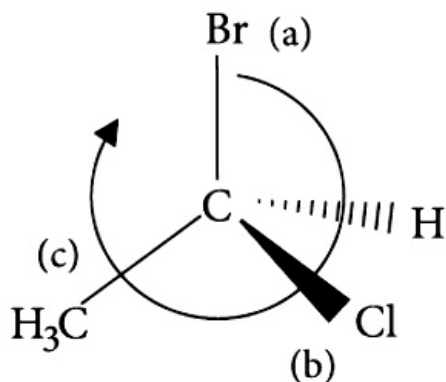
Options:

- A. R
- B. S
- C. E
- D. Z

Answer: A

Solution:

Solution:



Lowest priority atom is always away from the viewer. Priority is seen on the basis of atomic no. and if atomic no. are same then on the basis of atomic mass.

If clockwise then it is R, if anticlockwise then it is S.

Name of the molecule is, (R) 1 -bromo- 1 -chloroethane.

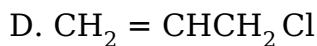
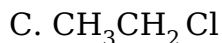
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## Question47

Which of the following is least reactive in a nucleophilic substitution reaction?  
(2004)

Options:

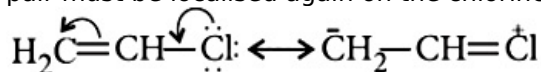
- A.  $(\text{CH}_3)_3\text{C} - \text{Cl}$
- B.  $\text{CH}_2 = \text{CHCl}$



**Answer: B**

### Solution:

The non-reactivity of the chlorine atom in vinyl chloride can be explained from the molecular orbital point of view as follows. If the chlorine atom has  $sp^2$  hybridisation, the C – Cl bond will be a  $\sigma$ -bond and the two lone pairs of electrons would occupy the other two  $sp^2$  orbitals. This would leave a p orbital containing a lone pair, and this orbital could now conjugate with the  $\pi$ -bond of the ethylenic link. Thus two M.O's will be required to accommodate these four  $\pi$ -electrons. Furthermore, since chlorine is more electronegative than carbon, the electrons will tend to be found in the vicinity of the chlorine atom. Nevertheless, the chlorine atom has now lost full control of the lone pair and so, is less negative than it would have been had there been no conjugation. Since two carbon atoms have acquired a share in the lone pair, each carbon atom acquires a small negative charge. Hence, owing to delocalisation of bonds (through conjugation), the vinyl chloride molecule has an increased stability. Before the chlorine atom can be displaced by some other group, the lone pair must be localised again on the chlorine atom. This requires energy, and so the chlorine is more firmly bound.

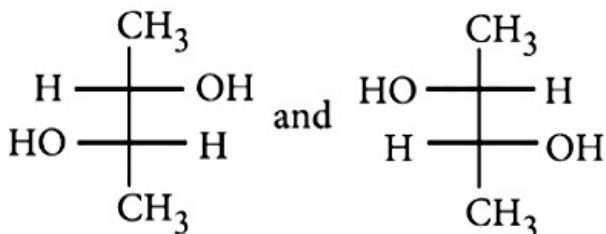


## Question 48

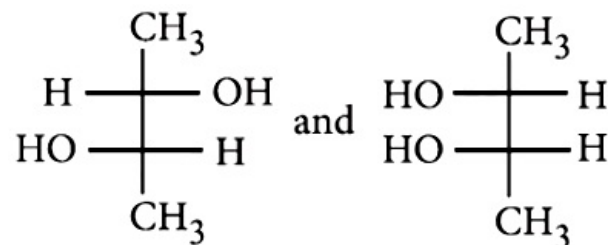
**Which of the following pairs of compounds are enantiomers? (2003)**

**Options:**

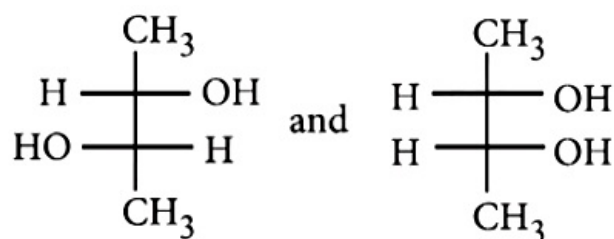
A.



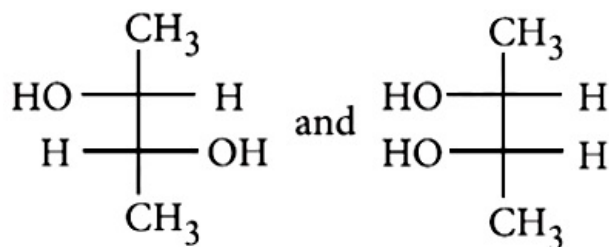
B.



C.



D.



**Answer: A**

**Solution:**

**Solution:**

These two are non-superimposable mirror images of each other, so they are enantiomers.

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## Question49

**Reactivity order of halides for dehydrohalogenation is (2002)**

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**Options:**

- A.  $\text{R} - \text{F} > \text{R} - \text{Cl} > \text{R} - \text{Br} > \text{R} - \text{I}$
- B.  $\text{R} - \text{I} > \text{R} - \text{Br} > \text{R} - \text{Cl} > \text{R} - \text{F}$
- C.  $\text{R} - \text{I} > \text{R} - \text{Cl} > \text{R} - \text{Br} > \text{R} - \text{F}$
- D.  $\text{R} - \text{F} > \text{R} - \text{I} > \text{R} - \text{Br} > \text{R} - \text{Cl}$

**Answer: B**

**Solution:**

$\text{I} > \text{Br} > \text{Cl} > \text{F} \rightarrow$  atomic radii

F, Cl, Br, I belongs to the same group orderly.

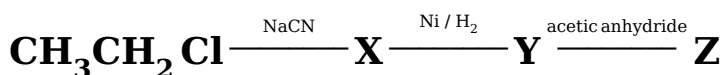
Atomic radii go on increasing as the nuclear charge increases in preceeding downwards in a group. The decreasing order of bond length  $\text{C} - \text{I} > \text{C} - \text{Br} > \text{C} - \text{Cl} > \text{C} - \text{F}$ .

The order of bond dissociation energy  $\text{R} - \text{F} > \text{R} - \text{Cl} > \text{R} - \text{Br} > \text{R} - \text{I}$ .

During dehydrohalogenation  $\text{C} - \text{I}$  bond breaks more easily than  $\text{C} - \text{F}$  bond. So reactivity order of halides is,  $\text{R} - \text{I} > \text{R} - \text{Br} > \text{R} - \text{Cl} > \text{R} - \text{F}$ .

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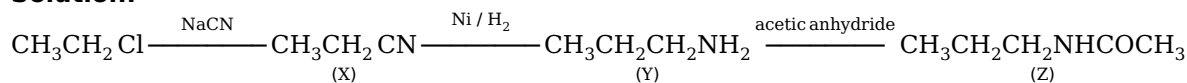
## Question50



**Z in the above reaction sequence is (2002)**

**Options:**

- A.  $\text{CH}_3\text{CH}_2\text{CH}_2\text{NHCOCH}_3$
- B.  $\text{CH}_3\text{CH}_2\text{CH}_2\text{NH}_2$
- C.  $\text{CH}_3\text{CH}_2\text{CH}_2\text{CONHCH}_3$
- D.  $\text{CH}_3\text{CH}_2\text{CH}_2\text{CONHCOCH}_3$

**Answer: A****Solution:****Solution:****Question 51**

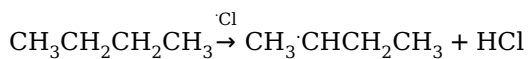
$\text{CH}_3 - \text{CH}_2 - \underset{\text{Cl}}{\underset{|}{\text{CH}}} - \text{CH}_3$  obtained by chlorination of n -butane will be (2001)

**Options:**

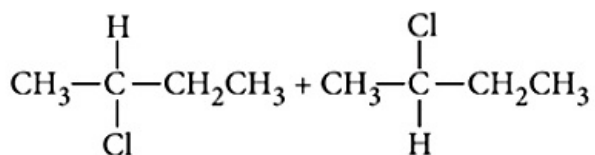
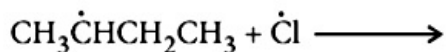
- A. meso form
- B. racemic mixture
- C. d -form
- D. l -form.

**Answer: B****Solution:**

Chlorination of n -butane takes place via free radical formation. i.e.  $\text{Cl}_2 \rightarrow \cdot\text{Cl} + \cdot\text{Cl}$



$\text{sp}^2$  - hybrid planar shape intermediate and  $\cdot\text{Cl}$  may attack from either side to give



Racemic mixture

(Mixture of 50% d-form and 50% l-form)

## Question52

An organic compound A( $C_4H_9Cl$ ) on reaction with Na/diethyl ether gives a hydrocarbon which on monochlorination gives only one chloro derivative then, A is (2001)

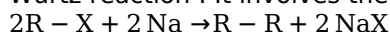
Options:

- A. t -butyl chloride
- B. s -butyl chloride
- C. iso-butyl chloride
- D. n -butyl chloride.

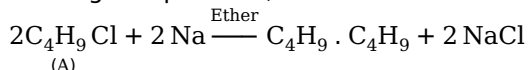
Answer: A

Solution:

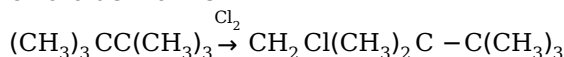
Wurtz reaction : It involves the reaction of alkyl halides with Na in ether to form higher alkanes.



In the given problem,



Compound A is t -butyl chloride, in this compound all -  $CH_3$  groups have primary hydrogen only and able to give only, one chloro derivative.



## Question53

A compound of molecular formula  $C_7H_{16}$  shows optical isomerism, compound will be (2001)

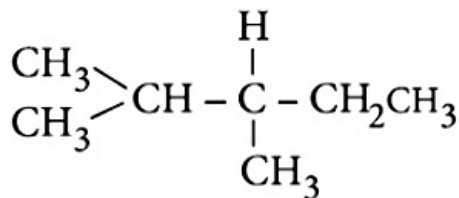
Options:

- A. 2,3 -dimethylpentane
- B. 2,2 -dimethylbutane
- C. 2 -methylhexane
- D. none of these.

Answer: A

Solution:

Organic compounds exhibit the property of enantiomerism (optical isomerism) only when their molecules are chiral. Most chiral compounds have a chiral centre, which is an atom bonded to four different atoms or groups.



2,3-Dimethylpentane has one chiral C-atom and do not have any symmetric element.

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## Question54

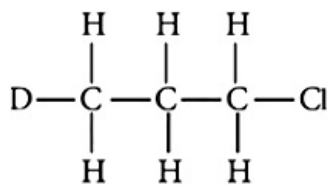
**Which of the following compounds is not chiral?  
(1998)**

**Options:**

- A.  $\text{CH}_3\text{CHDCH}_2\text{Cl}$
- B.  $\text{CH}_3\text{CH}_2\text{CHDCl}$
- C.  $\text{DCH}_2\text{CH}_2\text{CH}_2\text{Cl}$
- D.  $\text{CH}_3\text{CHClCH}_2\text{D}$

**Answer: C**

**Solution:**



The above compound has no chiral 'C'-atom. All the 'C' atoms are attached to two identical 'H' atoms, so they are not asymmetrical.

-----

## Question55

**Replacement of Cl of chlorobenzene to give phenol requires drastic conditions. But chlorine of 2,4 -dinitrochlorobenzene is readily replaced because  
(1997)**

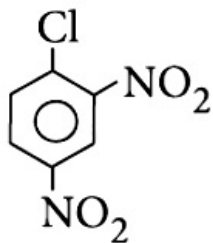
**Options:**

- A.  $\text{NO}_2$  donates  $e^-$  at meta position
- B.  $\text{NO}_2$  withdraws  $e^-$  from ortho/para positions
- C.  $\text{NO}_2$  make ring electron rich at ortho and para

D.  $\text{NO}_2$  withdraws  $e^-$  from meta position.

**Answer: B**

**Solution:**



Withdrawal of electrons by -  $\text{NO}_2$  groups from ortho/para positions cause easier removal of - Cl atom due to the development of positive charge on o - positions.

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## Question56

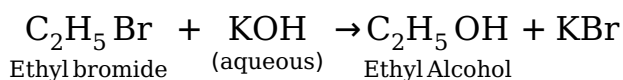
**The alkyl halide is converted into an alcohol by (1997)**

**Options:**

- A. elimination
- B. dehydrohalogenation
- C. addition
- D. substitution.

**Answer: D**

**Solution:**



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## Question57

**Reaction of t -butyl bromide with sodium methoxide produces (1994)**

**Options:**

- A. sodium t -butoxide
- B. t -butyl methyl ether



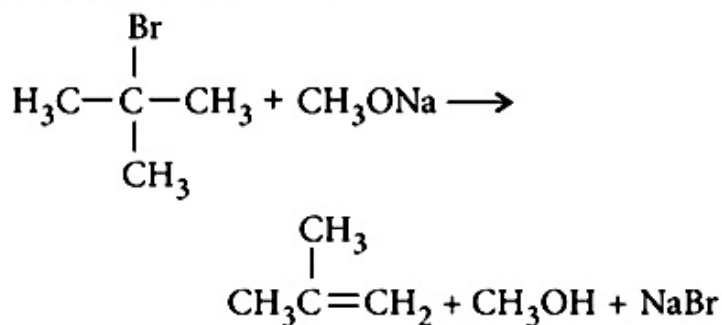
C. isobutane

D. isobutylene.

**Answer: D**

**Solution:**

Isobutylene is obtained.



Thus, the reaction produces isobutylene.

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## Question58

**Grignard reagent is prepared by the reaction between (1994)**

**Options:**

A. magnesium and alkane

B. magnesium and aromatic hydrocarbon

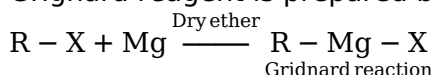
C. zinc and alkyl halide

D. magnesium and alkyl halide.

**Answer: D**

**Solution:**

Grignard reagent is prepared by heating an alkyl halide with dry magnesium powder in dry ether.

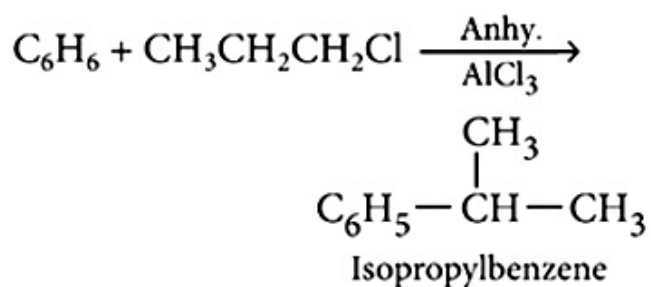


## Question59

**Benzene reacts with n -propyl chloride in the presence of anhydrous  $\text{AlCl}_3$  to give (1993)**

**Options:**

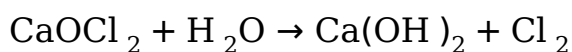
- A. 3 -propyl-1-chlorobenzene
- B. n -propylbenzene
- C. no reaction
- D. isopropylbenzene.

**Answer: D****Solution:****Question60**

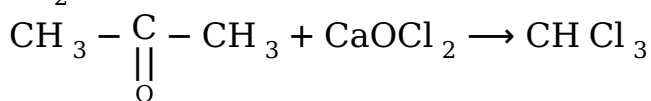
**Industrial preparation of chloroform employs acetone and (1993)**

**Options:**

- A. phosgene
- B. calcium hypochlorite
- C. chlorine gas
- D. sodium chloride.

**Answer: B****Solution:**

$\text{Cl}_2$  so obtained acts as a mild oxidising as well as chlorinating agent.

**Question61**

**Chlorobenzene reacts with Mg in dry ether to give a compound (A)**

**which further reacts with ethanol to yield (1993)**

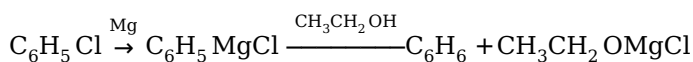
©

**Options:**

- A. phenol
- B. benzene
- C. ethyl benzene
- D. phenyl ether.

**Answer: B**

**Solution:**



## Question62

**When chlorine is passed through propene at 400°C, which of the following is formed? (1993)**

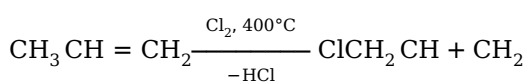
©

**Options:**

- A. PVC
- B. Allyl chloride
- C. Propyl chloride
- D. 1,2 -Dichloroethane

**Answer: B**

**Solution:**



At 400°C temperature, substitution occurs instead of addition.

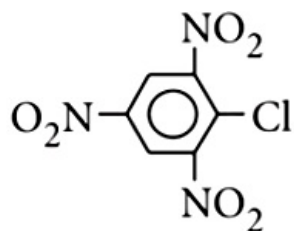
## Question63

**Which chloro derivative of benzene among the following would undergo**

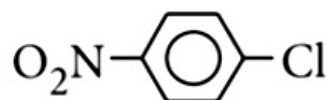
**hydrolysis most readily with aqueous sodium hydroxide to furnish the corresponding hydroxy derivative?  
(1989)**

**Options:**

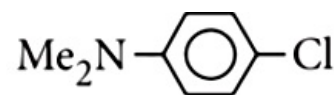
A.



B.



C.



D.  $\text{C}_6\text{H}_5\text{Cl}$

**Answer: A**

**Solution:**

Cl in 2, 4, 6-trinitrochlorobenzene is activated by three  $-\text{NO}_2$  groups at o -and p -positions and hence undergoes hydrolysis most readily.

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## Question64

**Which of the following is an optically active compound?  
(1989)**

**Options:**

A. 1 -Butanol

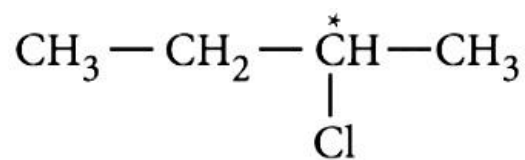
B. 1 -Propanol

C. 2 -Chlorobutane

D. 4 -Hydroxyheptane

**Answer: C**

**Solution:**



2-Chlorobutane contains a chiral carbon atom and hence it is optically active compound.

---

## Question65

**Phosgene is a common name for (1988)**

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**Options:**

- A. phosphoryl chloride
- B. thionyl chloride
- C. carbon dioxide and phosphine
- D. carbonyl chloride.

**Answer: D**

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