

## CHEMISTRY

### BOOKS - CENGAGE CHEMISTRY (HINGLISH)

#### AROMATIC COMPOUNDS AND ALKYL AND ARYL HALIDES

##### Illustration

1. Name of the following groups:

a.  $C_6H_5-$  , b.  $C_6H_5CH_2-$

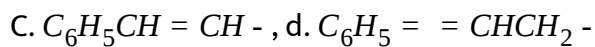
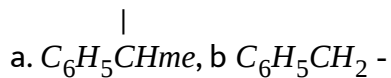
c.  $(C_6H_5)_2CH-$ , d.  $(C_6H_5)_3C-$  e.  $C_6H_5CH-$ , f.  $C_6H_5-$

overset{()underset{()}{C}-



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2. Write the name of phenyl-substituted groups:



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3. Write the structures of the following compounds:

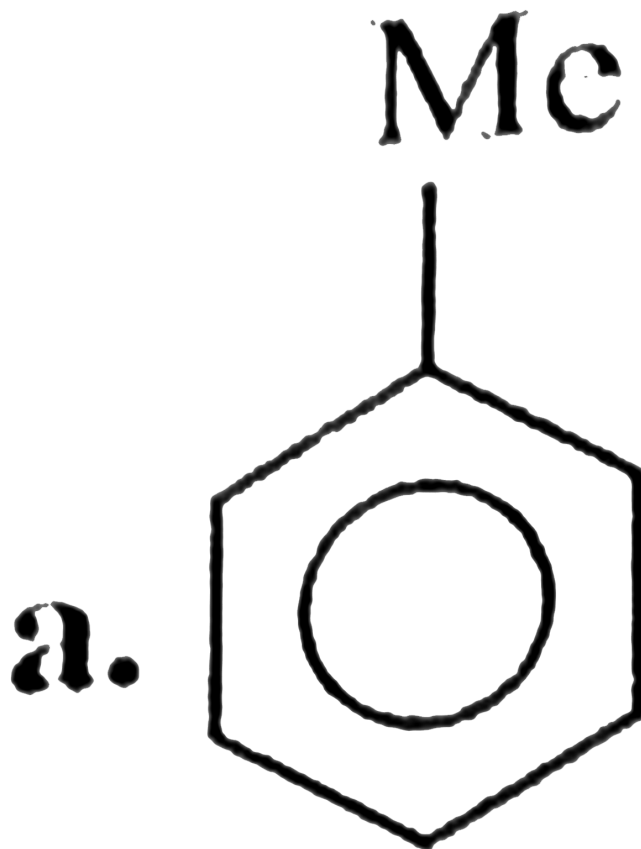
a. Adjacent tribromobenzene

b. Unsymmetrical trimethyl benzene

c. Symmetrical triethyl benzene

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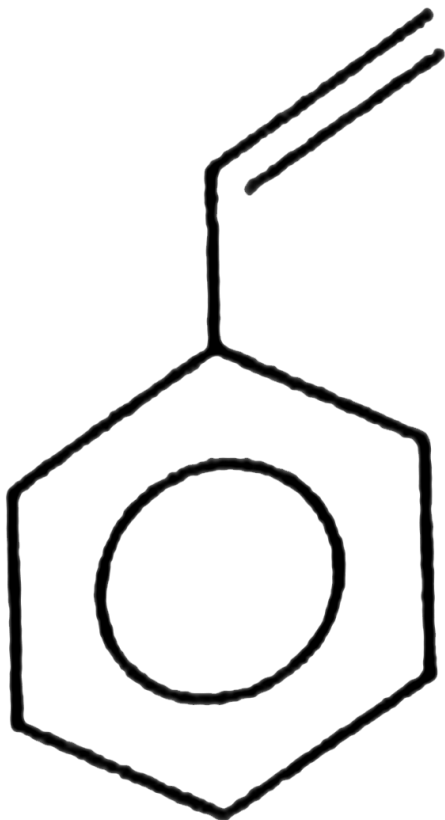
4. Name of the following compounds:

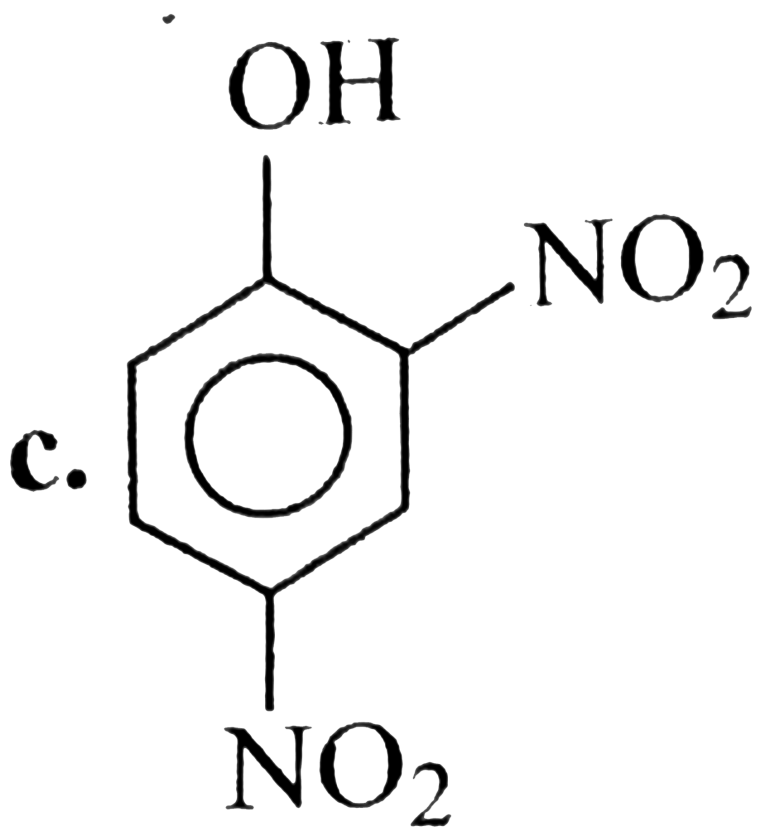


a.

, b.

**b.**

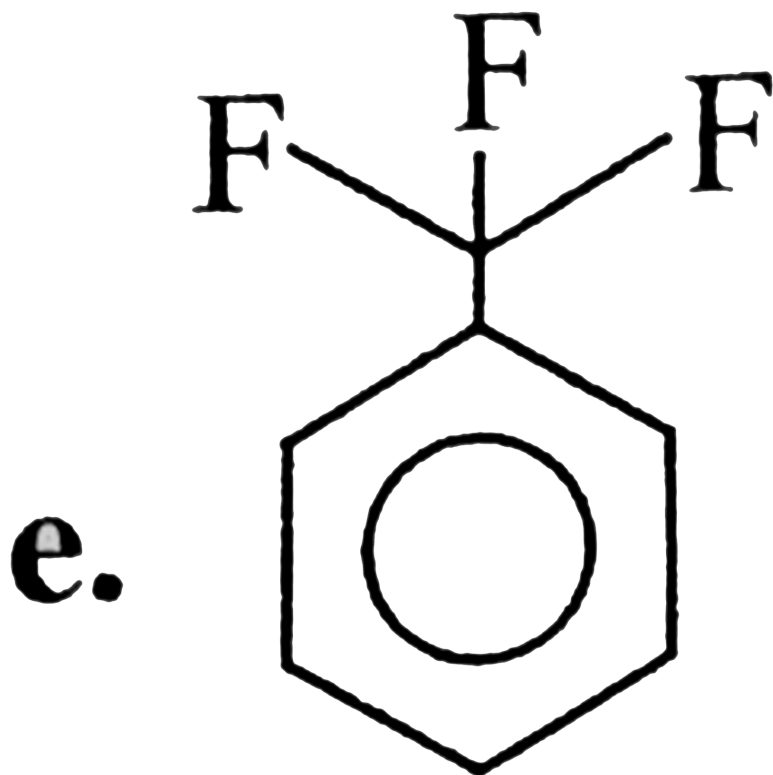
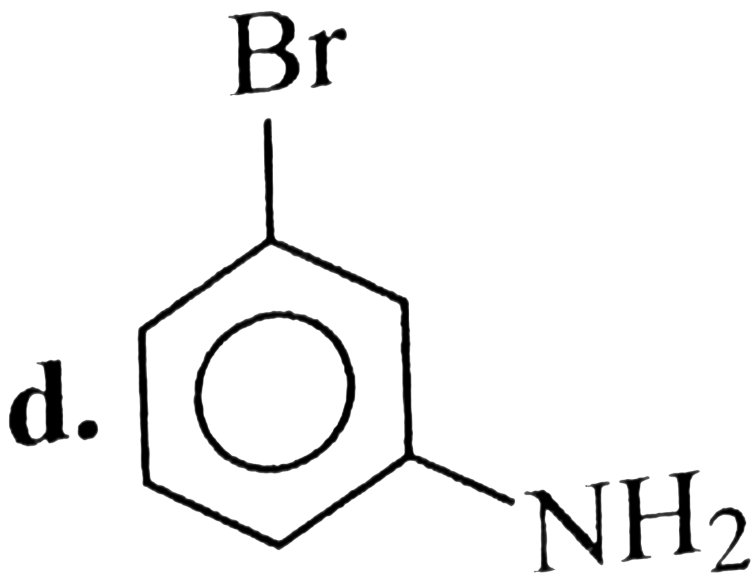




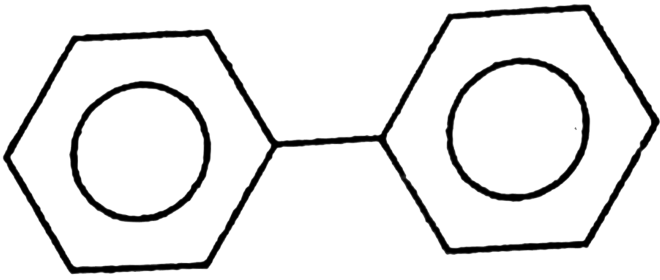
c.

,

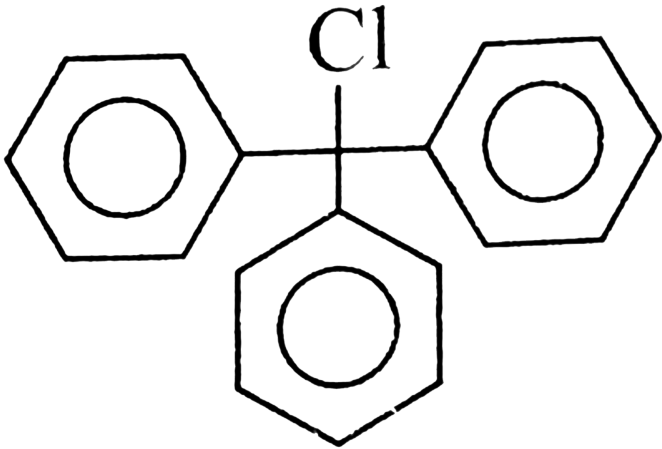
d.



**f.**



**g.**

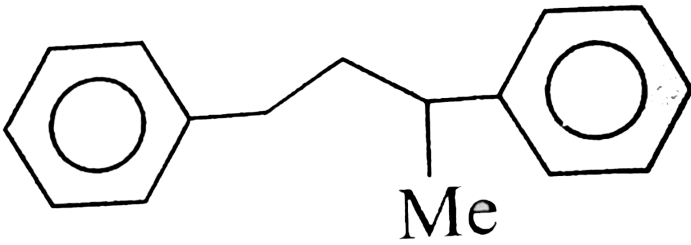


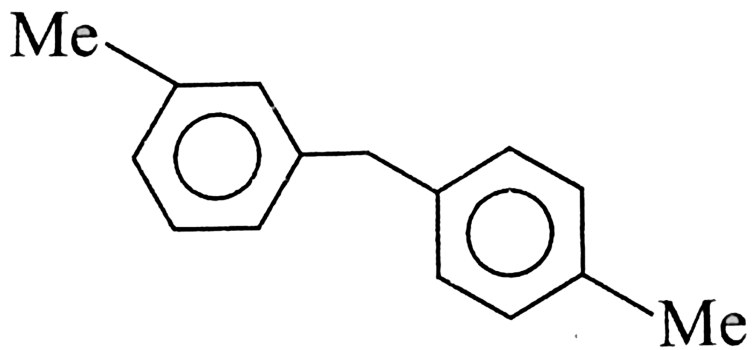
g.

,

h.

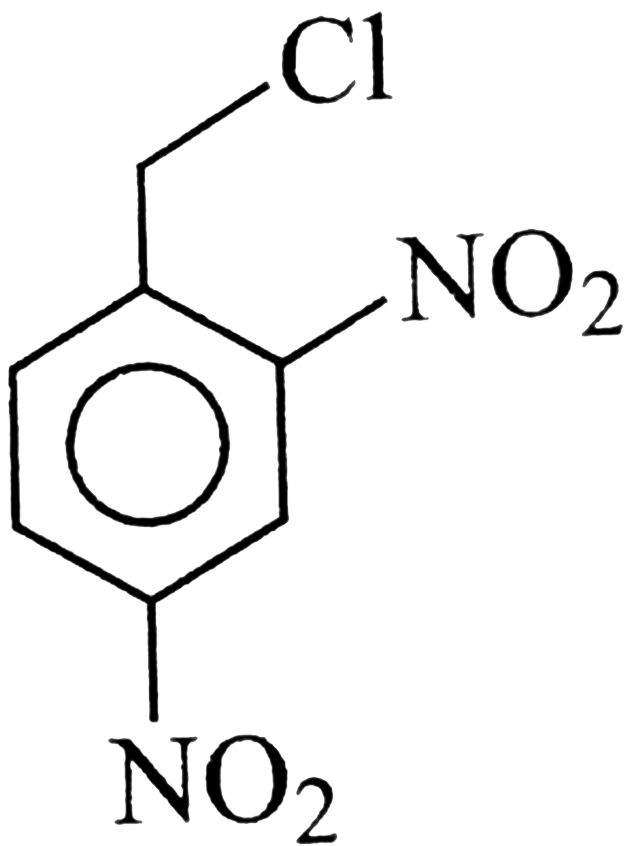
**h.**



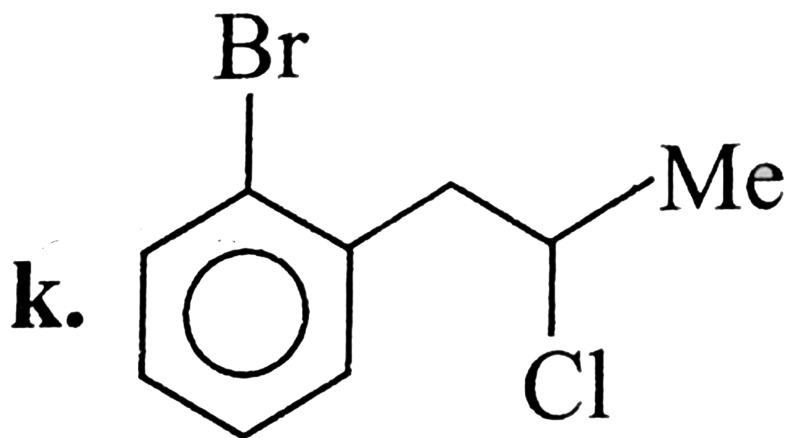


i.

j.



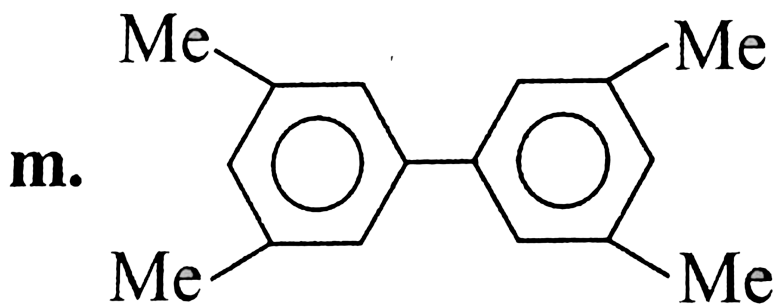
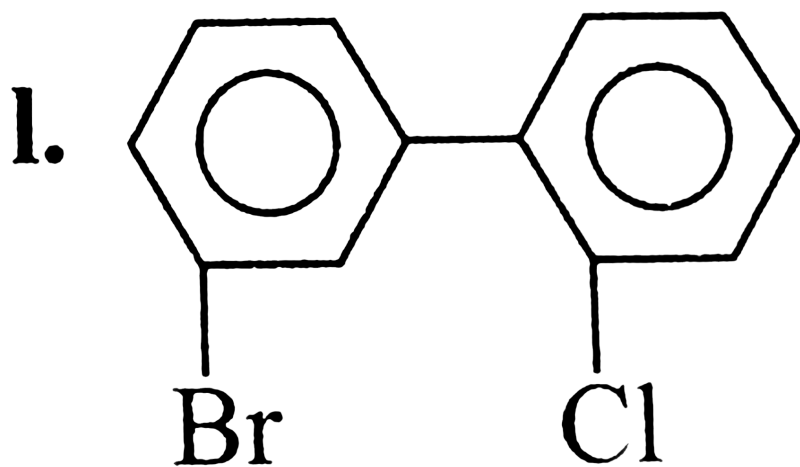




k.

,

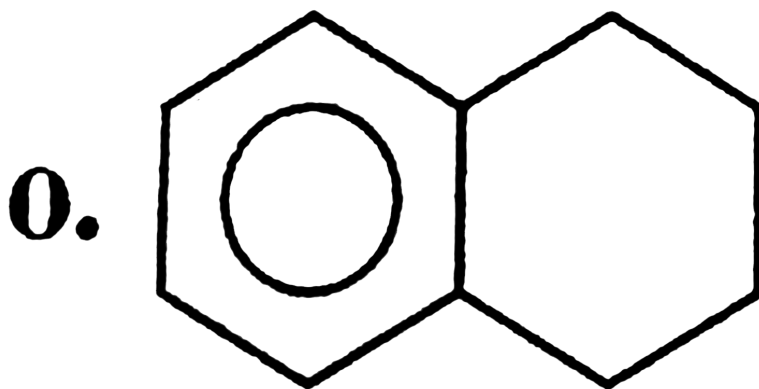
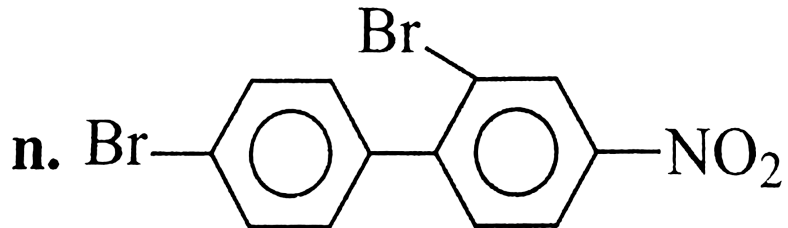
l.



m.

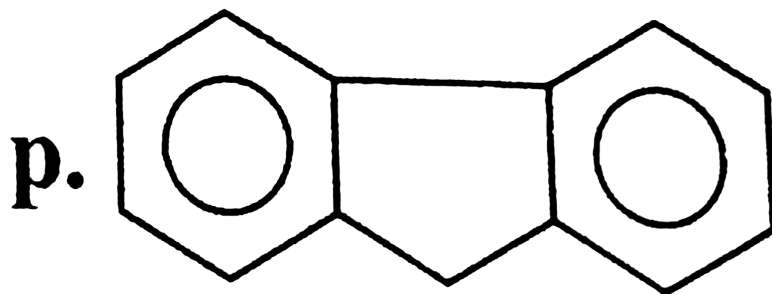
,

n.

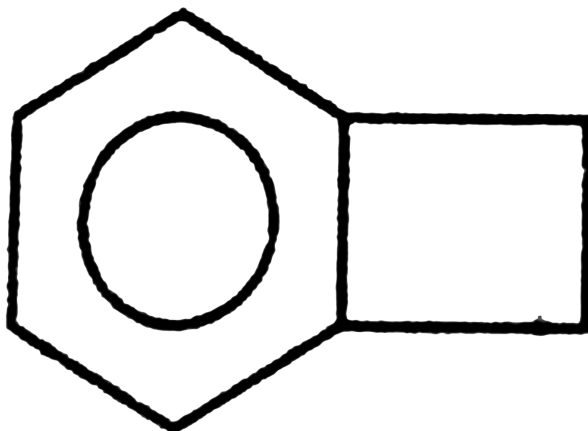


o.

, p.



q.

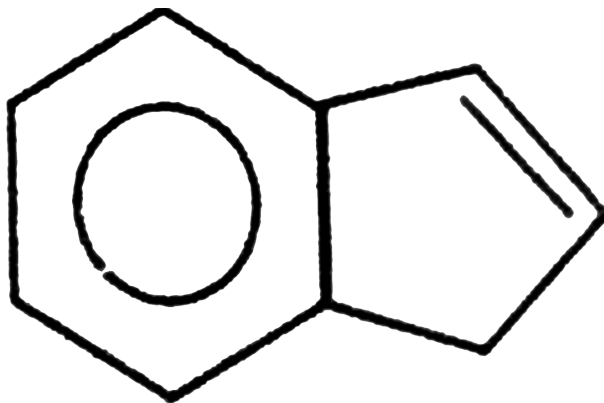


q.

,

r.

r.



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5. Write the names and structures of the following compounds:

a. *TNB*, b. *TNT*

c. *PABA*, d. *DDT*

e. *DES*, f. Cumene

g. Mesityphenone, h. Acetophenone

i. Benzoic acid, j. Benzyl chloride

k. Cinnamic acid, l. Cinnamyl alcohol

m.  $\alpha$  - Phenylethanol, n.  $\beta$  - Phenylethanol

o. Benzyl chloride

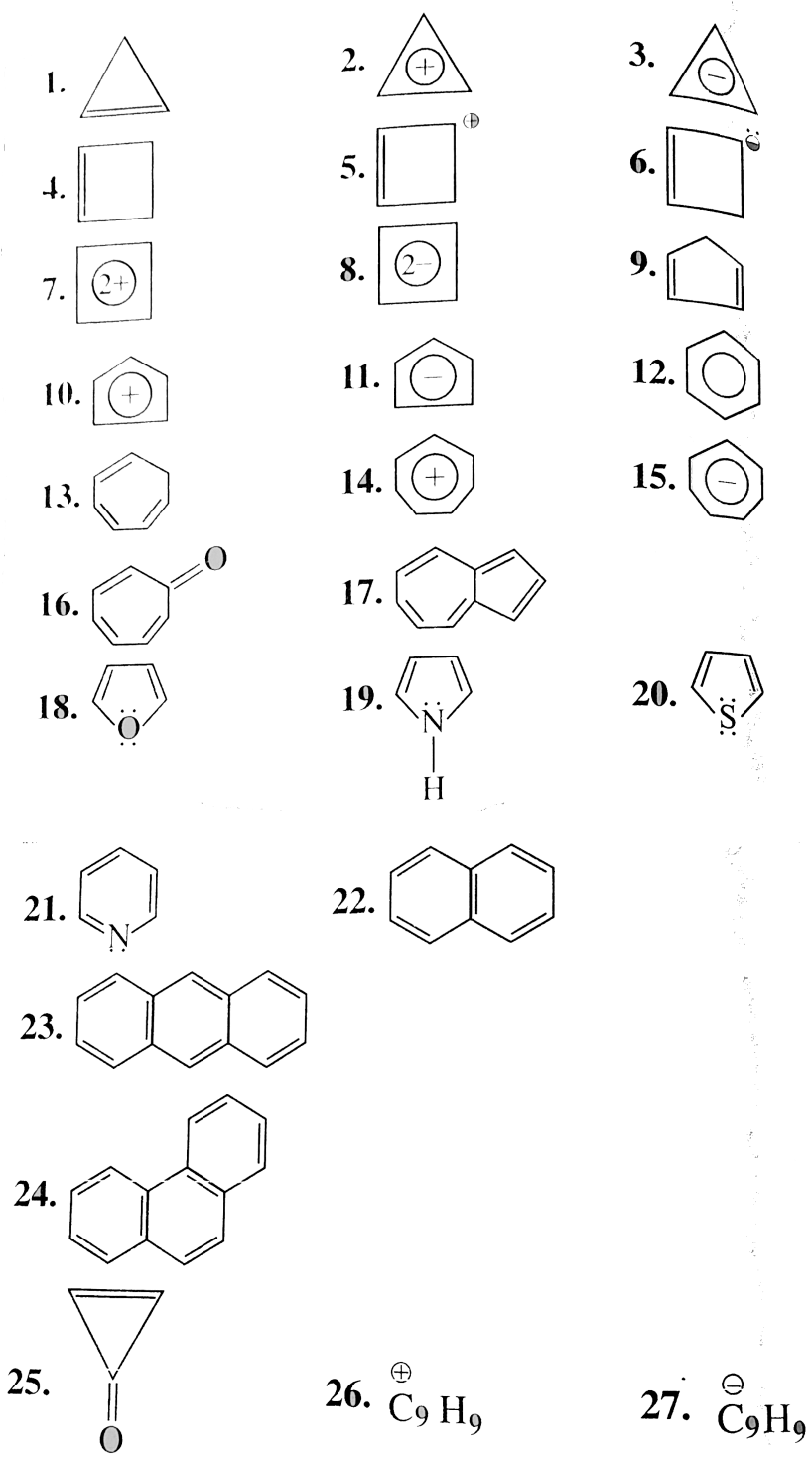


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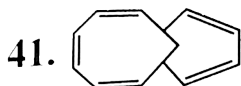
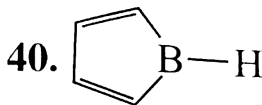
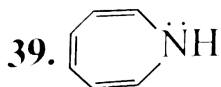
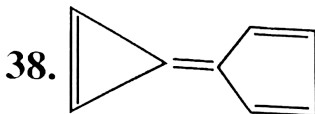
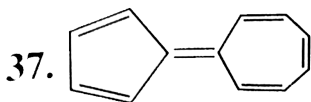
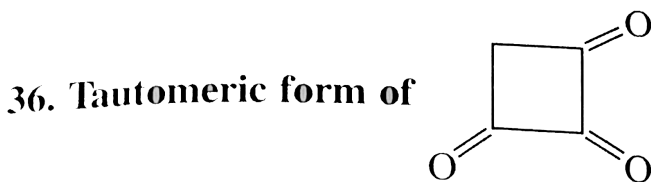
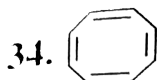
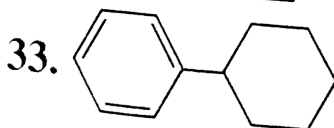
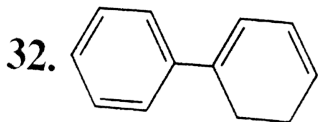
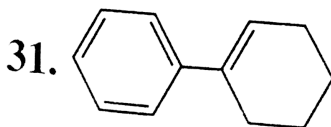
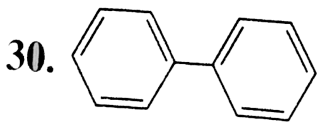
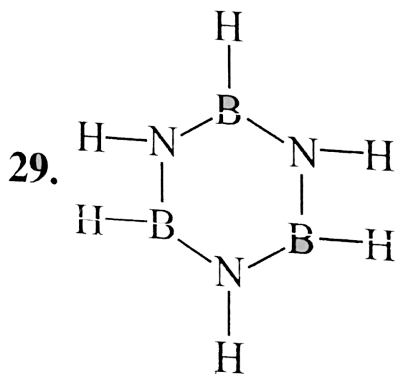
6. Select the aromatic, anti-aromatic, and non-aromatic compounds.

1, 2, 3.

4, 5



28.  $C_9H_{10}$



7. a.

7 - Bromocycloheptatriene (tropylium bromide) completely dissociates in water and gives a yellow precipitate of  $AgBr$  with  $AgNO_3$ . Why?

(b) Why is cyclopentadiene ( $K_a = 10^{-15}$ ) much more acidic than 1,3-cyclohexadiene?

C. Explain the following 1,3,5-cycloheptatriene through a cycle, planar triene with six  $\pi e^-$ 's is not aromatic, whereas tropolone behaves like a phenol.



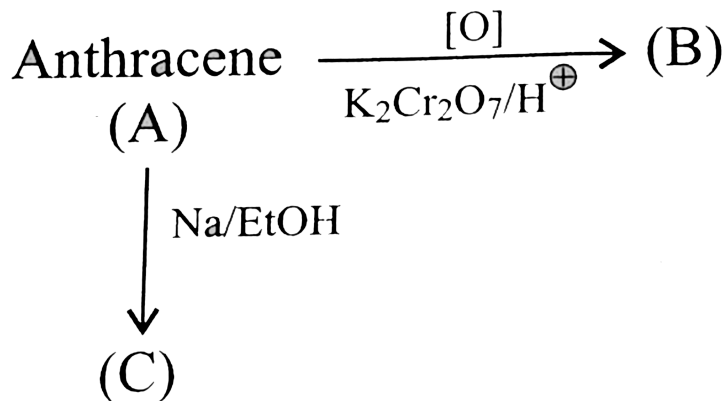
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8. Anthracene has a resonance energy of  $351 \text{ kJ mol}^{-1}$  and the resonance energy of benzene is  $150.5 \text{ kJ mol}^{-1}$ . Which of the two is more easily oxidised or reduced?



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9. a. Complete the following reaction.



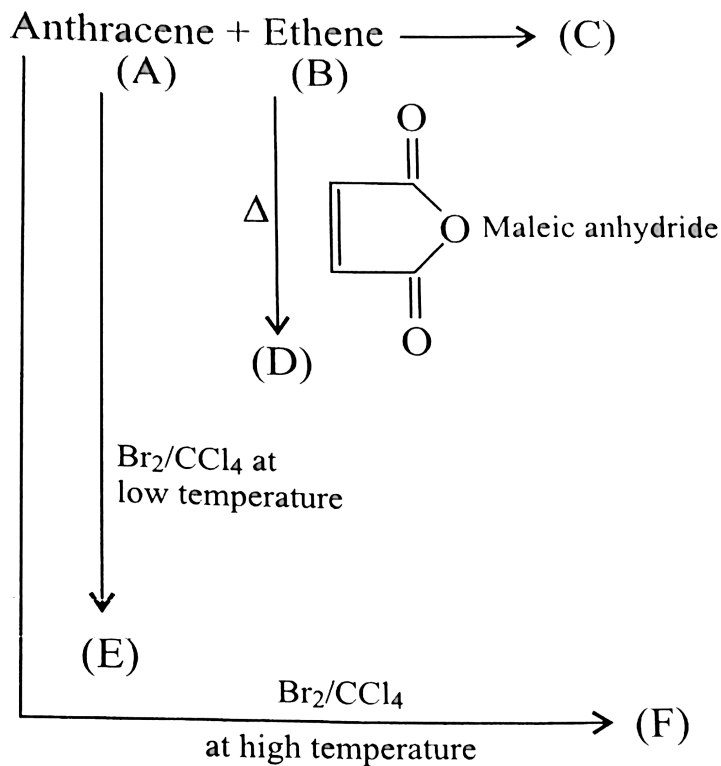
b. Why do the oxidation and reduction of anthracene take place at positions 9 and 10, respectively?



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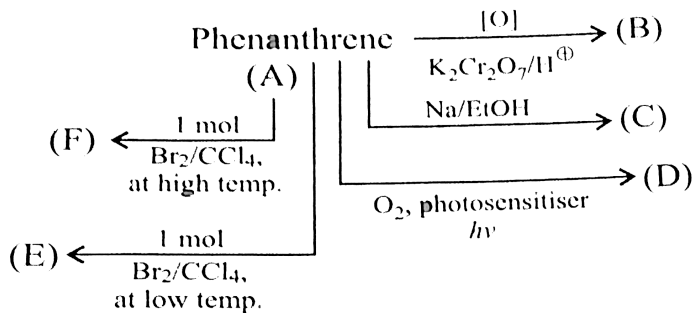


10. Complete the reactions.



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11. Complete the reactions.

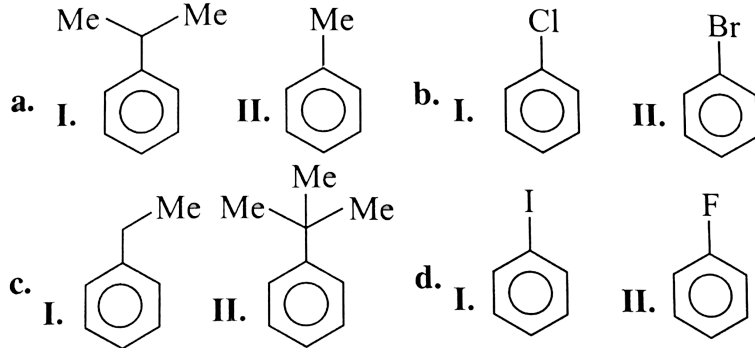


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12. Out of benzene naphthalene anthracene, and phenantrene, which is more reactive in the Diels - Alder reaction?

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13. Among the following in which cases is the *o/p* ratio on nitration more the first compound?

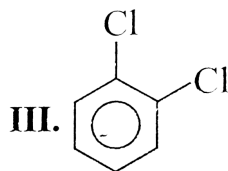
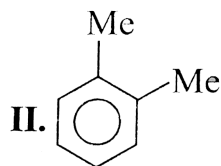
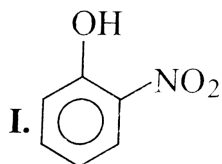


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14. Why are  $(o + p)$  ratios for the *SE* reaction (e.g. nitration) of toluene and *PhCF*, 32.3 and 0.14, respectively?

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15. Arrange the following in the decreasing of  $\mu$ .



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16. Write resonance-contributing structures for the intermediate carbocation (arenonium ion) of

(i) Phenol

(ii) Nitrobenzene with electrophile  $Cl^{\oplus}$  at the

a. ortho-,

b. para-, and

c. meta-positions.

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17. a. Why are  $\bar{e}$  donating groups, such as  $(-NH_2)$ ,  $(-OR)$ ,  $(-R)$  *o*- and *p*- directing towards *SE* reaction? Explain by an example.

b. Why are  $\bar{e}$  withdrawing groups, [such as  $(-NO_2)$ ,  $(-SO_3H)$ ,  $(-CN)$ ,  $(-COOH)$ , etc], *m*- direction towards *SE* reaction? explain by example.

c. Although halogens are  $\bar{e}$  withdrawing yet they are *o*- and *p*- directing towards *SE* reaction. Why?

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18. Arrange the following compounds in decreasing order of their property as indicated:

a. Reactivity and orientation of (I)  $\text{Ph} - \overset{\oplus}{\text{S}} \dots - \text{Et}_2$  and (II)  $\text{PhNO}$ .

Activating effects of the following *o*, *p* - directions:

I.  $-\text{O}^\ominus$  . II,  $-\text{OH}$ , III.  $-\text{NH}_2$  IV.  $-\text{NHCOCH}_3$  V.  $-\text{OCOCH}_3$

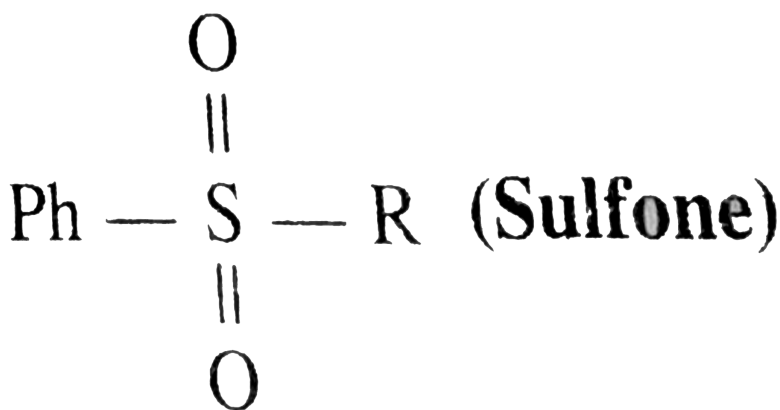
c. Respectivity and orientation of I.  $\text{PhOEt}$  and II.  $\text{PhSEt}$

d. Categories the following substituents as:

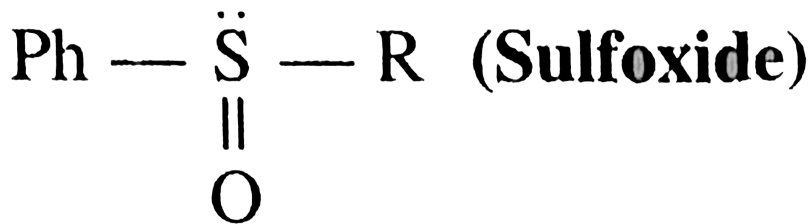
i. Activating,

ii. Deactivating,

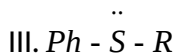
iii. *o*, *p*, or *m* - directing.



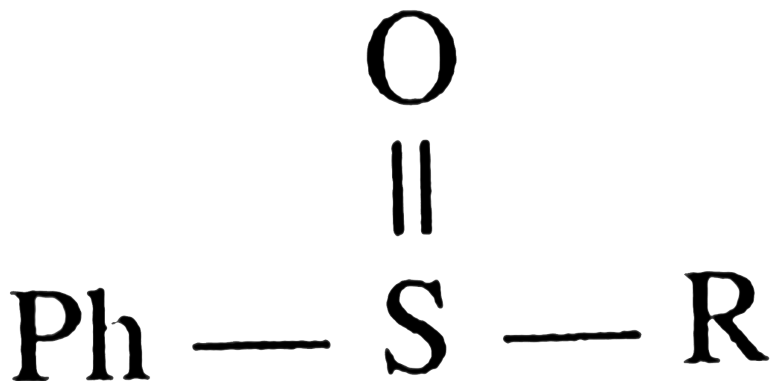
i.



II.



SE reactivity of:



I.  $\text{PhNO}$ , II.

f. SE reactivity and orientation of:

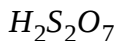
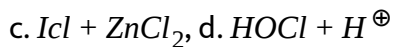
1.  $\text{PhF}$ , II.  $\text{PhCl}$ , III.  $\text{PhBr}$ , IV.  $\text{PhI}$



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19. Identify the electrophile and base for each for the following:

a.  $\text{HNO}_2 + \text{H}^\oplus$ , b.  $\text{Ibr} + \text{ZnCl}_2$



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20. Give the decreasing order of the relative reactivity towards *SE* reaction of the following compounds.

a. I. Benzene, II. Phenol, III. Aniline, IV. Chlorobenzene

b. I. Acetanilide, II. Aniline, III. Acetep[jempme, IV. Benzene.

c. I. 1,3- Dimryhyl benzene, II. 1, 4 - Dimenthyl-benzene, III. Tolune, IV. Trimehtyl benzene, V. 1, 3, 5 - Trimethhtyl benzene

d. I. *p* - Mehtyl benzoic acid, II. Terephtalic acid, III. *p* - Xylene, IV. Toluene.

e. I. Benzene, II. Toulene, III. *p* - Nitrotoulene, IV. *m* - Nitroluence, V. Benzence

g. I. Flurobenzene, II. Chlorobenzene, III. Bromobenzene, IV. Iodobenzene

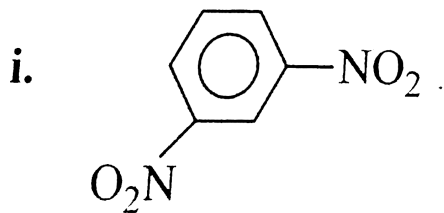
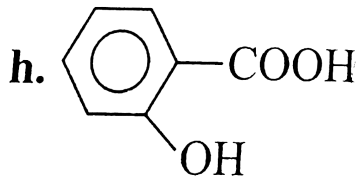
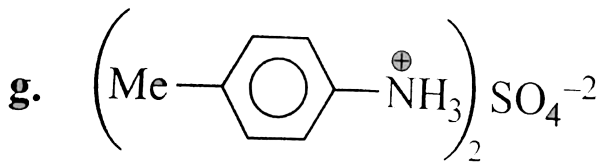
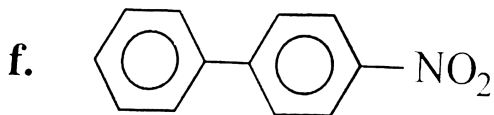
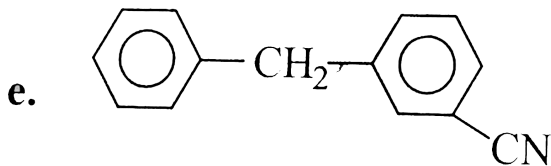
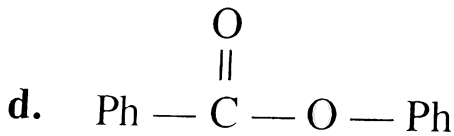
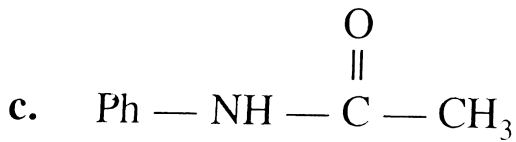
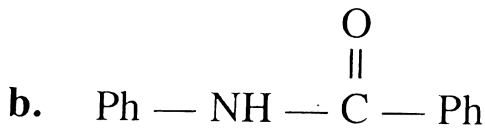
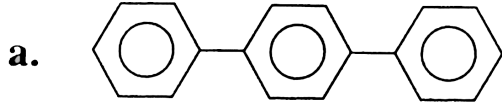
h. I. *p* - Nitrochlorobenze II. Chlorobenzene, III. 2.4 - Dintrochlorobenzene

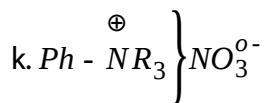
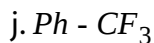


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21. Indicate by an arrow the position(s) where *SE* reaction takes place in the following:



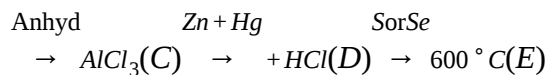
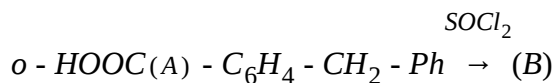




l. *o*, *m* - and *p* - Ethoxyacetanilide

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22. Complete the following reaction:



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23. a. Which of the following shows primary isotope effect?

i. Nitration of benzene. ii. Halogenation of  $CH_4$ . iii. Alkene prepared by *E.1cB* mechanism.

b. Which of the following shows secondary isotope effect?

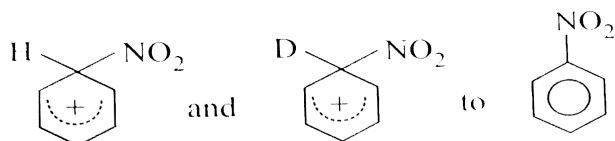
i. An alkene obtained by *E.2* mechanism

ii. An alkene obtained by *E. I* mechanism

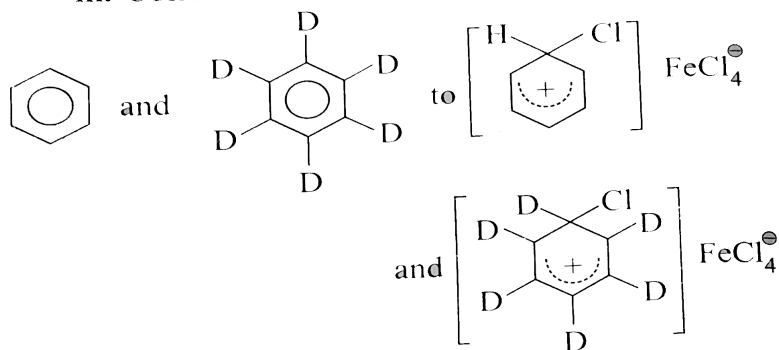
c. Which of the following examples of  $K_H/K_D$  (ratio of rate constant) is greater than 1.

i.  $PhCH_2CH_2Br$  and  $PhCD_2CH_2Br$  towards de-hydrohalogenation by strong base.

ii. Conservation of:



iii. Conversion of:



iii. Conservation of:



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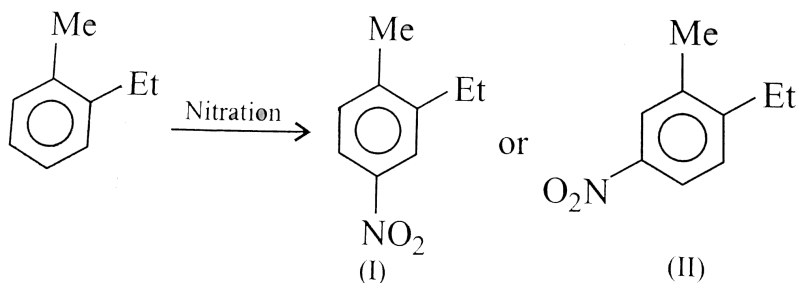
24. Convert benzene to adjacent trimethyl benzene.

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25. a. In the *F. C.* Alkylation with alkene, why is *HF*, not *HBr*, used? How does *HF* at high temperature effect the dealkylation process?

In Illustration 11.24 in the conversion of (I) to (II), why dows alkylation take place at 2 and 6 positions rather than at 3 and 5 positions?

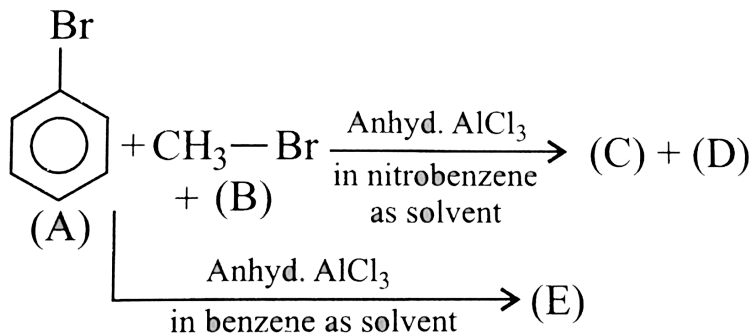
c. Explain which product (I) or (II) is formed and why.



d. What is the depression in freezing point of 1*m* soultion of nitrating mixture ( $K_f = xK, kgmol^{-1}$ )?

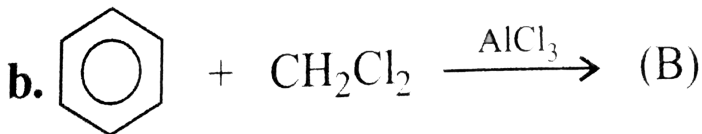
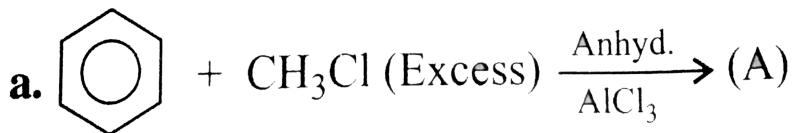
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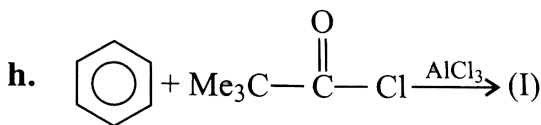
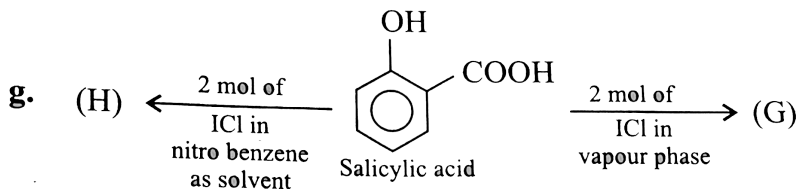
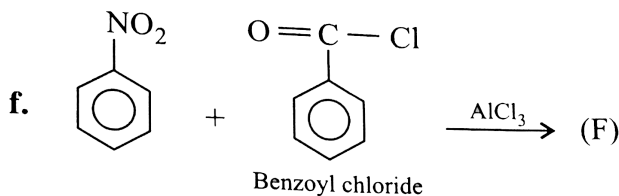
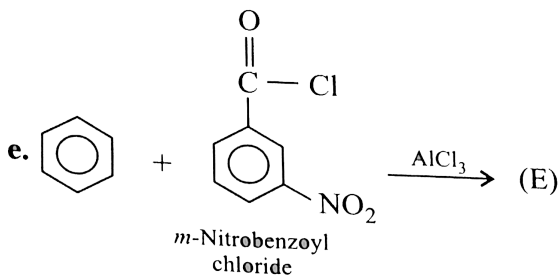
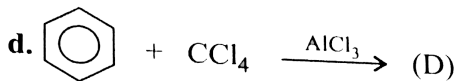
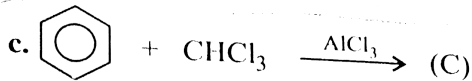
26. Complete the reaction:



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27. Complete the following reaction:





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28. For different reactions of  $\text{H}_2\text{SO}_4$  with (i)  $\text{HNO}_3$  and (ii)  $\text{HClO}_4$ ,

a. Write the equations for the reactions and identify the conjugate acids

and bases:

b. Explain the different behaviours of  $H_2SO_4$

II. Nitrobenzene can be prepared from benzene by using a mixture of conc.

$HNO_3$  and conc.  $H_2SO_4$  In the nitrating mixture,  $HNO_3$  acts as:

a. Base, b. Acid

c. Reducing agent, d. Catalyst

III. Among the following statements on the nitration of aromatic compounds, the false one is:

a. The rate of nitration of benzene is almost the same as that of hexadeuterobenzene.

b. The rate of nitration of toluene is greater than that of benzene.

c. The rate of nitration of benzene is greater than that of hexadeuterobenzene.

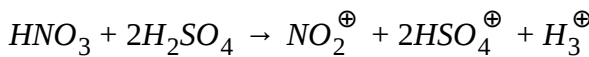
c. The rate of nitration of benzene is greater than that of hexadeuterobenzene.

d. Nitration is an electrophilic substitution reaction.

IV. Select the correct alternative(s).

The following reaction occurs in a mixture of conc.  $HNO_3$  and conc.  $H_2SO_4$

as:



Which of the following statements about this reaction is correct?

a. Nitric acid acts as a base.

b. Sulphuric acid acts as a base.

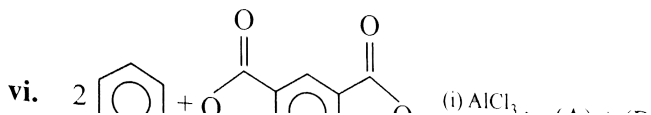
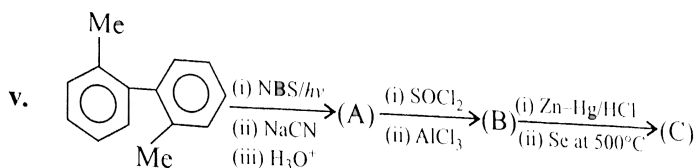
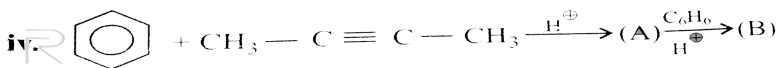
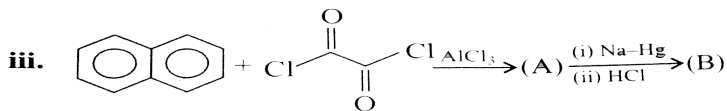
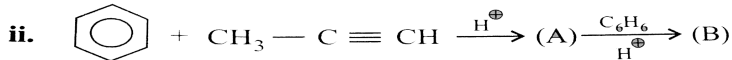
c. Sulphuric acid acts as a dehydrating agent.

d. Addition of  $\text{H}_2\text{O}$  will reduce the  $\text{NO}_2^{\oplus}$  concentration.

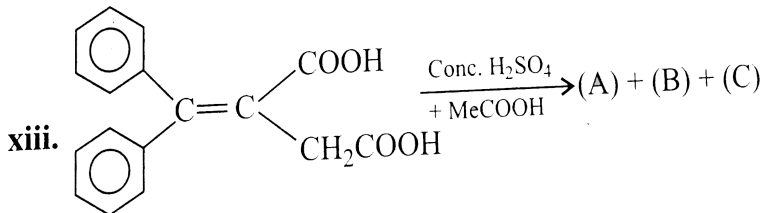
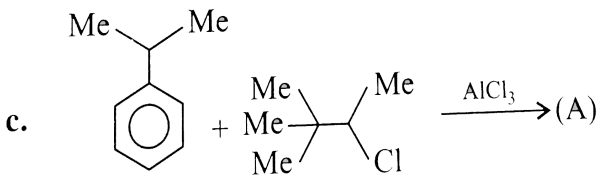
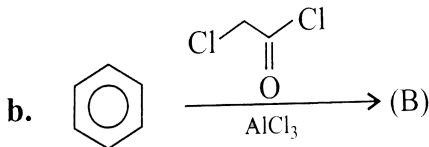
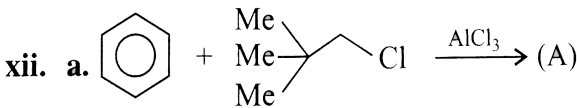
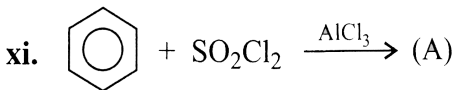
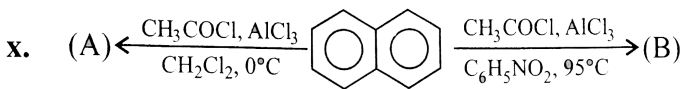
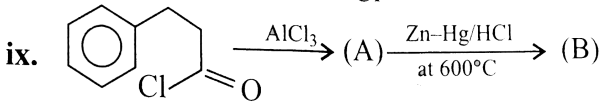
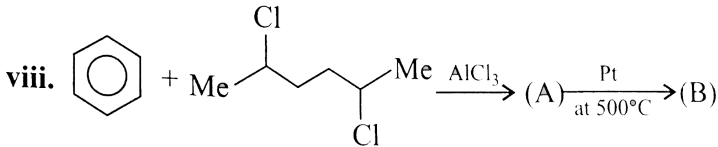
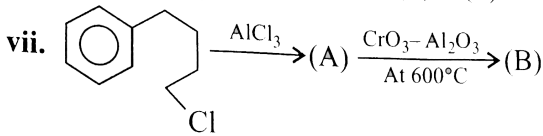
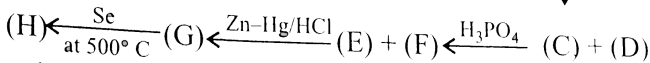
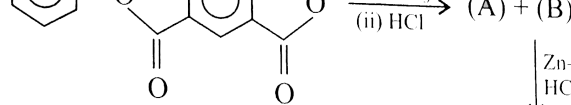
e.  $\text{HNO}_3$  and  $\text{NO}_2^{\oplus}$  are conjugate acid-base pair.

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29. Complete the following reaction:

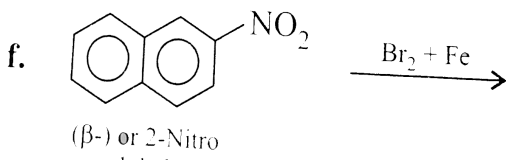
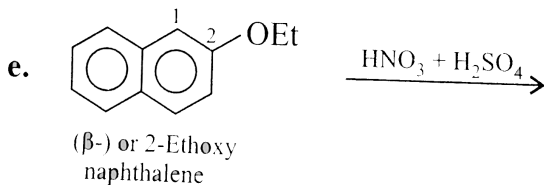
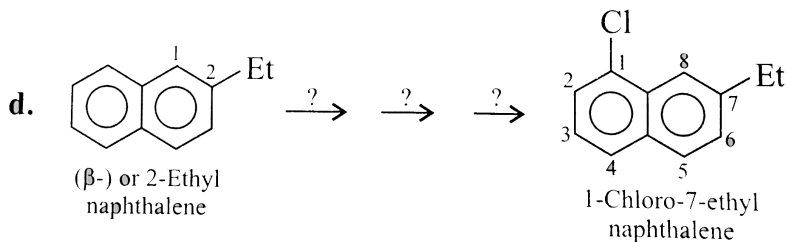
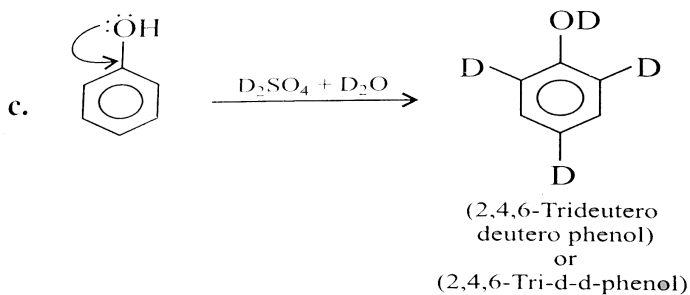
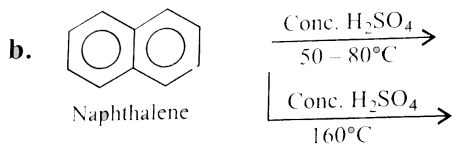
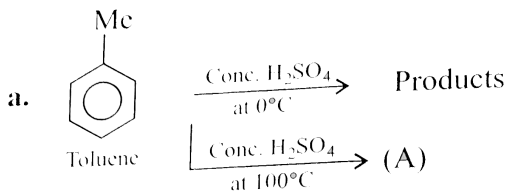


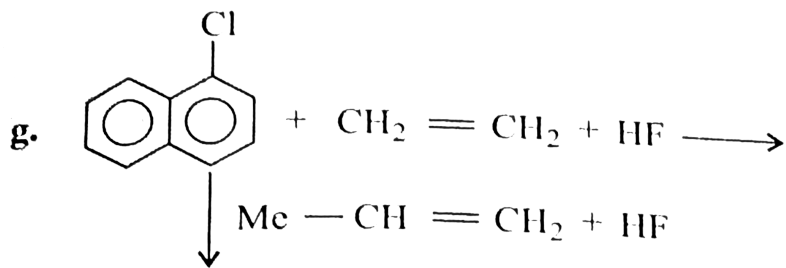






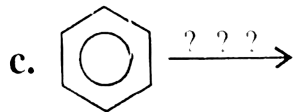
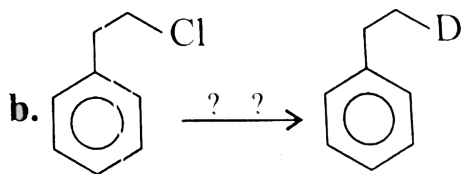
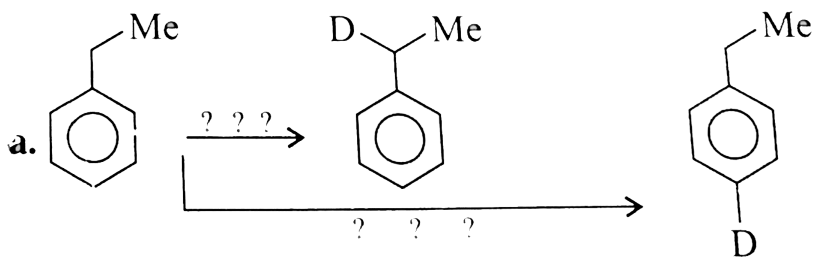
30. Complete the following reactions:



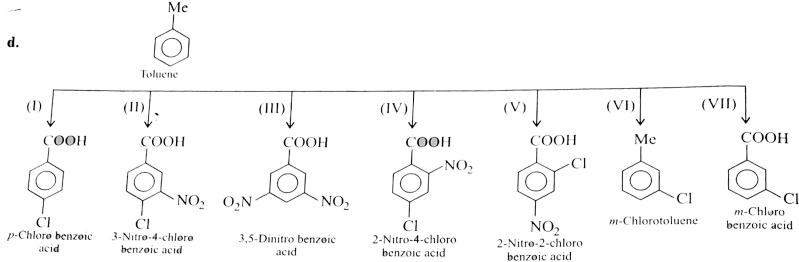


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31. Convert the following:

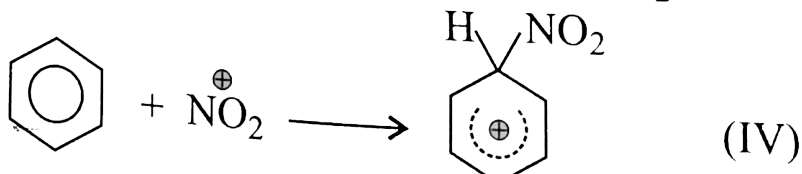
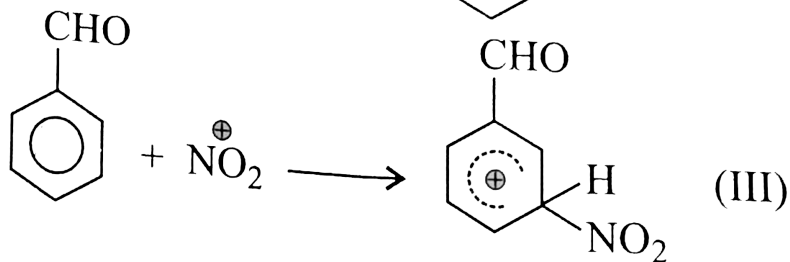
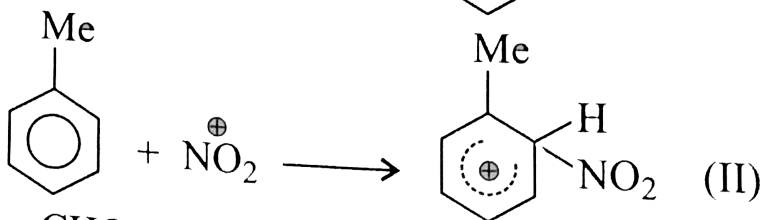
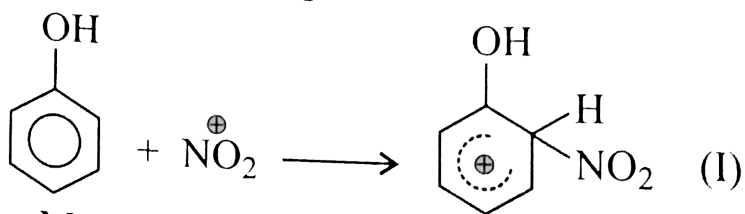


*p*-, *o*-, and *m*-Bromonitro benzene

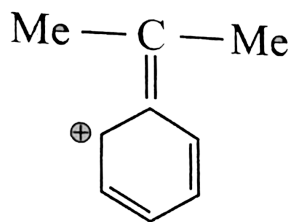


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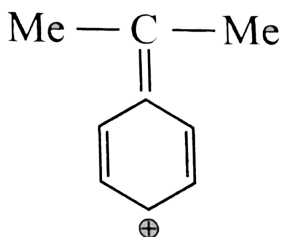
32. Answer the following species (I - IV) according to the decreasing order of potential energy.



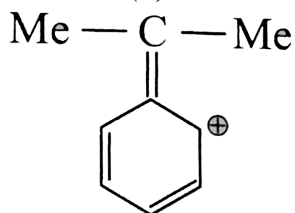
b. Which of the following resonance structures is/are wrong?



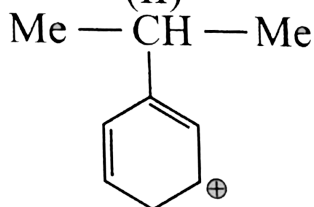
(I)



(II)

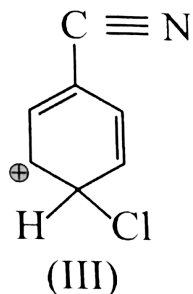
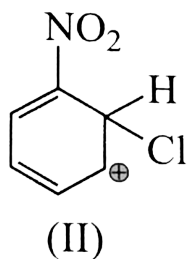
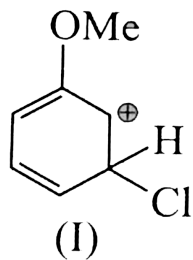


(III)

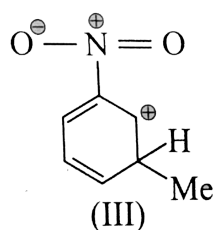
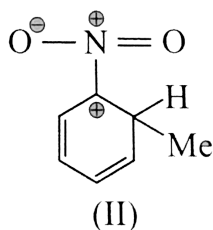
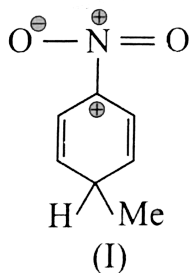


(IV)

c. Which of the following is/are stable resonating structures?



d. Which of the following is/are least stable arenium ion?



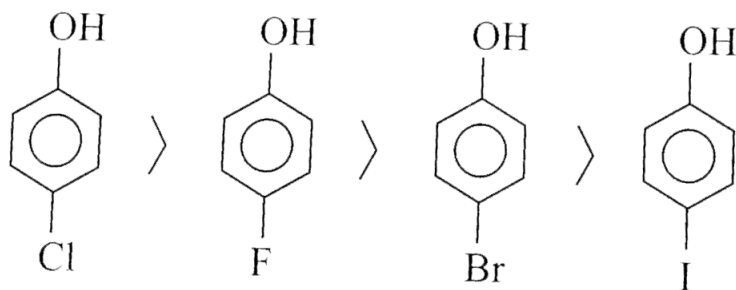
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33. Explain the following:

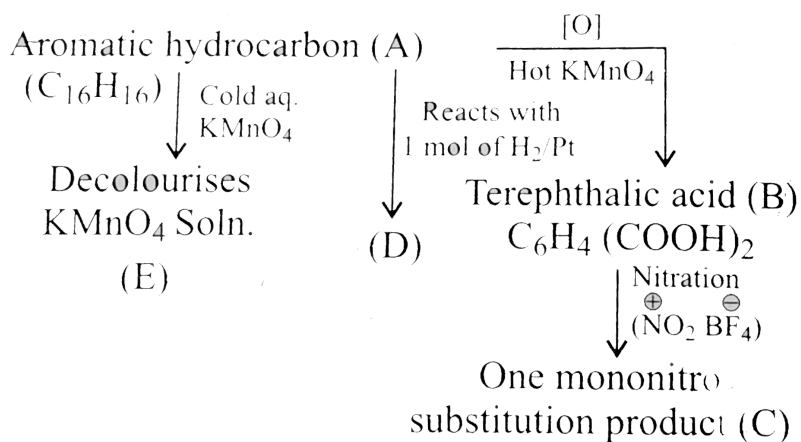
a. Reactivity of halogens in the presence of halogen carrier ( $FeX_3$ ) with benzene is  $F_2 > Cl_2 > Br_2 > I_2$ .

b. The reactivity of halogen-substituted benzene towards  $SE$  reaction is  $PhI > PhBr > PhCl > PhF$ .

C. The acidic character fo halogen-substituded pehnol is:



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34. Identify(A)to(E). Write the number of possible stereoisomers of(A).

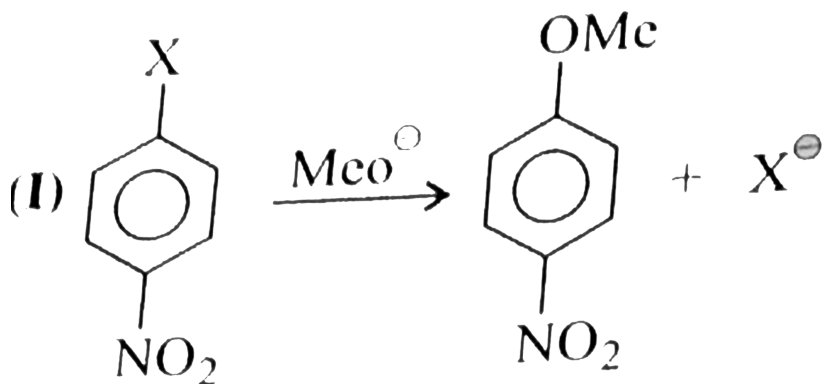
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35.

For

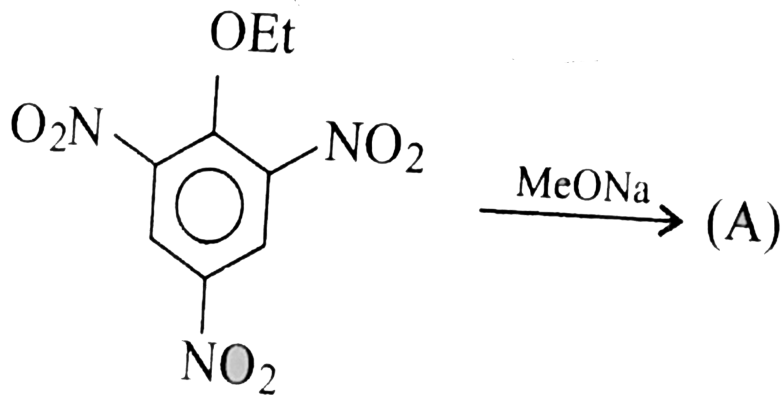
reaction(I)



,why is the

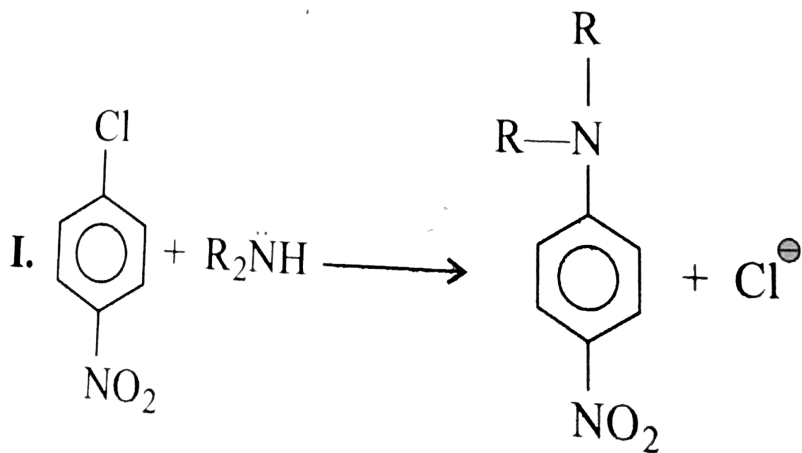
order of reactivity of halides  $\text{ArF} > \text{ArCl} > \text{ArBr} > \text{ArI}$ , although  $\text{F}^-$  is a poor leaving group? Explain.

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Write the structure of (A) and (B) and its resonance structures.

37. Give the decreasing order of reactivity of the following  $ArSN$  (addition-elimination reaction).



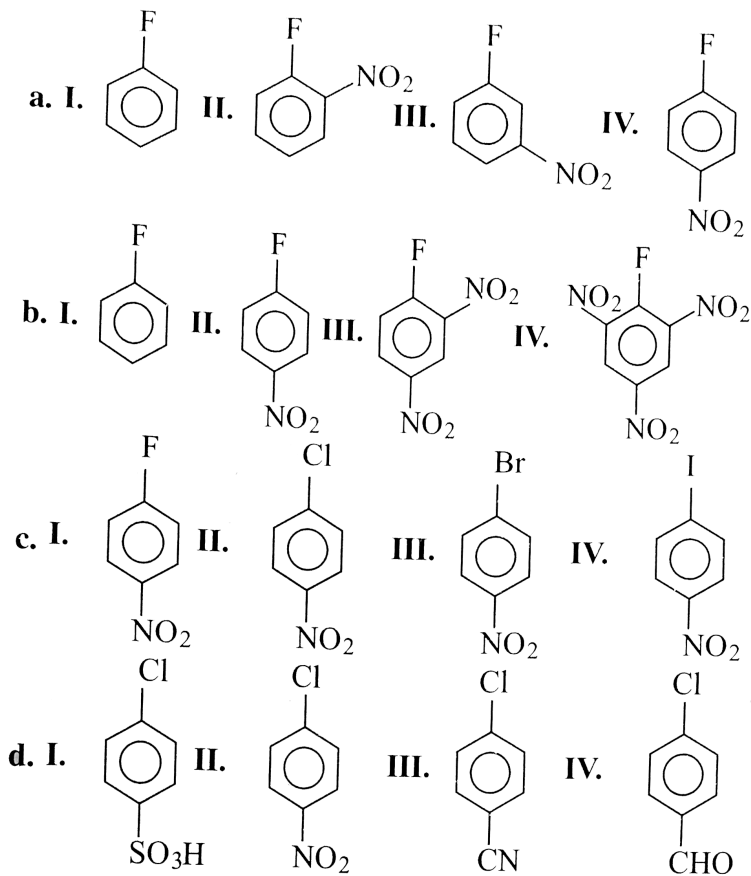
(a) I.

(II) 

(b)(III) 

(IV) 

38. Give the decreases order of  $ArSN$  reaction of the following with  $NaOMe$ :



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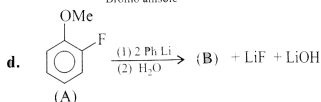
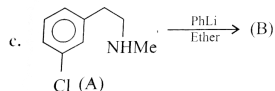
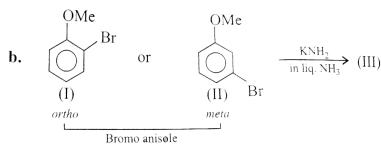
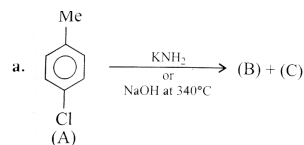
39. (a) Azulene has an appreciable dipole moment. Write the resonance structure that explains the dipole moment and its aromaticity.

(b) In the case of halogen-substituted azulenes, a halogen atom on C - 6 can

be displaced by nucleophile while one on C-1 is unreactive towards nucleophile. Rationalise this difference in behaviour.

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**40.** Give the reaction and explain the formation of the products.



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**41.** (a)  $\text{Me}_3\text{CCH}_2\text{CH}(\text{Cl})\text{Ph}$ ., (b)  $\text{MeC}(\text{Et})_2\text{CH}_2\text{Cl}$

(c)  $\text{MeCH} = \text{C}(\text{Br})\text{CH}_2\text{Me}_2$ , (d)  $m\text{-ClC}_6\text{H}_4\text{CH}_2\text{CHMe}_2$

(e)  $p\text{-I-C}_6\text{H}_4\text{CH}(\text{Me})\text{Et}$  (f)  $p\text{-FCH}_2\text{C}_6\text{H}_4\text{CH}_2\text{Cme}_3$

(g)  $\text{MeCH} = \text{CHC}(\text{Cl})\text{Me}_2$ , (h)  $\text{MeC}(\text{Br})(\text{et})\text{Et}$

(i)  $\text{MeCH} = \text{CHc}(\text{Cl})\text{Me}_2$ , (j)  $\text{EtC}(\text{Me})_2\text{CH}_2\text{Br}$

(k)  $\text{Me}_2\text{CHCH}(\text{Br})\text{Me}$  , (l)  $\text{BRCH}_2\text{C} \equiv \text{CH}_2\text{Cl}$

(m)  $\text{CHF}_2\text{BrClF}$  , (n)  $(\text{CBr}_3)_3\text{CBr}$  (o)  $\text{MeC}(\text{p-BrC}_6\text{H}_4)_2\text{CH}(\text{Cl})\text{Me}$   
, (p)  $\text{Me}_3\text{CCH}=\text{ClC}_6\text{H}_4\text{Cl-m}$

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**42.** Write the stick, condensed and bond line structure of the following compounds:

a. 4 - tertButyl-3 - chloroheptane      , b. 2 - (2 - Bromophenyl)-1 - chloro octane

c m-Bromochlorobenzene d Perchlorobenzene

e. 1 - Bromo-4 - ethyl cyclohexane . f. 2-Fluoro-3 - methyl pentane

1-Chloro-4sec butyl-2methyl benzene , 1, 4 - Dichloro but-2ene

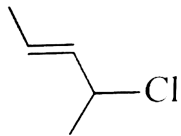
i. 4 - tert-Butyl-3 - fluoro heptane , j. 1-iodo-4 - ethyl cyclohexane

k. 2 - Fluoro-3 - methyl pentane

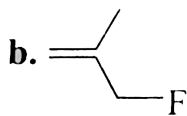
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43. Write IUPAC name of the following with (cis or trans) or (E and Z) if any.

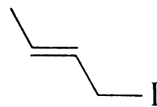
a.



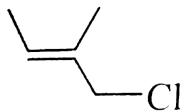
b.



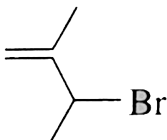
c.



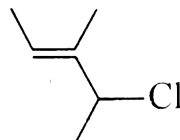
d.



e.



f.



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44. Arrange the following compounds in decreasing order of dipole moment values Explain the order

(a)  $CBr_4$ , (b)  $CHBr_3$ , (c)  $CH_2Br_2$ , (d)  $CH_3Br$ .

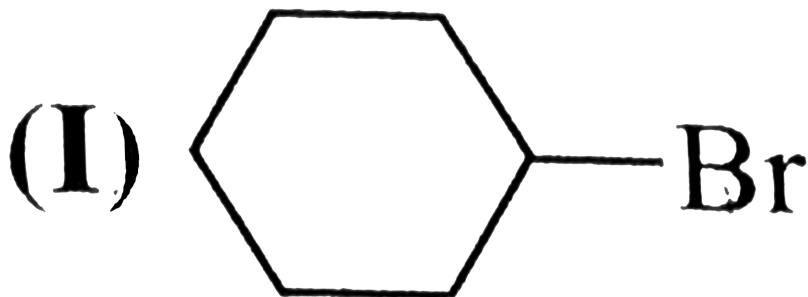


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45. a. Assign orientation to the three chlorotounes with  $\mu = 1.3, 1.78,$  and  $1.9D$ .

b. Assign orientation to the three chloronitrobenzenes with  $\mu = 2.5, 3.1,$  and  $4.3D$

c. Which has higher  $\mu$  : (1)

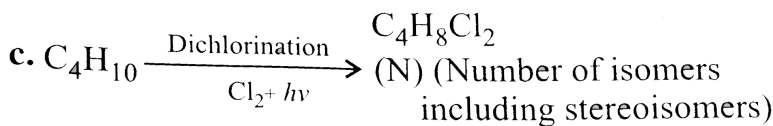
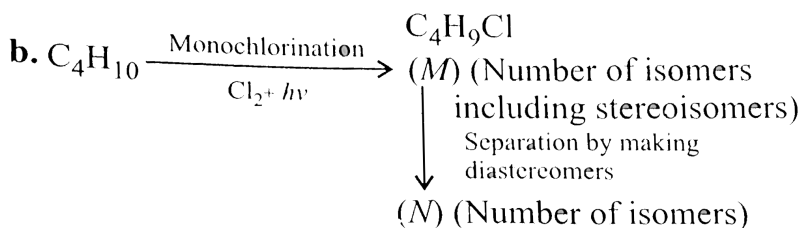
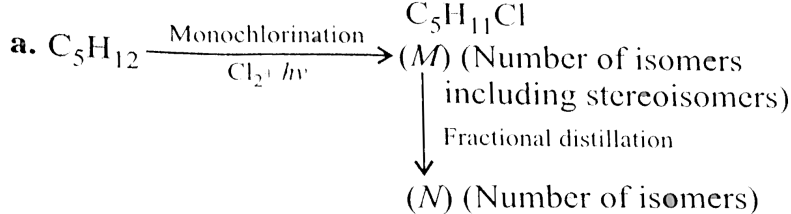


or II. *PhBr*?

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46. What are the numerical values of (*M*) and (*N*) in the following reaction

?



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47. a. Prepare the following ethers via Williamson's synthesis.

I. Di - n - propyl ether (A)

II. Benzyl methyl ether (B)

III. Phenylethyl ether (C)

IV. t - Butyl ethyl ether (D)

b. Which compound in the above problem, can be prepared by alternative Williamson's reaction?

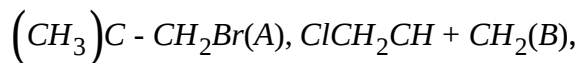
c. Explain the inability of (A), (C), (D) in the above problem to be



prepared via alternate Williamson's synthesis.

d. Give six types of ethers that cannot be synthesised by the typical williamson's synthesis.

e. Rank the following alkyl halides in the decreasing order of reactivity in Willamson's reaction.



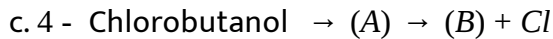
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48. a. An aromatic compound (A)  $(C_7H_8O)$  on reaction with  $Br_2 + H_2O$  gives a white ppt. of compound (B)  $(C_7H_5OBr_3)$ . Compound (C), an isomer of (A) also gives the same reaction and gives a white ppt. of compound (D)  $(C_7H_5OBr_3)$ . Compound (C) is insoluble in  $NaOH$ . identify (A), (B), (C) and (D).

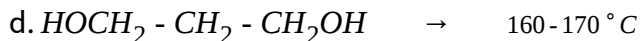
b. I. Starting from  $C_6H_6$  and  $C_6H_5OH$ , synthesise phenyl 1 - 2, 4 - dinitrophenyl ether (B).

II. Could we have first prepared and then nitrated  $Ph_2O$  ?

e.  
OH



Conc.  $H_2SO_4$

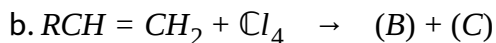


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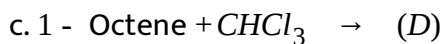
49. Identify (A) to (E).



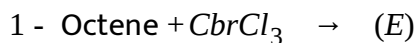
Peroxide



Peroxide

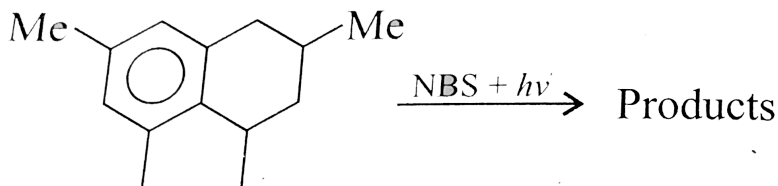
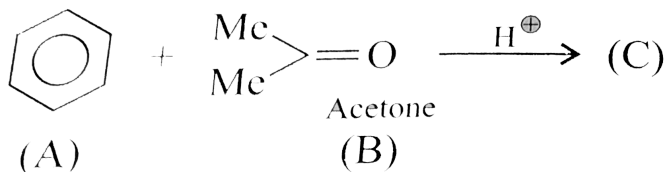


Peroxide



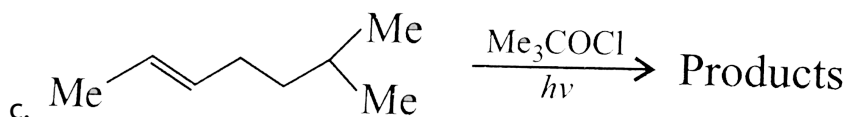
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50. a. Complete the following:



b.

Identify all the possible products. Give the major product and rank the products in decreasing order of reactivity with *NBS*.



Identify all the possible products. Give the major products and list them in decreasing order of reactivity with *Me<sub>3</sub>CoCl*.

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51. a. Explain the following

a. Dipole moment of *CH<sub>3</sub>F* is 1.85D and that of *CDF* is 1.86D.

b. 8 - Hydroxy quinoline can be separated from 4 - hydroxy quinoline by

steam distillation:

c. The boiling point of  $(C_2H_5)_2S > (C_2H_5)_2O$ .

d. The  $pK_a$  of *o*-fluorobenzoic acid (I) is 4.14, whereas that of *p*-chlorobenzoic acid (II) is 3.99.

e. Glycine exists as zwitterion, but PABA does not.

f. Write the structure of carbocation produced on treatment of a compound (A)  $(Ph_2CHC(OH)Me_2)$  with  $SbF_5/SO_2$ .



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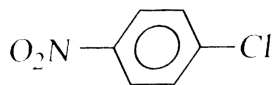
52. a. The dipole moment of  $HBr$  is  $2.6 \times 10^{-30} C.m$  and interatomic spacing is  $1.41 \text{ \AA}$ . What is the percent ionic character of  $HBr$ ?

b. A diatomic molecule has  $\mu = 1.2D$ . Its bond distance is  $1.0 \text{ \AA}$ . What fraction of electronic charge exists on each atom?

c. In water, ( $H - O - H$ ) bond angle is  $105^\circ$ . The distance between ( $O - H$ ) is  $0.94 \text{ \AA}$ .  $\mu$  of  $H_2O = 1.85D$ . Determine the magnitude of the charge on the oxygen atom in water molecule and hydrogen atom.

d.  $BI_3$  is a symmetrical planar molecule, all the ( $B - I$ ) bonds lie at  $120^\circ$  of each other. The distance between the  $I$  atoms is  $3.54 \text{ \AA}$ . The Estimate

the covalent radius of boron. e. Calculate the dipole moment of the following compound:



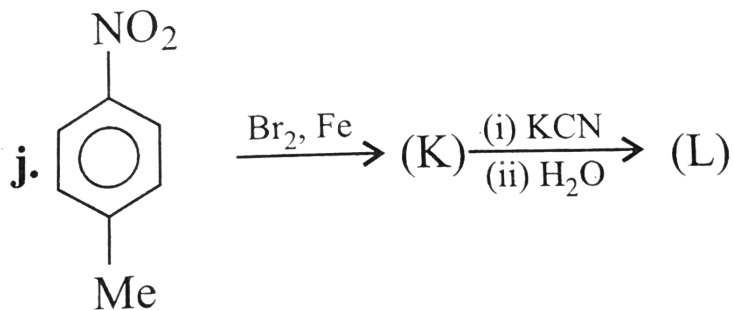
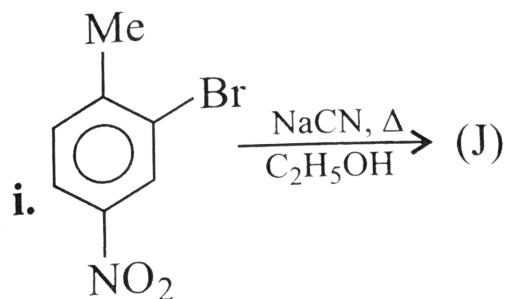
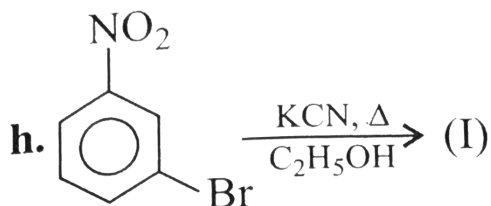
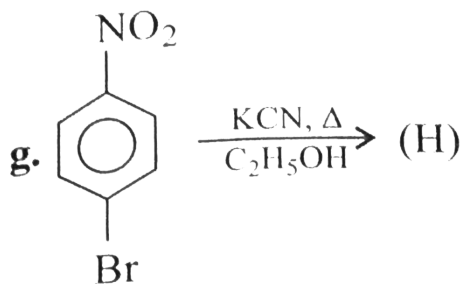
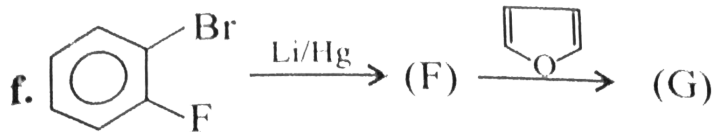
$$\text{Given : } \mu_{\text{C}-\text{Cl}} = 1.55 \text{ D}$$

$$\mu_{\text{C}-\text{NO}_2} = 3.95 \text{ D}$$

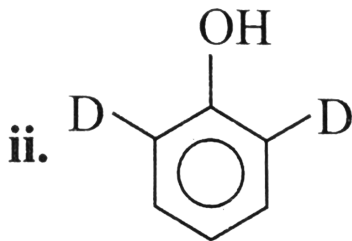
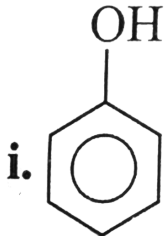
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53. Compler the following:

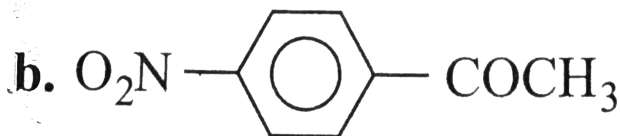
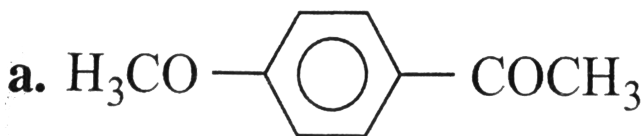




II. Which of the following has the greater  $K_a$  value?



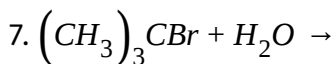
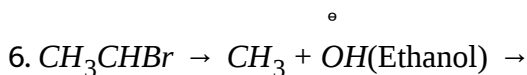
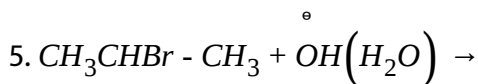
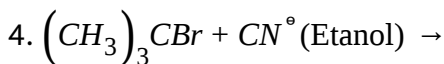
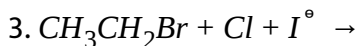
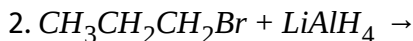
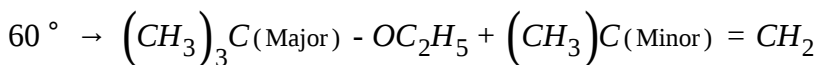
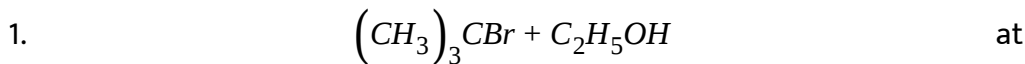
III. Which of the carbonyl groups in (a) and (b) protonate more readily in acid solution and why?



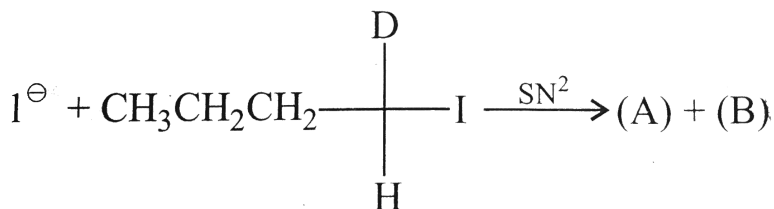
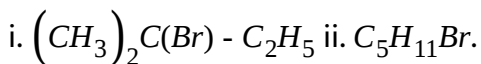
IV. When a trace of  $\text{KNH}_2$  is added to a solution of chlorobenzene and potassium triphenyl methide ( $\text{Ph}_3\text{C}^\ominus\text{K}^\oplus$ ) in liquid  $\text{NH}_3$ , a rapid reaction takes place to yield a product of formula  $\text{C}_{25}\text{H}_{20}$ . What is the product? What is the role of  $\text{KNH}_2$  and why is it needed?

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54. a. Indicate whether the following are  $SN^{-1}$ ,  $SN^{-1}$ ,  $E1$ , or  $E2$



b. Give the decreasing order of reactivity of the alkaline hydrolysis of the following alkyl halides towards (1)  $SN^{-1}$  (2)  $SN^2$  and (3) in the presence of  $AgNO_3$ :

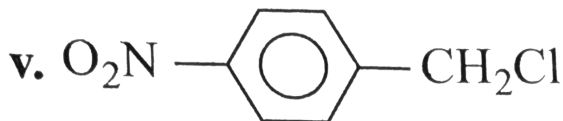
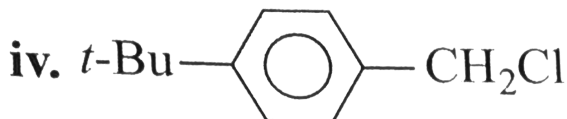
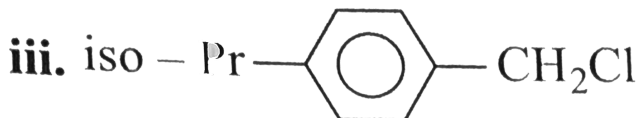
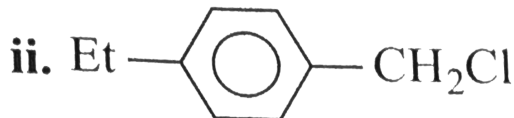
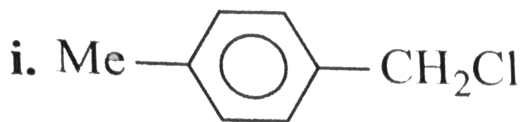


d. Give the decreasing order of reactivity of alkaline hydrolysis of the



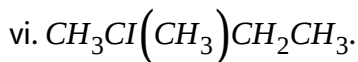
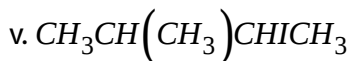
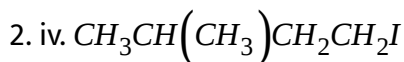
following halides towards  $SN^{-1}$ :

1.  $MeCl$ , ii.  $PhCH_2Cl$ , iii.  $Oh_2CHCl$ , iv.  $Ph_3Cl$



2.

e. Determine the relative  $SN^{-2}$  reactivity with  $NaCN$  in aprotic  $DMSO$  of the following alkyl halides:



3. vii. 4 - Bromo 1, 1 - dimethylcyclohexane

viii. 1 - Bromo -cis or trans-1, 4 - dimethylcyclohexane

ix. cis or trans -2 - Bromo-cis-1, 3 - Dimethyl-cyclohexane

f. Optically pure ( + ) - 2 - bromooctane,  $[\alpha] = +36^\circ$  reacts with aq.  $\text{NaOH}$  in acetone to give optically pure ( - ) - 2 - octanol,  $[\alpha] = -10.3^\circ$ . With partially racemised bromo compound whose  $[\alpha] = +30^\circ$ , the  $[\alpha]$  of alcohol product is  $-6.0^\circ$ . Calculate:

1. The percentage optical purity of partially racemised bromo compound and alcohol,
2. the percentage of inversion and racemisation, and
3. the percentage of frontside and backside attack.

g. Compare the rates of  $\text{SN}^{-1}$  and  $\text{SN}^{-2}$  reactions of:

1. Cyclopropyl and cyclopentyl chloride
2. Vinyl chloride

h. Arrange the following in the order of decreasing basicity:

1.  $\text{F}^-$ , 2.  $\text{O}^2-$ , 3.  $\text{NH}_2^-$ , 4.  $\text{CH}_3^-$

a.  $4 > 3 > 2 > 1$ , b.  $1 > 2 > 3 > 4$ ,

c.  $2 > 1 > 3 > 4$ , d.  $3 > 2 > 1 > 4$

i. Arrange the following in the order of decreasing nucleophilic character:

1.  $H_2O$ , 2.  $OH^-$ , 3.  $CH_3O^-$ , 4.  $CH_3COO^-$

i.  $2 > 3 > 4 > 1$ , ii.  $3 > 2 > 1 > 4$

iii.  $1 > 4 > 2 > 3$ , iv.  $4 > 1 > 2 > 3$

j. Arrange the following in the decreasing order as a leaving group in substitution reaction.

1.  $CH_3CO^-$ , 2.  $C_6H_5SO_3^-$ , 3.  $CH_3O^-$ , 4.  $CH_3COO^-$ . a.  $3 > 1 > 4 > 2$ , b.  $3 > 1 > 2 > 4$ , c.  $3 > 2 > 1 > 4$ , d.  $4 > 2 > 1 > 3$

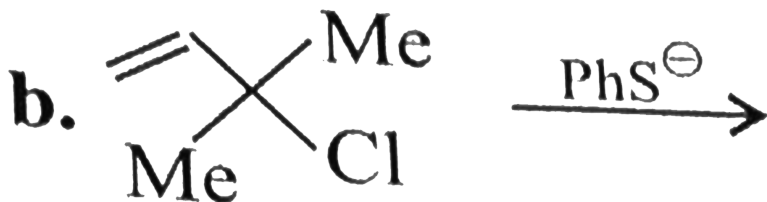
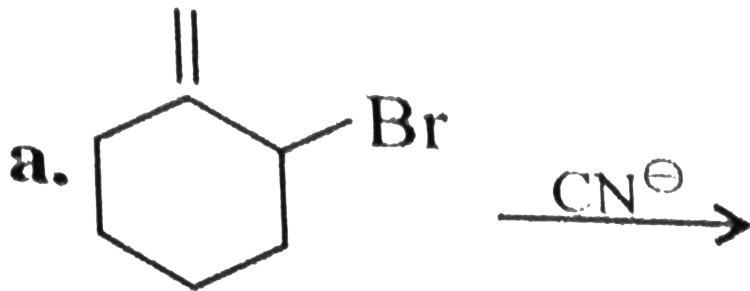
k. The order of leaving group ability of the following is: 1.  $-OAc$ , 2.  $-OMe$ , 3.  $-OSO_2Me$ , 4.  $-OSO_2CF_3$ . a.  $1 > 2 > 3 > 4$ , b.  $4 > 3 > 1 > 2$ , c.  $3 > 2 > 1 > 4$ , d.  $2 > 3 > 4 > 1$



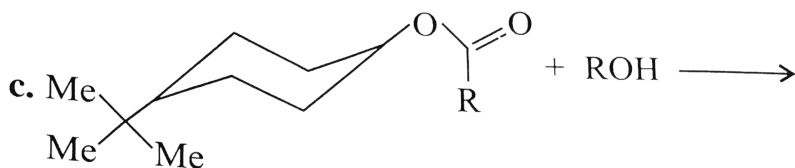
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55. Explain why solvolysis (where the solvent is nucleophilic) appears to follow a first-order rate law, rather than a second order.

II. Complete the following:



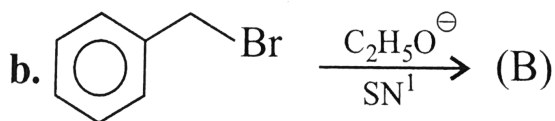
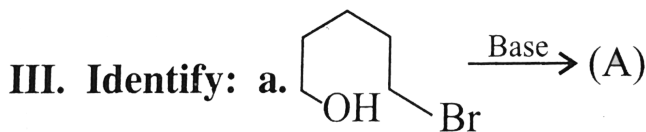
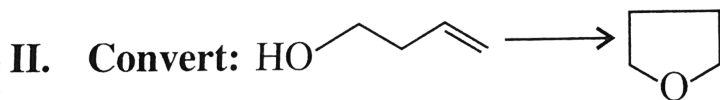
III. Predict the product of each  $S_N^2$  reaction. Indicate stereochemistry where appropriate.



IV. Why is  $PhO^-$  a weaker nucleophile than  $RO^-$ ?

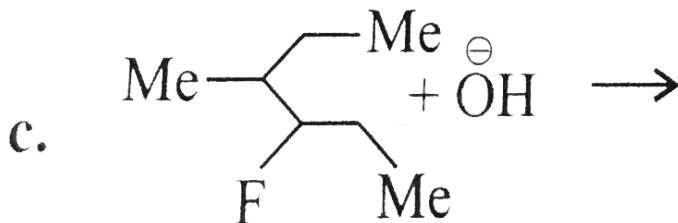
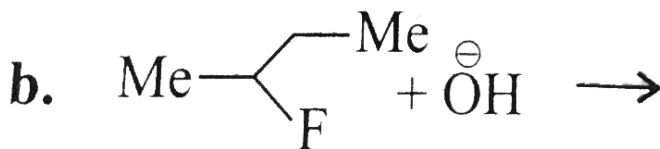
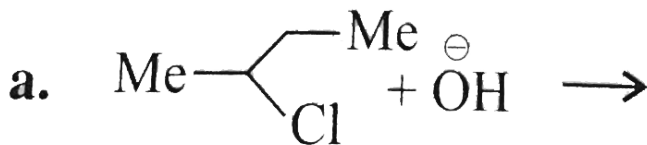
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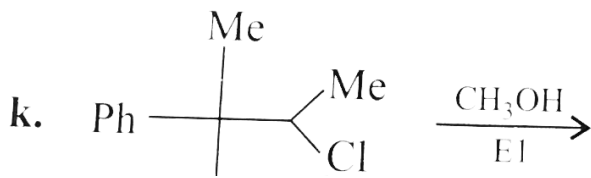
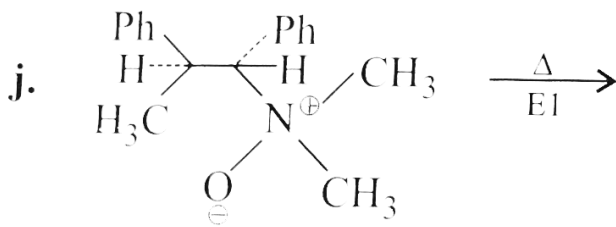
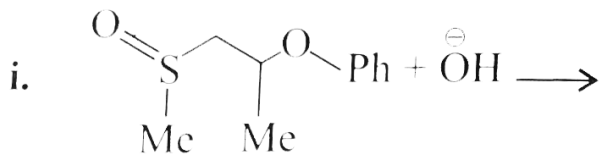
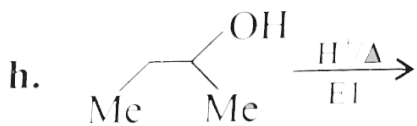
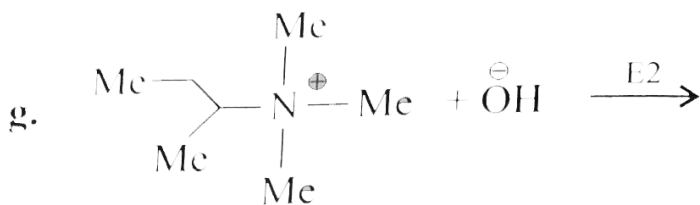
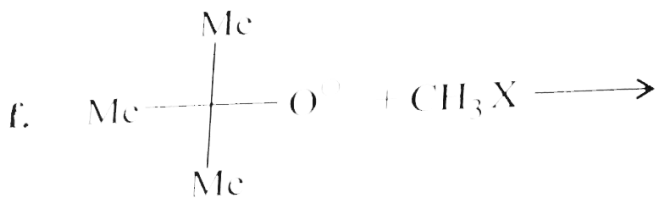
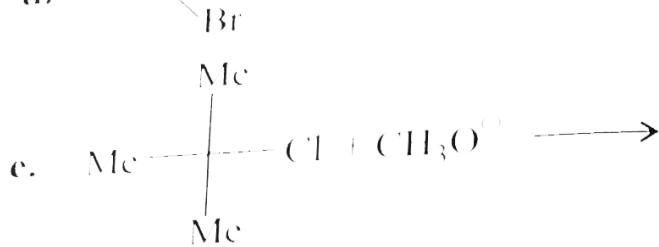
56. 1. 5-Chloro-1,3-cyclopentadiene undergoes  $S_N^1$  solvolysis in the presence of  $Ag^+$  ion slowly, even though chlorine is doubly allylic, and allylic halides normally ionise readily. Explain.



IV. 1-Chloro-2-(dimethylamino)propane (A) after standing for several weeks in a sealed container changes to 2-chloro-1-(dimethylamino)propane (B). Explain.

V. Give the major products of the following elimination reactions:

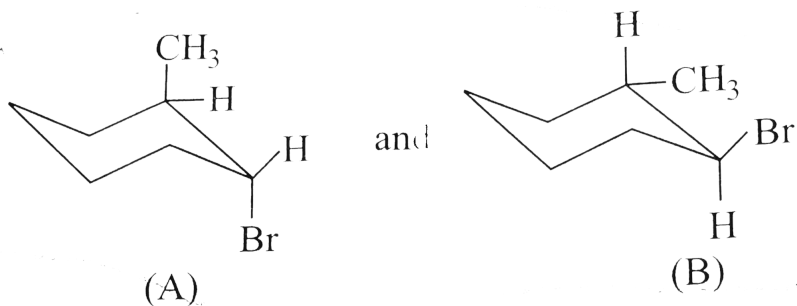




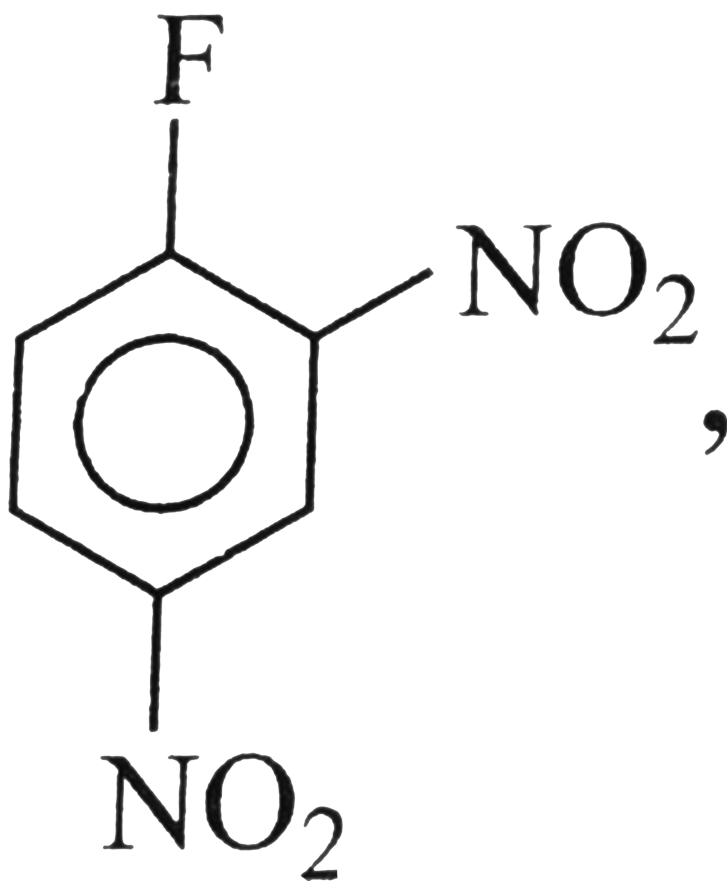
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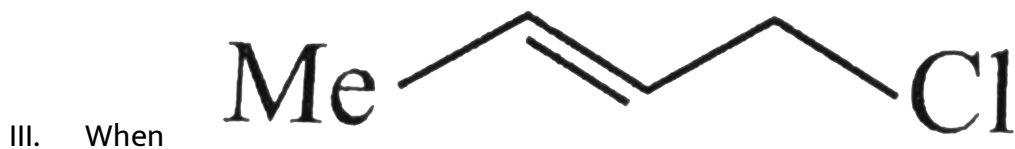
57. I. Which of the following will solvolyse faster is  $SN^{-1}$  and why?



II.  $F^{\ominus}$  works as a good leaving group in  $ArSN$  reaction



even though it is a poor leaving group in aliphatic  $S_N^{-1}$  and  $S_N^{-1}$  mechanisms. Explain.



reacts with alc.  $KCN$ , a mixture of isomeric products is obtained. Explain.



## Solved Examples

1. Benzene, toluene, xylene (*o*, *m*, *p*) and mesitylene dissolve in  $\text{HBF}_4$  to form salts. Explain the order of basicity:

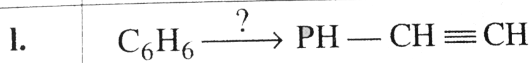
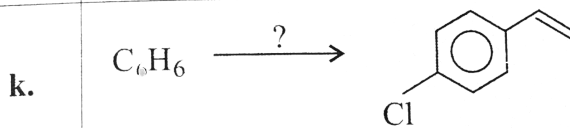
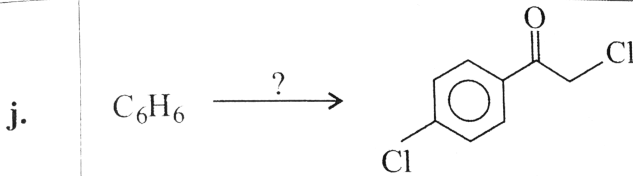
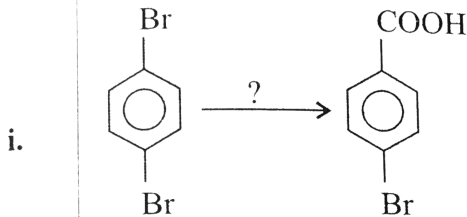
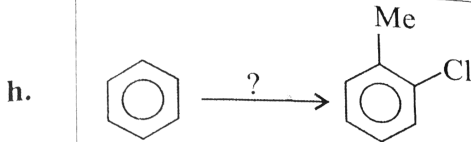
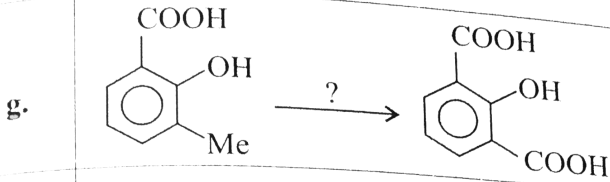
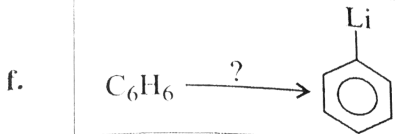
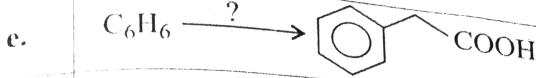
Mesitylene > *m*-Xylene > *o*- and *p*-Xylenes > Toluene > Benzene



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2. Complete the following :

a.	$2 \text{ } \begin{array}{c} \text{Me} \\   \\ \text{C}_6\text{H}_3 \\   \\ \text{NO}_2 \end{array} \text{I} + \text{Cu} \xrightarrow{\Delta} \text{B} + \text{C}$ <p>(A)</p>
b.	$2 \text{ } \begin{array}{c} \text{C}_6\text{H}_4 \\   \\ \text{NO}_2 \end{array} \text{I} + \text{Cu} \xrightarrow{\Delta} \text{B}$ <p>(A)</p>
c.	$\text{C}_6\text{H}_5\text{CH}_2\text{Me} \xrightarrow{??} \text{C}_6\text{H}_6$
d.	$\text{C}_6\text{H}_6 \xrightarrow{??} \text{C}_6\text{H}_4(\text{Me})_2$



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3. a. The treatment of  $RX$  with aqueous  $KOH$  leads to the formation of alcohols but in the presence of alcoholic  $KOH$  or  $NaOH$ , alkenes are the major products. Explain why.
- b.  $CHF_3$  is less acidic than  $CHCl_3$ . Explain.
- c. Wurtz reaction in case of tert-alkyl halide fails. Explain.
- d. Dipole moment of  $C_6H_5Cl$  is lower than that of  $C_6H_{11}Cl$  (cyclohexyl chloride). Explain why.
- e. Why should Grignard reagent be prepared under anhydrous conditions?
- f. Why does *p*-dichlorobenzene have higher melting point and lower solubility than *o*- and *m*-isomers?



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4. Give the decreasing order of following with their properties as indicated.

A. Decreasing basic, nucleophilic and fugacity orders:

a. I,  $H_2O$ , II.  $CH_3OH$ , III.  $O$  over  $(o)(H)$ , IV.  $CH_3O^-(o)$  b. I.  $H_2O$

$\oplus$ 

, II.  $C_2H_5OH$ , III.  $OH^-$ , IV.  $C_2H_5O^-$

c. I.  $HCO_3^-$ , II.  $F^-$

III.  $F_3C-COO^-$ , IV.  $NO_3^-$

B. The decreasing order of *ArSN* reaction:

a. I. *PhCl*

II. *p*- $NO_2-C_6H_4-Cl$

III. 2, 4, 6 - Trinitro chlorobenze

IV. 2,4 - Dinitro chlorobenze

I. *PhF*

II. *p*- $NO_2-C_6H_4-F$

III. *p*- $HOOC-C_6H_4-F$

IV. *p*- $NO_2-C_6H_4-F$

C. I. *PhCl*

II. *p*- $NO_2-C_6H_4-Cl$

III. *o*- $NO_2-C_6H_4-Cl$

IV. *m*- $NO_2-C_6H_4-Cl$

C. The decreasing order of *SE* reaction:

a. I. *PhCl*, II.  $C_6H_6$

III. *PhCH<sub>3</sub>*, IV. *PhOMe*

b. I. PhCH<sub>3</sub> II. o-MeO-C<sub>6</sub>H<sub>4</sub>-Me III. m-MeOC<sub>6</sub>H<sub>4</sub>-Me IV. p-MeO-C<sub>6</sub>H<sub>4</sub>-Me  
 V. PhNH<sub>2</sub> VI. PhNHCHO VII. PhNHCOPh VIII. PhNHTsd.  
 I. PhNH<sub>2</sub> II. o-NP<sub>2</sub>-C<sub>6</sub>H<sub>4</sub>-NH<sub>2</sub> III. m-NUO<sub>2</sub>-C<sub>6</sub>H<sub>4</sub>-NH<sub>2</sub> IV. p-NO<sub>2</sub>-C<sub>6</sub>H<sub>4</sub>-NH<sub>2</sub>



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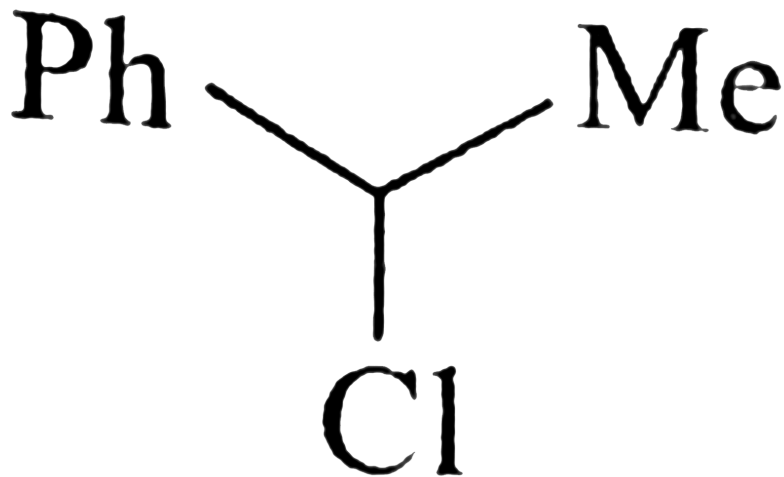
5. Distinguish between the following compounds:

a. I. *m*-Toluene and II. Benzyl iodide.

b. I. *p*-Bromotoluene and II. *p*-nitrotoluene,

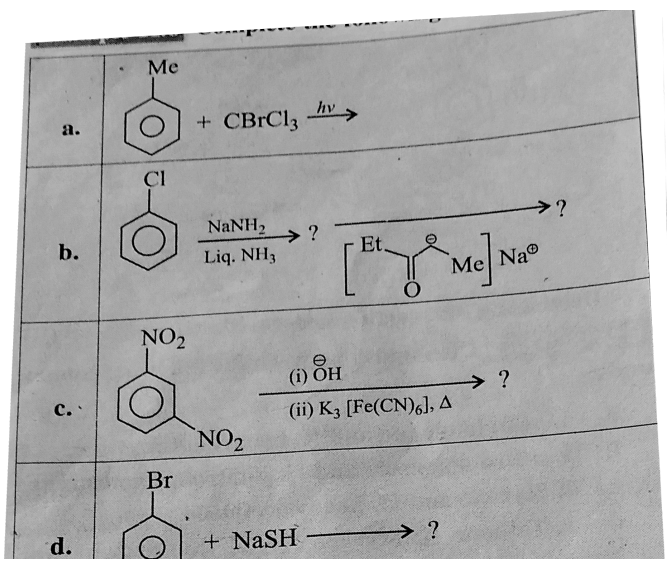
c. I. Styrene and II. Phenylacetylene,

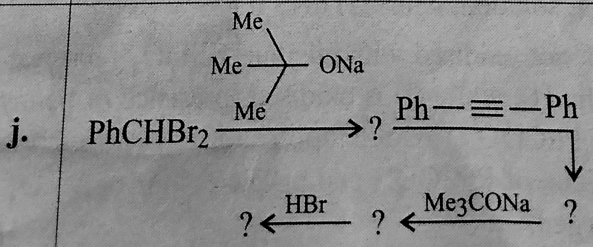
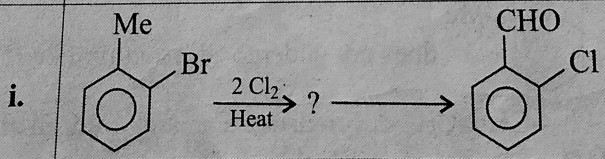
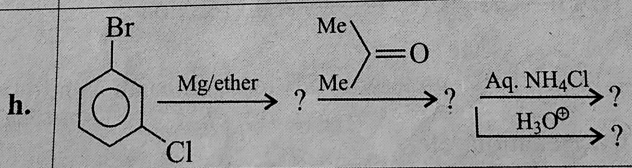
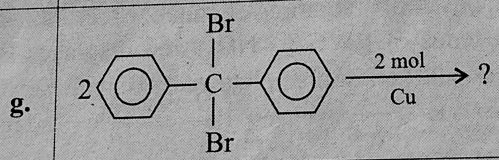
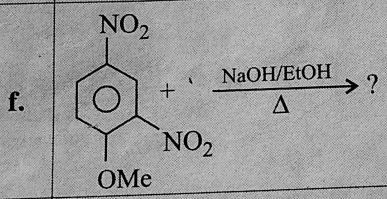
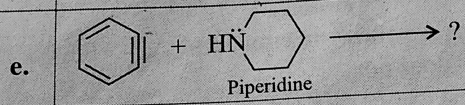
d. I. Touene II. Stryene III. Methyl cyclohexane, IV.  $\text{PhCMe}_3$  V.



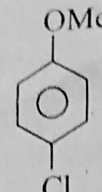
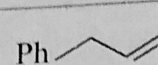
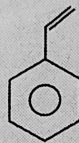
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6. Complete the following reactions:



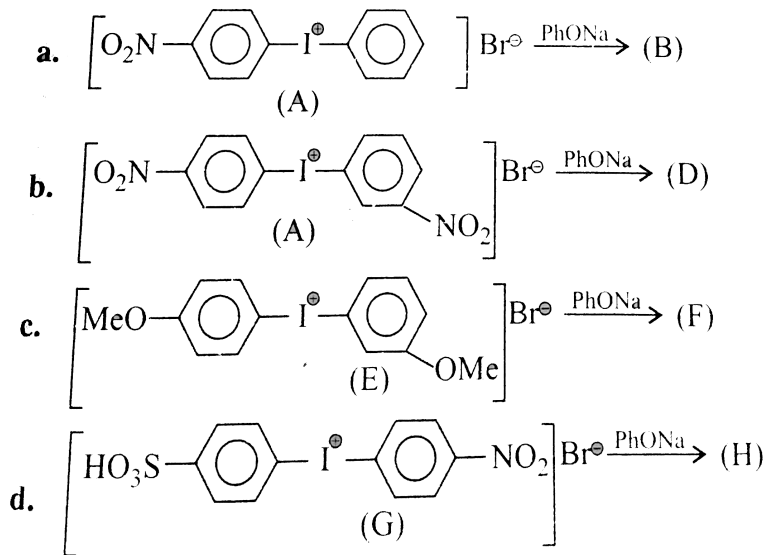




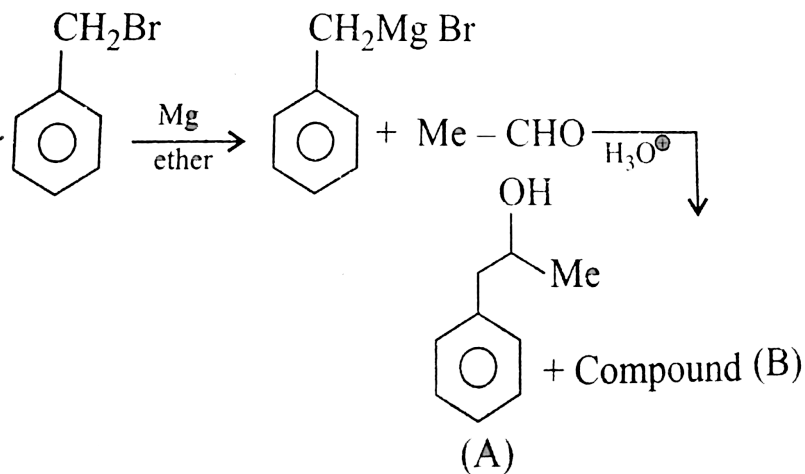
k.	 $\xrightarrow[\text{Ether}]{\text{LiNMe}_2} ? + ?$
l.	 $\xrightarrow[\Delta]{\text{KOH}} ?$
m.	$\text{Ph}_3\text{C}-\text{OH} + \text{HBF}_4 \xrightarrow[\text{Solvent}]{(\text{CH}_3\text{CO})_2\text{O}}$ <p>Triphenyl carbinol or Triphenyl methanol</p> <p>Bright orange ppt.</p> <p>Colour is discharged <math>\xleftarrow[\text{or ROH}]{\text{H}_2\text{O}}</math></p>
n.	$\text{Ph}_3\text{C}-\text{Cl} \xrightarrow{2\text{Na/Hg in ether}}$ <p>Chlorotriphenyl methane</p> <p>Deep red colour</p> <p>Colour is discharged <math>\xleftarrow[\text{or ROH}]{\text{H}_2\text{O}}</math></p>
o.	 $\xrightarrow{\text{Br}_2 + \text{FeBr}_3} (\text{B}) + (\text{C}) + (\text{D})$ <p>Styrene (A)</p>

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7. Indicate the position where  $ArSN$  reaction will take place and explain why.

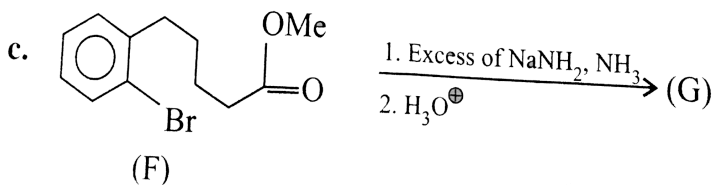
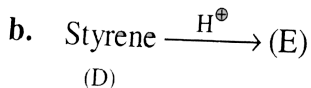
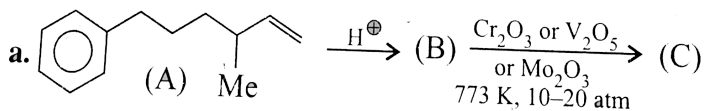


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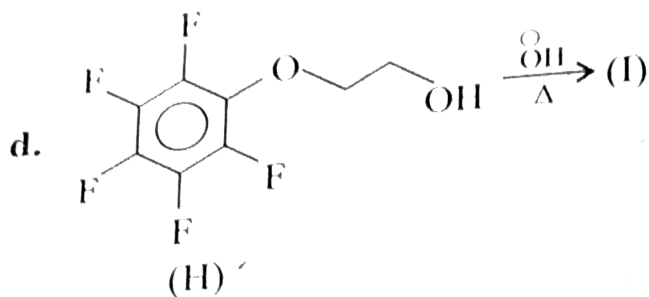


8.

Compound (B) is an isomer of (A). Compound (B) shows positive iodoform test and gives *o*-toluic acid. What is (B)? Explain its formation.



9.



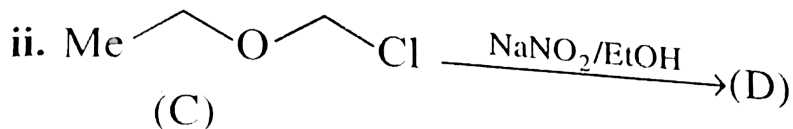
10. a. Account for the rapid rate of ethanoyl chloride



, although it

is  $1^\oplus$  halide.

Explain the following reactions:

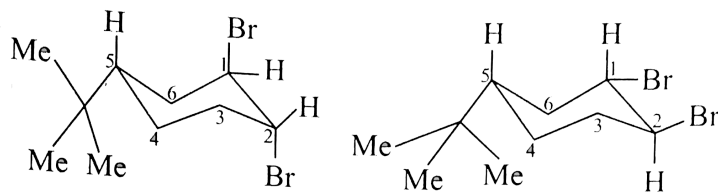


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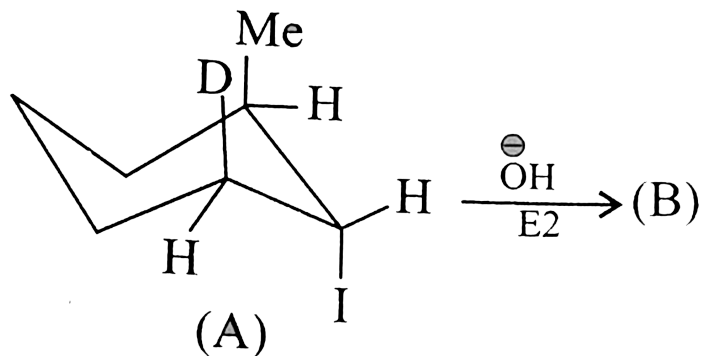
11. I. Give the product of debromination of with  $KI$  in acetone solution of the following.

a. Erythro or mes-2, 3 - dibromobutane

II. Explain why (A) readily forms alkene with  $1^\ominus$  but (B) does not.



III. Give the major product of the following reaction:

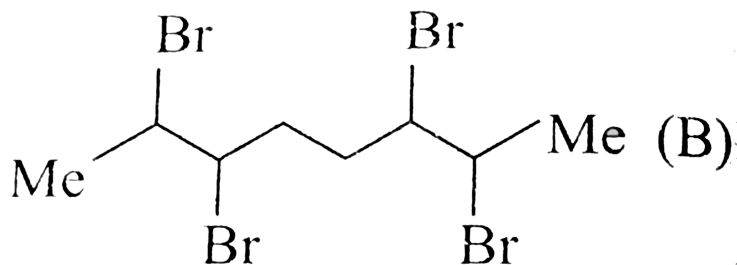


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12. Synthesise the following:

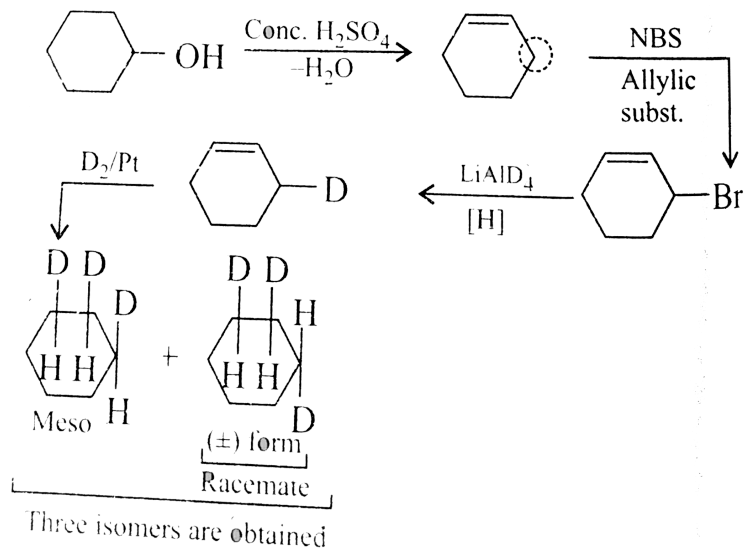
a. Cyclohexanol (A) to 1, 2, 3 - tridenuterocyclohexane (B).

b.  $E$ -but-2-ene to  $Z$ -but-2-ene



i. Which diastereomer of (B) is obtained?

ii. If  $Z$ -but-2-ene is used, which diastereomer of (B) is obtained?

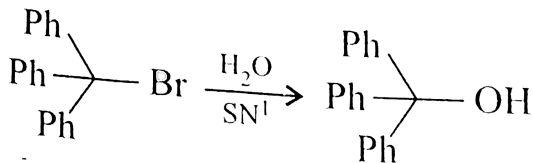
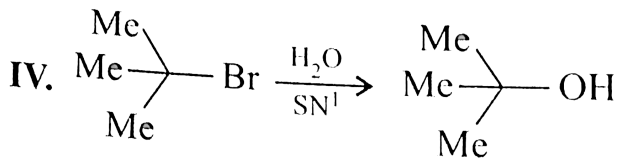
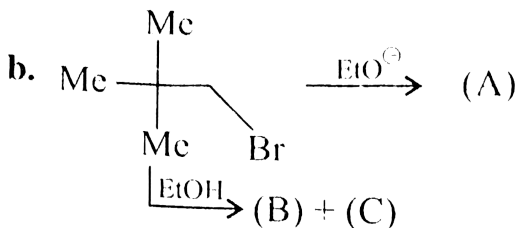
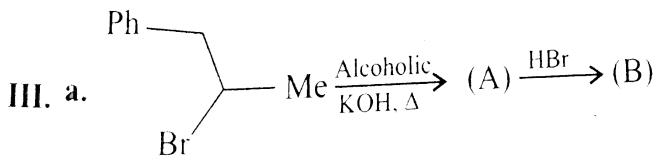
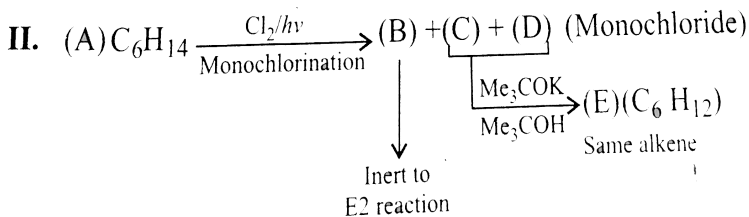
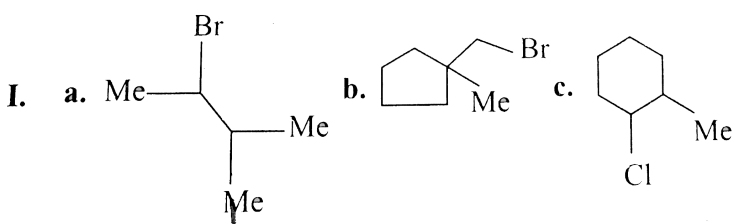


c.

What is product (C)? Which reaction [(1) or (2)] is faster and why?

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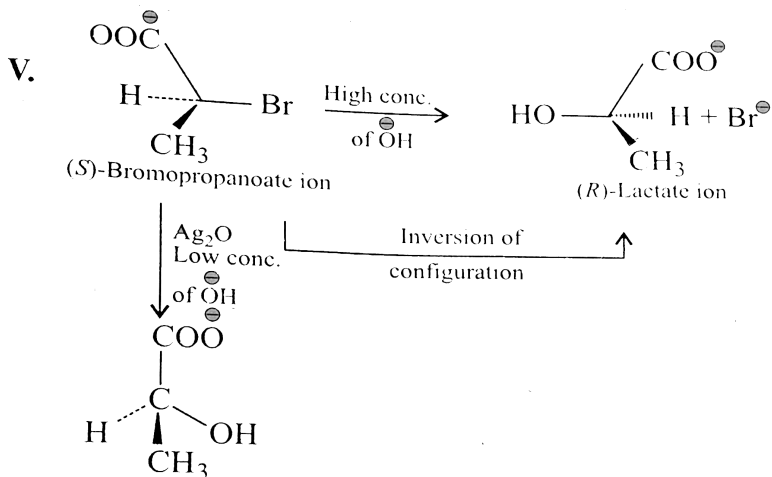
13. Give the major product, when the following compounds are treated with sodium methoxide.



Equation (1) shows no effect with the addition of  $Br^-$  to the reaction but

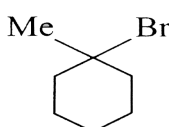
Eq. (2) shows the common ion effect of  $Br^-$  if supplied from external

source. Why?



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14. I. Which of the following solvents is the reaction faster?

- a.  $\text{Me} - \text{CH}_2 - \text{CH}_2 - \text{Cl} + \text{NH}_3$  in 20%  $\text{CH}_3\text{OH}$  and 80%  $\text{H}_2\text{O}$  or 40%  $\text{CH}_3\text{OH}$  and 60%  $\text{H}_2\text{O}$ .
- b.  in  $\text{CH}_3\text{OH}$  or  $\text{EtOH}$ .
- c.  $\text{Me} - \text{CH}_2 - \text{CH}_2 - \text{CH}_2 - \text{I} + \text{OH}^-$  in  $\text{CH}_3\text{OH}$  or 50%  $\text{CH}_3\text{OH}$  + 50%  $\text{H}_2\text{O}$ .
- d.  $\text{Me} - \text{CH}_2 - \text{CH}_2 - \text{Cl} + \text{:C} \equiv \text{N:}^-$  in  $\text{EtOH}$  or  $\text{DMSO}$ .

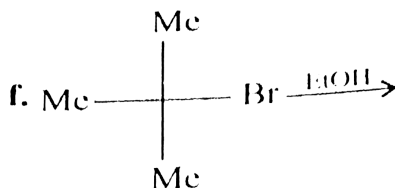
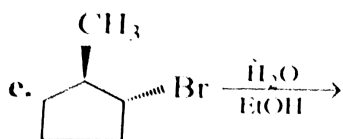
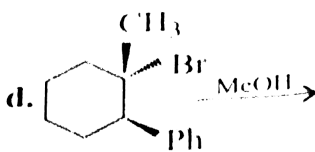
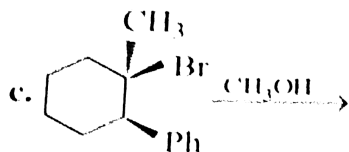
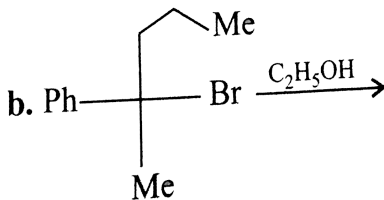
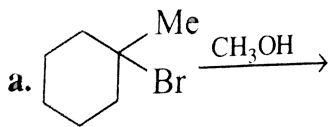
II. Indicate  $\text{SN}^{-1}$  or  $\text{SN}^2$ .



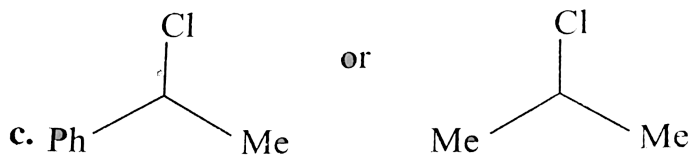
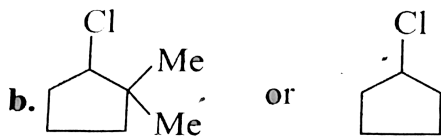
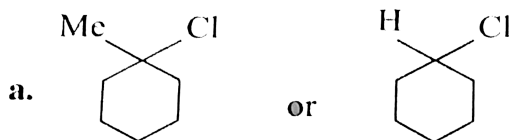
- a.  $\text{Me}-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{OTs} + \text{Cl}^- \xrightarrow{\text{CH}_3\text{OH}}$
- b.  $\text{Me}-\text{C}(\text{Me})_2-\text{Br} + \text{CH}_3\text{COO}^- \xrightarrow{\text{CH}_3\text{COOH}}$
- c.  $\text{Me}-\text{CH}_2-\text{CH}(\text{Cl})-\text{Me} + \text{H}_2\text{O} \xrightarrow[\text{EtOH}]{\text{H}_2\text{O}}$
- d.  $\text{Cyclopentane ring with OMe} + \text{SH}^- \xrightarrow{\text{DMF}}$
- e.  $\text{CH}_3\text{I} + \text{Me}-\text{C}(\text{Me})_2-\text{O}^- \xrightarrow{t\text{-BuOH}}$
- f.  $\text{CH}_2=\text{CH}-\text{CH}_2-\text{Cl} + \text{CH}_3\text{OH} \xrightarrow{\text{CH}_3\text{OH}}$
- g.  $\text{CH}_2=\text{CH}-\text{CH}_2-\text{Cl} + \text{CH}_3\text{COO}^- \xrightarrow{\text{CH}_3\text{COOH}}$
- h.  $\text{C}_6\text{H}_5-\text{CH}(\text{Br})-\text{Me} + \text{CH}_3\text{CH}_2\text{OH} \xrightarrow{\text{EtOH}}$

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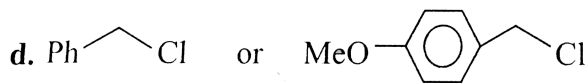
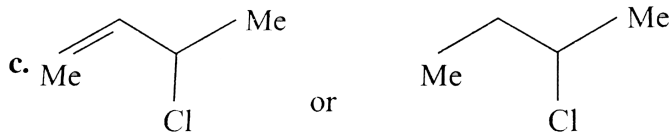
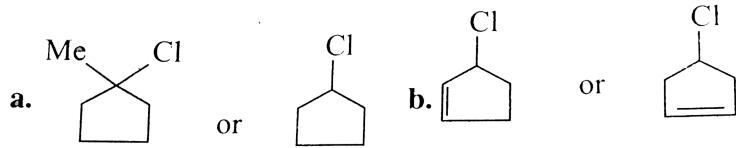
15. I. Give the products of the following:



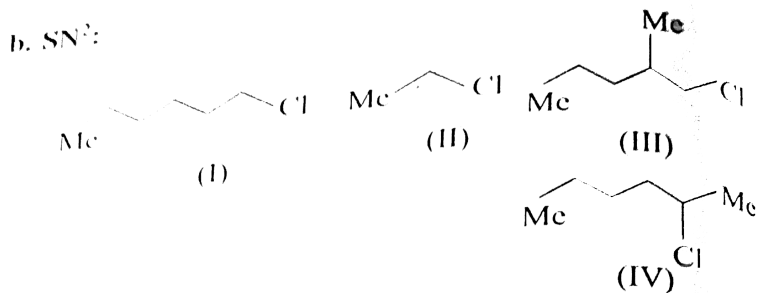
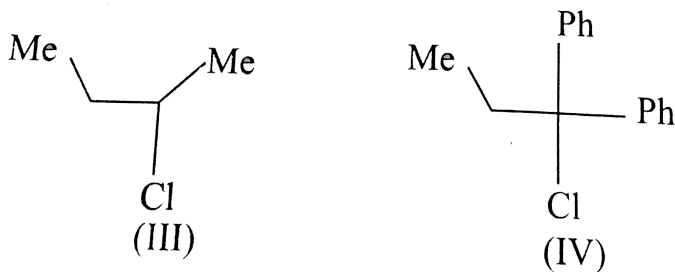
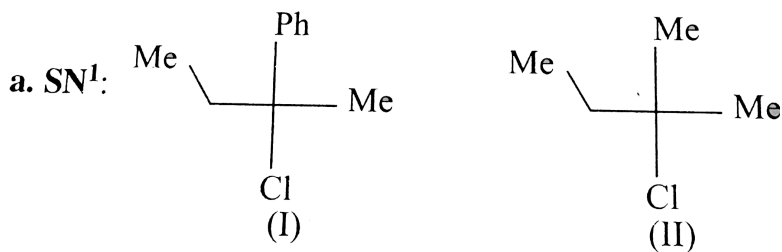
II. Which has faster rate of  $S_N1$ ?



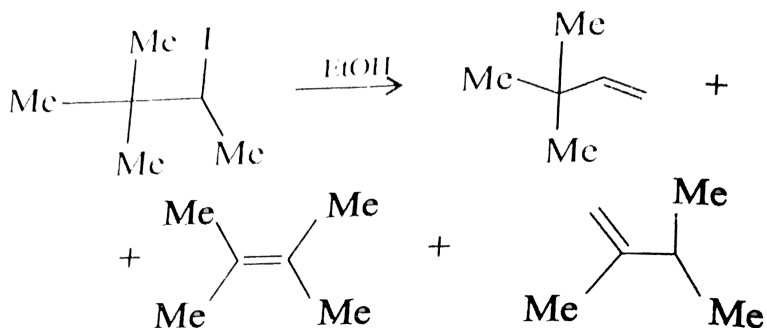
III. Which has faster rate of  $S_N1$ ?



IV. Give the decreasing order of:

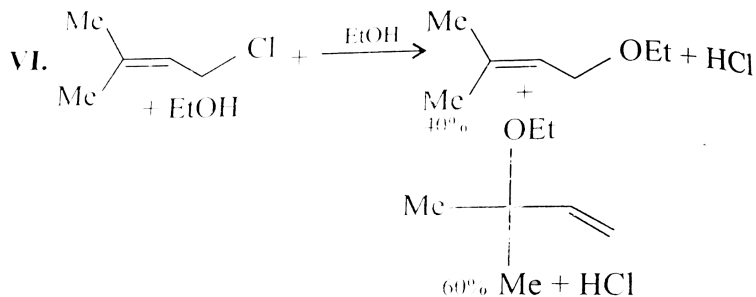


Show the mechanism for the formation of these products and sub-products.

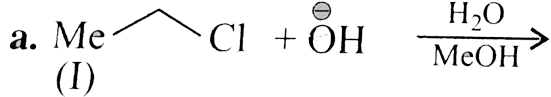


Explain the formation and mechanism.

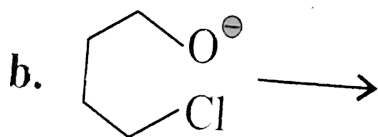
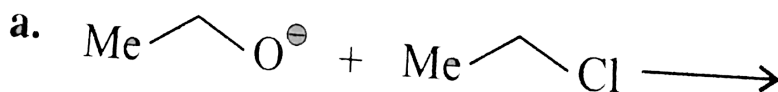
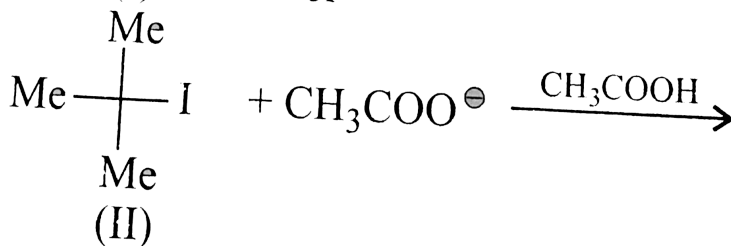
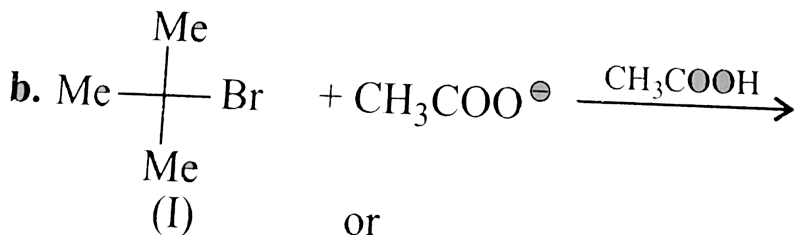
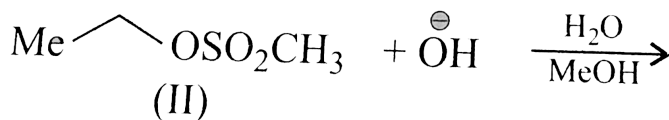
VII. Explain whether these reactions would follow the  $S_N^{-1}$  or  $S_N^{-2}$  and which is faster.



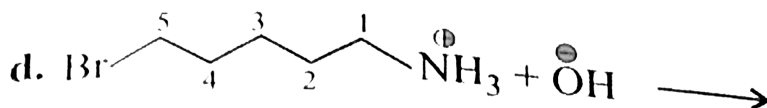
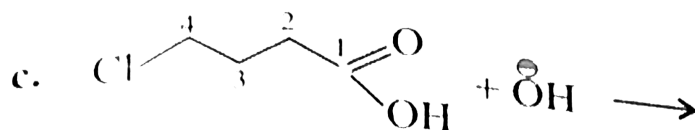
VIII. What are the products of the following reactions?



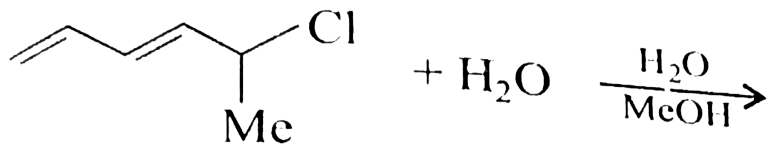
or



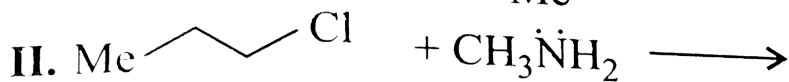
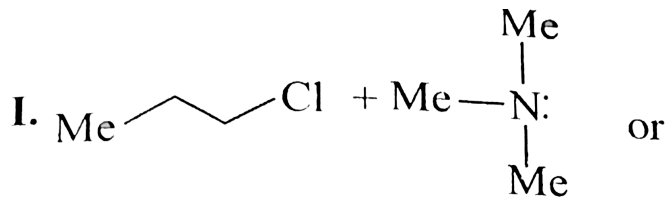
IXgt a. This reaction gives three substitution products. Show the structure.



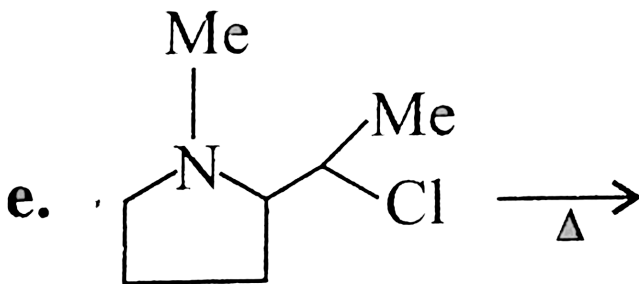
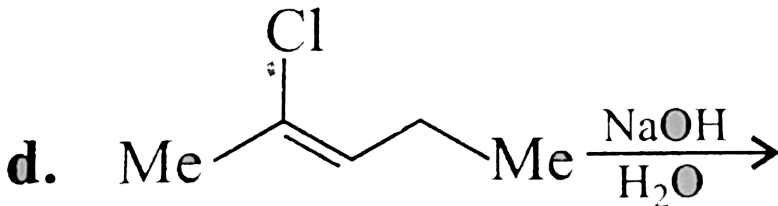
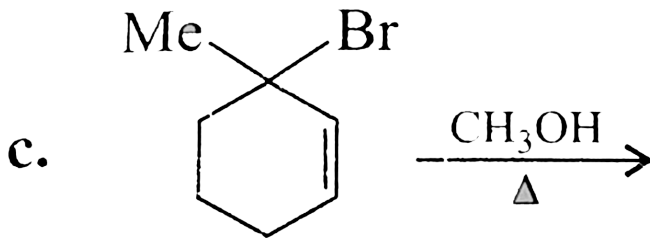
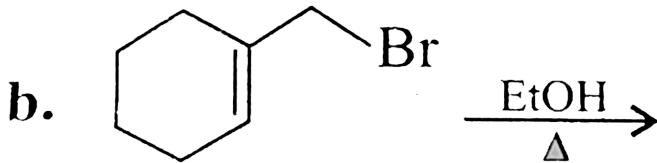
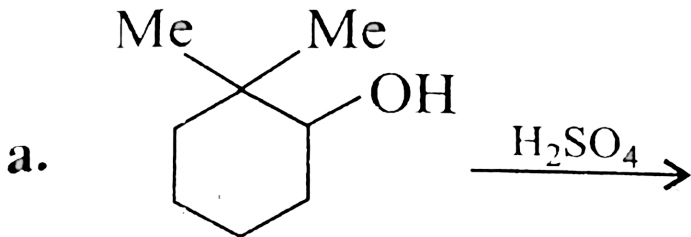
b. Which has faster rate?



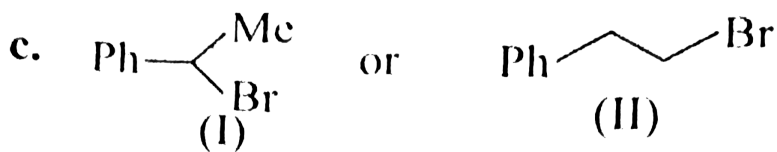
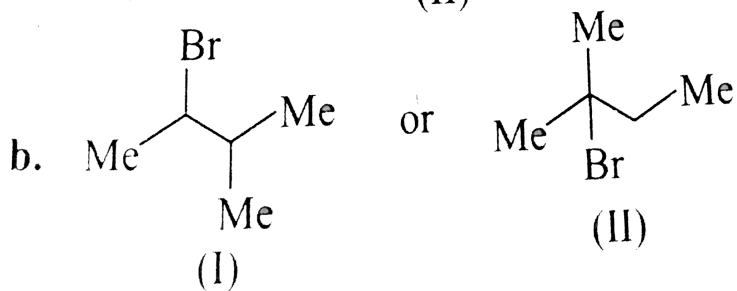
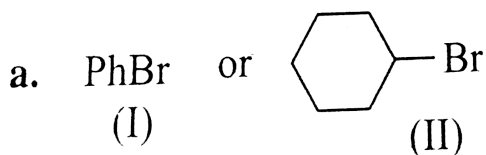
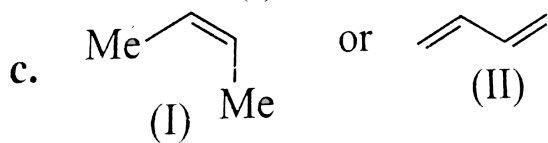
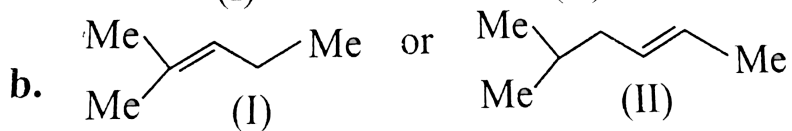
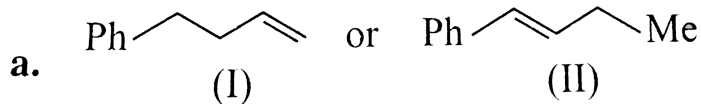
X. Give the products of the following:



XI. Which is more reactive towards acid-catalysed hydration and explain the regiochemistry of the alcohol formed.



XII. Which hydrolyses at faster rate?



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16. A Grignard reagent (A) and a haloalkene (B) react together to give (C) compound (C) on heating with  $KOH$  yields a mixture of two geometrical isomers, (D) and (E) of which (D) predominates. (C) gives 1-bromo-3-phenylpropane on reaction with  $HBr$  in the presence of a peroxide. Give the structures of (A), (B), and (C) and the configurations of (D) and (E).

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17. Catalytic dehydrogenation of methylcyclohexane, obtained from petroleum, gives a liquid which on treatment with chlorosulphonic acid at  $370K$  yields a mixture of two isomers (A) and (B),  $C_7H_7SO_2Cl$ . The major isomer (A) reacts with ammonia to form (C), which on oxidation with permanganate gives compound (D) gives a well-known sweetening agent. (E). The major isomer (B) also reacts with ammonia to give a compound (F) which on treatment with  $NaCl \frac{\text{O}}{N} aOH$  gives an antiseptic (G). Identify (A) to (G).

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18. An organic compound (A) on analysis was found to contain  $C = 16.271\%$ ,  $H = 0.677\%$ , and  $Cl = 72.203\%$ . It reduced Fehling's solution and on oxidation gave a monocarboxylic acid (B), having  $C = 14.679\%$ ,  $H = 0.612\%$ , and  $Cl = 65.137\%$ . On distillation with soda lime, (B) gave a sweet-smelling liquid (C), containing 89.12% chlorine. (C) can also be obtained by heating (A) with alkali. What structural formulae would you assign to (A), (B), and (C)?

Explain the above reactions:.



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19. Benzene reacts with  $CH_3I$  in the presence of  $AlCl_3$  to give compound (A) ( $C_7H_8$ ) which undergoes oxidation with alkaline  $KMnO_4$  to give an acidic compound (B). (B) reacts with  $SOCl_2$  to give (C) which gives (D) with ammonia. The compound (D) on reaction with bromine and alkali forms (E) which reacts with bromine water to give (F) ( $C_6H_4NBr_3$ ). (F) reacts with  $NaNO_2$  and  $HCl$  and forms (G) ( $C_6H_2N_2Br_3Cl$ ). Addition of  $KI$  forms (H). Identify the compounds (A) to (H).

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20. Two isomeric mononitro derivatives (B) and (C) are obtained by the nitration of an organic compound (A),  $C_7H_8O$ . Treatment of (A) with acetyl chloride produces (D) which on reaction with  $CrO_2Cl_2$  gives (E) whose oxidation with neutral  $KMnO_4$  followed by acidification gives (F). Compound (F) on heating gives phenol. (A) on treatment with alkaline  $C_6H_5SO_2Cl$  produces (G) which on oxidation with  $KMnO_4$  gives (H). Hydrolysis of (H) also gives (F). Give the structures of (A) to (H).

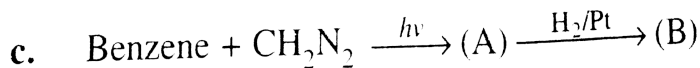
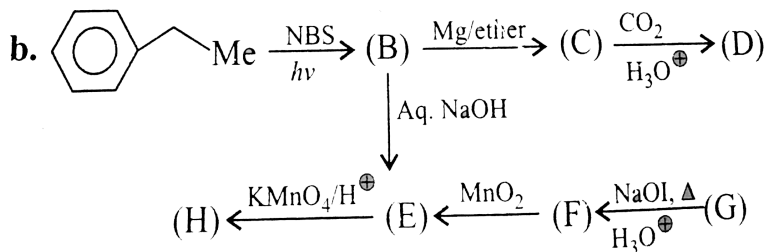
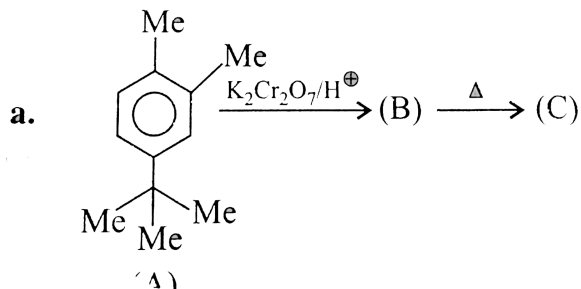
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21. An organic compound (A) ( $C_9H_{12}$ ) gave (B) ( $C_8H_6O_4$ ) on oxidation by alkaline  $KMnO_4$ . (B) on heating does not form anhydride. Also, (B) reacts with  $Br_2$  in the presence of iron to give only one monobromo-derivative (C) ( $C_8H_5BrO_4$ ). What are (A), (B), and (C) ?

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## Solved Problems

1. Complete the following reactions:



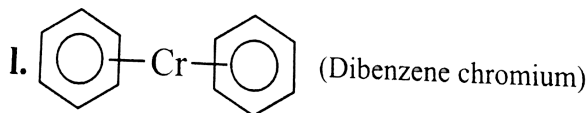
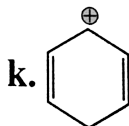
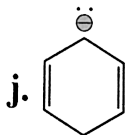
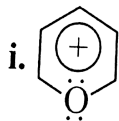
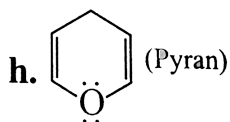
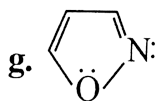
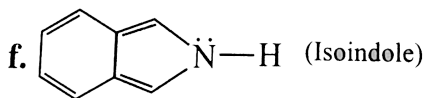
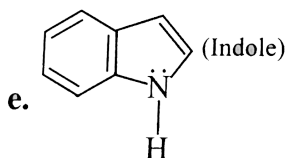
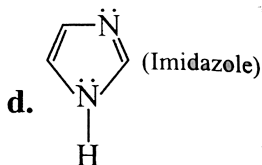
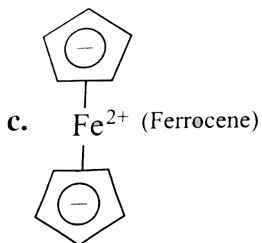
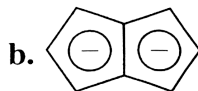
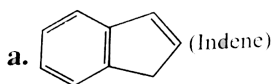
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2. Cycloocta -1, 3, 5, 7 - tetraene (*A*), a non-aromatic compound reacts with 2 mol potassium to form a stable compound (*B*) without producing  $H_2$ . Write the structure of (*B*) and explain the reason for its stability.



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3. Select the aromatic, anti-aromatic, and non-aromatic compounds.

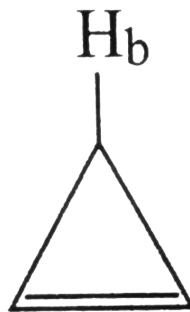


4. Answer the following :

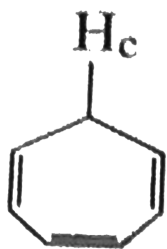
a. Which of the following  $H$  will have lower  $pK_a$  value in the given compound?



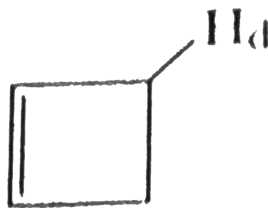
(I)



(II)

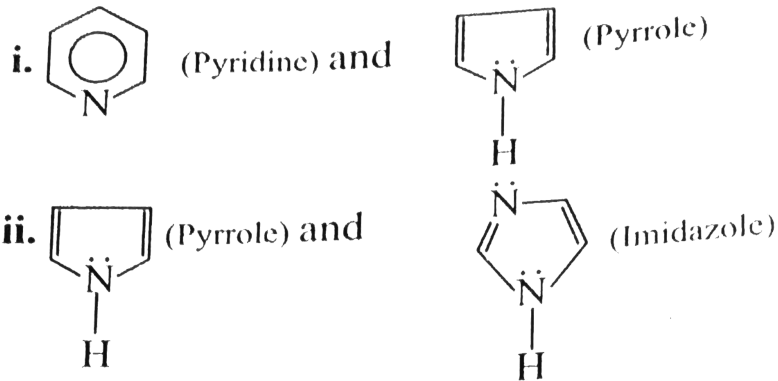


(III)

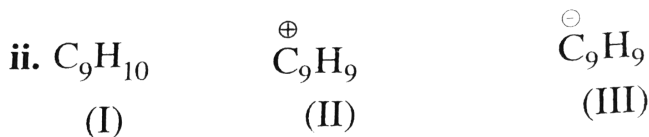
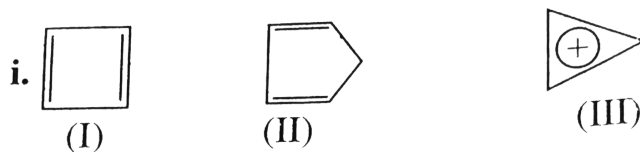


(IV)

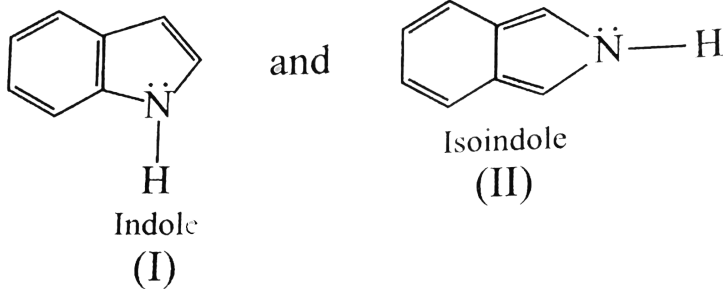
b. Which of the following have lower  $pK_b$  value?



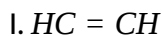
c. Give the decreasing order of the stability of the following:



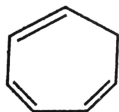
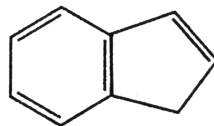
d. Which of the following is more stable?



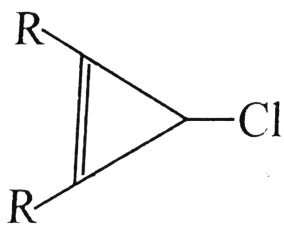
e. What of the decreasing order of  $pK_a$  values of the following?





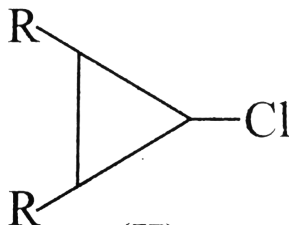
**II.****III.**

f. Which of the following has highest  $K_a$  value?

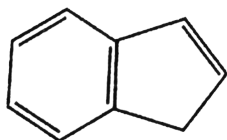
**i.**

(I)

or

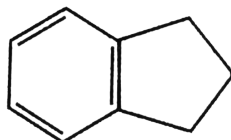


(II)

**ii.**

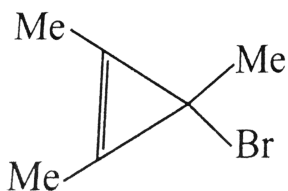
(I)

or



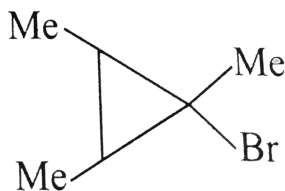
(II)

g. Which of the following reacts faster with aq.  $\text{NaOH}$  by  $\text{SN}^{-1}$  mechanism,?



(I)

or



(II)


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5. Distinguish chemically between the following:

i. (a) Cyclohexane, (b) Cyclohexane, (c) Benzene

ii. (a) Benzene, (b) Naphthalene

iii. (a) Anthracene and (b) Phenanthrene.

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6. Explain the following observations:

a. Benzene is soluble in water while both 1,3- and 1,4-cyclohexadiene are insoluble.

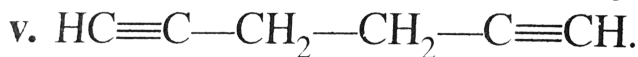
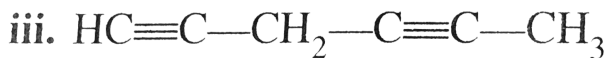
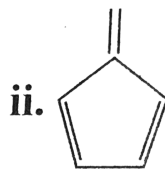
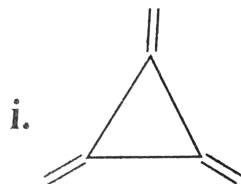
b. Melting point of benzene (32K) is much higher than both 1,3- and 1,4-cyclohexadienes.

c. *p*-Xylene has higher melting point than *o*- and *m*-xylenes.

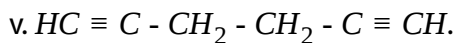
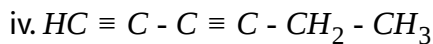
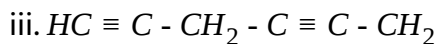
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7. Some alternate structures proposed for benzene are given. Indicate for each of them how many (i) mono and (ii) di-substituted products are

possible. Which structures fits in the isomer number observed for

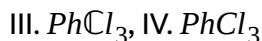
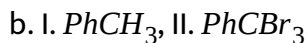
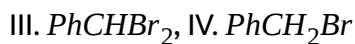
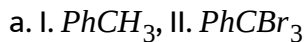


benzen?



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8. Arrange the following in decreasing order of  $\left(\frac{m}{o+p}\right)$  ratio.



c. I.  $ArN^{\oplus}R_3$ , II.  $ArCH_2N^{\oplus}R_3$

III.  $ArCH_2CH_2N^{\oplus}R_3$

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9. Give the principal product on bromination with  $Br_2/FeBr_3$  of the following.

I.  $Ph - OCOR$ , II.  $Ph - CH = CH - NO_2$

III.  $PhCOONH_2$ , IV.  $Ph - O - Ph$

V.  $PhCH_2OH$ , VI.  $PhCoEt$

VII.  $Ph - CH = CH - COOH$

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10. Give the decreasing order of  $\left(\frac{o+p}{m}\right)$  ratio for the nitration of compounds in Problem 11.9.

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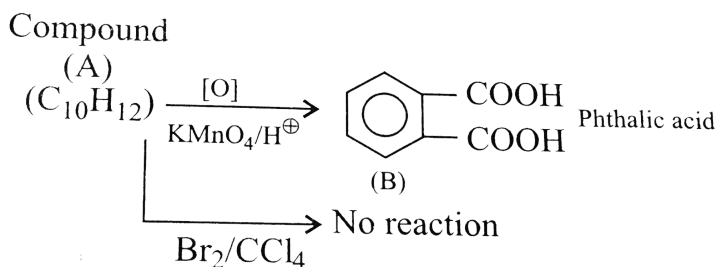
11. Give major products of the following:



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12. Give the decreasing order of  $\frac{\oplus p}{m}$  ratio of the reactions or decreasing order of the reactivity given is problem 11.11.

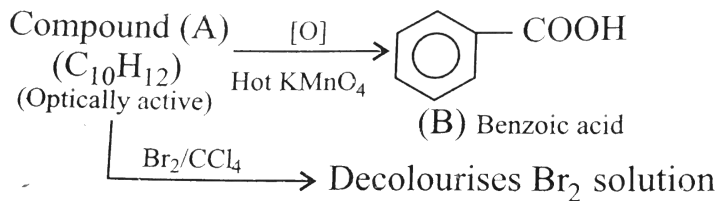
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Identify compound (A) How many isomers including geometrical isomers are possible for (A) ?

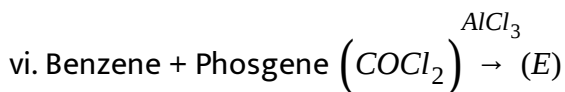
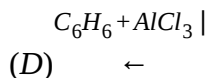
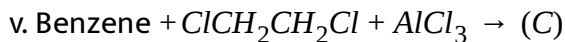
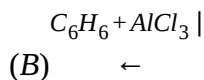
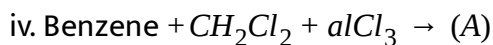
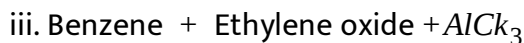
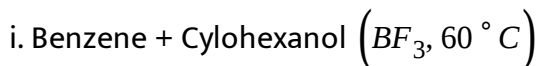
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14. Identify (A).



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15. Give the products of the following:



overset(H<sub>3</sub>O<sup>(o+)</sup>)(rarr)

(F)vii. Toe ≠ + Maloncyanhydride

overset(AlCl<sub>5</sub>)(rarr) (A)underset(H<sub>3</sub>PO<sub>4</sub>)(rarr) (B) underset(HCl)

overset(Zn-Hg)(rarr) (C)

viii. Expla ∈ howalkylbenze ≠ napha ≤ ≠ , anthrace ≠ and phenanthre ≠ canbe

F.C.` reaction ?

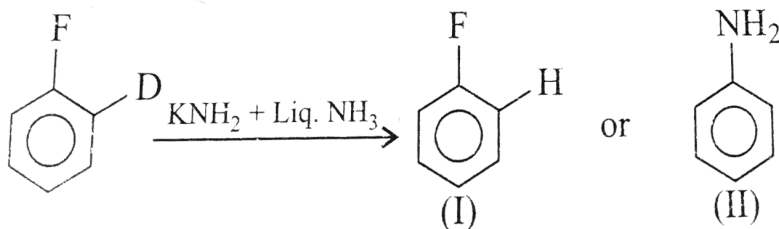
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16. a. Show that reaction of *o*- bromogluro benzene with *Mg* produces benzyene.

b. Why does *I*- cj,prp-2.6 - dimerhyl benzene not undergo elimination-addition reaction with *NaNH<sub>2</sub>* in liq. *HN<sub>3</sub>*?

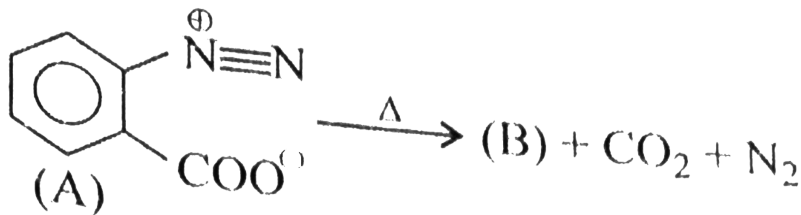
c. Why does bromobenzene react faster than 2.6 - dideuteriobromonbenzene with *NaNH<sub>2</sub>* in liw. *NH<sub>3</sub>*?

In the following reactions:



Explain why the formation of (I) is faster than that of (II).

c. Why does compound (A) on heating give benzene intermediate which dimerizes to give product (B).



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17. When three isomeric chlorotouenes are treated with  $\text{KNH}_2$  in liquid  $\text{NH}_3$ , they yield different toluidines as shown below. Explain these observations. It rgt a. *o*-Chlorotouene  $\rightarrow$  (*o*:*m*) - Toluidine

b. *m*-Chlorotouene  $\rightarrow$  (*o* + *m* + *p*) - Toidine

c. *p*-Chlorotouene  $\rightarrow$  (*m* + *p*) - Toidine

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18. Give the order of reactivity towards  $SN^2$  reaction of the following:

a. i. 1 - Bromopentane

ii. 2 - Bromopentane

iii. 2 - Bromo -2 methyl butane

b. i. *n* - Butyl bromide ( $C_4H_9Br$ )

ii. Isobutyl bromide  $Me_2CHCH_2Br$

iii. sec - Butyl bromide  $CH_3 - CH(Me)CH_2Br$

iv. tert- Butyl bromide  $Me_3C - Br$



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19. Give the order of reactivity towards  $SN^1$  reaction of the following:

a. i. 1 - Bromopentane

ii. 2 - Bromopentane

iii. 2 - Bromo -2 methyl butane



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20. Give the order of  $SN^{-1}$  of  $SN^{-1}$  and  $SN^2$  displacement of halogen  $CH_3X$ ,  $1^\circ X$ ,  $2^\circ X$ ,  $3^\circ X$ .

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21. Give the order of reactivity towards  $E2$  dehydrohalo-genation of the following.

a. i.  $C_2H_5Br$

ii.  $C_3H_7Br$

iii. Isobutyl bromide

iv. Neopentyl bromide

b. i. 2 - Bromo -2 - methybutane

ii. 1 - Bromo pentane

iii. 2 - Bromo pentane

iv. 3 - Bromo pentane

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22. Give the order of reactivity towards  $SN^{-1}$  solvolysis of the following:

i. Benzyl chloride

ii. *p* - Chlorobenzyl chloride

iii. *p* - Methoxybenzyl chloride iv. *p* - Methyl benzyl chloride

v. *p* - Nitrobenzyl chloride



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23. Give the order of reactivity towards  $SN^1$  reaction of the following:

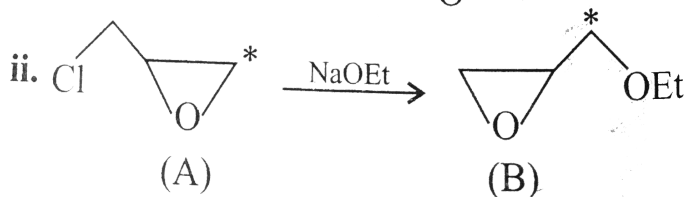
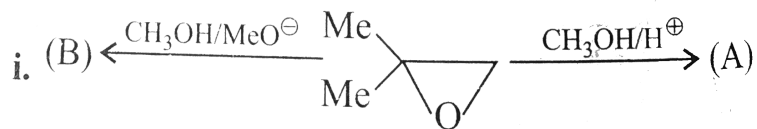
i.  $ClCH_2CH = CH_2$

ii.  $CH_3CH_2CH_2Cl$

iii.  $CH_3CH = CHCl$

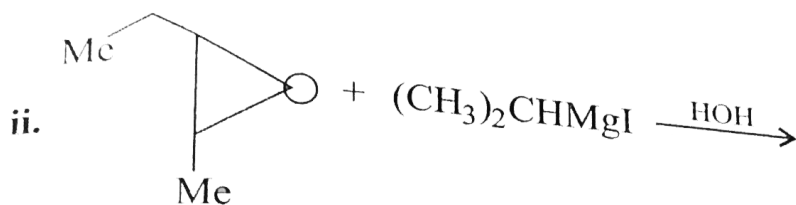
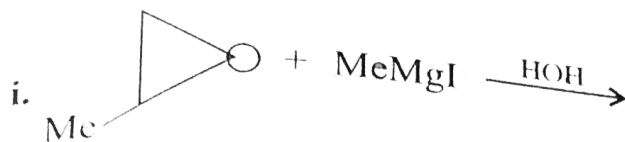


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24.

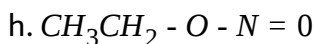
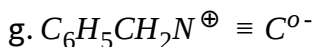
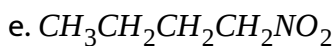
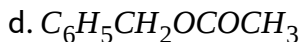
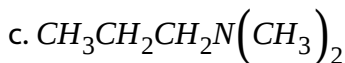
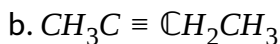
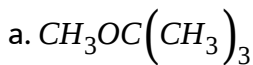
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25.

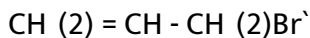
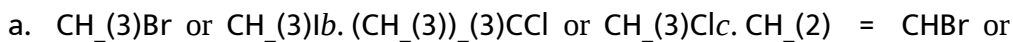
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26. Outline the preparation of the following compounds using a nucleophilic substitution reaction.



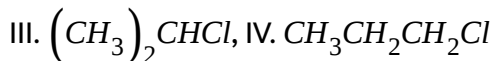
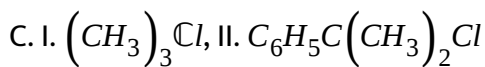
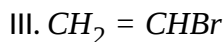
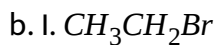
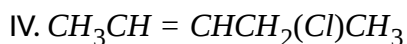
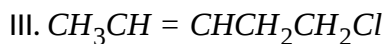
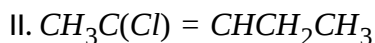
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27. Which compound in each of the following pairs will react faster in  $\text{S}_\text{N}2$  reaction with  $\text{HO}^\ominus$  ?



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28. Arrange the following compounds in increasing order of  $S_N^2$  reactivity.



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29. Predict all the alkenes that would be formed by dehydrohalogenation of the following halides with sodium ethoxide in ethanol and identify

major alkene.

a. 1 - Bromo - 1 - methylcyclohexane

b. Cyclohexylmethyl bromide

c. 2 - Chloro - 2 - butane

d. 3 - Bromopentane

e. 2, 2, 3 - Trimethyl 3 - bromopentane.



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30. Predict all order of reactivity of the following compounds in dehydrohalogenation.

a. i.  $CH_3CH_2CH_2CH_2Cl$

ii.  $(CH_3)_2CHCH_2Cl$

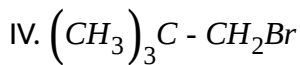
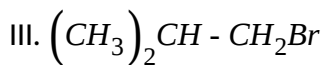
iii.  $(CH_3)_2CH - CH_2Br$

iv.  $CH_3CH(Br)CH_2CH_3$

v.  $(CH_3)_3C - Br$

b. i.  $CH_3CH(Br)CH_3$

ii.  $CH_3CH_2CH_2Br$



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31. Explain :

- Vinyl chloride is unreactive in nucleophilic substitution reactions.
- Neopentyl bromide undergoes nucleophilic substitution reactions very slowly.
- 3 - Bromocyclohexane is more reactive than 4 - bromocyclohexane in hydrolysis with aqueous  $\text{NaOH}$ .
- ter* - Butyl chloride reacts with aqueous sodium hydroxide by  $\text{SN}^1$  mechanism while *n* - butyl chloride reacts by  $\text{SN}^2$  mechanism.

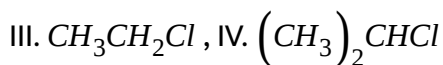
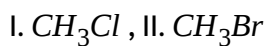
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32. The nucleophilic substitution of primary alkyl chlorides with sodium acetate is catalysed by sodium iodide. Explain why.

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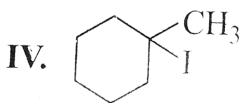
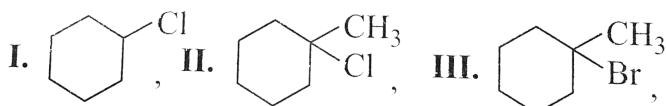


33. Arrange the following halides in order of increasing  $SN^2$  and  $SN^1$  reactions.



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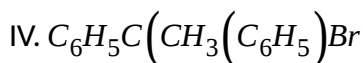
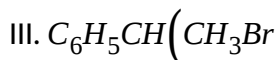
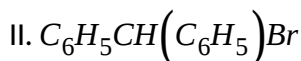
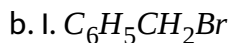
34. Predict the order of reactivity of the following compounds in  $SN^1$  and  $SN^2$  reactions.



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35. Predict the order of reactivity of the following compounds in  $SN^1$  and  $SN^2$  reactions.

a. The four isomeric bromobutanes



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36. Identify all the possible alkenes that would be formed on the dehydrohalogenation of the following organic halides with alcoholic  $KOH$

. Also, identify the major alkene.

a. 1-Chloropentane

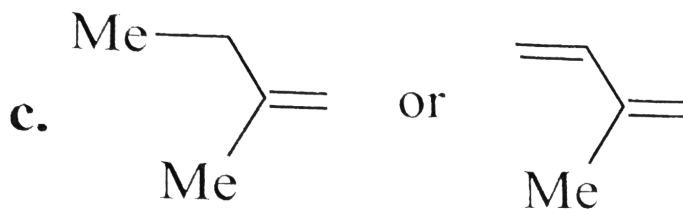
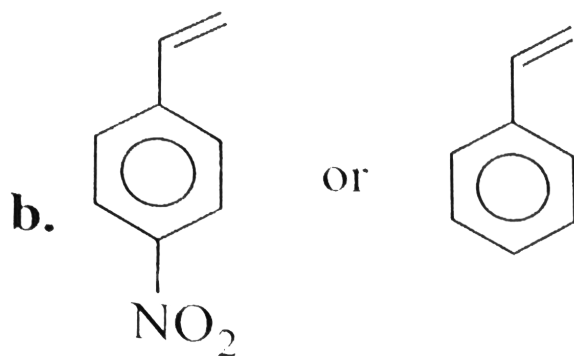
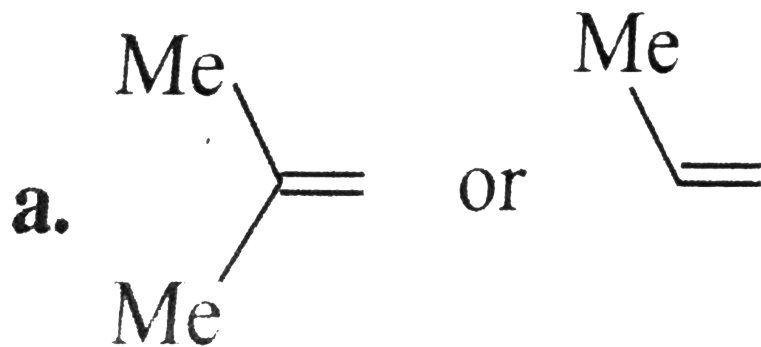
b. 2-Chloropentane

c. 2-chloro-2-methyl butane

d. 2-Chloro-3-methyl butane

e. 3-Chloro-2-methyl pentane

37. I. Which compound has the faster rate of reaction with  $HCl$  ?



II. Two compounds ( $C_8H_{15}Br$ ) are formed when 1.6-dimethylcyclohexene reacts with  $HBr$  in dark and are formed from 1.2-dimethylcyclohexene. What are these two compounds?

III. A hydrocarbon ( $M_w = 70$ ) gave only one monochloro-derivative on photochemical chlorination. Name both the compound.

IV. Among the isomeric alkanes of  $MF(C_5H_{12})$  identify the one that on photochemical chlorination yields:

- A single monochloride
- Three isomeric derivatives
- Four isomeric derivatives

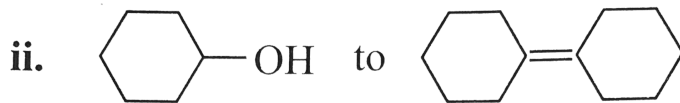


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## Exercises Subjective Type

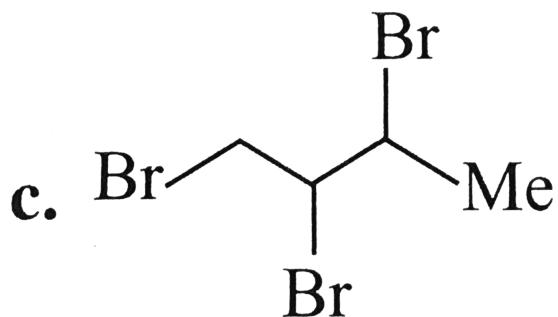
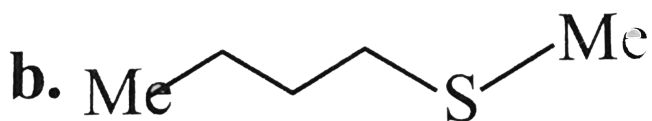
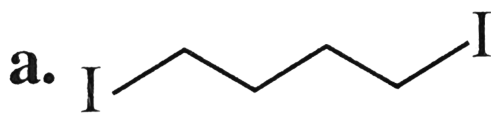
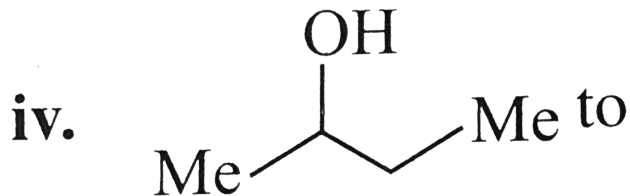
1. Synthesise the following:

- Propane to a. 1-fluoropropane, b. cyclopropane, and c. 2,3-dimethylbutane



ii.

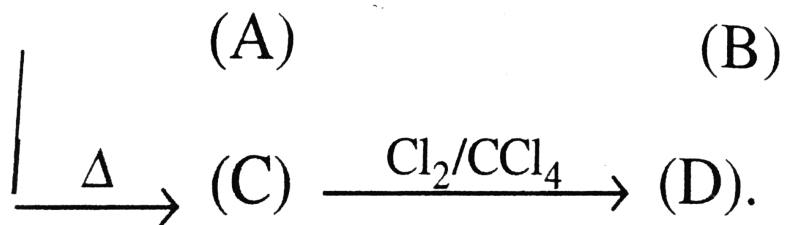
iii.  $C_2H_5Br$  to but - 1 - ene



iv.

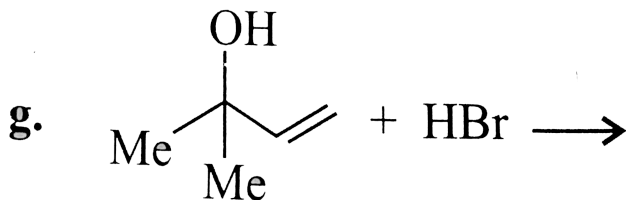
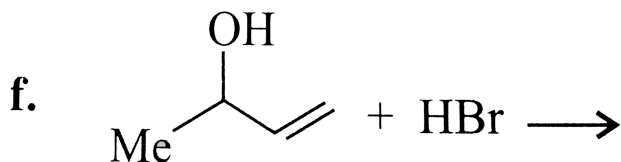
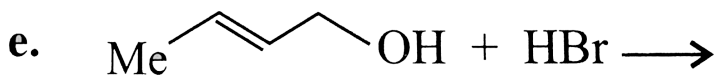
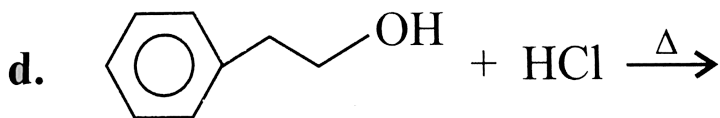
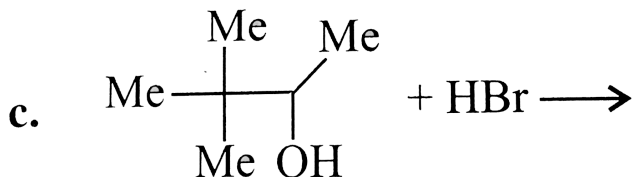
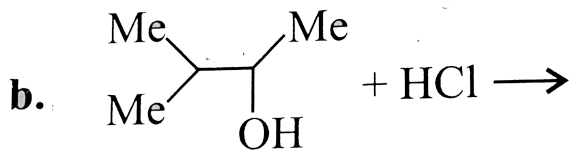
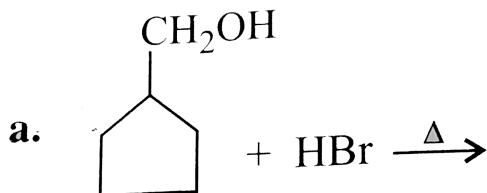
v. Cyclopentane to hexachloropentadiene

(vi) Cyclopentadiene + Hexachlorocyclopentadiene



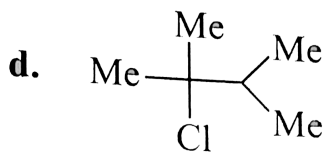
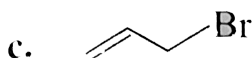
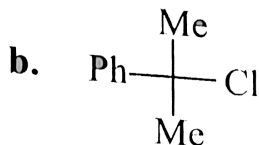
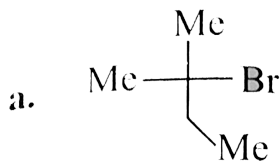
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2. (i) Identify the major and minor products.



1-Butanol + HBr → 1-Bromobutane  
2-Butanol + HBr → 2-Bromobutane

ii. Suggest the reagent that could be used to prepare these alkyl halides from alcohols.



iii. Identify the products in the following reactions:

(A) iv. Identify

(B), (C) and (D) in the following:

(B)

v. Identify the products in the following reaction:

(C)

vi. Identify the products when (A) reacts with:

(D)

a.  $Br_2/Fe$ , b.  $Br_2/Cl_4$

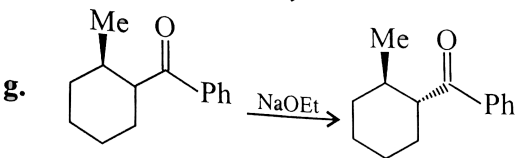
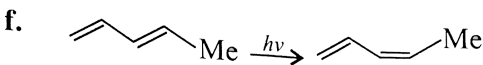
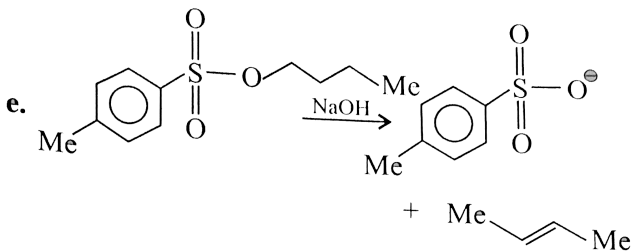
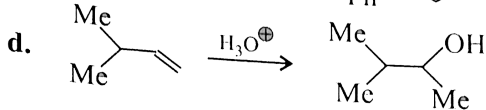
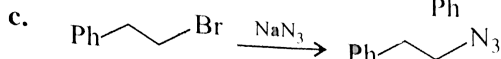
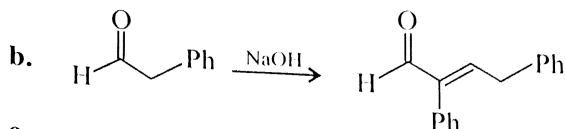
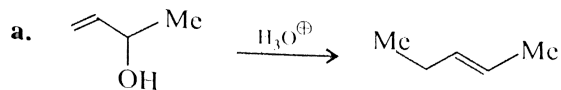
c. NBS, d. HBr



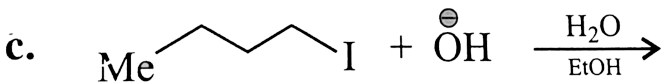
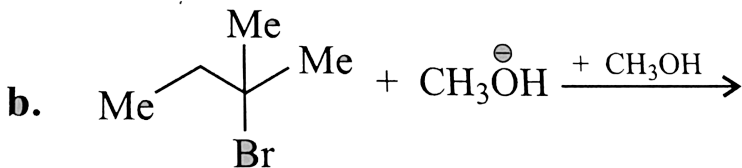
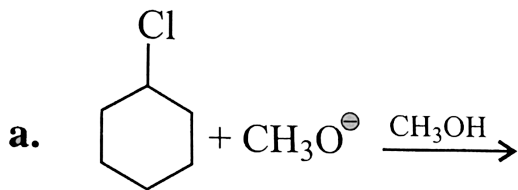
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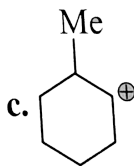
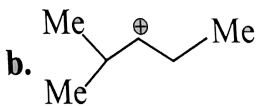
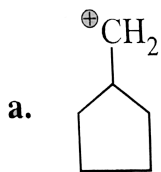
3. i. Classify the following reactions as an addition elimination substitution condensation rearrangement geometrical isomerisation, or oxidation/reduction.



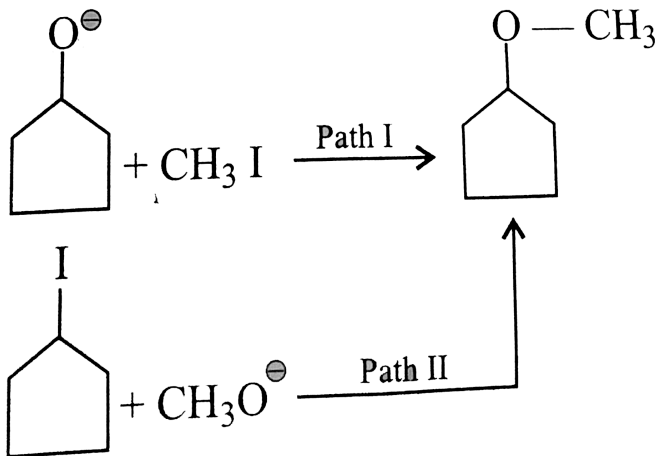
ii. Show both substitution and elimination products in these reactions.



iii. Show the rearranged carbocations that are expected from these carbocations.



iv. Which path is better for the reaction ?



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4. i. sec-Butyl alcohol will undergo alkaline hydrolysis in the polar solvent

by

a.  $\text{SN}^2$  , b.  $\text{SN}^1$  c.  $\text{SN}^1$  and  $\text{SN}^2$

, d. No  $\neq$  ii. Which of the following statements is correct? (A)  $\text{C}_2\text{H}_5\text{Br}$  reacts with  $\text{OH}^-$  in  $\text{C}_2\text{H}_5\text{OH}$  to form  $\text{C}_2\text{H}_5\text{O}^-$  and  $\text{H}_2\text{O}$ .

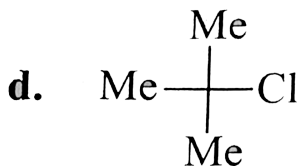
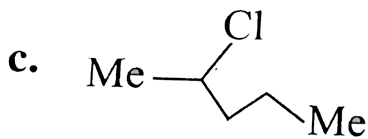
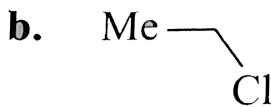
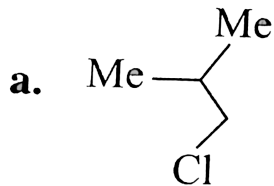
(B)  $\text{C}_2\text{H}_5\text{Br}$  reacts with  $\text{OH}^-$  in  $\text{C}_2\text{H}_5\text{OH}$  to form  $\text{C}_2\text{H}_5\text{O}^-$  and  $\text{H}_2\text{O}$ .

(C)  $\text{C}_2\text{H}_5\text{Br}$  reacts with  $\text{OH}^-$  in  $\text{C}_2\text{H}_5\text{OH}$  to form  $\text{C}_2\text{H}_5\text{O}^-$  and  $\text{H}_2\text{O}$ .

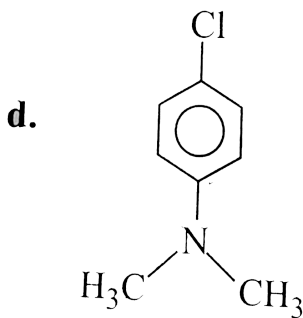
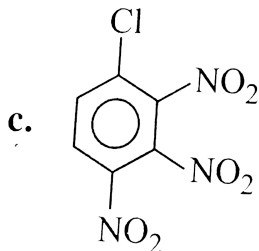
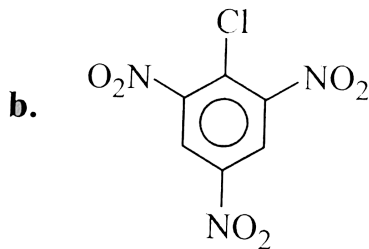
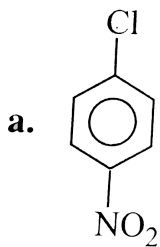
(D)  $\text{C}_2\text{H}_5\text{Br}$  reacts with  $\text{OH}^-$  in  $\text{C}_2\text{H}_5\text{OH}$  to form  $\text{C}_2\text{H}_5\text{O}^-$  and  $\text{H}_2\text{O}$ .

. d. (A) is isomer of dimethyl ether (B) is dehydrated compound of (A)

. iii. Which of the following will undergo  $S_N2$  readily?

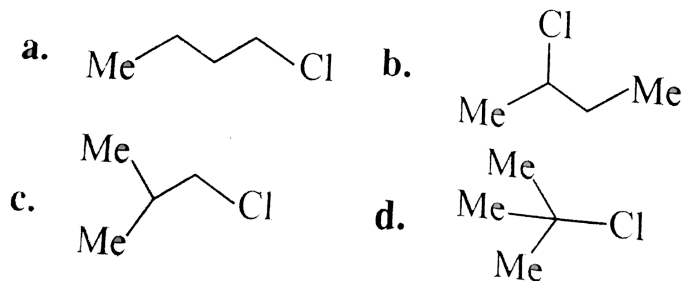


iv. Which of the following will undergo hydrolysis most readily with aq. KOH?



v. In two experiments equal quantities of an alkyl halide,  $(C_4H_9Cl)$  were

treated at the same temperature with equal volumes of 0.1M and 0.2M solution of  $\text{NaOH}$ , respectively. In the two experiments the time taken for the reaction of exactly 50% of the alkyl halide was the same. The most likely structures of the halides is:



vi. Which of the following methods is not suitable for the preparation of  $\text{RX}$ ?

- Halogenation of alkane
- $\text{ROH}$  and  $\text{PX}_3$
- $\text{ROH}$  and  $\text{HX}$
- Alkene and  $\text{HX}$

vii. The heterolytic bond dissociation energy of the  $(\text{C} - \text{Cl})$  bond in vinyl chloride is  $207 \text{ kcal mol}^{-1}$  as compared to  $191 \text{ kcal mol}^{-1}$  in the case of  $(\text{C} - \text{Cl})$  bond in  $\text{C}_2\text{H}_5\text{Cl}$ . One explanation for this observation is:

a. In  $\text{C}_2\text{H}_5\text{Cl}$  the bond energy is decreased through resonance involving lone

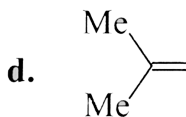
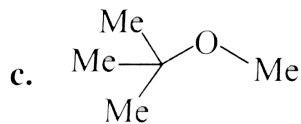
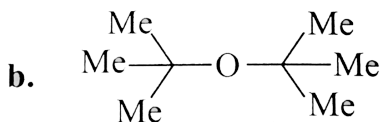
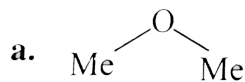
pair of  $\bar{e}$ 's on chlorine.

b. The hyperconjugation stabilisation in  $C_2H_5Cl$ .

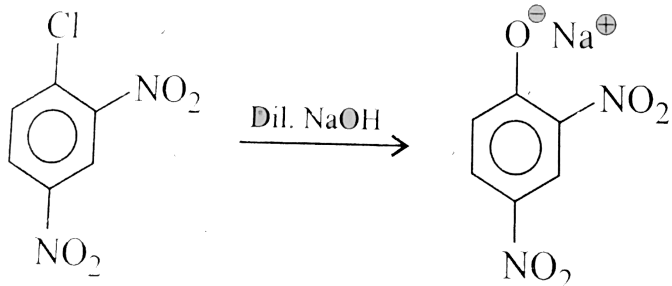
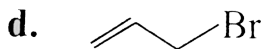
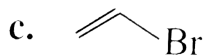
c. In vinyl chloride, ( $C - Cl$ ) bond gets partial double bond character through resonance.

d. In vinyl chloride the  $Cl$  atom is attached to  $sp$  - hybridised  $C$ , whereas in  $C_2H_5Cl$  it is attached to  $sp^3$ -hybridised  $C$  atom.

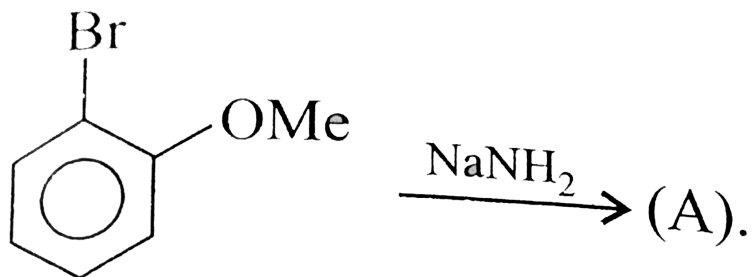
viii. When  $(Me_3C - Cl)$  is reacted with  $NaOMe$  the major product is



ix. Which of the following would be hydrolysed most readily?

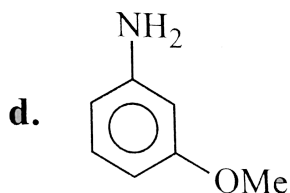
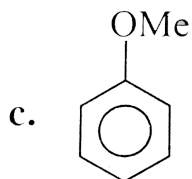
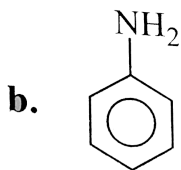
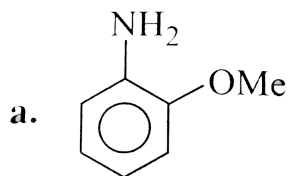


x.



The above reaction proceeds through:

- a. Electrophilic addition
- b. Benzyne intermediate
- c. Activated nucleophilic substitution
- d. Oxirane



xi.

## Exercises Subjective

1. Give the chemical test to distinguish between

(a) Hexane (I),  $MeCH = CHCl$ (II),

Propylchloride (III), and  $CH_2 = CHCH_2Cl$ (IV)

(b) (V) $CH_3Cl$ , (VI)  $CH_3Br$ , and (VII)  $CH_3I$



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2. Give the product of  $E_2$  reaction of the following compounds with  $atc.$

$KOH$ .

a. *Ertho*-and *therno* form of *I*- bromo 1,2 - diphenyl propane

b. *Eryhgro*- and *thero*-from of 2,2,3,5,5 - penta methyl 4 - bromo hexane



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3. Draw the diastereomers of 2-chloro-1,3-dimethylcyclohexane and indicate the diastereomers that cannot undergo  $E2$  elimination and give the product from these that react.

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4. Draw the diastereomers of 2-chloro-1,3-dimethylcyclohexane and indicate the diastereomers that cannot undergo  $E2$  elimination and give the product from these that react.

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5. (W) and (X) are optically active isomers of  $C_5H_9Cl$  (W) on treatment with 1 mol of  $H_2$  is converted to an optically inactive compound (Y), but (X) gives an optically active compound (Z) under the same conditions. Give the structure of (Y) and the configurations of (W), (X) and (Z) in Fischer projections.

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6. An organic compound (A),  $C_8H_4O_3$  in dry benzene in the presence of anhydrous  $AlCl_3$  gives compound (B). Compound (B) on treatment with  $PCl_5$  followed by reaction with  $H_2/Pd(BaSO_4)$  gives compound (C) which on reaction with hydrazine gives a cyclised compound (D),  $(C_{14}H_{10}N_2)$ . Identify (A), (B), (C) and (D). Explain the formation of (D) from (C).

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7. Three isomeric hydrocarbons  $C_9H_{12}$   $\begin{bmatrix} A \\ B \\ C \end{bmatrix}$  oxidise to  $C_9H_6O_6$   $\begin{bmatrix} D \\ E \\ F \end{bmatrix}$  while (D) may give three different monoethyl esters, (E) may form only two and (F) only one. Identify compounds (A) to (F).

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8. An organic compound (A) contains  $C = 92.3\%$  and  $H = 7.7\%$ . Its vapour density has been found to be 52. (A) is oxidised by  $KMnO_4$  solution in cold and when fully oxidised gives an acid (B) which contains  $C = 68.8\%$  and  $H = 4.9\%$ . (A) reacts with bromine and yields a compound (C) containing  $60.8\%$  bromine. What is the structure of (A)?

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9. An aromatic hydrocarbon (A) containing  $C = 91.3\%$  and  $H = 8.7\%$  on treatment with chlorine gave three isomeric monochloro compounds (X), (Y) and (Z), each having  $28\%$  chlorine. On oxidation with  $KMnO_4$  all the three gave monobasic acids. The acid from (X) on distillation with soda lime gave benzene while those from (Y) and (Z) gave monochlorobenzene. What formula would you assign to the various compounds?

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10. Compound  $C_8H_9Cl(A)$  on treatment with  $KICN$  followed by hydrolysis gives  $C_9H_{10}O_2(B)$ . Ammonium salt of  $(B)$  on dry distillation yields  $(C)$ , which reacts with alkene solution of bromine to give  $C_8H_{14}N(D)$ . Another compound  $C_8H_{10}O(E)$  is obtained by the action of nitrous acid of  $(D)$  or by the action of aqueous potassium on  $(A)$ ,  $(E)$  on oxidation gives the inner anhydride  $C_8H_6O_4(F)$  which gives the inner anhydride  $C_8H_4O_3(G)$  on heating. Identify the various compounds  $(A)$  to  $(G)$ .

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11. On disulfonation followed by fusion with  $NaOH$  an acidificant, an aromatic compound  $C_8H_{10}(A)$  gives two isomeric compounds  $(B)$  and  $(C)$  (molecular formula  $C_8H_{10}O_2$ ). Oxidation of esters of  $(B)$  and  $(C)$  followed by hydrolysis gives two isomeric compounds  $(D)$  and  $(E)$  ( $C_7H_6O_4$ ) which may be decarboxylated to form  $C_6H_6O_2(F)$ . Identify compounds  $(A)$  to  $(F)$ .

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12. The compound (A) is xylene. On sulphonation, it gives only one product (B) which on alkaline fusion followed by acidification yields (C), a phenol. On treatment with chloroform and alkali, (C) gives  $C_9H_1O_2$  (D). Identify the various organic compounds (A) to (D).

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13. An organic compound  $C_6H_4O_2NCl$  (A) reacts with  $NaOH$  to give  $C_6H_5O_3N$  (B). On reduction with tin and  $HCl$ , (B) gives  $C_6H_7ON$  (C) whereas on milder reduction it gives (D)  $C_6H_7O_2N$ , which rearranges in the presence of a mineral acid to give a new compound (E). Identify the various compounds from (A), to (E).

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14. A hydrocarbon  $C_9H_{10}$  (A) rapidly decolourises cold aqueous permanganate and rapidly absorbs one molecule of bromine. Vigorous

hydrogenation of (A) gives a compound (A) gives a compound (B) of the formula  $C_9H_{18}$ . Strong oxidation of (A) yields phthalic acid. Propose a structure for (A) and give the basis for your answer.

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15. On nuclear chlorination,  $C_8H_{10}$ (A) gives a product (B) which may be oxidised to  $C_8H_5O_4Cl$ (C). (A) may also be chlorinated to give  $C_8H_8Cl_2$ (D),  $C_8H_6Cl_4$ (E), and  $C_8H_4Cl_6$ (F). (D), (E) and (F) on hydrolysis give halogen-free compounds (G), (H), and (I), respectively. With  $CrCO_2Cl_2$ , (A) gives  $C_8H_6O_4$  which is identical with (I). Identify with (I). Identify the various compounds (A) to (J).

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16. When bromobenzene is monochlorinated. Two isomeric compounds (A) and (B) are obtained. Monochlorination of (A) gives several isomeric products of molecular formula  $C_6H_3ClBr_2$ . While monochlorination of (B) yields only two isomers (C) and (D). Compound (C) is identical to

one of the compounds obtained from the bromination of (A). However, (D) is totally different from any of the isomeric compounds obtained from the bromination of (A). Give the structures of (A), (B), (C) and (D), and also the structures of the four isomeric monobrominated products of (A). Support your answer with reasoning.

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17. 0.450 gm of an aromatic organic compound (A) on ignition gives 0.905 gm  $CO_2$  and 0.15 gm  $H_2O$ . 0.350 gm of (A) on hydrolysis with  $Ca(OH)_2$  yields (B) which on mild reduction gives an optically active compound (C). On heating (C) with  $I_2$  and  $NaOH$ , iodoform is produced along with (D). With  $HCl$ , (D) gives a solid, which is more soluble in hot water than in cold. Identify (A) to (D) with proper explanation.

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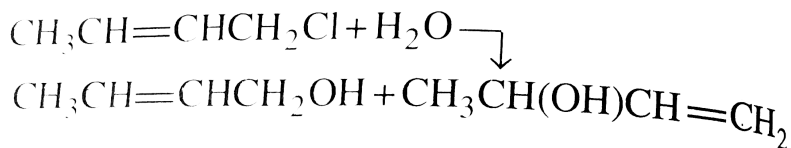
1. In each of the following reactions, a new ring is formed. Use curved arrow notation to explain the formation of the new ring.

a. 1, 4 - Dibromobutane ( $0.1\text{mol}$ ) is treated with sodium sulphide ( $0.1\text{mol}$ ) in aqueous ethanol.

b. 1 - Chloro - 2 (2 - hydroxyphenyl) ethane is treated with aqueous sodium hydroxide.

c. *o* - Methoxybenzyl bromide reacts faster than *p* - nitrobenzyl bromide with ethanol to form an ether product. Explain why.

d. Explain the formation of the two products in the following reactions:



e. What mass of propene obtained from  $34.0\text{gm}$  of *I* - iodopropane on heating with ethanolic *KOH*, if the yield is  $36\%$  ?

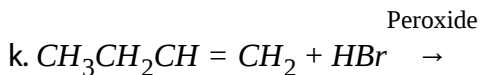
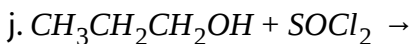
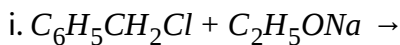
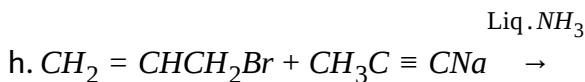
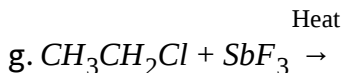
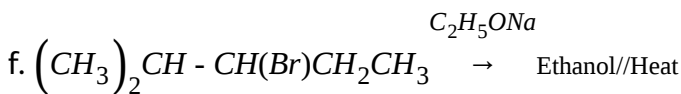
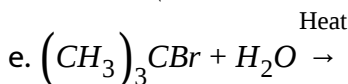
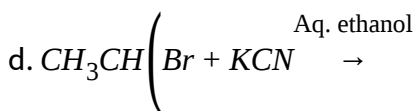
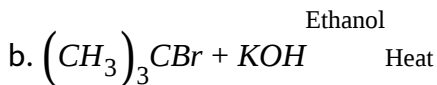
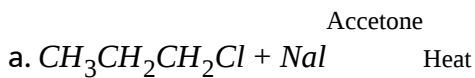
f. The nucleophilic substitution of primary alkyl chlorides with sodium acetate is catalysed by sodium iodide. Explain why.

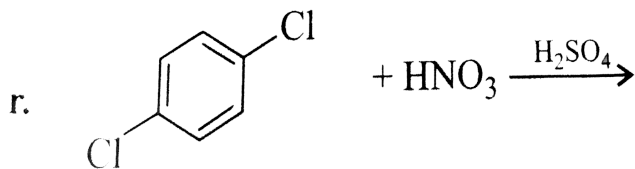
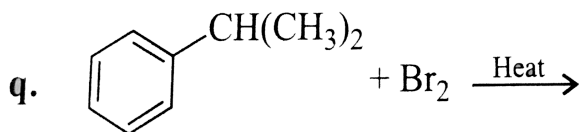
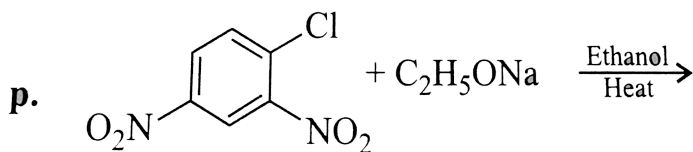
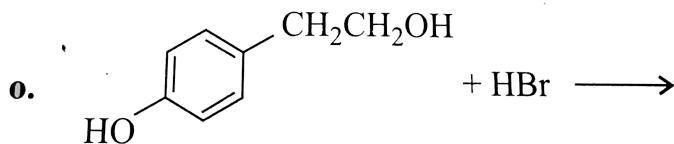
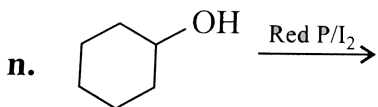
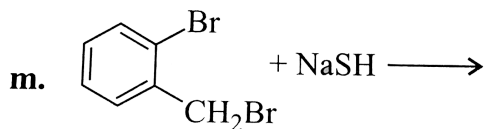
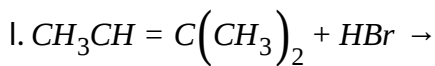


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2. Write the structure of the major organic product in each of the following reactions:





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1. How will you bring about the following conversations in not more than two steps?

a. Ethanol to but -1 yne

b. Ethene to bromoethene

c. Propene to 1 - Nitropropane

d. Toluene to benzyl alcohol

e. Propane to propyne

f. Ethanol to ethyl fluoride

g. Bromomethane to propanone

h. But -1 - e  $\neq$  to but -2 - e  $\neq$

i. 1 - Chlorobutane to *n* - octane

j. Bromoethane to cis-hex -3 - ene

k. Benzyl alcohol to phenylethanenitrile

l. Benzene to biphenyl

m. Cyclopentene to cyclopentadiene -1, 3 - diene

n. Aniline to phenyl isocyanide



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2. a. A sample of 0.50g of an organic compound was treated according to Kjeldahl's method. The ammonia evolved was absorbed in 50ml of 0.5M  $H_2SO_4$ . The residual acid required 60mL of 0.5M solution of  $NaOH$  for neutralisation. Find the percentage composition of nitrogen in the compound.

b. On analysis, 0.2g of a monobasic acid gave 0.505gm of  $CO_2$  and 0.0864gm  $H_2O$ . 0.305gm of this acid required 25ml of  $M/10NaOH$  solution for neutralisation. Find the molecular formula of the acid.

c. A liquid aromatic organic compound (A) containing carbon (92.3%) and hydrogen (7.7%) decolourised  $KMnO_4$  and on ozonolysis gave methanal and another compound (B). The molecular mass of (A) is 104. On treatment with a suitable catalyst, (A) gave a high molecular mass solid product (C) having the same empirical formula as that of compound (A). Compound (C) is used in making toys and household goods. Identify (A), (B), and (C) and explain the reactions.

d. A sample of 0.246gm of an organic compound gave 0.198gm of  $CO_2$  and 0.104gm of  $H_2O$  on complete combustion. 0.37gm of the compound gave

0.638gm of silver bromide in Carius method. What is the molecular formula of the compound if its molecular mass is 109.

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**3. Convert the following:**

a, *t* - Butyl chloride to isobutyl bromide

b. Chloroethane to butane

c. 2 - Chloropropane to 1 - bromopropane

d. Bromo benzene to *p* - nitrophenol

e. Isopropyl alcohol to iodofrom

f. 2 - Chloropropane to 1 - Propanal

g. Butene to *n* - butyl iodide

h. Ethyl bromide to propanoic acid.

i. 2 - Methyl prop - 1 ene to 2 - Chloro - 2 - methyl propane

j. 2 - Bromobutane to 3, 4 - dimethyl hexane

k. Aniline to propane nitrile

m. Benzyl alcohol to phenyl ethanoic acid

n. Benzene to 4 - bromo nitro benzene

o. Toluene to benzyl alcohol

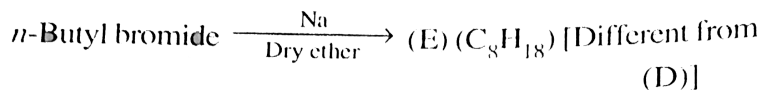
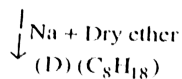
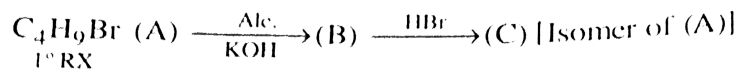
p. *I*-Bromopropane to 2-bromopropane

q. Ethanol to but-1-yne

r. Propane to propan-1-ol

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4. Identify (A) to (E):

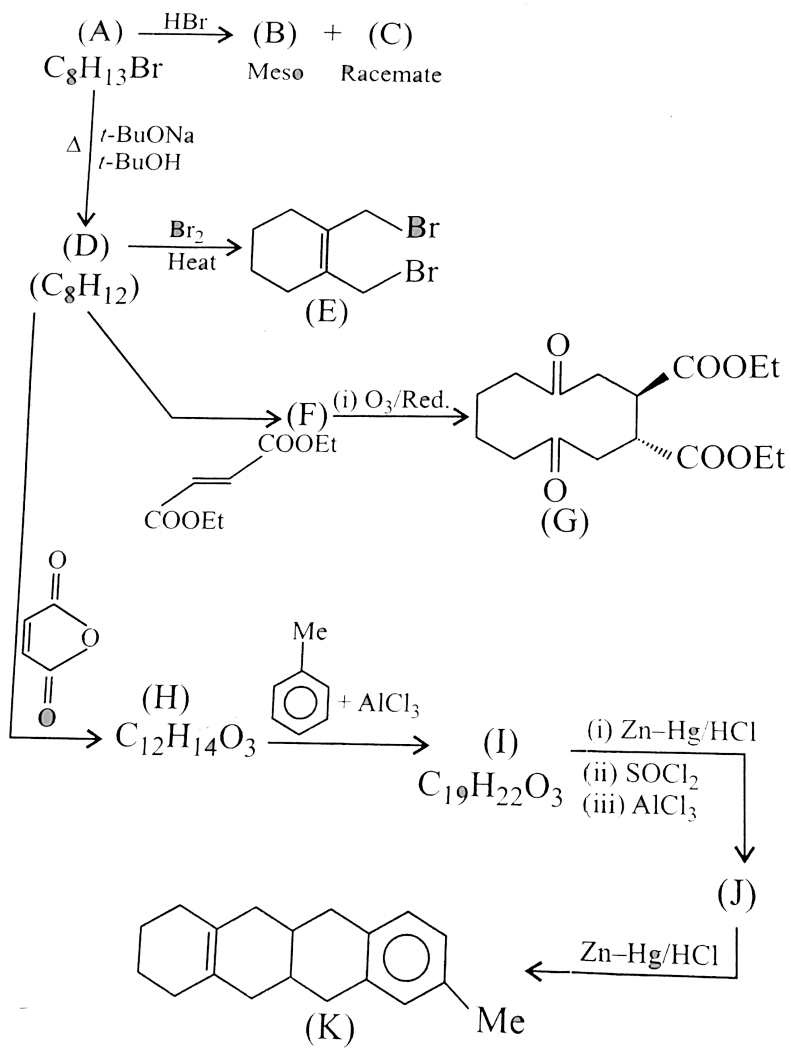


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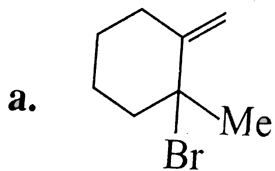
Exercises Linked Comprehension

1. This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

Paragraph for Problems 1 - 8

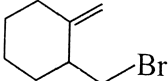
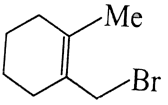
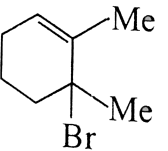


Compound (A) is:



A.



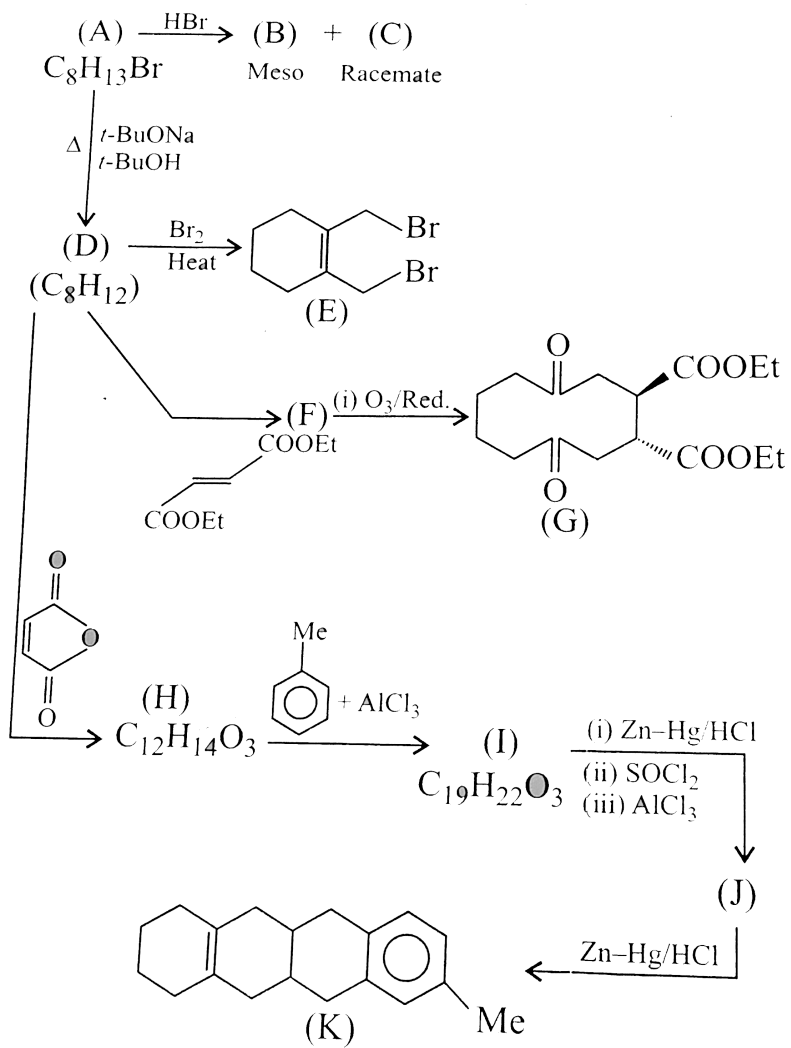
- b. 
- B.
- c. 
- C.
- d. 
- D.

**Answer: a**

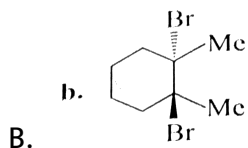
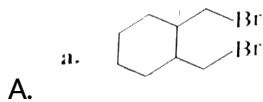
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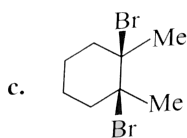
2. This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

Paragraph for Problems 1 - 8

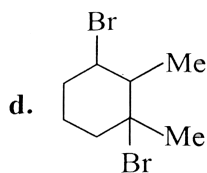


Compound (B) is:





C.



D.

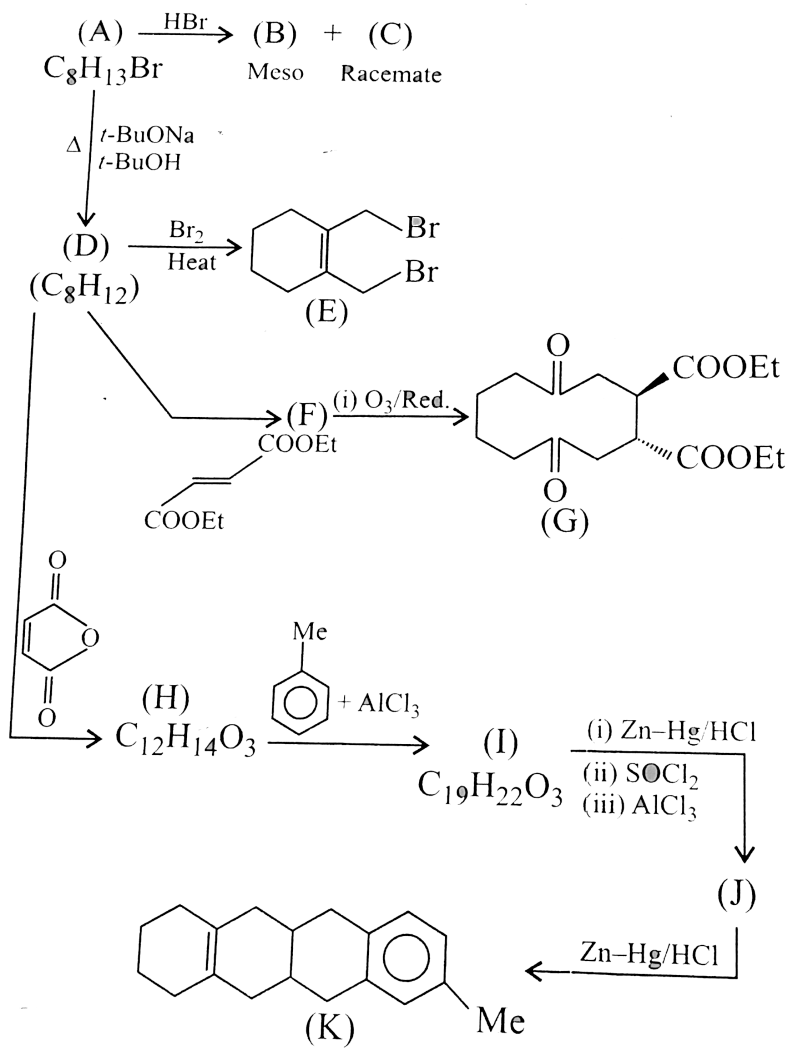
**Answer: c**



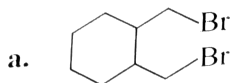
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**3.** This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

Paragraph for Problems 1 - 8



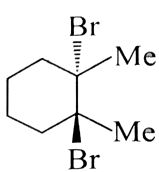
Compound (C) is:



A.

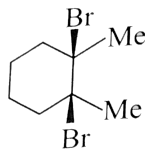
B.

b.



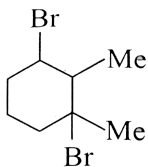
C.

c.



D.

d.



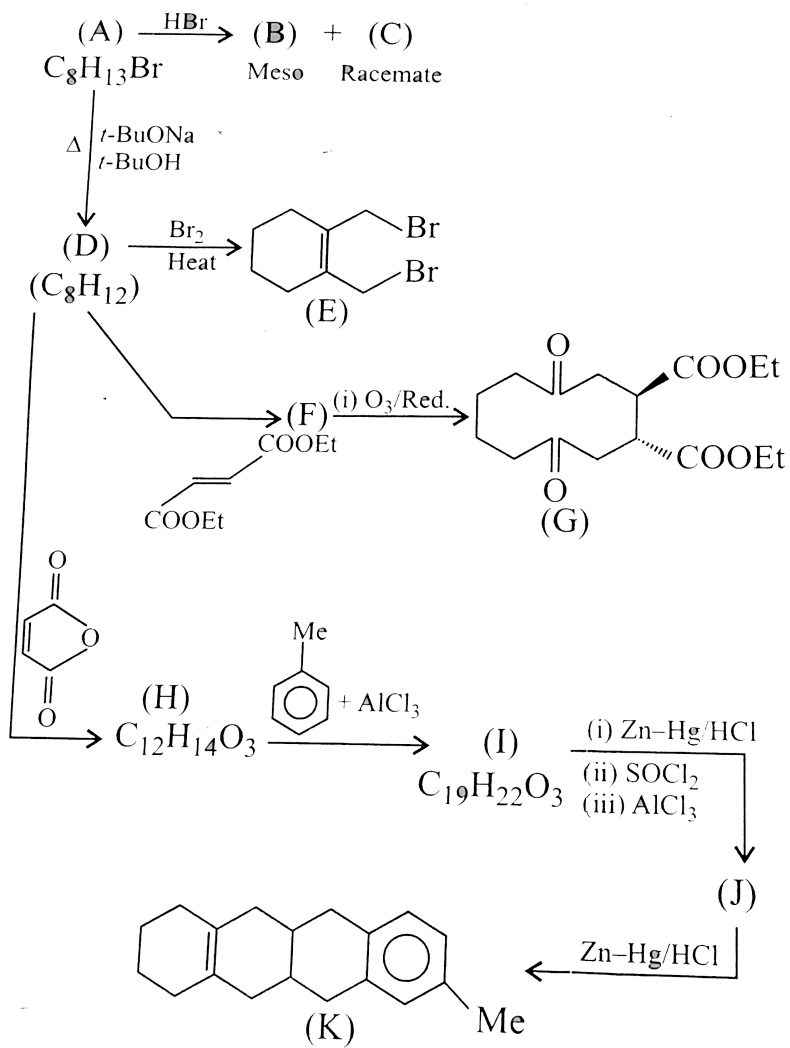
**Answer: b**



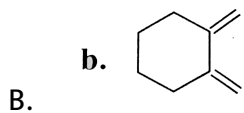
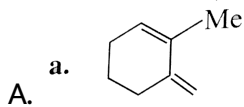
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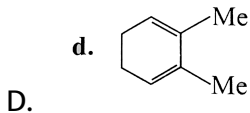
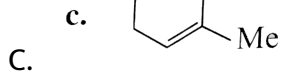
4. This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

Paragraph for Problems 1 - 8



Compound (D) is:



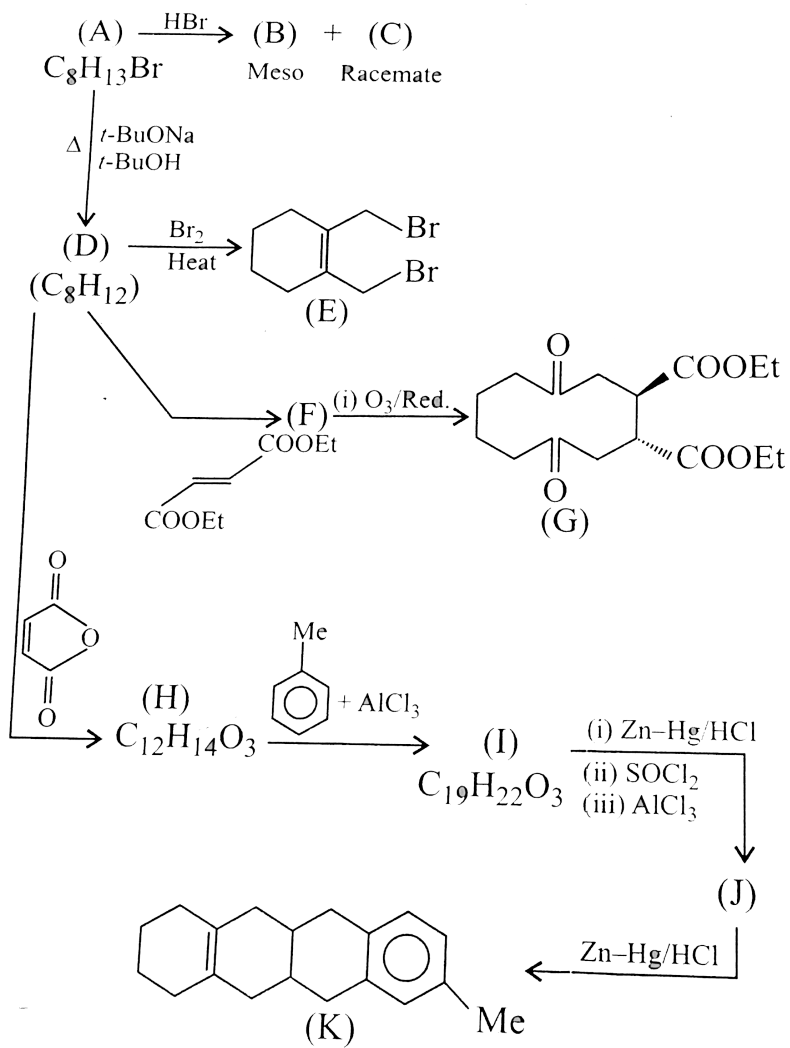


**Answer: b**

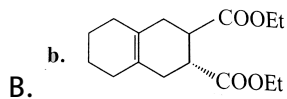
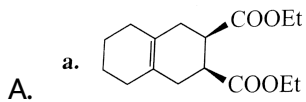
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5. This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

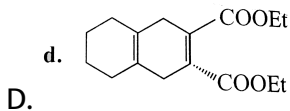
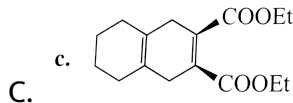
Paragraph for Problems 1 - 8



Compound (F) is:





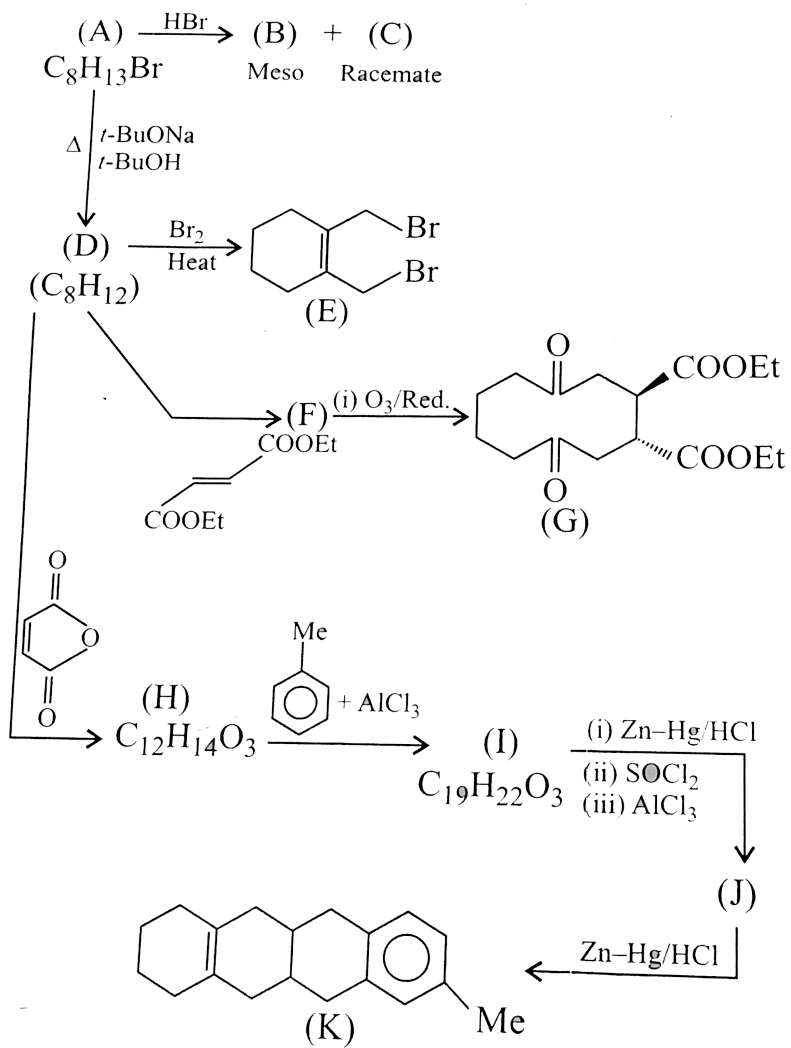


**Answer: b**

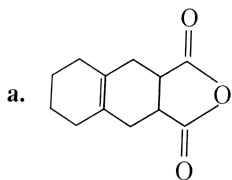
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6. This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

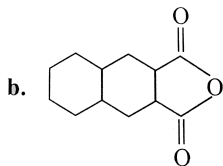
Paragraph for Problems 1 - 8



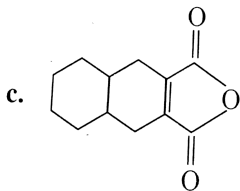
Compound (H) is:



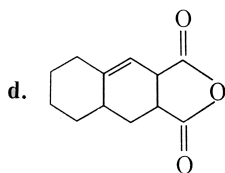
A.



B.



C.



D.

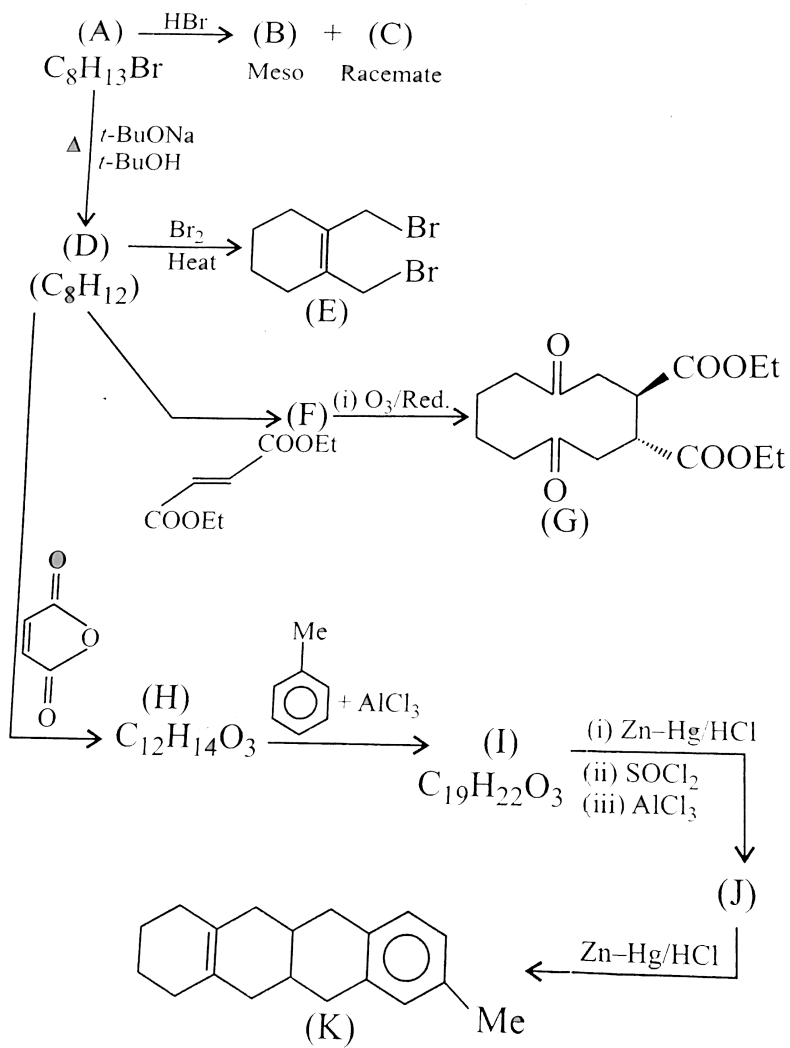
**Answer: a**



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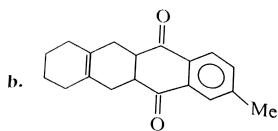
7. This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

Paragraph for Problems 1 - 8

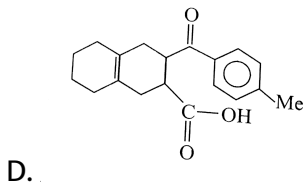
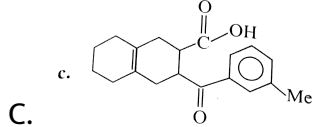


Compound (I) is:

A.



B.

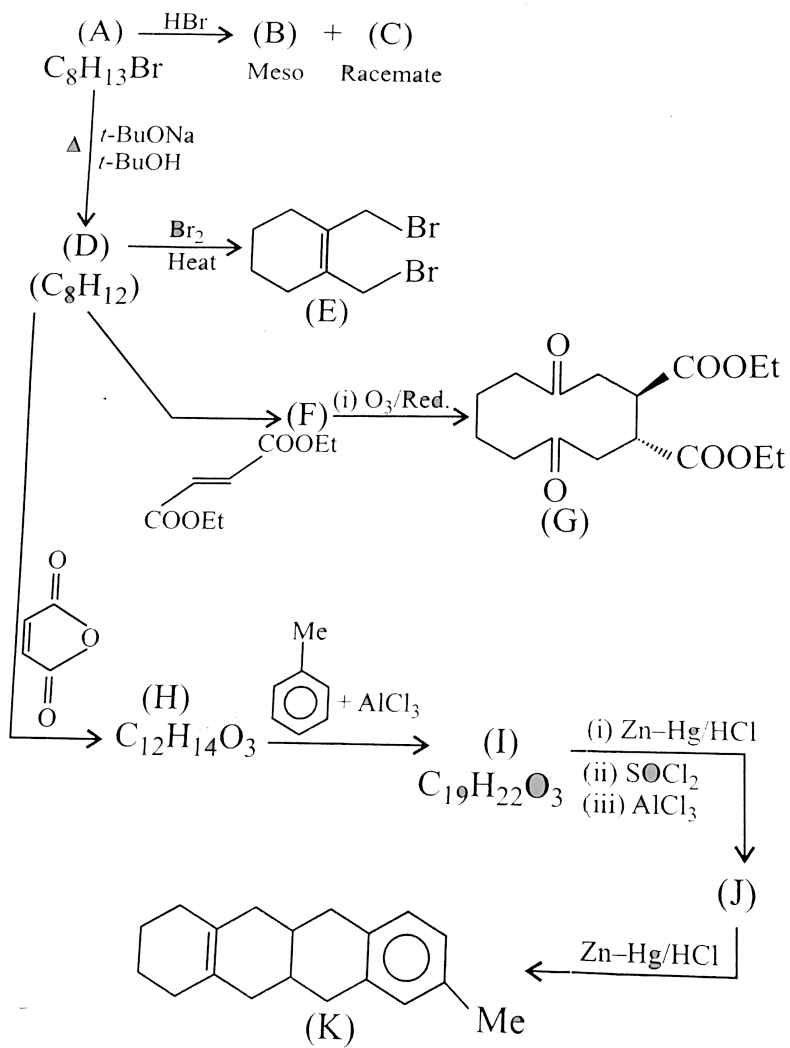


**Answer: d**

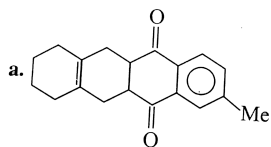
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**8.** This section contains 11 paragraphs, Based on each paragraph, four to nine multiple choice questions have to be answered. Each question has four multiple choice questions have to be answered. Each question has four multiple choices (a), (b), (c) and (d) out of which only one is correct or wrong.

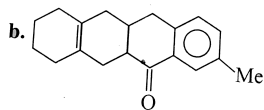
Paragraph for Problems 1 - 8



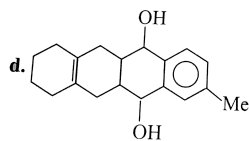
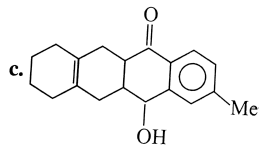
Compound (J) is:



A.



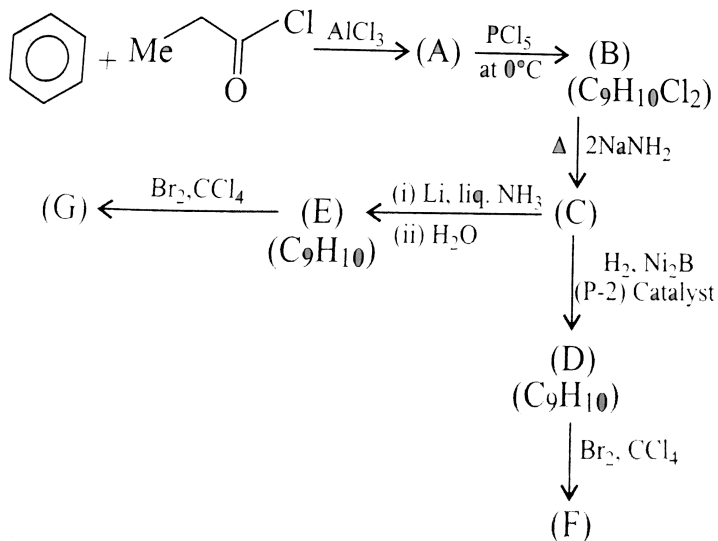
B.



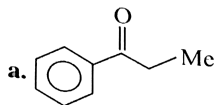
Answer: b

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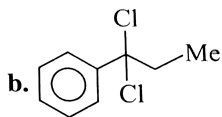
9. Paragraph for problems 9 - 15



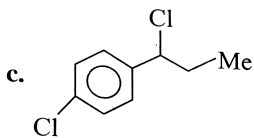
Compound (A) on reaction with  $NH_2NH_2 + OH^-$  is:



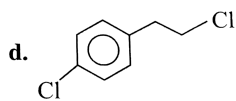
A.



B.



C.



D.

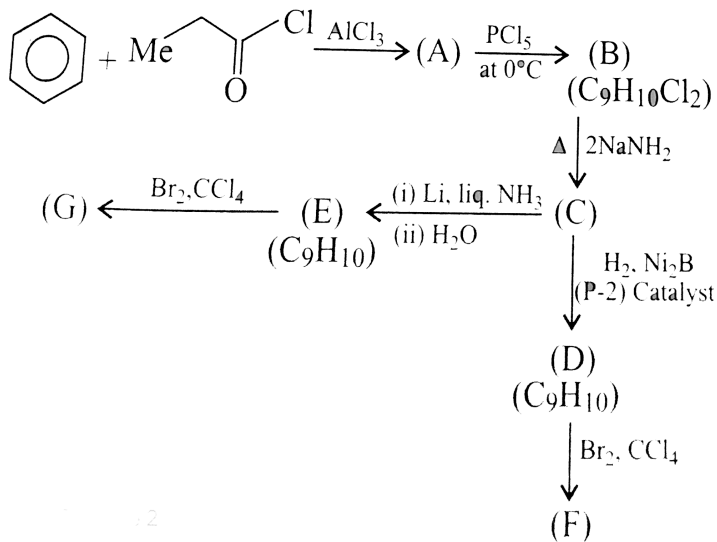
Answer: c



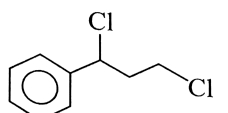
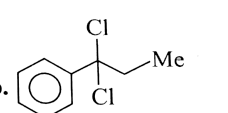
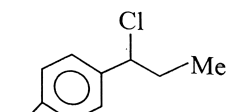
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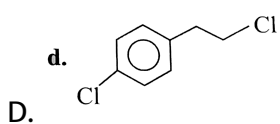


10. Paragraph for problems 9 - 15



Compound (B) is:

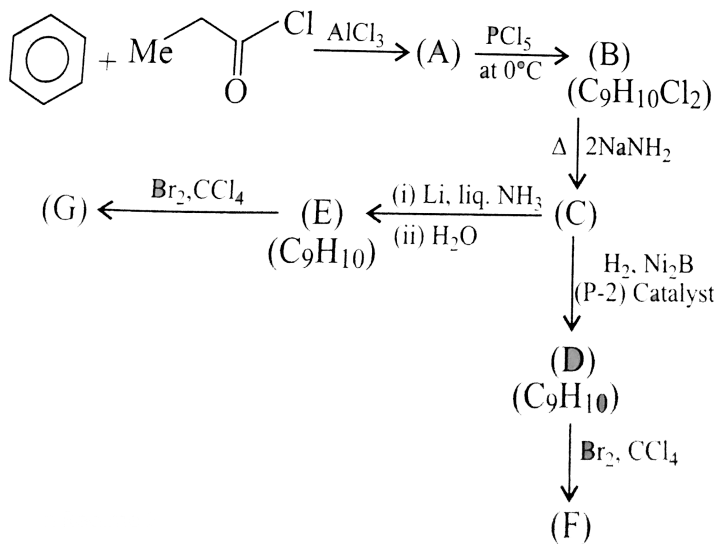
- A. 
- B. 
- C. 



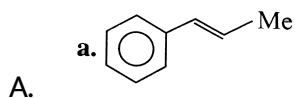
Answer: *b*

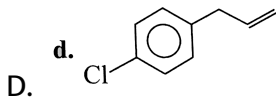
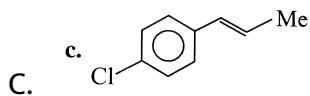
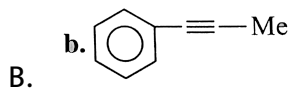
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11. Paragraph for problems 9 - 15



Compound (C) is:

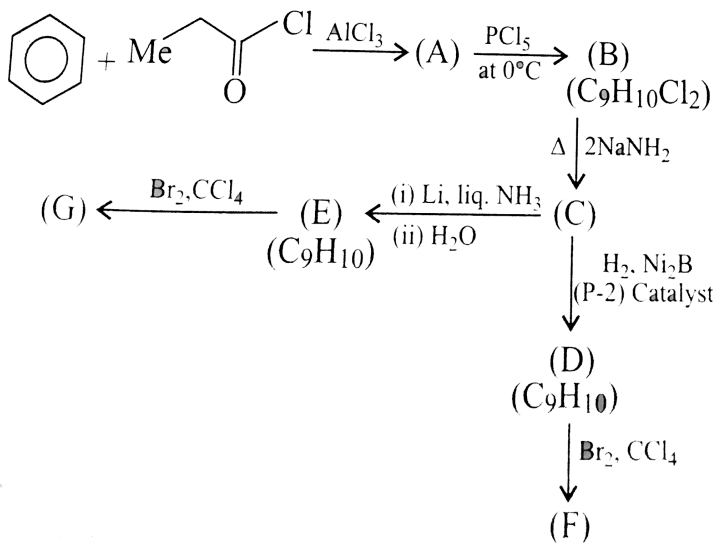




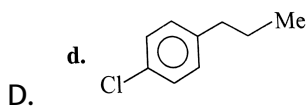
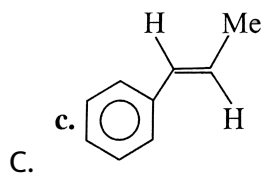
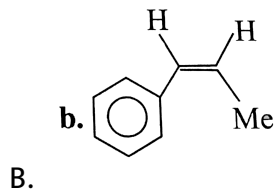
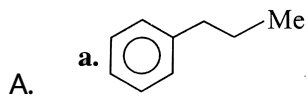
Answer: *b*

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## 12. Paragraph for problems 9 - 15



Compound (D) is:

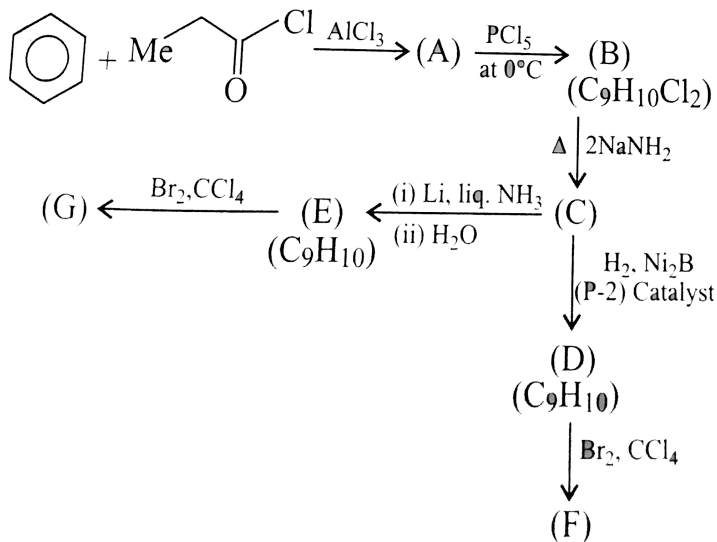


**Answer: b**

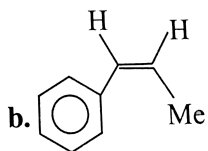
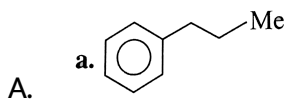


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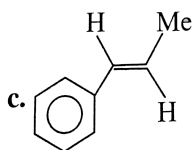
13. Paragraph for problems 9 - 15



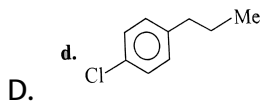
Compound (E) is:



B.



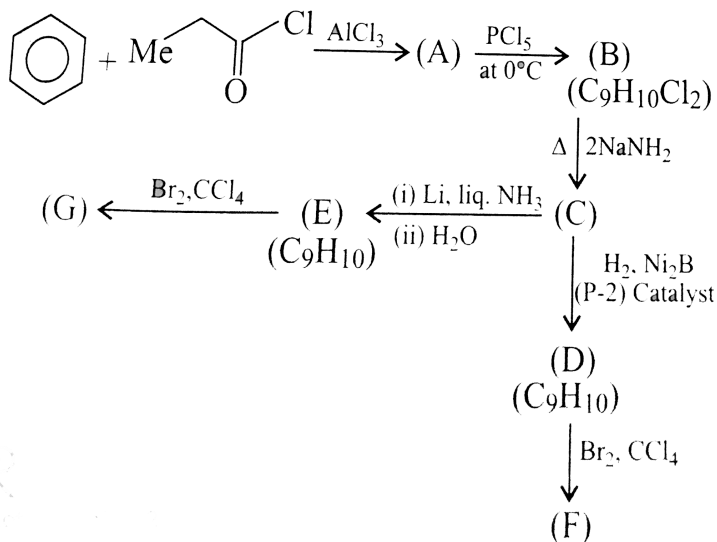
C.



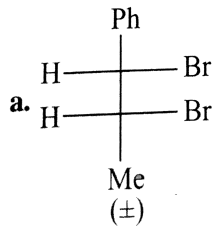
Answer: c

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#### 14. Paragraph for problems 9 - 15

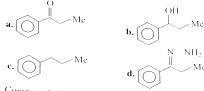


Compound (F) is:

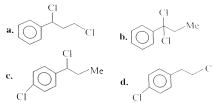


A.

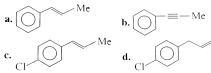
9. Compound (A) on reaction with  $\text{NH}_4\text{SH}$ , +  $\text{OH}^-$  is:



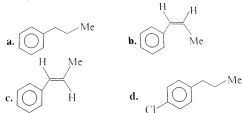
10. Compound (B) is:



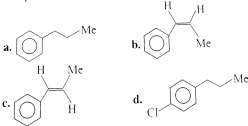
11. Compound (C) is:



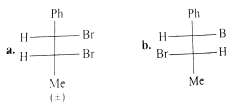
12. Compound (D) is:



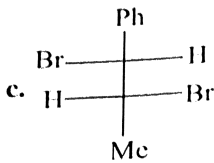
13. Compound (E) is:



14. Compound (F) is:



B.

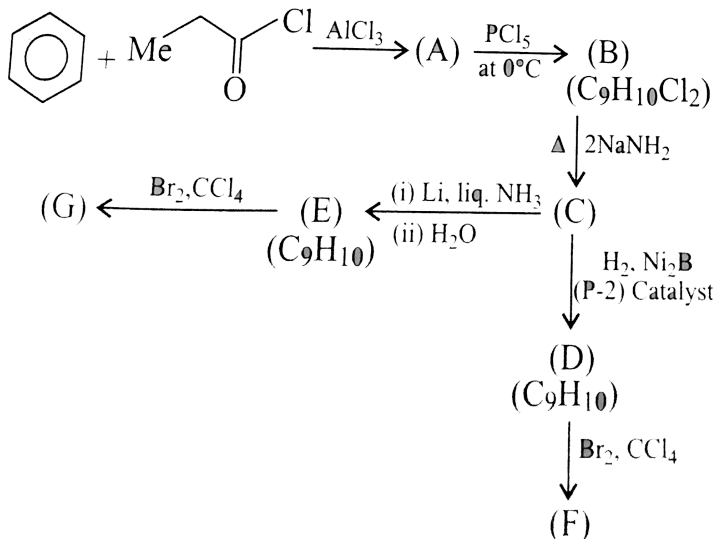


C.

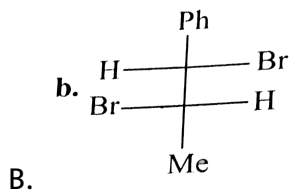
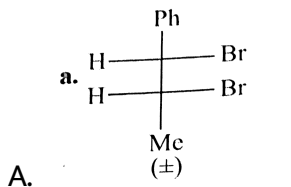
D. 

Answer: d

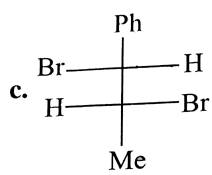
15. Paragraph for problems 9 - 15



Compound (G) is:







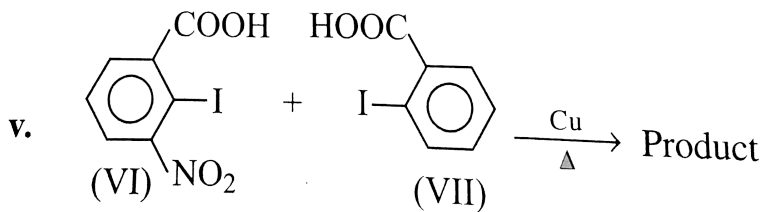
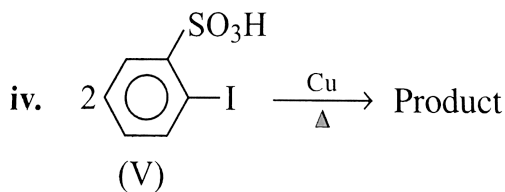
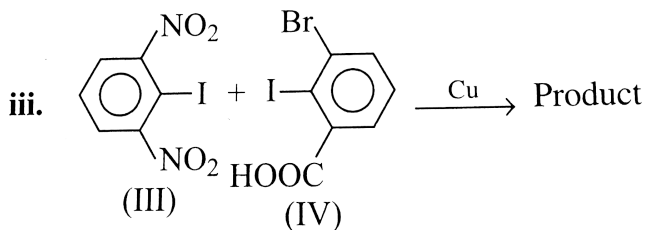
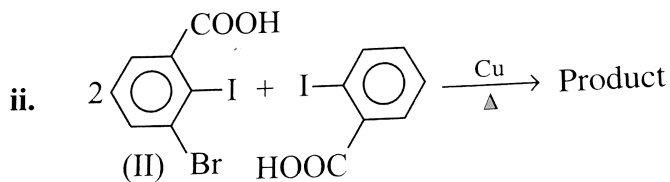
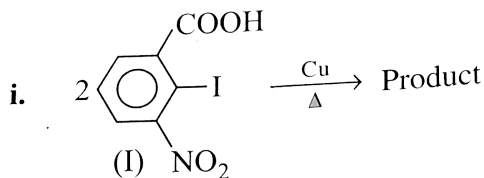
C.

D. Both (b) and (c)

**Answer: a**



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16.

All the reactions *i*) to (*v*) are examples of:

A. Wurtz reaction

B. Wurtz-Fitting reaction

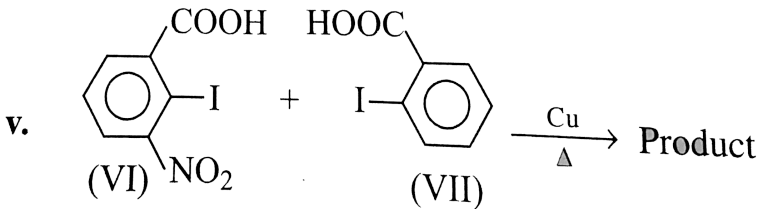
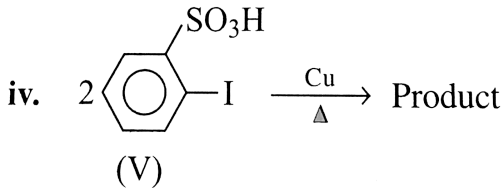
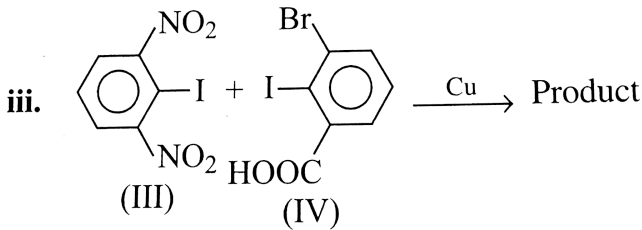
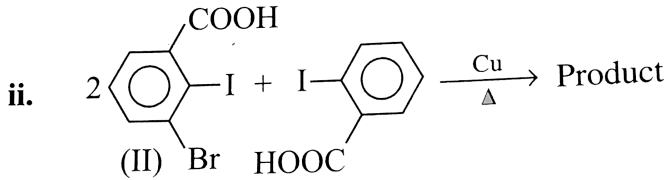
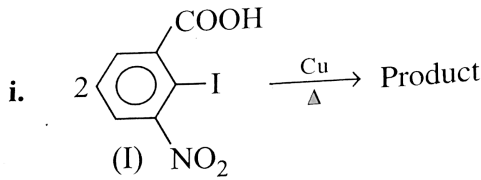
C. Ullmann reaction

D. Frankland reaction

**Answer:** *c*

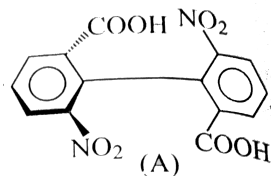


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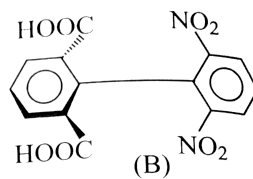
17.

The compound(s) in the reaction (i) is/are:



A. Resolve (+) and (-) forms of

B. Non-resolvable form of (A)

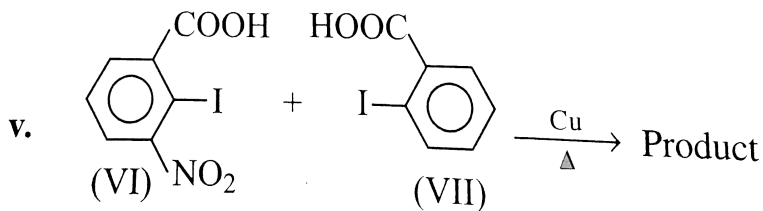
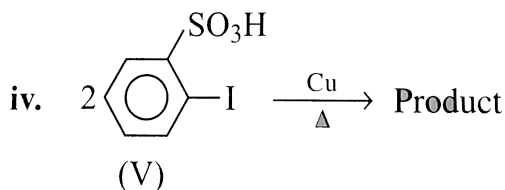
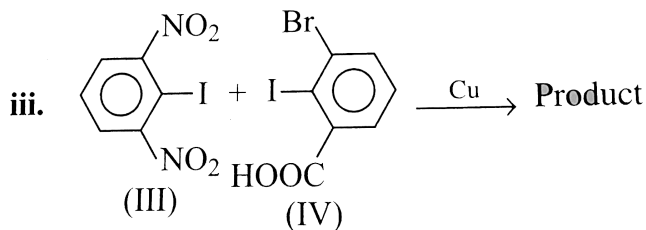
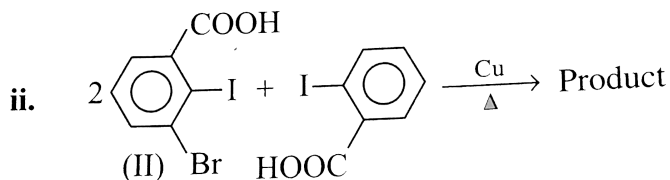
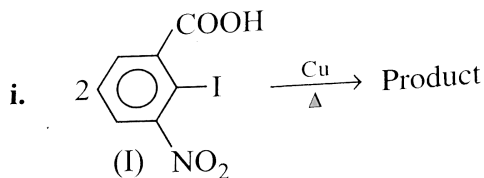


C. Resolvable ( + ) and ( - ) forms of

D. Non-resolvable form to (B).

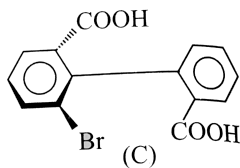
**Answer: a**

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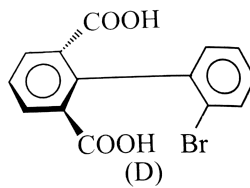
18.

The compound(s) in the reaction (ii) is/are:



A. Resolve (+) and (-) forms of

B. Non-resolvable form of (C)



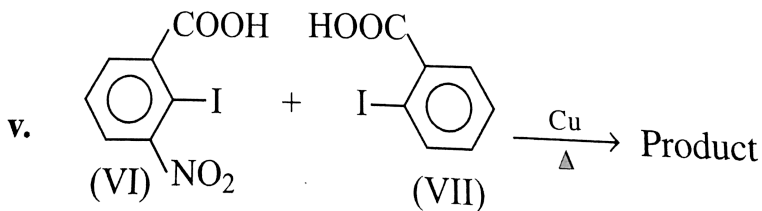
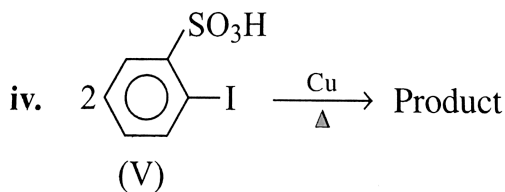
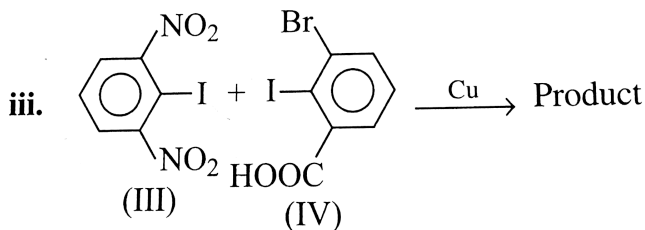
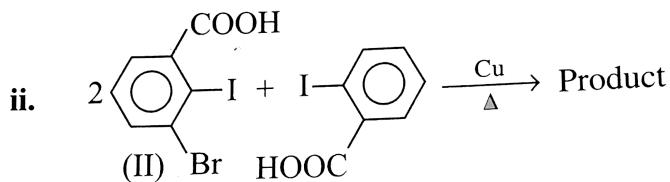
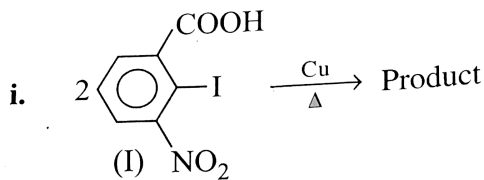
C. Resolvable (+) and (-) forms of

D. Non-resolvable form to (D).

**Answer: a**

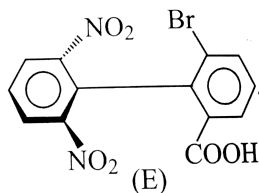


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19.

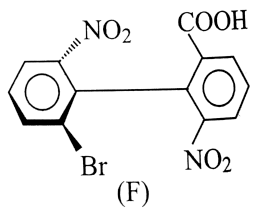
The compound(s) in the reaction (iii) is/are:



A. Resolve (+) and (-) forms of



B. Non-resolvable form of (*E*)



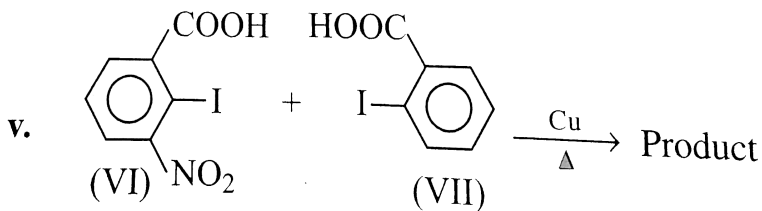
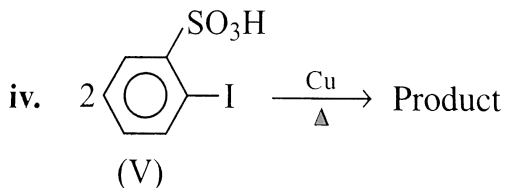
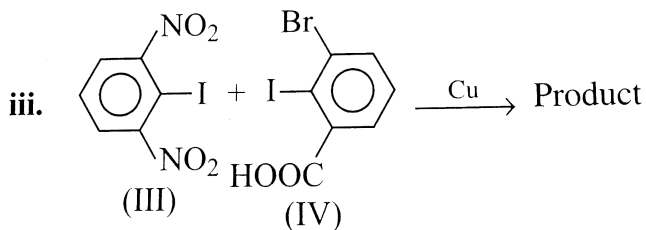
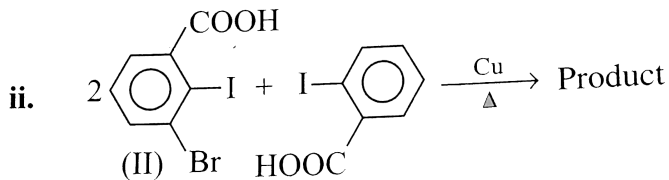
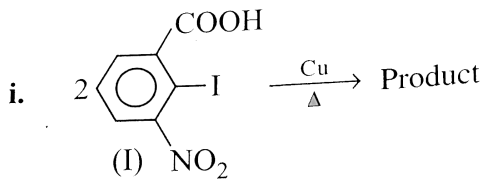
C. Resolvable (+) and (-) forms of

D. Non-resolvable form to (*F*).

**Answer: b**

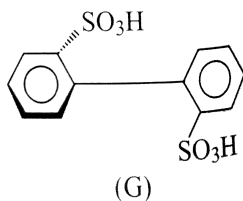


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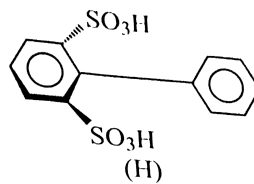
20.

The compound(s) in the reaction (iv) is/are:



A. Resolve (+) and (-) forms of

B. Non-resolvable form of (*G*)



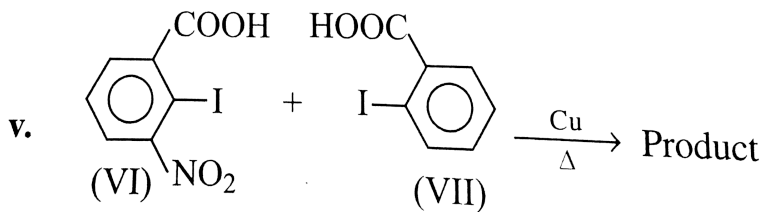
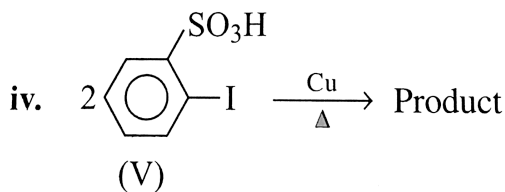
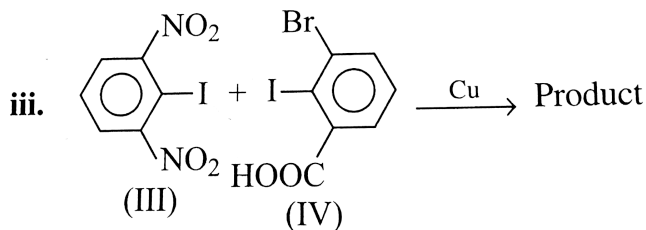
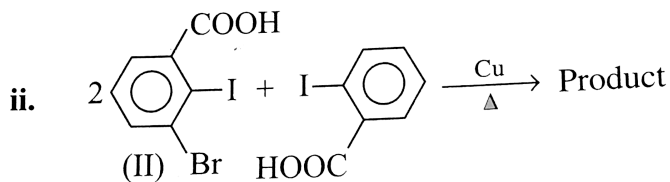
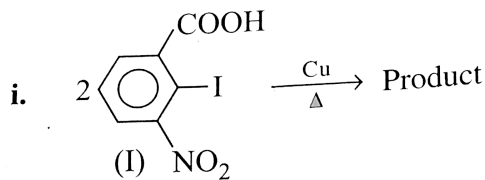
C. Resolvable (+) and (-) forms of

D. Non-resolvable form to (*H*).

**Answer: a**

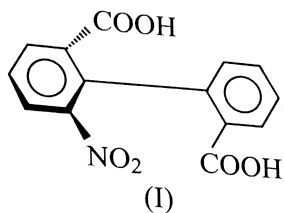


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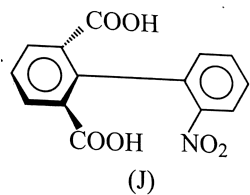
21.

The compound(s) in the reaction (v) is/are:



A. Resolve (+) and (-) forms of

B. Non-resolvable form of (I)



C. Resolvable (+) and (-) forms of

D. Non-resolvable form to (J).

**Answer: a**

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22. Compound (A), an alkene with molecule formula ( $C_5H_{10}$ ) exists in various structures. On monochlorination and dichlorination, it again shows various structures and stereoisomers.

A. 5

B. 6

C. 7

D. 8

Answer: c

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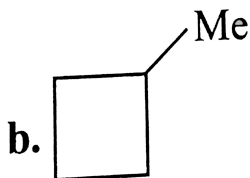
23. Compound (A), an alkene with molecule formula ( $C_5H_{10}$ ) exists in various structures. On monochlorination and dichlorination, it again shows various structures and stereoisomers.

One of the cyclic isomers of (A) on monochlorination gives one product.

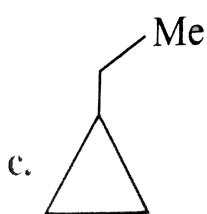
The isomer (I) is:



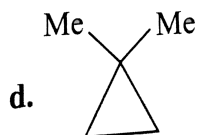
A.



B.



C.



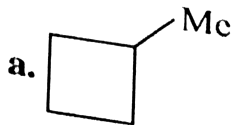
D.

**Answer: a**

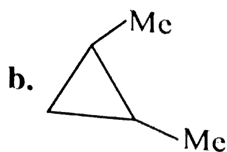
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**24.** Compound (A), an alkene with molecule formula ( $C_5H_{10}$ ) exists in various structures. On monochlorination and dichlorination, it again shows various structures and stereoisomers.

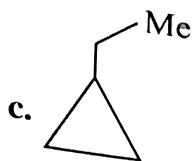
Another isomer (II) on monochlorination gives two products. The isomer (II) is:



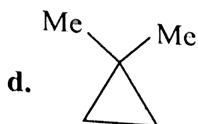
A.



B.



C.



D.

**Answer: d**

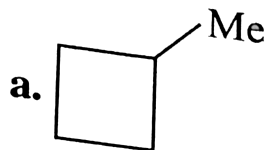
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25. Compound (A), an alkene with molecule formula ( $C_5H_{10}$ ) exists in various structures. On monochlorination and dichlorination, it again

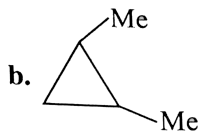


shows various structures and stereoisomers.

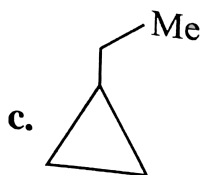
Isomer (III) on monochlorination gives three products. The isomer (III) is:



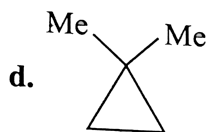
A.



B.



C.



D.

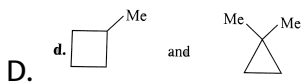
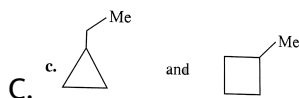
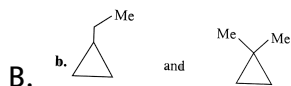
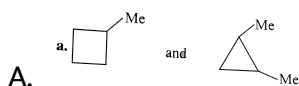
Answer: b



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26. Compound (A), an alkene with molecule formula ( $C_5H_{10}$ ) exists in various structures. On monochlorination and dichlorination, it again shows various structures and stereoisomers.

Two isomers (IV) and (V) on monochlorination give four products. The isomers (IV) and (V) are:



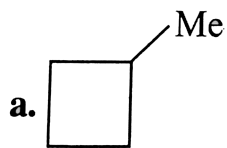
**Answer: c**

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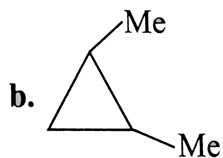
27. Compound (A), an alkene with molecule formula ( $C_5H_{10}$ ) exists in various structures. On monochlorination and dichlorination, it again

shows various structures and stereoisomers.

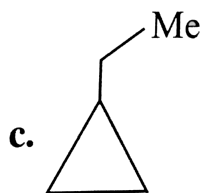
Which isomer of (A) shows stereoisomerism ?



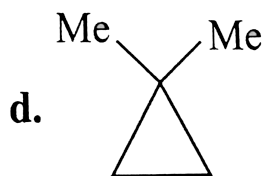
A.



B.



C.



D.

**Answer: b**



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28. Compound (A), an alkene with molecule formula ( $C_5H_{10}$ ) exists in various structures. On monochlorination and dichlorination, it again shows various structures and stereoisomers.

The total number of stereoisomerism shown by isomer in Q.No 26 is:

A. 2

B. 3

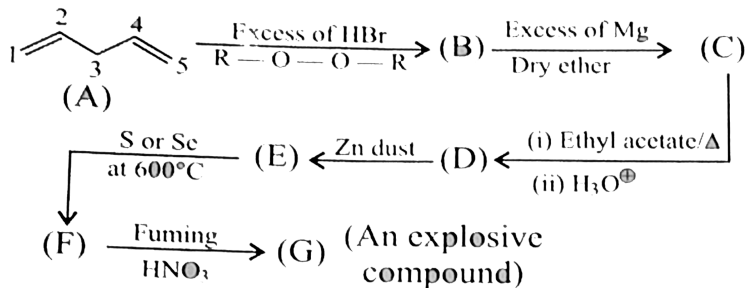
C. 4

D. 5

**Answer: b**

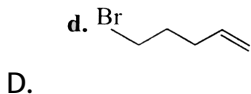
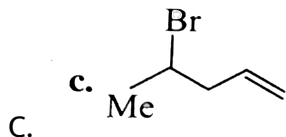
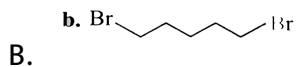
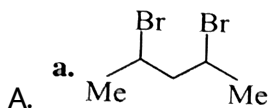


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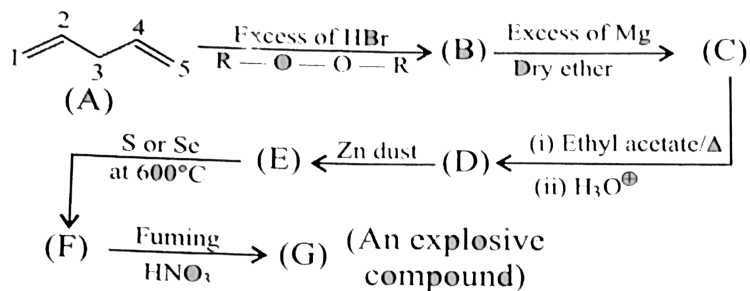
29.

Compound (B) is:



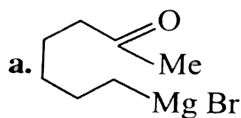
Answer: b

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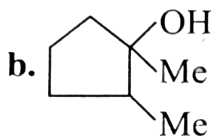


30.

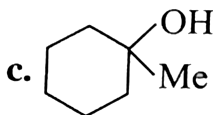
Compound (D) is:



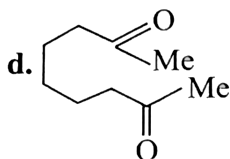
A.



B.



C.

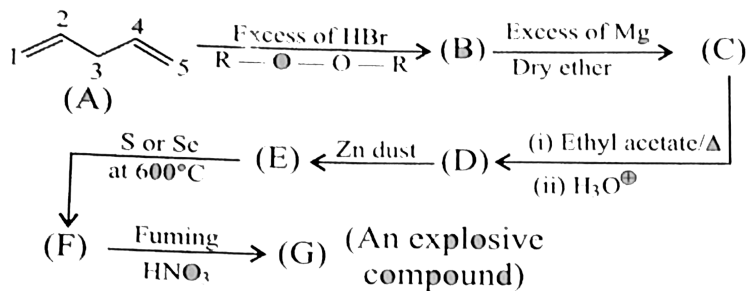


D.

Answer: c

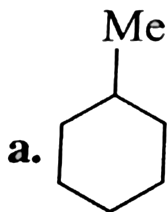


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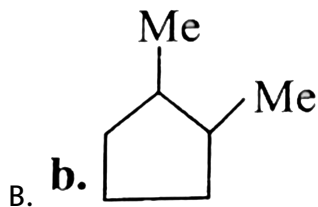


31.

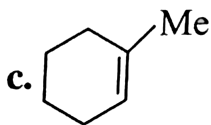
Compound (E) is:



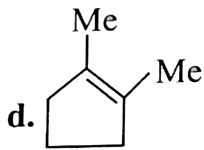
A.



B.



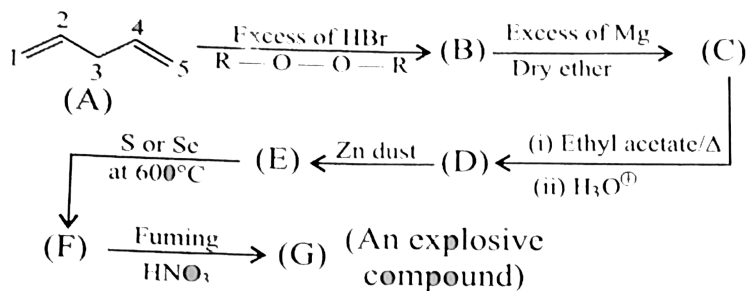
C.



D.

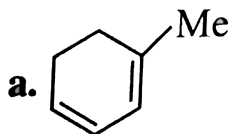
Answer: a

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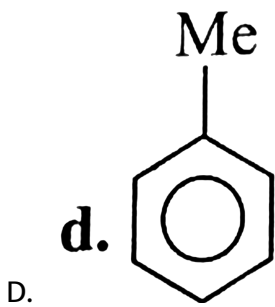
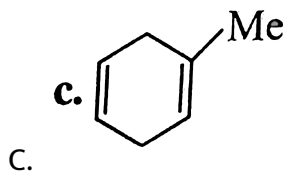
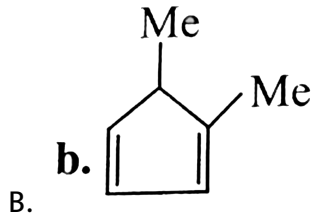
32.

Compound (F) is:



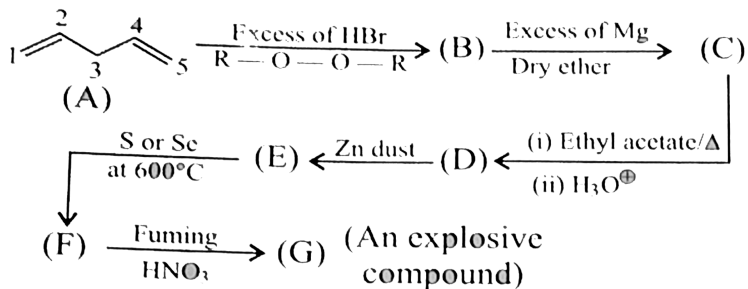
A.





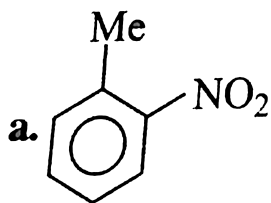
Answer: *d*

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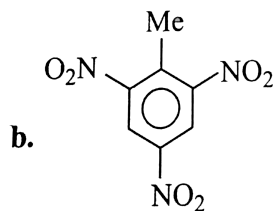


33.

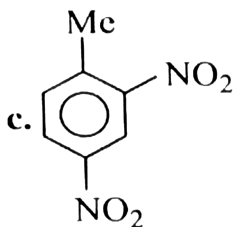
Compound (G) is:



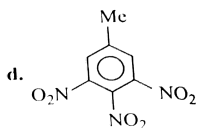
A.



B.



C.



D.

**Answer: b**

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34. A hydrocarbon (A)  $(C_8H_{10})$  is a stream volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.

The name of (A) is:

- A. Dibenzocyclobutane
- B. Dibenzocyclobutadiene
- C. Dibenzocyclobutane
- D. Benzocyclobutane

**Answer: b**

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35. A hydrocarbon (A)  $(C_8H_{10})(E)$  is a steam volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.

Degree of unsaturation ( $DU$ ) in (A) is:

A. 7

B. 8

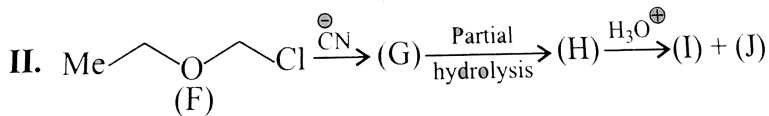
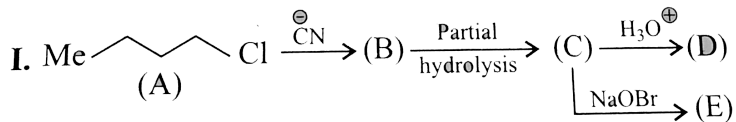
C. 9

D. 10

**Answer: c**



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36.

Compound (B) is:

A.  $n\text{-BU}-\text{C} \equiv \text{N}$

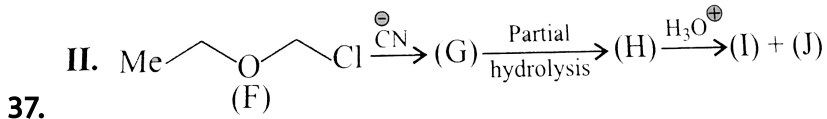
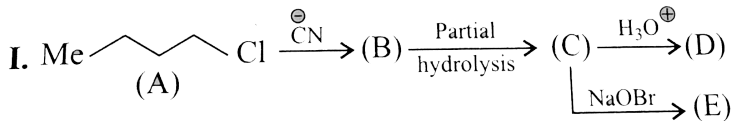
B.  $n\text{-Bu}-\overset{\oplus}{\text{N}} \equiv \text{overser}(o-)(\text{C})$

C. Both

D. None

Answer: a

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Compound (C) is:

A.  $\text{Bu}-\text{COOH}$

B.  $\text{Bu}-\text{NH}-\text{CHO}$

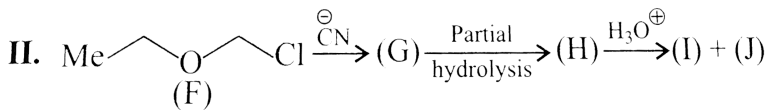
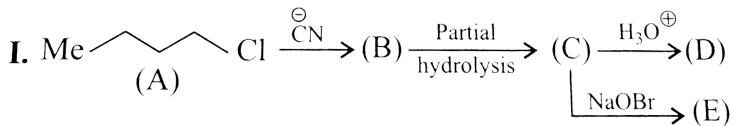
C.  $\text{Bu}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}_2$

D.  $\text{Bu}-\text{NH}_2$

Answer: c



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38.

Compound (D) is:

A.  $\text{Bu}-\text{COOH}$

B.  $\text{Bu}-\text{NH}-\text{CHO}$

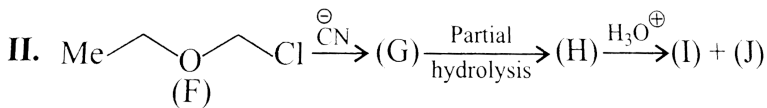
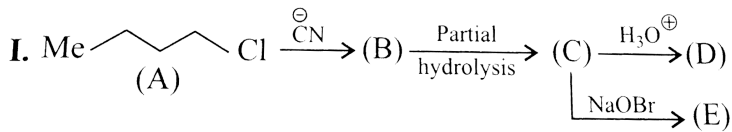
C.  $\text{Bu}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}_2$

D.  $\text{Bu}-\text{NH}_2$

Answer: a



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39.

In the conversion of (C) to (E), (E) is  $\text{Bu}-\text{NH}_2$ . This reaction is called:

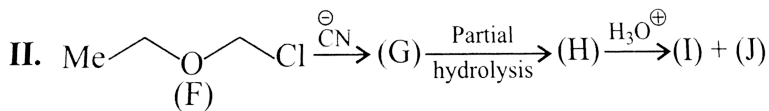
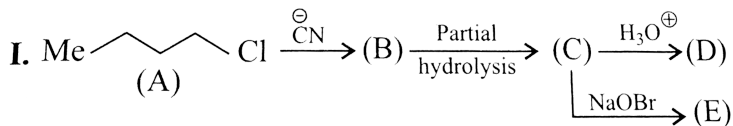
- A. Halofrom reaction
- B. Halfrom reaction
- C. Hofmann bromide reaction
- D. Hofmann bromide rearrangement reaction

Answer: d



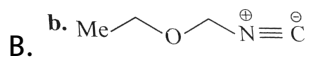
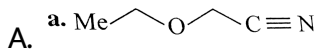
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40.

Compound (G) is:



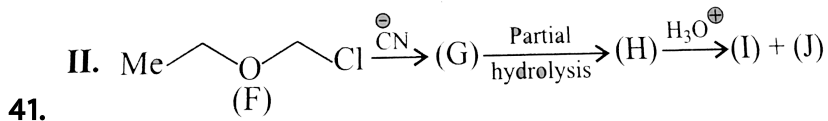
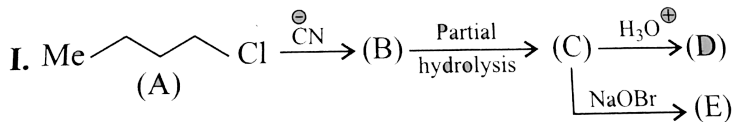
C. Both

D. None

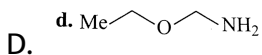
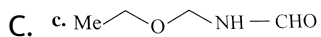
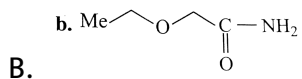
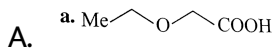
Answer: *b*



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Compound (H) is:

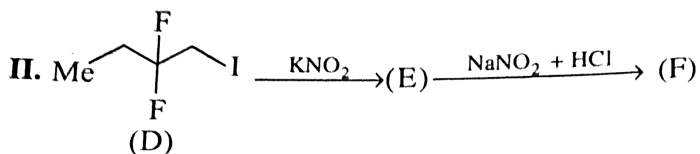
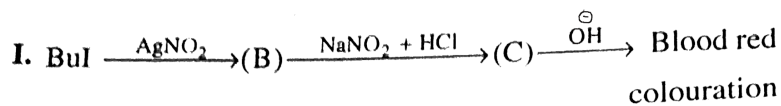


Answer: c

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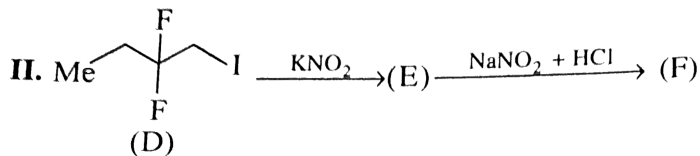
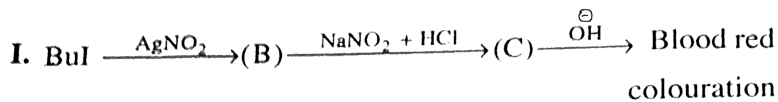
44.

Compound (B) is:

- A.  $\text{Bu} - \text{NO}_2$
- B.  $\text{Bu} - \text{O} - \text{N} = \text{O}$
- C. Both
- D. None

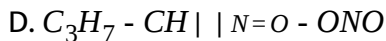
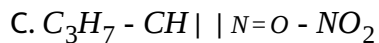
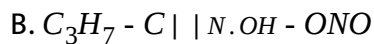
Answer: a

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45.

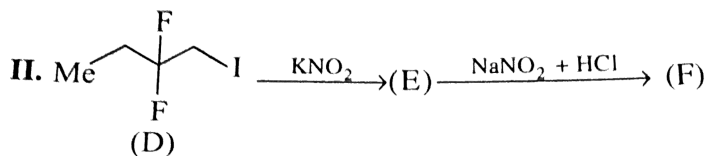
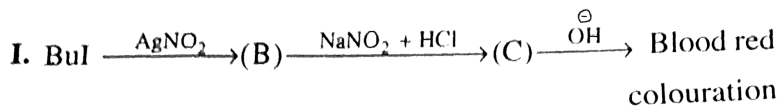
Compound (C) is:



Answer: c

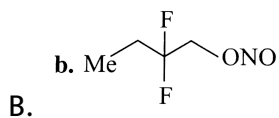
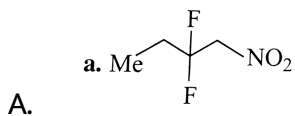


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46.

Compound (E) is:

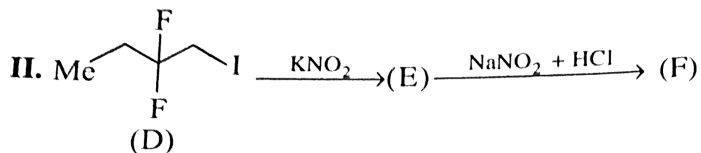
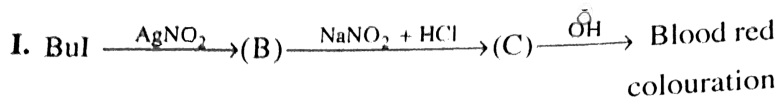


C. Both

D. None

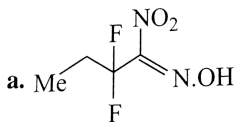
Answer: b

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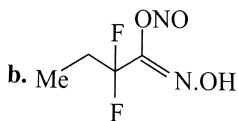


47.

Compound (F) is:



A.



B.

C. No reaction

D. Both (a) and (b)

Answer: c

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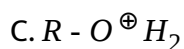
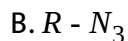
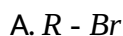


**48.**  $SN$  reaction is given by these compounds, which have a nucleophilic group and a good leaving  $EWG$ . It should be stable after leaving with bonding pair of  $\bar{e}$ 's and it should have high polarisability.

Nucleophilic aliphatic substitution reaction is mainly of two types  $SN^{-1}$  and  $SN^2$ .  $SN^{-1}$  mechanism is a two step process. Reaction velocity of  $SN^{-1}$  depends only on the concentration of the substrate. It proceeds via the formation of carbocation, optically active substrate gives (  $\oplus$  ) and (  $o -$  ) forms of the product.

In most of the cases, the product usually consists of 5 - 20 % inverted and (95 - 80 % ) racemised species. The more stable is the carbocation, the greater is the proportion of racemisation. In solvolysis reaction, the more nucleophilic is the solvent, the greater is the proportion of inversion.

Which of the following will give  $SN$  reaction?



D. All

Answer: d



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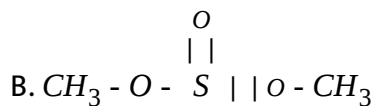
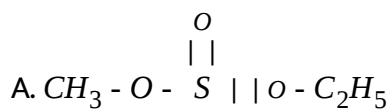
49.  $S_N$  reaction is given by these compounds, which have a nucleophilic group and a good leaving  $EWG$ . It should be stable after leaving with bonding pair of  $\bar{e}$ 's and it should have high polarisability.

Nucleophilic aliphatic substitution reaction is mainly of two types  $S_N^{-1}$  and  $S_N^2$ .  $S_N^{-1}$  mechanism is a two step process. Reaction velocity of  $S_N^{-1}$  depends only on the concentration of the substrate. It proceeds via the formation of carbocation, optically active substrate gives (  $\oplus$  ) and (  $o -$  ) forms of the product.

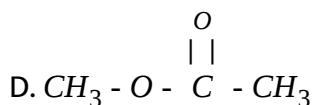
In most of the cases, the product usually consists of 5 - 20 % inverted and (95 - 80 %) racemised species. The more stable is the carbocation, the greater is the proportion of racemisation. In solvolysis reaction, the more nucleophilic is the solvent, the greater is the proportion of

inversion.

Fugacity power of which group will be maximum?



C. 



Answer: c

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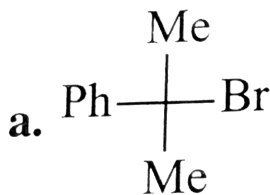
50.  $\text{S}_{\text{N}}$  reaction is given by these compounds, which have a nucleophilic group and a good leaving  $\text{EWG}$ . It should be stable after leaving with bonding pair of  $\bar{e}$ 's and it should have high polarisability.

Nucleophilic aliphatic substitution reaction is mainly of two types  $\text{S}_{\text{N}}^{-1}$  and  $\text{S}_{\text{N}}^2$ .  $\text{S}_{\text{N}}^{-1}$  mechanism is a two step process. Reaction velocity of  $\text{S}_{\text{N}}^{-1}$  depends only on the concentration of the substrate. It proceeds via the

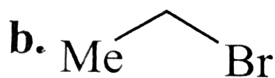
formation of carbocation, optically active substrate gives (  $\oplus$  ) and (  $\ominus$  ) forms of the product.

In most of the cases, the product usually consists of 5 - 20 % inverted and (95 - 80 % ) racemised species. The more stable is the carbocation, the greater is the proportion of racemisation. In solvolysis reaction, the more nucleophilic is the solvent, the greater is the proportion of inversion.

Which of the following gives  $SN^1$  reaction?



A.



B.

C.  $MeBr$

D. All

**Answer: a**



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51.  $SN^2$  reaction is a bimolecular reaction which takes place by the formation of  $T.S.$  Velocity of the reaction depends on the concentration of the substrate as well as the nucleophile.

The reaction is favoured by strong  $Nu^{o-}$  and in the presence of polar aprotic solvent, optically active halides give Walden inversion by  $SN^2$  mechanism. The presence of hetero group (atom) at  $\beta$ -C atom,

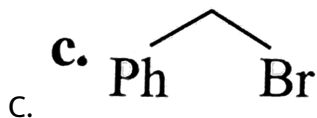
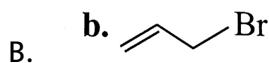
unsaturation at  $\beta$ -C and  $\left( \begin{array}{c} O \\ | | \\ - C - \end{array} \right)$  group at  $\alpha$ -C atom favor  $SN^2$

mechanism.

Allyl halides and benzyl halides give  $SN^1$  and  $SN^2$  reactions. Allyl halides also give  $SN^2$  mechanism.  $EDG$  at ortho- and para- positions in benzyl halides favors  $SN^1$  mechanism, whereas  $EWG$  favors  $SN^2$  mechanism.

Which of the following will give  $SN^2$  mechanism?

A.  $MeBr$



D. All

Answer: d

 [View Text Solution](#)

52.  $SN^2$  reaction is a bimolecular reaction which takes place by the formation of  $T.S.$  Velocity of the reaction depends on the concentration of the substrate as well as the nucleophile.

The reaction is favoured by strong  $Nu^{o-}$  and in the presence of polar aprotic solvent, optically active halides give. Walden inversion by  $SN^2$  mechanism. The presence of hetero group (atom) at  $\beta$ -C atom,

unsaturation at  $\beta$ -C and  $\left( \begin{array}{c} O \\ || \\ - C - \end{array} \right)$  group at  $\alpha$ -C atom favor  $SN^2$

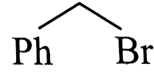
mechanism.

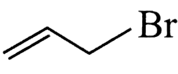
Allyl halides and benzyl halides give  $SN^1$  and  $SN^2$  reactions. Allyl halides also give  $SN^2$  mechanism  $EDG$  at ortho- and para- positions in benzyl

halides favors  $SN^1$  mechanism, whereas  $EWG$  favors  $SN^2$  mechanism.

Which of the following will give Walden inversion ?

A.  $C_4H_9 - CHD - Br$ ,

B. **b.**  CC(Br)C

C. **c.**  C=CCBr

D. All

Answer: a



[View Text Solution](#)

53.  $SN^2$  reaction is a bimolecular reaction which takes place by the formation of  $T.S.$  Velocity of the reaction depends on the concentration of the substrate as well as the nucleophile.

The reaction is favoured by strong  $Nu^{o-}$  and in the presence of polar aprotic solvent, optically active halides give Walden inversion by  $SN^2$  mechanism. The presence of hetero group (atom) at  $\beta$ -C atom,

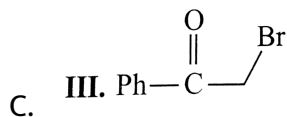
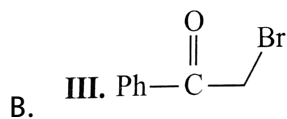
unsaturation at  $\beta$ -C and  $\left( \begin{array}{c} \text{O} \\ | | \\ - \text{C} - \end{array} \right)$  group at  $\alpha$ -C atomn favor  $SN^2$

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Which of the following will gives  $SN^2$  mechanism gt

A. *EtBr*



D. All

**Answer: d**



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54.  $SN^2$  reaction is a bimolecular reaction which takes place by the formation of  $T.S.$  Velocity of the reaction depends on the concentration of the substrate as well as the nucleophile.

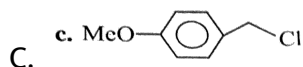
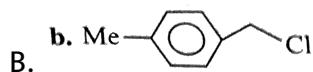
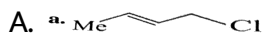
The reaction is favoured by strong  $Nu^{O-}$  and in the presence of polar aprotic solvent, optically active halides give Walden inversion by  $SN^2$  mechanism. The presence of hetero group (atom) at  $\beta$ -C atom,

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Allyl halides and benzyl halides give  $SN^1$  and  $SN^2$  reactions. Allyl halides also give  $SN^2$  mechanism.  $EDG$  at ortho- and para- positions in benzyl halides favors  $SN^1$  mechanism, whereas  $EWG$  favors  $SN^2$  mechanism.

Which of the following will give  $SN^1$  reaction?



D. All

Answer: d



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55. Isopropyl bromide was treated separately with sodium ethoxide under two different conditions.

Reaction I:

Treatment of isopropyl bromide with  $(Me_3CONa)$  at  $40^\circ C$  gave almost exclusively compound (A)  $(C_3H_6)$ .

Reaction II:

Treatment of  $(i-PrBr)$  with  $NaOC_2H_5$  at  $30^\circ C$  yielded compound (A)  $(C_3H_6)$  along with a small amount of an ether (B)  $(C_5H_{12}O)$ .

Compound (A) was readily oxidised by a neutral solution of cold dil.  $KMnO_4$  to give a brown precipitate.

The formations of (A) and (B) are best explained by:

A.  $SN^2$  reaction and  $E2$  reaction, respectively

B.  $E2$  reaction and  $SN^2$  reaction, respectively

C.  $E1$  reaction and  $SN^1$  reaction, respectively

D.  $E2$  reaction and  $SN^1$  reaction, respectively

**Answer: b**

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56. Isopropyl bromide was treated separately with sodium ethoxide under two different conditions.

Reaction I:

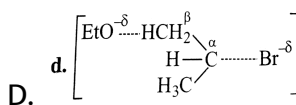
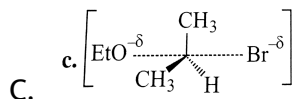
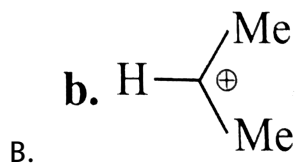
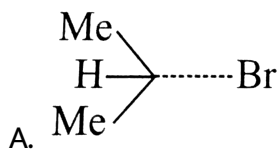
Treatment of isopropyl bromide with  $(Me_3CONa)$  at  $40^\circ C$  gave almost exclusively compound (A)  $(C_3H_6)$ .

Reaction II:

Treatment of  $(i - PrBr)$  with  $NaOC_2H_5$  at  $30^\circ C$  yielded compound (A)  $(C_3H_6)$  along with a small amount of an ether (B)  $(C_5H_{12}O)$ .

Compound (A) was readily oxidised by a neutral solution of cold dil.  $KMnO_4$  to give a brown precipitate.

Which of the following most accurately represents the activated complex formed in reaction (II) that leads to compound (A) ?



**Answer: d**



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57. Isopropyl bromide was treated separately with sodium ethoxide under two different conditions.

Reaction I:

Treatment of isopropyl bromide with  $(Me_3CONa)$  at  $40^\circ C$  gave almost exclusively compound (A)  $(C_3H_6)$ .

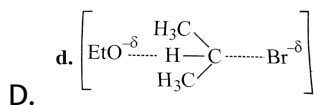
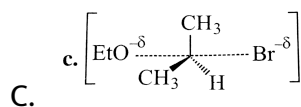
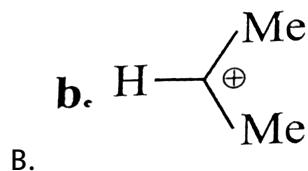
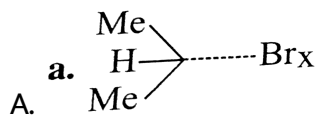
Reaction II:

Treatment of (*i*-PrBr) with  $\text{NaOC}_2\text{H}_5$  at  $30^\circ\text{C}$  yielded compound

(A) ( $\text{C}_3\text{H}_6$ ) along with a small amount of an ether (B) ( $\text{C}_5\text{H}_{12}\text{O}$ ).

Compound (A) was readily oxidised by a neutral solution of cold dil.  $\text{KMnO}_4$  to give a brown precipitate.

Referring to Q. No. 67 which of the following represents the intermediate *T.S* for the formation of compound (B) ?



**Answer: c**

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58. Isopropyl bromide was treated separately with sodium ethoxide under two different conditions.

Reaction I:

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Compound (A) was readily oxidised by a neutral solution of cold dil.  $KMnO_4$  to give a brown precipitate.

Which of the following is an accurate representation of compound (B)?

A.  $MeOMe$

B.  $Et - O - Et$

C.  $i - Pr - O - Et$

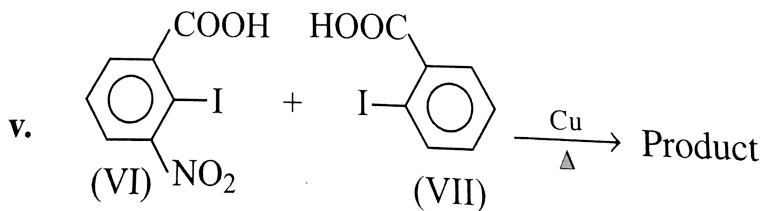
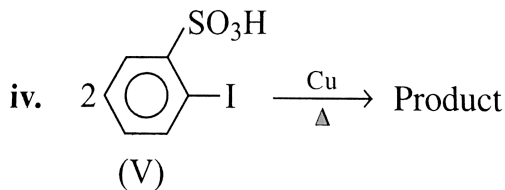
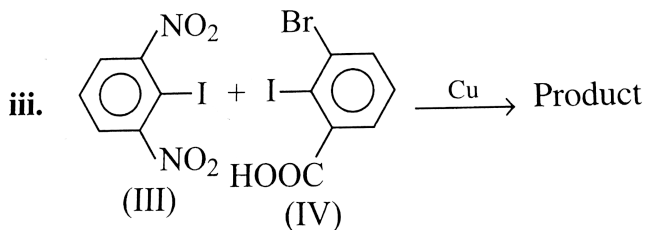
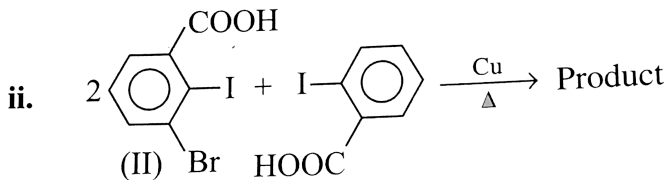
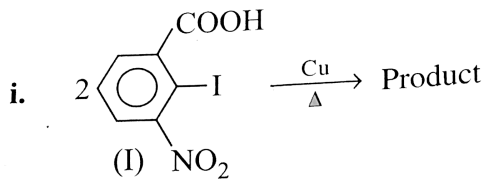
D.  $EtOMe$

Answer: c



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## Exercises Linked Comprehension Type



1. The type of stereoisomerism arising from restricted rotation about a single bond and where the stereoisomers can be isolated is called.

A. Kryptomerism



B. Atropisomerism

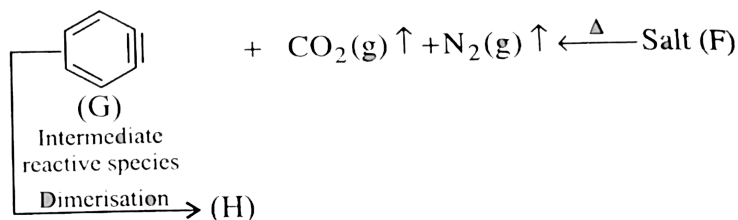
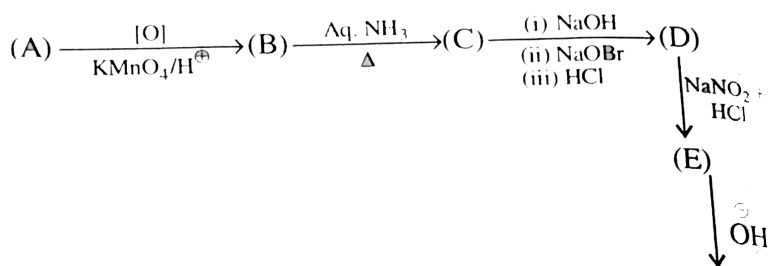
C. Allotropism

D. Merotropy

Answer: b

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2. A hydrocarbon (A) ( $C_8H_{10}$ ) (E) is a steam volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.



Compound (A) is:

A. Ethylbenzene

B. p-Xylene

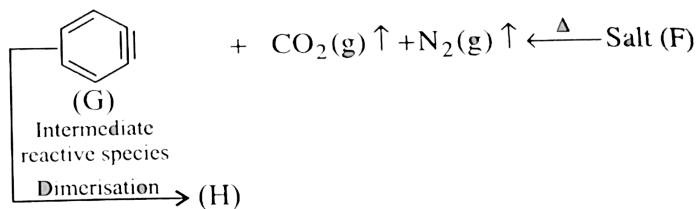
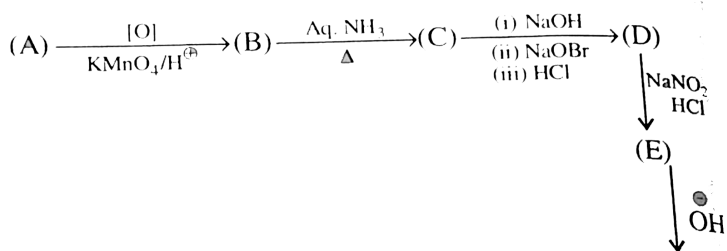
C. m-Xylene

D. o-Xylene

**Answer: d**

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3. A hydrocarbon (A) ( $C_8H_{10}$ ) (E) is a steam volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.



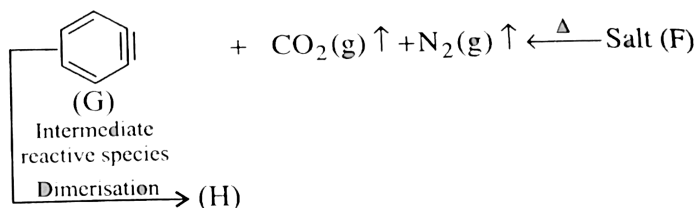
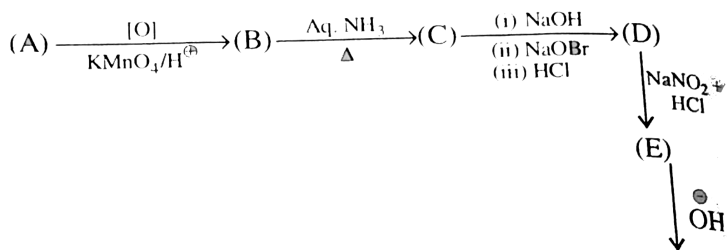
Compound (B) is:

- A. Phthalic acid
- B. Isophthalic acid
- C. Terephthalic
- D. Benzoic acid

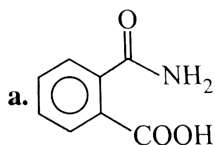
Answer: a

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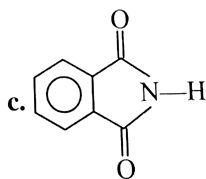
4. A hydrocarbon (A) ( $C_8H_{10}$ ) (E) is a steam volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.



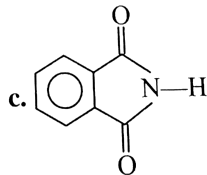
Compound (C) is:



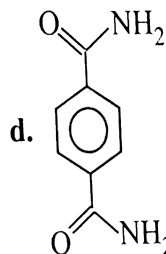
A.



B.



C.

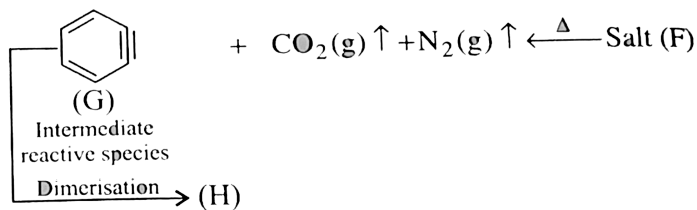
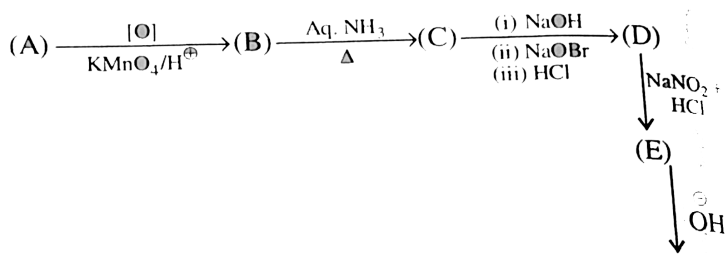


D.

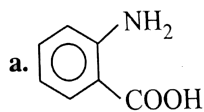
**Answer: c**

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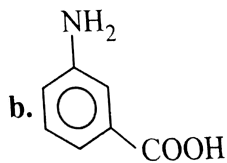
5. A hydrocarbon (A) ( $C_8H_{10}$ ) (E) is a stream volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.



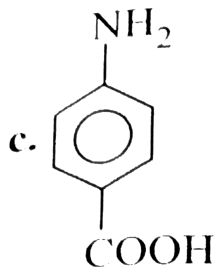
Compound (D) is:



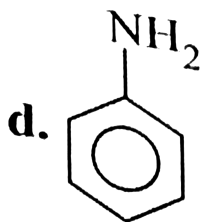
A.



B.



C.

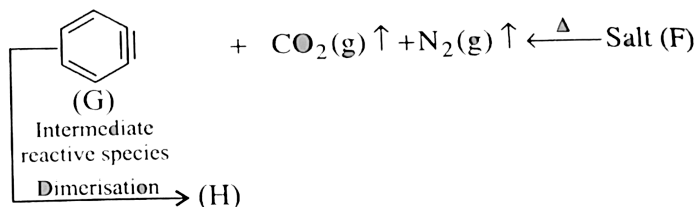
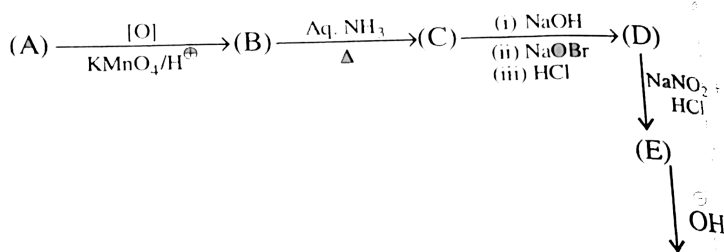


D.

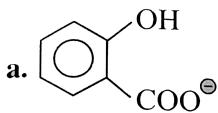
Answer: a

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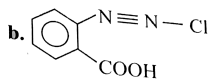
6. A hydrocarbon (A) ( $C_8H_{10}$ ) (E) is a steam volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.



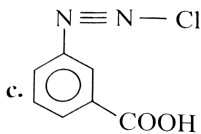
Compound (E) is:



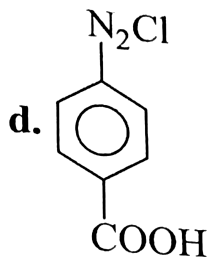
A.



B.



C.



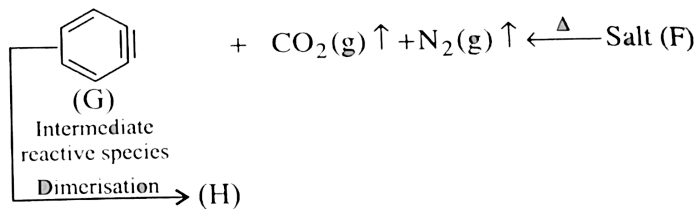
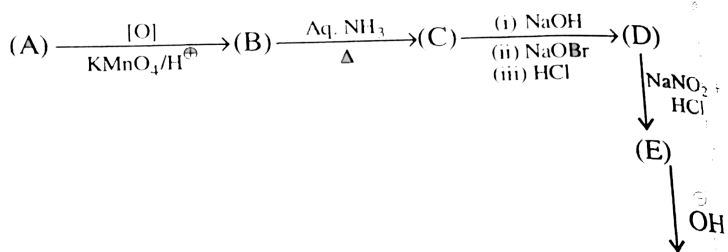
D.

**Answer: b**

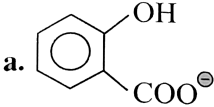
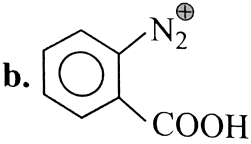
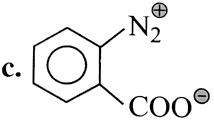
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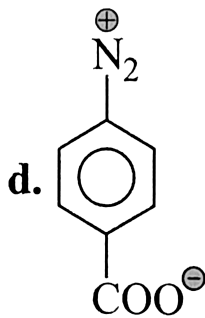
7. A hydrocarbon (A) ( $C_8H_{10}$ ) (E) is a steam volatile compound and on nitration gives two mononitro derivatives. (A) gives the following reactions.





Compound (F) is:

- A. 
- B. 
- C. 



D.

Answer: c

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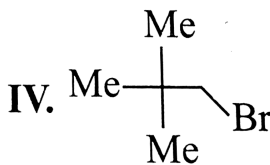
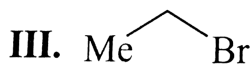
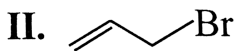
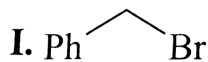
8.  $S_N$  reaction is given by these compounds, which have a nucleophilic group and a good leaving  $EWG$ . It should be stable after leaving with bonding pair of  $e^-$ 's and it should have high polarisability.

Nucleophilic aliphatic substitution reaction is mainly of two types  $S_N^{-1}$  and  $S_N^2$ .  $S_N^{-1}$  mechanism is a two step process. Reaction velocity of  $S_N^{-1}$  depends only on the concentration of the substrate. It proceeds via the formation of carbocation, optically active substrate gives  $(\oplus)$  and  $(o-)$  forms of the product.

In most of the cases, the product usually consists of 5 - 20 % inverted

and (95 - 80 %) racemised species. The more stable is the carbocation, the greater is the proportion of racemisation. In solvolysis reaction, the more nucleophilic is the solvent, the greater is the proportion of inversion.

Which of the following gives  $SN^1$  reaction?



Select the correct answer.

A. (I), (II) and (III)

B. (I) and (II)

C. (II), (III), and (IV)

D. (I), (III), and (IV)

**Answer: b**



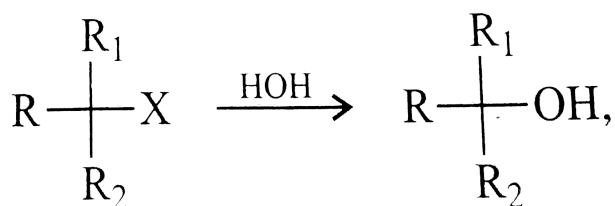
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9.  $S_N$  reaction is given by these compounds, which have a nucleophilic group and a good leaving  $EWG$ . It should be stable after leaving with bonding pair of  $e^-$ 's and it should have high polarisability.

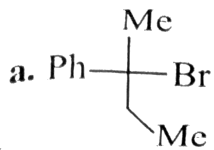
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In most of the cases, the product usually consists of 5 - 20 % inverted and (95 - 80 % ) racemised species. The more stable is the carbocation, the greater is the proportion of racemisation. In solvolysis reaction, the more nucleophilic is the solvent, the greater is the proportion of inversion.

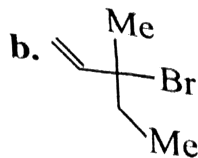
For the reaction



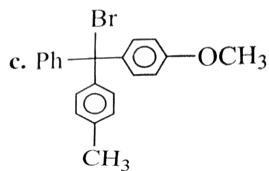
which substrate will give maximum racemisation ?



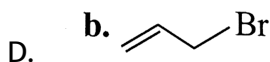
A.



B.



C.



Answer: c

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10.  $SN^2$  reaction is a bimolecular reaction which takes place by the formation of  $T.S.$  Velocity of the reaction depends on the concentration of the substrate as well as the nucleophile.

The reaction is favoured by strong  $Nu^{0-}$  and in the presence of polar

aprotic solvent, optically active halides give Walden inversion by  $S_N^2$  mechanism. The presence of hydroxyl group (atom) at  $\beta$ -C atom,

unsaturation at  $\beta$ -C and  $\left( \begin{array}{c} O \\ | | \\ - C - \end{array} \right)$  group at  $\alpha$ -C atom favor  $S_N^2$

mechanism.

Allyl halides and benzyl halides give  $S_N^1$  and  $S_N^2$  reactions. Allyl halides also give  $S_N^2$  mechanism. EDG at ortho- and para- positions in benzyl halides favors  $S_N^1$  mechanism, whereas EWG favors  $S_N^2$  mechanism.

Which of the following will give  $S_N^2$  reaction?



- A. (I) and (II)
- B. (I) and (III)
- C. (I), (II), and (III)
- D. All

**Answer: c**



**View Text Solution**

11.  $S_N2$  reaction is a bimolecular reaction which takes place by the formation of  $T.S.$  Velocity of the reaction depends on the concentration of the substrate as well as the nucleophile.

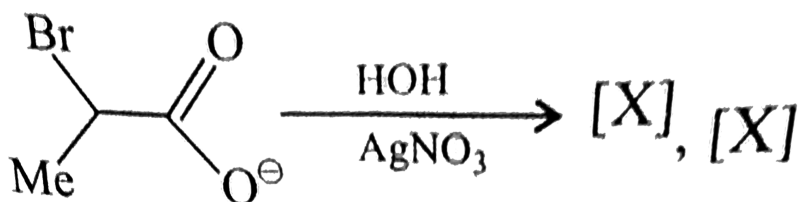
The reaction is favoured by strong  $Nu^{o-}$  and in the presence of polar aprotic solvent, optically active halides give Walden inversion by  $S_N2$  mechanism. The presence of hetero group (atom) at  $\beta$ -C atom,

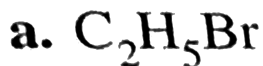
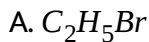
unsaturation at  $\beta$ -C and  $\left( \begin{array}{c} O \\ || \\ - C - \end{array} \right)$  group at  $\alpha$ -C atom favor  $S_N2$

mechanism.

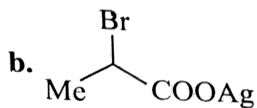
Allyl halides and benzyl halides give  $S_N1$  and  $S_N2$  reactions. Allyl halides also give  $S_N2$  mechanism.  $EDG$  at ortho- and para- positions in benzyl halides favors  $S_N1$  mechanism, whereas  $EWG$  favors  $S_N2$  mechanism.

In the reaction

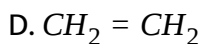




B.



C.



**Answer: c**

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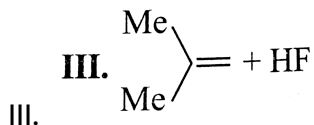
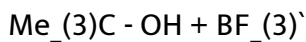
## Exercises Multiple Correct

1. Both *t*-butyl and ( $-SO_3H$ ) groups are used as a blocking group in certain synthesis of organic compounds. Which of the following statements are correct?

A. *t*-Butyl group is easily introduced by any of the variations of the Friedel-Crafts alkylation reaction.



B. *t*-Butyl group can be formed or deduced by using: I.  $\text{Me}_3\text{C} - \text{Cl} + \text{AlCl}_3$  II.



C. *t*-Butyl group can be easily removed under acidic conditions, because of the stability of tert-butyl cations.

D. *t*-Butyl group has advantage over a  $(-\text{SO}_3\text{H})$  group as a blocking group, because *t*-butyl group activates the ring to further *SE* reaction.

**Answer: (a, b, c, d)**

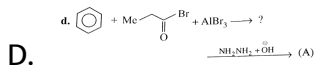
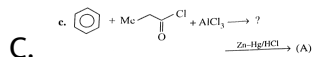
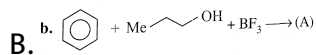
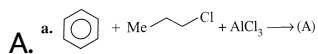


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2. Which compound in each of the following pairs will react faster in  $\text{SN}^2$  reaction with  $\text{HO}^-$  ?

a.  $\text{CH}_3\text{Br}$  or  $\text{CH}_3\text{I}$ .  $(\text{CH}_3)_3\text{CCl}$  or  $\text{CH}_3\text{Cl}$ .  $\text{CH}_2 = \text{CHBr}$  or

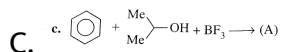
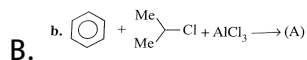
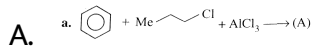
$\text{CH}_2 = \text{CH} - \text{CH}_2\text{Br}$

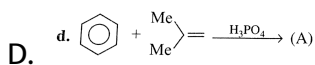


Answer: (c, d)

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3. Which of the following are the best methods for the preparation of cumene (A) ?

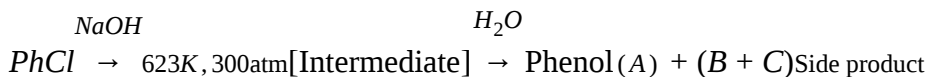




Answer: (b, c, d)

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4. In Dow's process for the process for the manufacture of phenol,  $PhCl$  is fused with  $NaOH$  at elevated temperature under pressure.



Which of the following statements are correct:

- A. Phenol is formed via the formation of benzyne intermediate.
- B. *p* - Phenyl phenol is also formed as by-product.
- C. Diphenylether is also formed as a by-product.
- D. Biphenylene is also formed as by-product.

Answer: (a,b,c)

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5. Which of the following statements are correct.

$o-$

A.  $OH$  is weaker nucleophile than  $H_2O$ .

B.  $R - \overset{\cdot\cdot}{S} \dots H$  is a stronger nucleophile than  $R - \overset{\cdot\cdot}{O} \dots H$  in polar protic solvent such as ethanol.

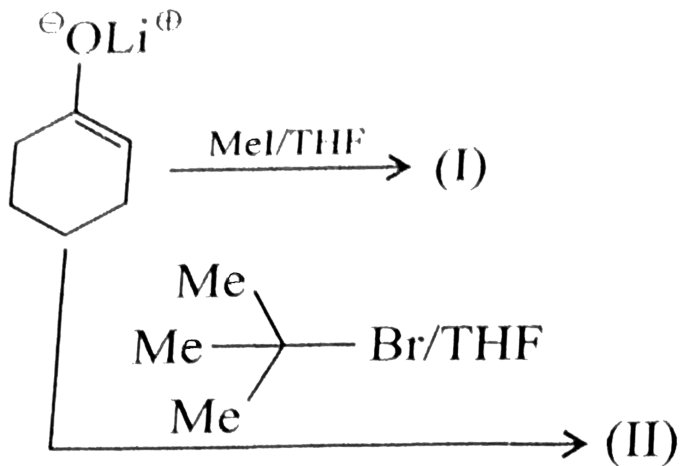
C.  $\overset{\cdot\cdot}{N}H_3$  is a weaker nucleophile than  $H_2\overset{\cdot\cdot}{H}$ .

D. a.  + Me-CH2-CH2-Cl + AlCl3 → (A) (I) is a stronger nucleophile than  $(C_2H_5)_3\overset{\cdot\cdot}{N}(II)$ .

**Answer: (b, c, d)**

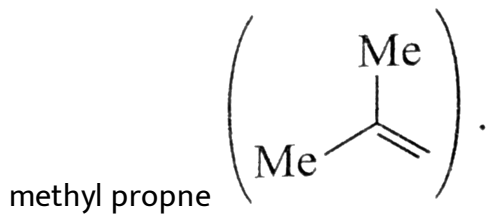
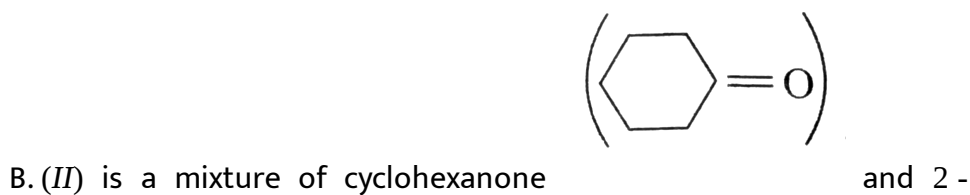
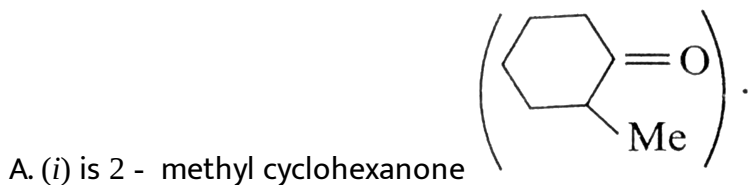


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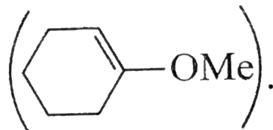


6.

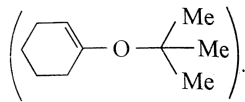
The products (I) and (II) are:



C. (I) is 1 - methoxy cyclohexene

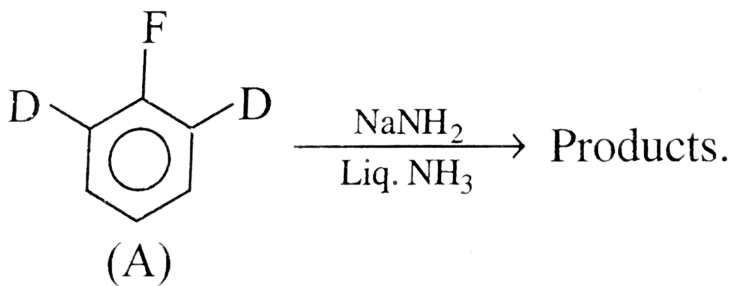


D. (II) is *t* - butoxy cyclohexene



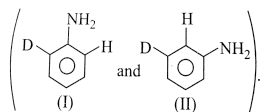
Answer: (a, b)

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Which of the following statements are correct for the above reaction?

A. The product is a mixture of



B. The product (II) is a cine-substitution product.

C. The reaction proceeds via benzene intermediate

D. The reaction is  $ArSN$  (addition-elimination).

**Answer: (a, b, c)**

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8. Refer to Q. No. (8) above. Which statements is/are correct ?

A. Both reactions (i) and (ii) proceed via  $SN^2$  mechanism.

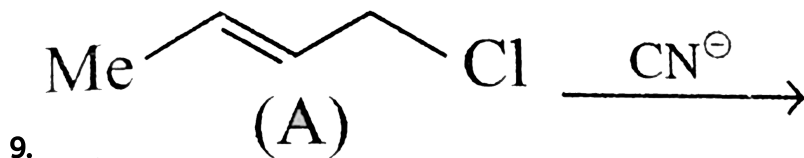
B. Both reactions (i) and (ii) proceed via  $SN^1$  mechanism.

C. Reaction (i) proceeds via  $SN^1$  and reaction (ii) via  $SN^2$  mechanism.

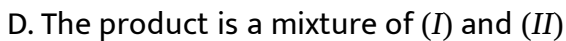
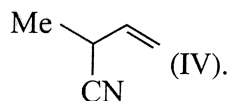
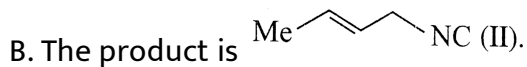
D. Reaction (i) proceeds via  $SN^2$  and reaction (ii) via  $SN^1$  mechanism.

**Answer: d**

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select the correct statements



**Answer: c**

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10. Refer to Q. No. 10. Which of the following statements are correct?



- A. Formation of (I) and (II) proceeds via  $SN^1$  mechanism.
- B. Formation of (I) and (II) proceeds via  $SN^2$  mechanism.
- C. Formation of (III) proceeds via  $SN^2$  mechanism.
- D. Formation of (IV) proceeds via  $SN^2$  mechanism with allylic rearrangement and is called  $SN^2$  - prime ( $SN^2$ ) mechanism.

**Answer: (c, d)**

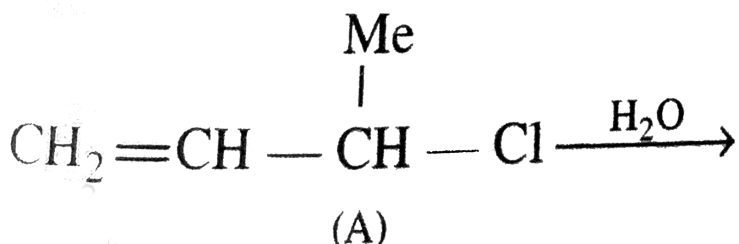
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11. Which of the following statements are correct about the reactivities of (I) *n*-propyl chloride and (II) allyl chloride.

- A. Rate of  $SN^1$  reaction of (I) > (II)
- B. Rate of  $SN^1$  reaction of (II) > (I)
- C. Rate of  $SN^2$  reaction of (I) > (II)
- D. Rate of  $SN^2$  reaction of (II) > (I)

Answer: (b, d)

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Which of the following statements are correct ?

- A. The product is  $\text{CH}_2=\overset{\text{Me}}{\underset{|}{\text{CH}}}-\text{OH}$  (I) (major)
- B. The product is a mixture of (I) and  $\text{HO}-\text{CH}_2-\text{CH}=\text{CH}-\text{Me}$  (II) (major).
- C. Allyl chloride is reactive both by  $\text{SN}^1$  and  $\text{SN}^2$  mechanism but more reactive by  $\text{SN}^1$  mechanism.
- D. Formation of (II) takes place by an allylic rearrangement.

Answer: (b, c, d)

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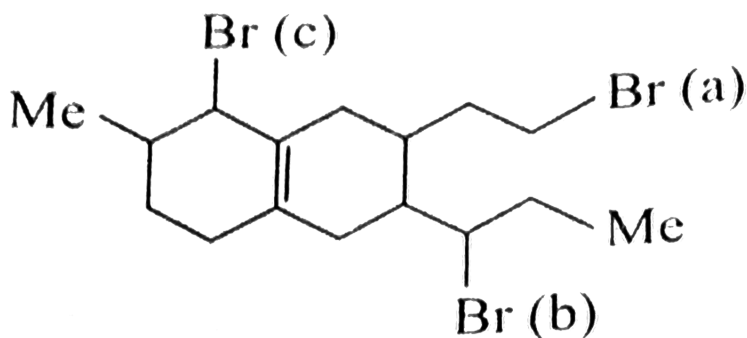
13. Which of the following statements are correct about Friedel-Crafts reaction?

- A. It is an aromatic electrophilic substitution reaction.
- B. The reaction intermediate is an  $\bar{e}$ 's deficient species.
- C. The reaction involves alkylation and acylation.
- D. A Lewis acid is used as a catalyst.

**Answer: (a, c, d)**

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14. Consider the following compound:



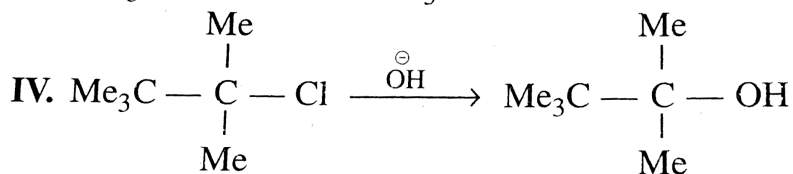
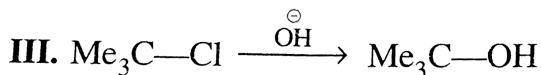
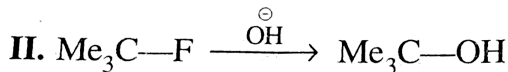
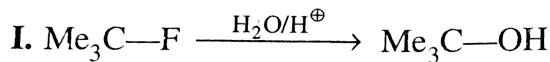
Which of the following statements are correct?

- A. Loss of *Br(a)* atom in dehydrobromination reaction results in the formation of the most reactive double bond towards hydrogentaion reaction.
- B. Removal of *Br(c)* atom results in the formaion of the most stable carbocation.
- C. The above compound conatins five asymmetric *C* atoms.
- D. The above compound does not show geometrical isomers.

**Answer: (a, b, c)**



15. Consider the following reactions:



Which of the statements are correct?

- A. Reaction (I) is faster than (II).
- B. Reaction (II) is faster than (I).
- C. Reaction (III) is faster than (IV).
- D. Reaction (IV) is faster than (III).

Answer: (a, d)

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16. Which of the following reactions are both stereospecific and stereoselective ?

A.  $SN^1$

B.  $SN^2$

C.  $E1$

D.  $E2$

**Answer: (b, d)**



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17. Which of the following reactions(s) is/are both non-stereospecific but stereoselective ?

A.  $SN^1$

B.  $E1$

C.  $E2$

D. *ElcB*

**Answer: b**



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**18.** Which of the following reactions(s) is/are stereospecific but non-stereoselective?

A.  $SN^2$

B.  $E1$

C.  $E2$

D. *ElcB*

**Answer: d**



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19. Which of the following reaction(s) is/are neither stereospecific nor stereoselective?

A.  $SN^1$

B.  $SN^2$

C.  $E2$

D.  $ElcB$

**Answer: a**



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20. The first steps of  $SN^1$  and  $SN^2$  reactions are, respectively

A. Both exothermic

B. Both endothermic

C. Endothermic and exothermic

D. Exothermic and endothermic



**Answer:** *c*

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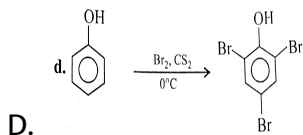
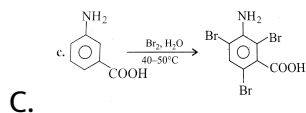
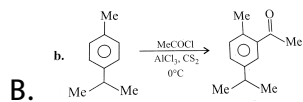
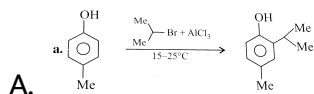
**21.** Which of the following statements are correct about *ElcB* reaction ?

- A. It proceeds via the formation of carbanion intermediate.
- B. Strong *EWG* and poor leaving groups favour the reaction.
- C. It is a unimolecular reaction with second order kinetics.
- D. When *D* is incorporated in the starting material by the solvent *EtOD* and the reaction is interrupted before completion, no *D* is found either in the substrate or in the product.

**Answer:** (*a, b, c*)

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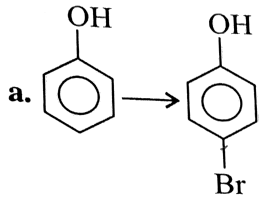
22. In which of the following reactions is the correct major product formed?



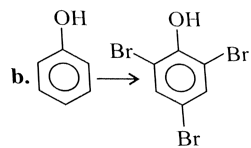
Answer: (a, b, c)

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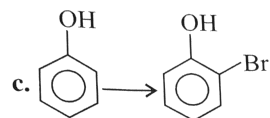
23. Which of the following syntheses could not be done without involving blocking positions on the ring?



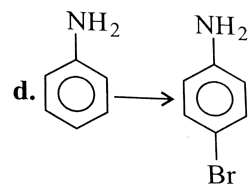
A.



B.



C.



D.

Answer: c



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24. Which of the following side chain reaction/s can be used to reduce the activating group such as ( - OH) or ( - NH<sub>2</sub>).

A. Benxolation

B. Acetylation

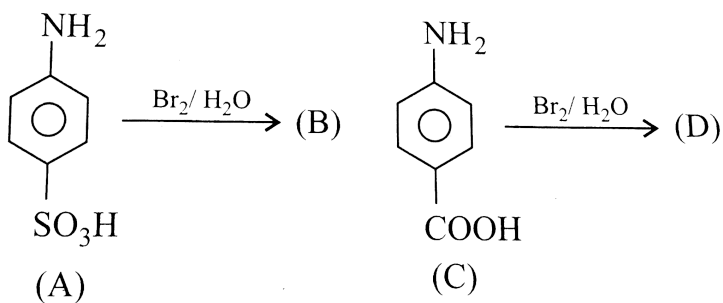
C. Tosylation

D. Sulphonation

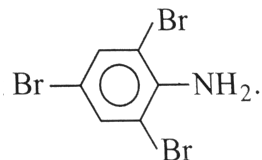
**Answer: (a, b, c)**

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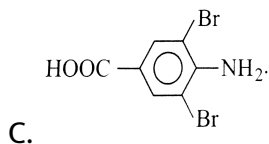
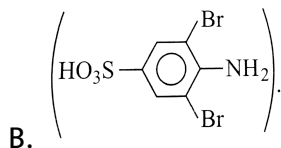
25. In the following reactions:



Which of the following statements are correct about the above reactions?



A. (B) and (D) are the same product



D. The above reaction is called ipso substitution.

**Answer: (a, b)**

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26. Which are the sources of phenol?

A. Cumene

B. Hydrolysi of benzene diaxonium salt

C. Middle oil of coal tar distillation

D. Reaction of diazonium salt with  $H_3PO_2$

**Answer:** (a, b, c)



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27. Which content (s) of middle oil separate on cooling?

A. Napthalene

B. Phenol

C. Benzene

D. Pyride

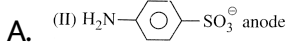
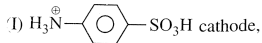
**Answer:** a



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28. Sulphanilic acid at  $pH = 2$  and  $12$  exists as....and migrates towards.....

Respectively.



B. (II) anode and (I) cathode

C. At both  $pH$ , only (I) and migrates towards the cathode

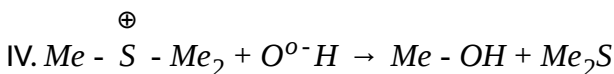
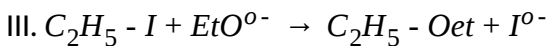
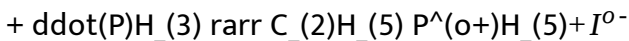
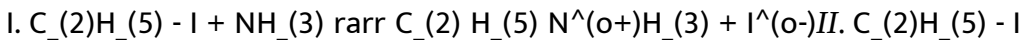
D. At both  $pH$ , only (II) and migrates towards the anode

Answer: a



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29. Consider the following reactions:



In which of the above reactions does not the rate of  $SN^2$  reactions increases in solvent polarity?

A. (I)

B. (II)

C. (III)

D. (IV)

**Answer: (a, b)**



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**30.** In which of the above reactions does the rate fo  $SN^2$  reactions decreases with an increases in solvent polarity?

A. (I)

B. (II)

C. (III)

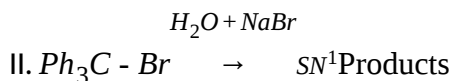
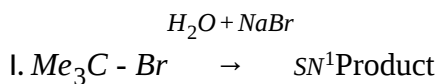


D. (IV)

Answer: (c, d)

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31. Consider the following reactions:



Which of the following statements are correct about the above reactions?

A. The products in reactions (I) and (II) are mixture of

$(\text{Me}_3 - \text{OH} + \text{Me}_3\text{Br})$  and  $(\text{Ph}_3\text{C} - \text{OH} + \text{Ph}_3\text{C} - \text{Br})$ , respectively.

B. The product in (I) is  $(\text{Me}_3\text{C} - \text{OH})$  and in (II) is

$(\text{Ph}_3\text{C} - \text{OH} + \text{Ph}_3\text{C} - \text{Br})$ .

C. The product in (I) is  $(\text{Me}_3\text{C} - \text{OH} + \text{Me}_3\text{C} - \text{Br})$  and in (II) is

$(\text{Ph}_3\text{C} - \text{OH})$ .

D.  $Ph_3C^{\oplus}$  is more stable than  $MeC^{\oplus}$ .

**Answer: (b, d)**



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**32.** When benzene sulphonic acid and *p*-nitrophenol are treated with  $NaHCO_3$ . The gases released, respectively, are:

A.  $SO_2, NO_2$

B.  $SO_2, NO$

C.  $SO_2, CO_2$

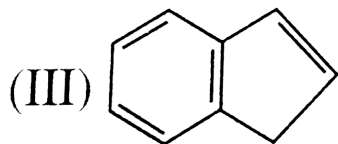
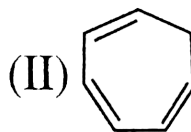
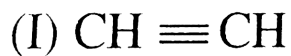
D.  $CO_2, CO_2$

**Answer: d**



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33. The decreasing order of  $pK_a$  value of the following is:



A. (III) > (I) > (II)

B. (II) > (I) > (III)

C. (I) > (III) > (II)

D. (I) > (II) = (III)

Answer: b

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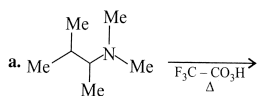
34. Among the following, which is/are correct?

- A. Both cyclopentadienyl anion and benzene are aromatic and have the same stability.
- B. Benzene is aromatic and more stable than cyclopentadienyl anion and it is non-aromatic
- C. Both cyclopentadienyl anion and benzene are aromatic but benzene is more stable than cyclopentadienyl anion.
- D. Cyclopentadienyl anion is more stable than benzene though both are aromatic

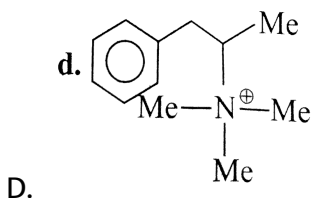
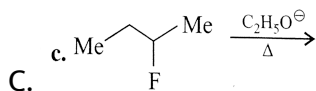
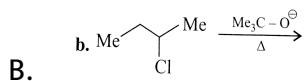
**Answer: c**

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**35. Which of the following will give Hofmann alkene?**



**A.**



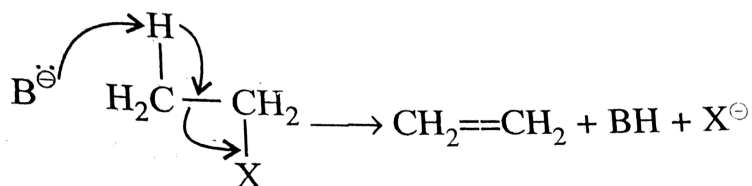
Answer: (a, b, c)

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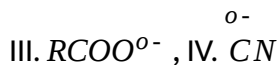
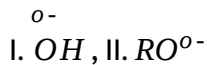
## Exercises Multiple Correct Answers Type

1.  $\beta$ - Elimination or anti-elimination reaction is carried out with base

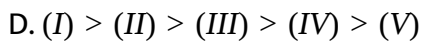
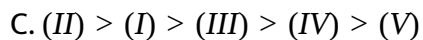
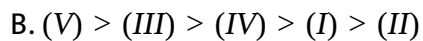
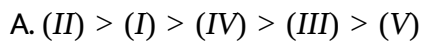
( $B^{o-}$ ) as shown below:



The following bases are used.



The decreasing order of reactivity for the above elimination is:

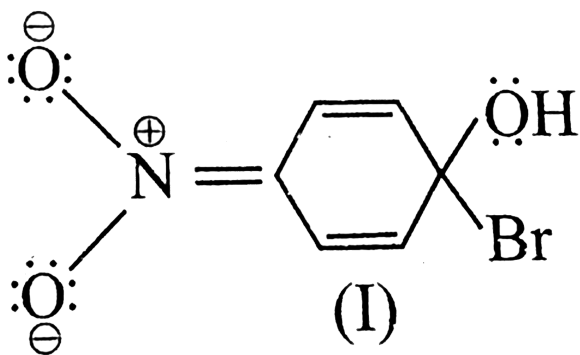


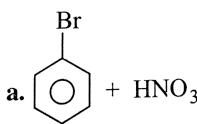
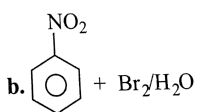
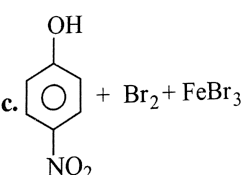
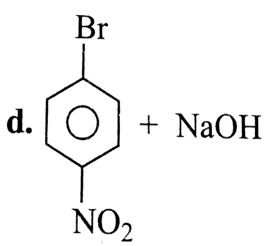
**Answer: a**



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2. Which combination of reactants will not give species (I) shown as a reactive intermediate?

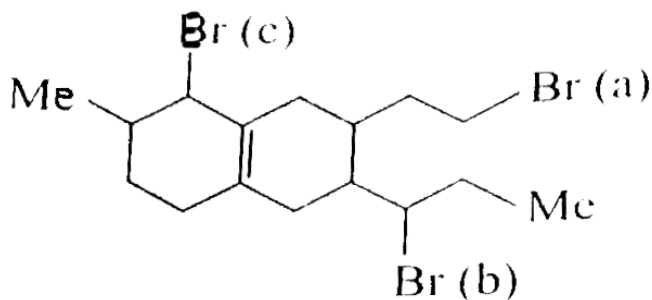


- A.  + HNO<sub>3</sub>
- B.  + Br<sub>2</sub>/H<sub>2</sub>O
- C.  + Br<sub>2</sub> + FeBr<sub>3</sub>
- D.  + NaOH

Answer: (a, b, c)

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3. Consider the following reactions:



Which of the statements are correct?

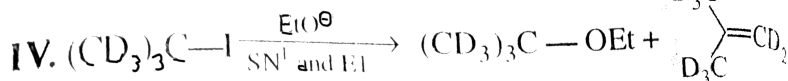
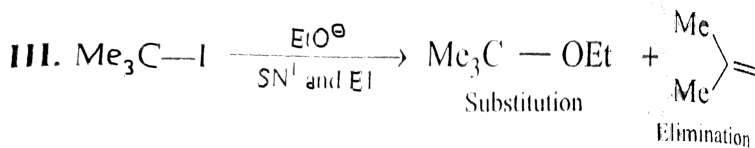
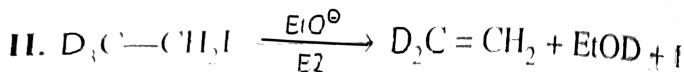
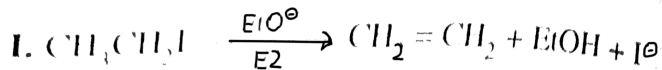
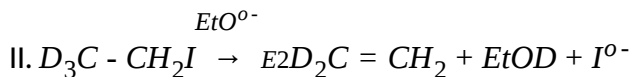
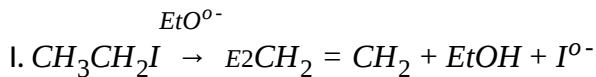
- A. Reaction (I) is faster than (II).
- B. Reaction (II) is faster than (I).
- C. Reaction (III) is faster than (IV).
- D. Reaction (IV) is faster than (III).

**Answer: (b, d)**

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4. Consider the following reactions:



Which of the following statement(s) is/are correct?

A. Reactions (I) and (II) show primary kinetic isotope effect, whereas reactions (III) and (IV) show 2° kinetic isotope effect.

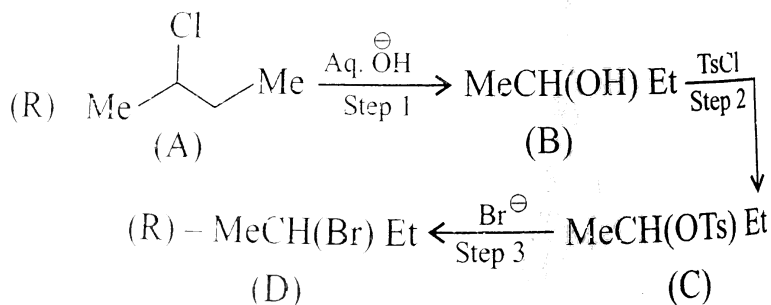
B. Reactions (I) and (II) show 2° kinetic isotope effect, whereas reactions (III) and (IV) show 1° kinetic isotope effect.

C. All reactions show 1° kinetic isotope effect.

D. All reactions show 2° kinetic isotope effect.



6. In the conversion of optically active (*R*) from of 2 - chlorobutane to (*R*) from of 2 - bromobutane the following sequence of reaction is carried out.



Direct conversion is not feasible since *Cl* cannot be replaced by *Br* in one step.

In which of the following steps does the inversion occur?

A. Step 1

B. Step 2

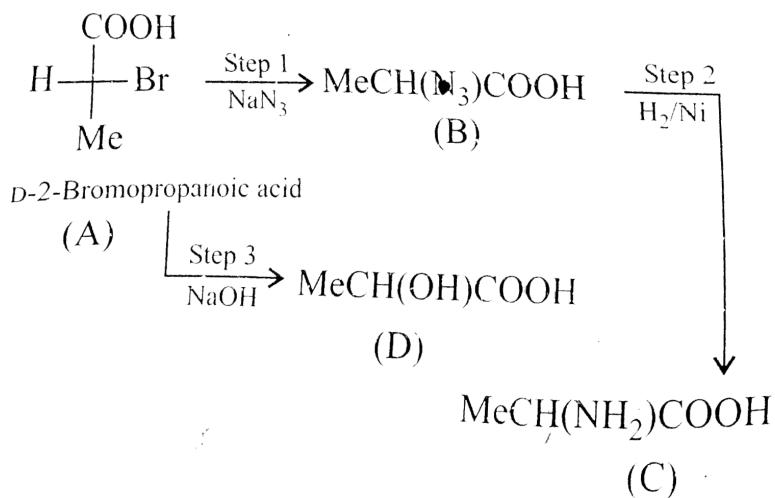
C. Step 3

D. Step 4

Answer: (a, c)

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7. In which of the following steps does inversion occur?



A. Step 1

B. Step 2

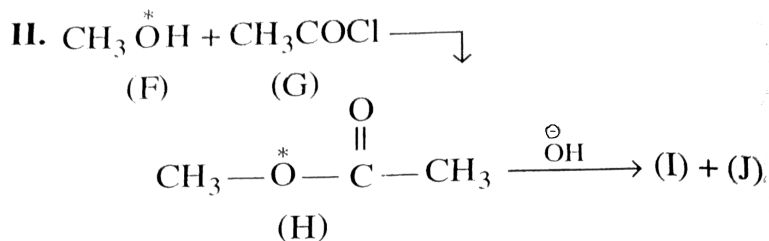
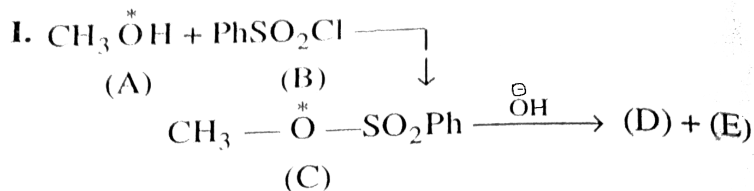
C. Step 3

D. None

Answer: (a, c)

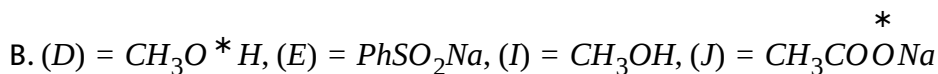
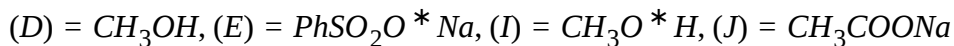
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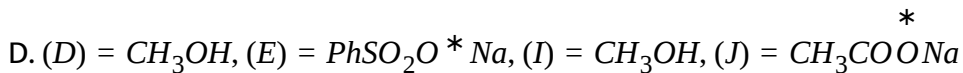
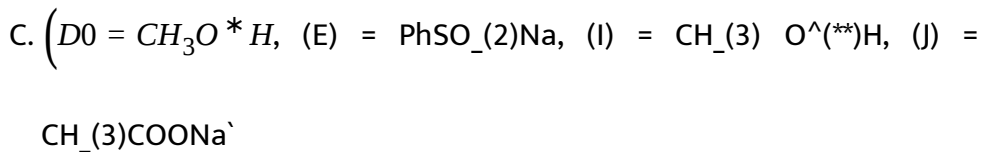
8. Consider the following reactions.



The products (D), (E), (I) and (J) are respectively.

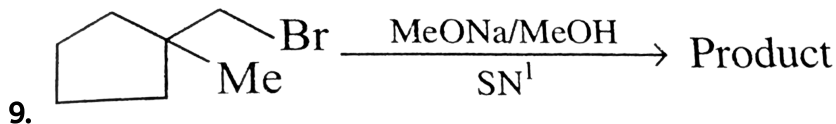
A.



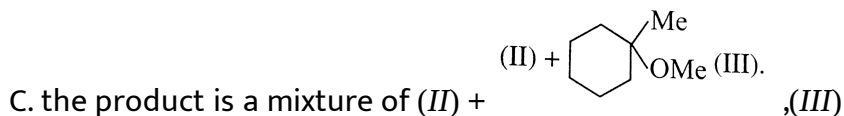
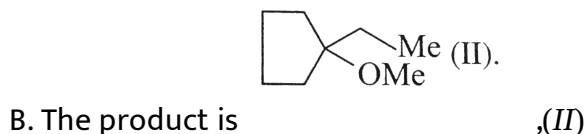


Answer: a

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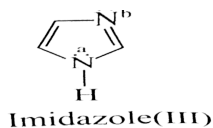
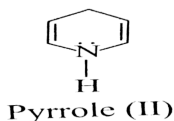
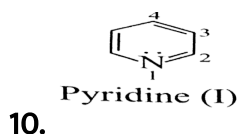
Which of the following statements are correct?



D. Product (II) is formed by 1,2Me shift and product (III) is formed by 1,2H<sup>+</sup> shift ring expansion.

Answer: (b, c, d)

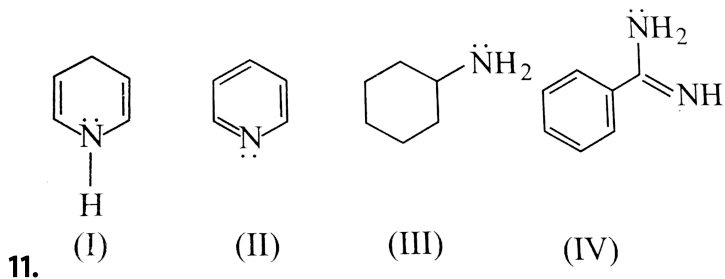
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Which of the following statements are true?

- A. (I) and (III) are modest Bronsted bases, whereas (II) is not.
- B. In (III), N<sup>a</sup> is more basic than N<sup>b</sup>.
- C. When (II) is protonated in the presence of a strong acid, protonation occurs at C - 2.
- D. All the nitrogen present in (I), (II) and (III) are sp<sup>2</sup> hybridised.

Answer: (a, c, d)



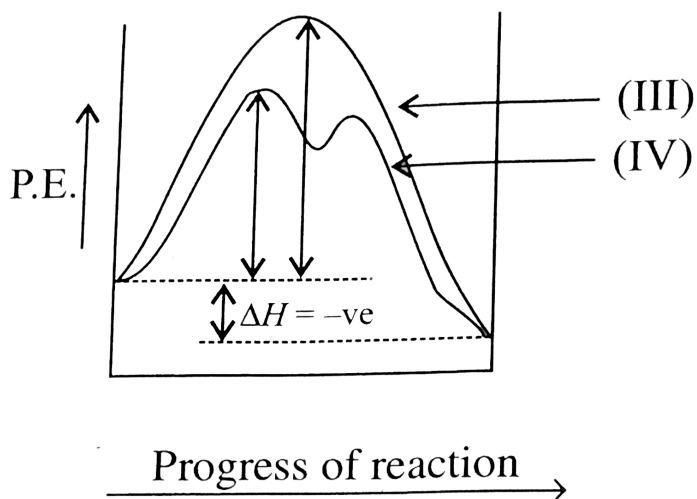
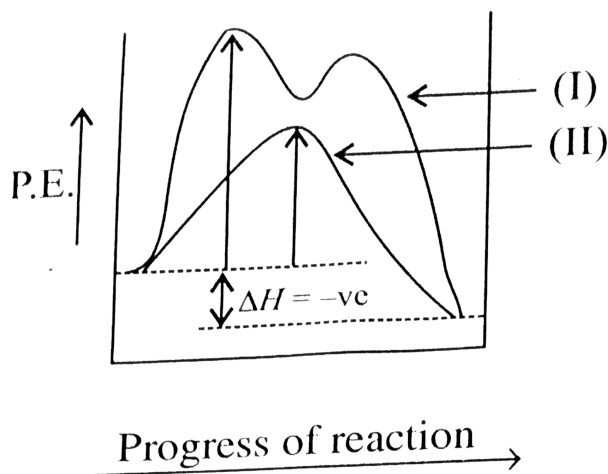
Which of the following statements are correct?

- A. (I) and (II) are aromatic and have equal basic strength.
- B. (I) is aromatic, (II) is anti-aromatic, but (II) is a stronger base than (I).
- C. The basicity order of above compounds is (IV) > (III) > (II) > (I)
- D. The conjugate acid of (IV) is more stabilized than the conjugate acid of (II).

**Answer: (c, d)**



12. Energy diagram of  $SN^1$  and  $SN^2$  reactions. The order of hydrolysis of  $RX$  by  $SN^1$  is  $3^\circ > 2^\circ > 1^\circ RX$  and by  $SN^2$  path is  $1^\circ > 2^\circ > 3^\circ RX$ .



A. (I) is Fig, (a) and (IV) in Fig, (b) represent  $SN^1$  reaction.

B. (II) in Fig. (a) and (III) in Fig(b) represent  $SN^2$  reaction.

C. Fig (a) and Fig (b) are the energy diagrams for  $1^\circ RX$  and  $3^\circ RX$ , respectively.

D. Fig (a) and Fig (b) are the energy diagram for  $3^\circ RX$  and  $1^\circ RX$ , respectively.

**Answer: (a, b, c)**

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## Exercises Single Correct

1. The decreasing order of the rate of nitration of the following compounds is

I. Benzene, II.  $C_6D_6$

III. Nitrobenzene, IV Chlorobenzene

A. (I) > (II) > (III) > (IV)

B. (I) > (II) > (IV) > (III)

C.  $(I) = (II) > (IV) > (III)$

D.

**Answer: c**

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2. The decreasing order of the rate of bromination of the following compounds is:

I.  $PhN^{\oplus}Me_3$ , II.  $Ph\overset{N}{CH_2}Me_3$

III.  $PhMe$ , IV.  $PhMe_2$

A.  $(I) > (II) > (III) > (IV)$

B.  $(IV) > (III) > (II) > (I)$

C.  $(III) > (IV) > (I) > (II)$

D.  $(III) > (IV) > (II) > (I)$

**Answer: b**





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3. The decreasing order of aromaticity for the following is

I. Benzene, II. Napthalene, III. Anthrance

A. (I) > (II) > (III)

B. (III) > (II) > (I)

C. (II) > (I) > (III)

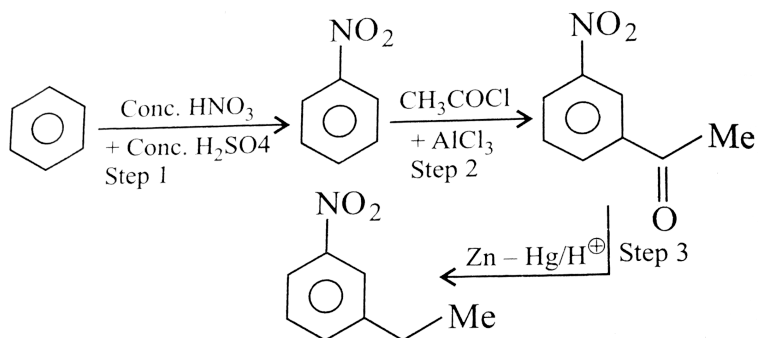
D. (II) > (III) > (I)

Answer: a



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4. In the following reaction, which of the following steps is wrong?



A. Step 1

B. Step 2

C. Step 3

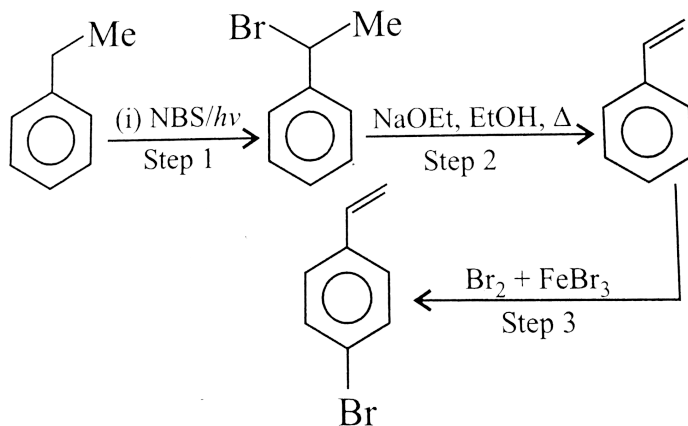
D. None

**Answer: b**



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5. In the following reaction, which of the following steps is wrong?



A. Step 1

B. Step 2

C. Step 3

D. None

**Answer: c**



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6. Nitration of xylene gives only one mono-nitro derivatives. Which xylene is it?

A. ortho

B. meta

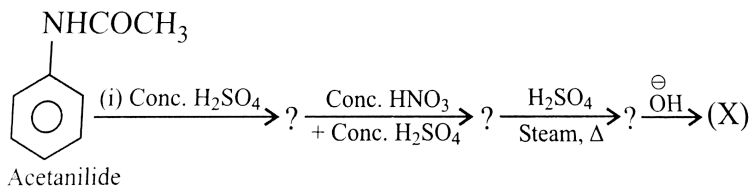
C. para

D. Both *o* and *p*

**Answer: c**

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7. The final product (X) in the following reaction is:  $\text{NHCOCH}_3$



A. 2 - Nitroaniline

B. 3 - Nitroaniline

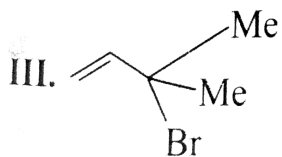
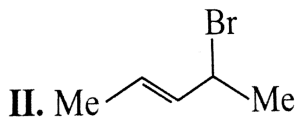
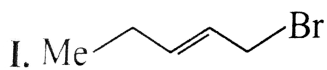
C. 4 - Nitroaniline

D. Sulphanilic acid

**Answer: a**

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8. Which of the following halides will be most reactive in  $S_N^2$  reaction and  $S_N^2$  reaction respectively?



A. (I), (II)

B. (II), (I)

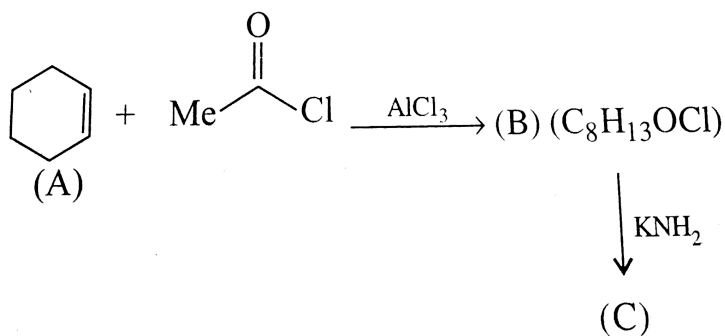
C. (I), (III)



D. (III), (I)

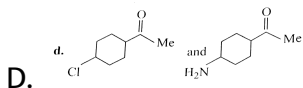
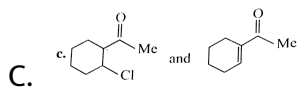
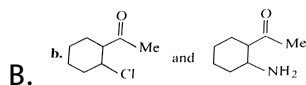
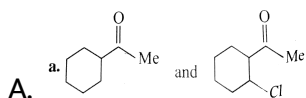
Answer: c

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9.

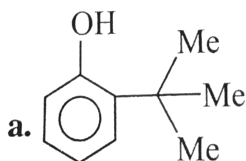
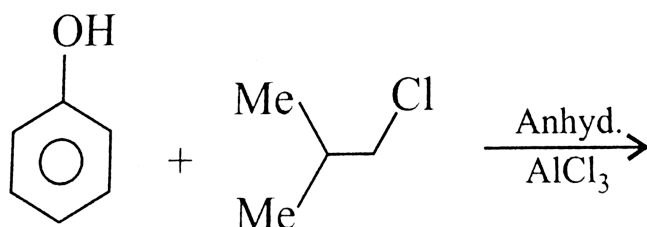
Compounds (B) and (C) respectively are:



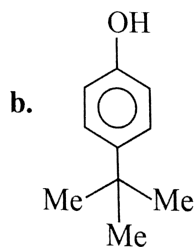
Answer: c

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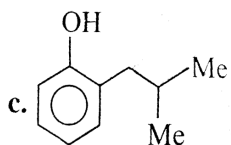
10. Give the major product of the following reaction.



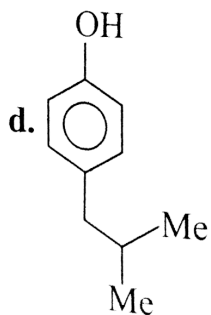
A.



B.



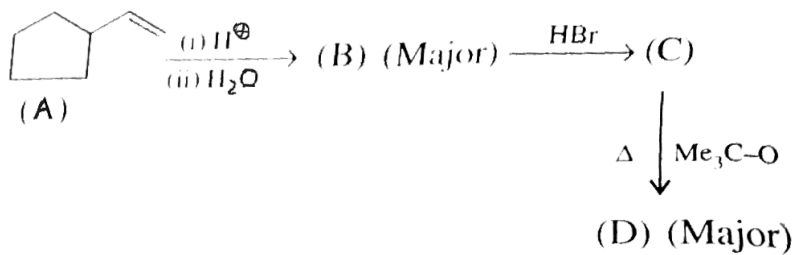
C.



D.

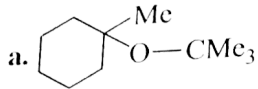
Answer: *b*

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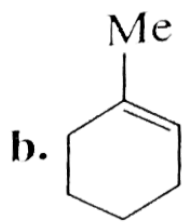


11.

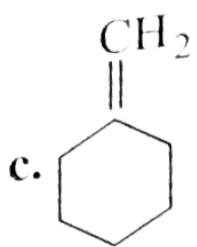
Compound (*D*) is:



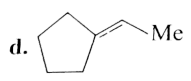
A.



B.



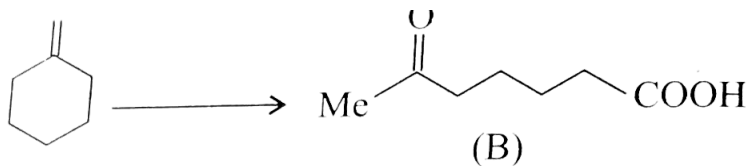
C.



D.

**Answer: c**

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12. (A)

The above conversion can be carried out by which process.

A. i.  $HBr$  + peroxide, ii.  $Me_3CO^{\ominus}$ ,  $\Delta$ , iii.  $O_3/H_2O$

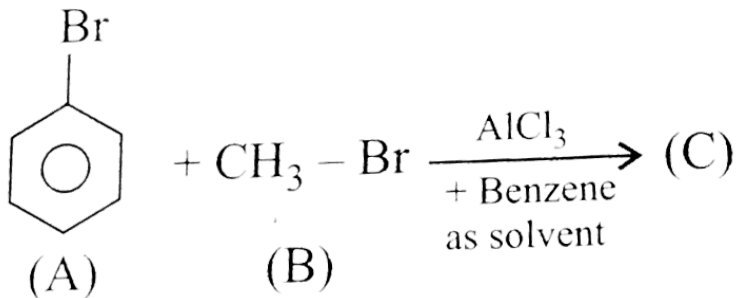
B. i.  $HBr$ , ii.  $C_2H_5O^{\ominus}$ ,  $\Delta$ , iii.  $O_3/H_2O$

C. i.  $HCl$ , ii.  $MeO^{\ominus}$ ,  $\Delta$ , iii.  $O_3/Zn$  acid

D.  $HCl$  + peroxide

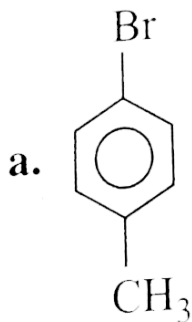
Answer: b

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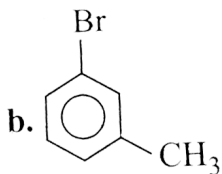


13.

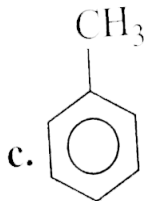
Product (C) is:



A.



B.



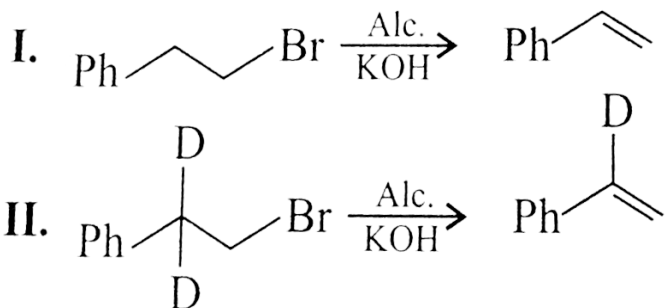
C.

D. 

Answer: c

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14. In the following reactions



the rate of reaction (I) and (II) are same. Both reactions proceed by which mechanism.

A. E1

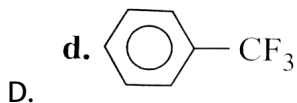
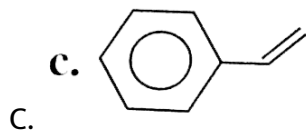
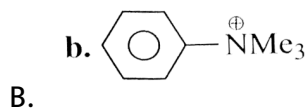
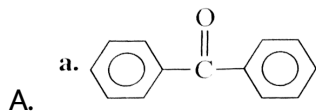
B. E2

C. E1cB

D. Anti-elimination

Answer: a

15. In which case will *SE* not be in *m* - position?

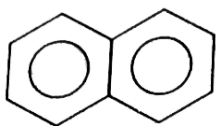


Answer: a

16. Which of the following has the highest dipole moment?

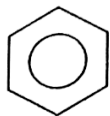


a.

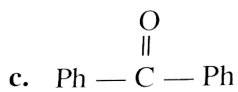


A.

b.

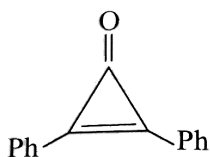


B.



C.

d.



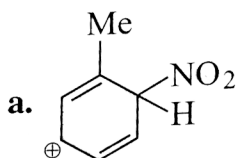
D.

**Answer: d**

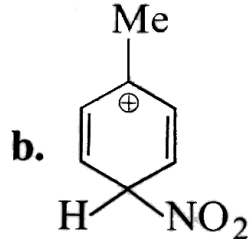


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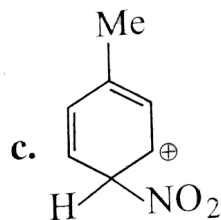
17. Which of the following is the most stable arenium or benzenium ion?



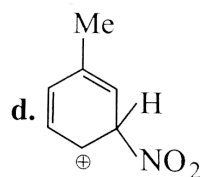
A.



B.



C.

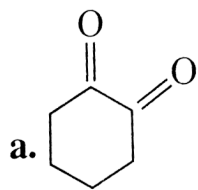


D.

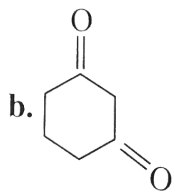
**Answer: b**

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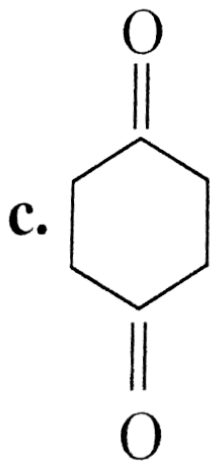
**18.** Which of the following ketonic compound is the least stable?



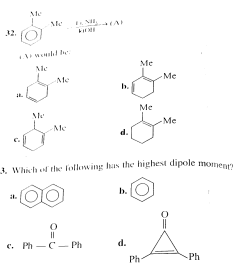
A.



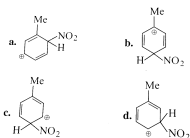
B.



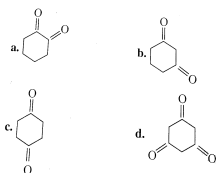
C.



34. Which of the following is the most stable arenium ion of benzene ring?



35. Which of the following ketonic compound is the least stable?

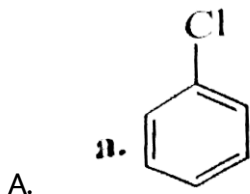


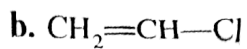
D.

**Answer: d**

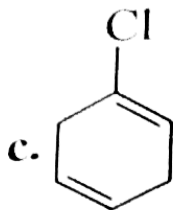
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19. Which of the following compounds will give curdy precipitate with  $AgNO_3$  solution?

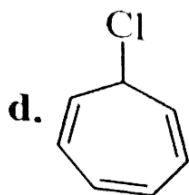




B.



C.



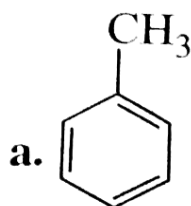
D.

Answer: *d*

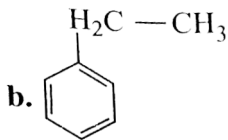


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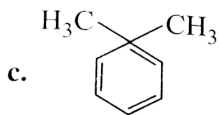
20. Which of the following compounds is the most reactive towards electrophilic substitution reaction?



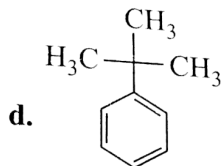
A.



B.



C.



D.

Answer: a

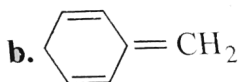


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21. Which of the following is the most stable species?



A.

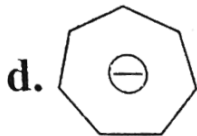


B.

C.



D.



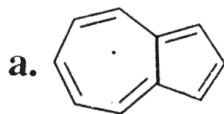
**Answer: a**



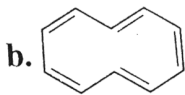
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22. Which of the following is not aromatic in nature?

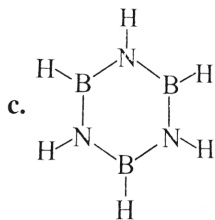
A.

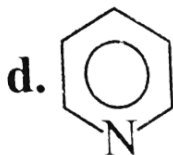


B.



C.



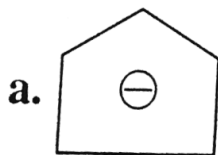


D.

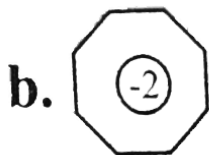
Answer: *b*

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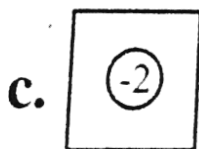
23. Which of the following species will be least stable?



A.



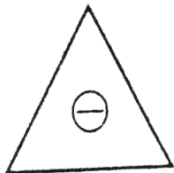
B.



C.



d.



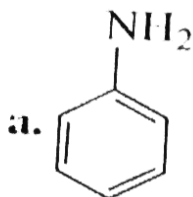
D.

Answer: *d*

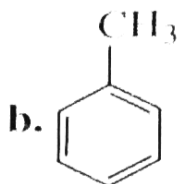


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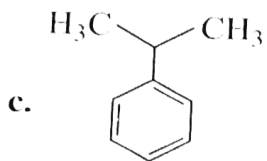
24. Which of the following compounds will undergo Friedel Crafts alkylation with faster rate?



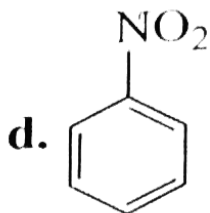
A.



B.



C.

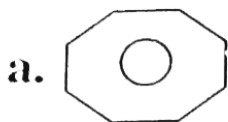


D.

Answer: *b*

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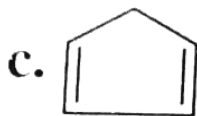
25. Which of the following is anti-aromatic in nature?



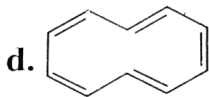
A.



B.



C.

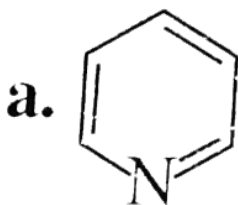


D.

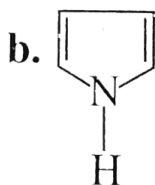
Answer: *b*

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26. Which of the following aromatic compounds is least reactive towards electrophilic substituions?

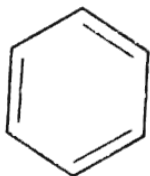


A.



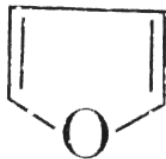
B.

c.



C.

d.



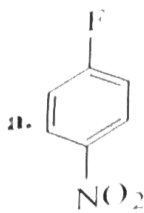
D.

Answer: a

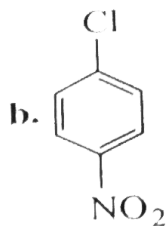


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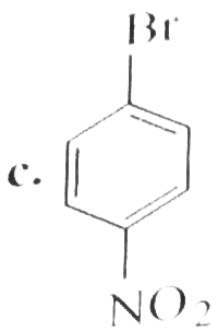
27. Which of the following compounds will show faster  $ArSN^2$  reactions?



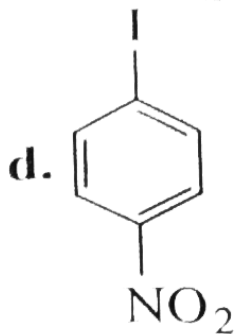
A.



B.



c.



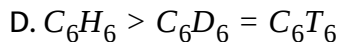
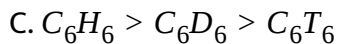
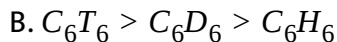
D.

Answer: a

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28. Which of the following is the order of the rate of reaction of  $C_6H_6$ ,  $C_6D_6$ , and  $C_6T_6$  towards sulphonation?

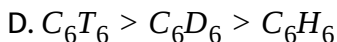
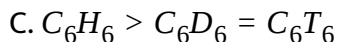
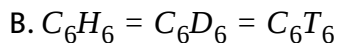
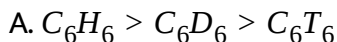
A. Same rates of reactions of  $C_6D_6$ ,  $C_6D_6$  and  $C_6T_6$ .



**Answer: c**

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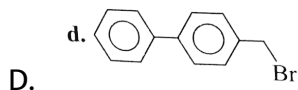
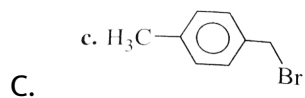
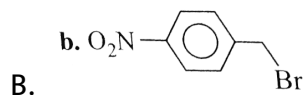
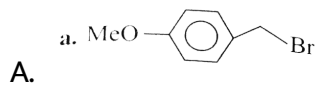
29. Which of the following is the correct order of the rate of reaction of  $C_6H_6$ ,  $C_6D_6$  and  $C_6T_6$  towards nitration?



**Answer: b**

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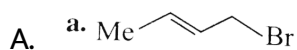
30. Which of the following will give  $SN^2$  mechanism?

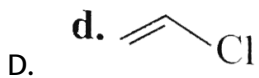
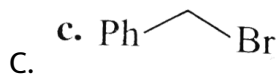
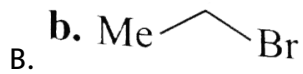


Answer: b

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31. Which of the following compounds gives  $SN^1$ ,  $SN^2$  and  $SN^2$  mechanisms?

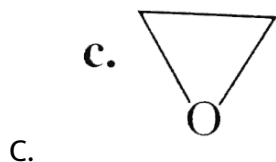
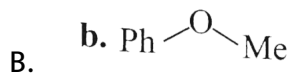
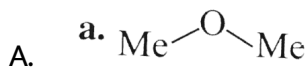




**Answer: a**

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**32.** Which of the following ether will always give  $S_N2$  mechanism in acidic as well as basic conditions?



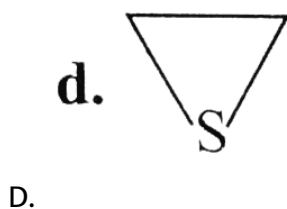
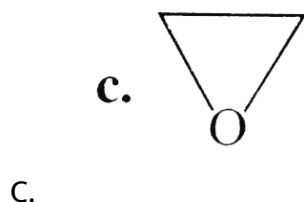
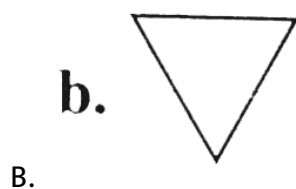
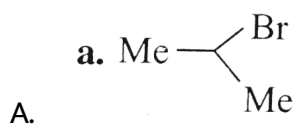
D. All



Answer: d

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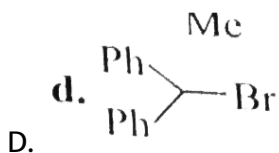
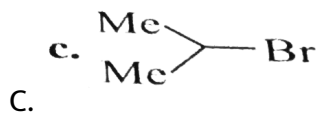
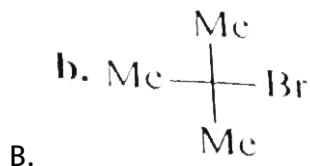
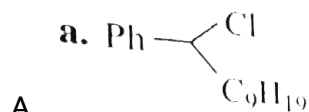
33. Which of the following ether will always give  $S_N2$  mechanism in acidic as well as basic conditions?



Answer: *d*

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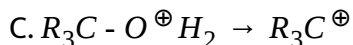
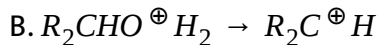
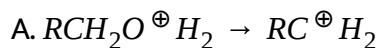
34. Which of the following substrates will give racemised product?



Answer: *a*

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35. The energy of activation is lowest for which reaction?



D. All have the same energy of activation.

Answer: c



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36. Isopentane on monochlorination gives....isomers and out of them .....

Are optically active.

A. 3, 1

B. 4, 2

C. 3, 2

D. 4, 1

**Answer: c**

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37. Propane on dichlorination gives.....isomers and out of them .....are optically active.

A. 4, 1

B. 3, 1

C. 4, 2

D. 3, 2

**Answer: a**

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38. A solution of (+)-2-chloro-2-phenylethane in toluene racemises slowly in the presence of a small amount of  $SbCl_5$ . Due to the formation

of:

A. Carbanion

B. Carbene

C. Free-radical

D. Carbocation

**Answer: d**



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**39.** The number of isomers for the compounds with molecular formula

$C_2BrClFI$  is:

A. 3

B. 4

C. 5

D. 6

**Answer: d**

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**40.** The decreasing basic order of the following compounds is:

i.  $NH_3$ , ii.  $PH_3$ , iii.  $AsH_3$ , iv.  $SbH_3$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

**Answer: a**

 [View Text Solution](#)

**41.** The decreasing nucleophilic order of the following compounds is:

i.  $NH_3$ , ii.  $PH_3$ , iii.  $AsH_3$ , iv.  $SbH_3$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

**Answer: b**

 [View Text Solution](#)

**42.** The decreasing nucleophilic order of the following compounds is:

i.  $NH_3$ , ii.  $PH_3$ , iii.  $AsH_3$ , iv.  $SbH_3$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

**Answer: b**

 [View Text Solution](#)

43. The decreasing basic order of the following is:

i.  $F^{O-}$ , ii.  $Cl^{O-}$ , iii.  $Br^{O-}$ , iv.  $I^{O-}$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

Answer: a

 [View Text Solution](#)

44. The decreasing nucleophilic order for the following compounds is:

i.  $F^{O-}$ , ii.  $Cl^{O-}$ , iii.  $Br^{O-}$ , iv.  $I^{O-}$

A. (i) > (ii) > (iii) > (iv)



B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

**Answer: b**

 [View Text Solution](#)

**45.** The decreasing leaving group order of the following is:

i.  $F^{0-}$ , ii.  $Cl^{0-}$ , iii.  $Br^{0-}$ , iv.  $I^{0-}$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

**Answer: b**

 [View Text Solution](#)

46. The decreasing basic order of the following is:

i.  $\overset{\ominus}{C}H_3$ , ii.  $NH_2^\ominus$ , iii.  $OH^\ominus$ , iv.  $F^\ominus$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

Answer: a



View Text Solution

47. The decreasing nucleophilic order of the following compounds is:

i.  $\overset{\ominus}{C}H_3$ , ii.  $NH_2^\ominus$ , iii.  $OH^\ominus$ , iv.  $F^\ominus$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

D. (ii) > (i) > (iv) > (iii)

**Answer: a**



[View Text Solution](#)

**48.** The decreasing leaving group order of the following compounds is:

i.  $\overset{\ominus}{C}H_3$ , ii.  $\overset{\ominus}{N}H_2$ , iii.  $\overset{\ominus}{O}H$ , iv.  $F^{\ominus}$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (i) > (iii) > (iv)

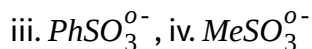
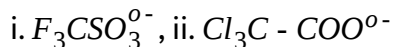
D. (ii) > (i) > (iv) > (iii)

**Answer: b**



[View Text Solution](#)

49. The decarising basic order of the following is:



A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (iv) > (iii) > (i)

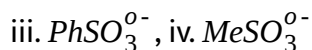
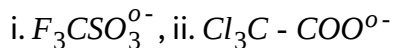
D. (iv) > (ii) > (i) > (iii)

Answer: c



View Text Solution

50. The decreasing nucleophilic order of the following is:



A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (ii) > (iv) > (iii) > (i)

D. (iv) > (ii) > (i) > (iii)

**Answer: c**

 [View Text Solution](#)

**51.** The decreasing leaving group order of the following is:

i.  $F_3CSO_3^{0-}$ , ii.  $Cl_3C - COO^{0-}$

iii.  $PhSO_3^{0-}$ , iv.  $MeSO_3^{0-}$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iii) > (i) > (ii) > (iv)

D. (i) > (iii) > (ii) > (i)

**Answer: d**



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52. The decreasing basic order of the following compounds is:

i.  $H_2O$ , ii.  $H_2S$ , iii.  $H_2Se$ , iv.  $H_2Te$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iii) > (i) > (ii) > (iv)

D. (iv) > (ii) > (i) > (iii)

**Answer: a**



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53. The decasing nucleophilic order of the following compounds is:

i.  $H_2O$ , ii.  $H_2S$ , iii.  $H_2Se$ , iv.  $H_2Te$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iii) > (i) > (ii) > (iv)

D. (iv) > (ii) > (i) > (iii)

**Answer: b**

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**54.** The decreasing leaving group order (fugacity order) of the following compounds is:

i.  $H_2O$ , ii.  $H_2S$ , iii.  $H_2Se$ , iv.  $H_2Te$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iii) > (i) > (ii) > (iv)

D. (iv) > (ii) > (i) > (iii)

**Answer: b**



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55. The decreasing basic order of the following is:

i.  $\text{O}^-(\text{C})\text{H}_3$ , ii.  $\text{O}^-(\text{O})\text{H}$ , iii.  $\text{CH}_3\text{COO}^-$ , iv.  $\text{H}_2\text{O}$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iii) > (i) > (ii) > (iv)

D. (iii) > (ii) > (iv) > (i)

**Answer: a**



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56. The decreasing nucleophilic order of the following compounds is:

i.  $\text{O}^-(\text{C})\text{H}_3$ , ii.  $\text{O}^-(\text{O})\text{H}$ , iii.  $\text{CH}_3\text{COO}^-$ , iv.  $\text{H}_2\text{O}$

A. (i) > (ii) > (iii) > (iv)



B. (iv) > (iii) > (ii) > (i)

C. (iii) > (i) > (ii) > (iv)

D. (iii) > (ii) > (iv) > (i)

**Answer: a**

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57. The decreasing fugacity order of the following compounds is:

i.  $\text{overset(o-)(C)H}_3$ , ii.  $\text{Overset(o - )(O)H}$ , iii.  $\text{CH}_3\text{COO}^{0-}$ , iv.  $\text{H}_2\text{O}$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iii) > (i) > (ii) > (iv)

D. (iii) > (ii) > (iv) > (i)

**Answer: b**

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58. The decreasing basic order of the following is:

i.  $\text{PhSO}_3^{0-}$ , ii.  $\text{C}_2\text{H}_5\text{SO}_3^{0-}$ , iii.  $\text{C}_2\text{H}_5\text{COO}^{0-}$ , iv.  $\text{CN}^{0-}$ , v.  $\text{OH}^{0-}$

A. (v) > (iv) > (iii) > (ii) > (i)

B. (i) > (ii) > (iii) > (iv) > (v)

C. (iv) > (v) > (iii) > (ii) > (i)

D. (i) > (ii) > (iii) > (v) > (iv)

Answer: a



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59. The decreasing nucleophilic order of the following compounds is:

i.  $\text{PhSO}_3^{0-}$ , ii.  $\text{C}_2\text{H}_5\text{SO}_3^{0-}$ , iii.  $\text{C}_2\text{H}_5\text{COO}^{0-}$ , iv.  $\text{CN}^{0-}$ , v.  $\text{OH}^{0-}$

A. (v) > (iv) > (iii) > (ii) > (i)

B. (i) > (ii) > (iii) > (iv) > (v)

C. (iv) > (v) > (iii) > (ii) > (i)

D. (i) > (ii) > (iii) > (v) > (iv)

**Answer: a**

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**60.** The decreasing fugacity order of the following compounds is:

i.  $\text{PhSO}_3^{0-}$ , ii.  $\text{C}_2\text{H}_5\text{SO}_3^{0-}$ , iii.  $\text{C}_2\text{H}_5\text{COO}^{0-}$ , iv.  $\text{CN}^{0-}$ , v.  $\text{OH}^{0-}$

A. (v) > (iv) > (iii) > (ii) > (i)

B. (i) > (ii) > (iii) > (iv) > (v)

C. (iv) > (v) > (iii) > (ii) > (i)

D. (i) > (ii) > (iii) > (v) > (iv)

**Answer: b**

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61. The decreasing basic order of the following is:

$\overset{\ominus}{\text{C}}\text{N}$ , ii.  $\overset{\ominus}{\text{O}}\text{H}$ , iii.  $\overset{\ominus}{\text{O}}\text{Me}$ , iv.  $\overset{\ominus}{\text{C}}\text{H}_3$ , v.  $\text{H}^{\ominus}$

A. (v) > (iv) > (iii) > (ii) > (i)

B. (i) > (ii) > (iii) > (iv) > (v)

C. (iv) > (v) > (iii) > (ii) > (i)

D. (i) > (ii) > (iii) > (v) > (iv)

Answer: c



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62. The decreasing nucleophilic order of the following compounds is:

$\overset{\ominus}{\text{C}}\text{N}$ , ii.  $\overset{\ominus}{\text{O}}\text{H}$ , iii.  $\overset{\ominus}{\text{O}}\text{Me}$ , iv.  $\overset{\ominus}{\text{C}}\text{H}_3$ , v.  $\text{H}^{\ominus}$

A. (v) > (iv) > (iii) > (ii) > (i)

B. (i) > (ii) > (iii) > (iv) > (v)

C. (iv) > (v) > (iii) > (ii) > (i)

D. (i) > (ii) > (iii) > (v) > (iv)

Answer: c



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63. The decreasing fugacity order of the following is:

$\overset{\ominus}{\text{C}}\text{N}$ , ii.  $\overset{\ominus}{\text{O}}\text{H}$ , iii.  $\overset{\ominus}{\text{O}}\text{Me}$ , iv.  $\text{CH}_3$ , v.  $\text{H}^{\ominus}$

A. (v) > (iv) > (iii) > (ii) > (i)

B. (i) > (ii) > (iii) > (iv) > (v)

C. (iv) > (v) > (iii) > (ii) > (i)

D. (i) > (ii) > (iii) > (v) > (iv)

Answer: d



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64. The decreasing basic order of the following is:

i.  $\text{Me}_2\text{N} - \text{Nme}_2$ , ii.  $\text{MeNH} - \text{NHMe}$

iii.  $\text{H}_2\text{N} - \text{NH}_2$ , iv.  $\text{NH}_3$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iv) > (iii) > (i) > (ii)

D. (iii) > (iv) > (ii) > (i)

**Answer: a**



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65. Decreasing nucleophilic order of the following is:

i.  $\text{Me}_2\text{N} - \text{Nme}_2$ , ii.  $\text{MeNH} - \text{NHMe}$

iii.  $\text{H}_2\text{N} - \text{NH}_2$ , iv.  $\text{NH}_3$

A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iv) > (iii) > (i) > (ii)

D. (iii) > (iv) > (ii) > (i)

**Answer: a**

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**66.** The decreasing fugacity order of the following is:

i.  $\text{Me}_2\text{N} - \text{Nme}_2$ , ii.  $\text{MeNH} - \text{NHMe}$

iii.  $\text{H}_2\text{N} - \text{NH}_2$ , iv.  $\text{NH}_3$

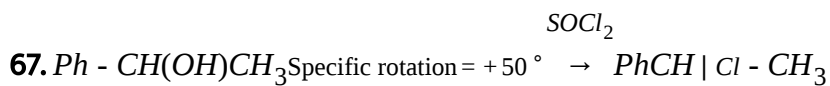
A. (i) > (ii) > (iii) > (iv)

B. (iv) > (iii) > (ii) > (i)

C. (iv) > (iii) > (i) > (ii)

D. (iii) > (iv) > (ii) > (i)

**Answer: b**

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What is the specific rotation of the product?

A.  $+50^\circ$

B.  $+60^\circ$

C.  $-60^\circ$

D. Zero

Answer: **b**

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68. Which of the following mechanisms does the reaction proceed?

A.  $\text{SN}^1$

B.  $\text{SN}^2$



C.  $SN^i$

D.  $E2$

**Answer: c**



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69. Which of the following acts as a nucleophilic?

A.  $Cl^{o-}$

B.  $OH^{o-}$

C.  $SO_2$

D. None

**Answer: a**



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70. The reaction is carried out in the presence of pyridine, Which of the following mechanisms does it follow?

A.  $SN^1$

B.  $SN^2$

C.  $SN^i$

D.  $E2$

**Answer: b**



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71. Which of the following acts as a leaving group?

A.  $OH^\theta$

B.  $Cl^\theta$

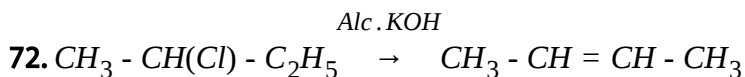
C.  $SO_2$

D.  $E2$

**Answer: c**



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The above reaction proceeds via *E1cB* mechanism. Which of the following statements is true about *E1cB* mechanism?

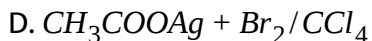
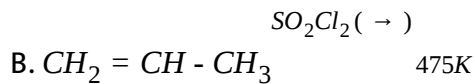
- A. It is second order and bimolecular.
- B. It is first order and unimolecular.
- C. It is first order and unimolecular.
- D. It is second order of unimolecular.

**Answer: d**



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73. Alkyl halides can be obtained by all methods except



**Answer: c**

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**74.** Neopentyl chloride on reaction with ethanolic KOH is likely to give

A. Neopentyl alcohol

B. Pentylene

C. 2 - Methyl - 2 butene

D. Undergo no reaction

**Answer: d**

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75. In the reaction  $CH_3C \equiv CNa^{\ominus} + (CH_3)_2CHCl \rightarrow$ , the product formed is:

- A. 4 - Methyl -2 - pentyne only
- B. Propyne
- C. Propyne and propylene
- D. Mixture of propene, propyne, and 4 - methyl -2 - pentyne

**Answer: d**

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76. In order to get ethanediol from bromoethane, the reagent used is :

- A. Sodium bisulphide
- B. Sodium sulphide

C. Potassium thiocyanate

D. Potassium sulphide

**Answer: a**

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77. For the preparation of chloroethane,

A.  $HCl$  gas is passed through ethanol.

B. Ethanol is treated with thionyl chloride in the presence of dimethyl amine or pyridine.

C. Ethyl sulphide is treated with hydrogen chloride.

D. Any of the above methods can be employed.

**Answer: b**

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78. X on treatment with sodium hydroxide followed by the addition of silver nitrate give white precipitate at room temperature which are soluble in  $NH_4OH$ . X can be :

- A. Chlorbenzene
- B. Ethyl bromide
- C. Benzyl chloride
- D. Vinyl chloride

**Answer: c**



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79. Which of the following cannot be used for the preparation of iodoform ?

- A. Acetone
- B. Methanol

C. Ethanol

D. Acetealdeyde

**Answer: b**



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**80.** Which of the following hladies is capable of exhibitin enantiomerism?

A. Ethyl chloride

B. Isopropyl bromide

C. sec-Butyl iodide

D. tert-Butyl chloride

**Answer: c**



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81. A suspension of  $\text{CaOCl}_2$  in water is heated with ethanol, the product formed is :

- A. Ethylene
- B. Ethanol
- C. Trichloromethane
- D. Chloroethane

**Answer: c**



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82. Pick up the correct statements about alkyl halides.

- A. They are associated with each other by  $H$  - bonds.
- B. They dissolve in water quickly.
- C. They dissolve easily in organic solvents.
- D. They do not contain any polar bonds in their molecules.

**Answer: c**

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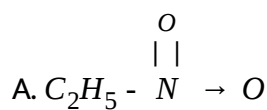
**83.** Which of the following is a geminal dihalide ?

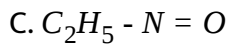
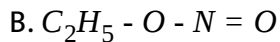
- A. Ethylene dibromide
- B. Propylidene chloride
- C. Isopropyl bromide
- D. None of the above

**Answer: b**

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**84.**  $C_2H_5I \xrightarrow{AgNO_2} X$ . Here X (major product) is :





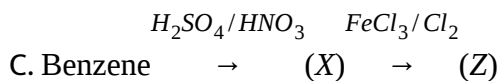
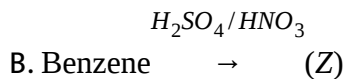
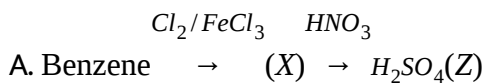
D. All of above

**Answer: c**



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85. Which of the following sequence would yield m-nitrochlorobenzene (Z) from benzene ?



D. All of these above will produce (Z)

**Answer: c**



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86.  $C_3H_7Cl \xrightarrow{KOH(alc.)} (A) \xrightarrow{Cl_2(g)} 770K(X)$ . (X) can be:

- A. Vinyl chloride
- B. Allyl chloride
- C. Ethyl chloride
- D. Ethyl isodide

**Answer: b**

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87. Which of the following alkyl halides undergoes the fastest  $S_N1$  reaction ?

- A. Methyl chloride
- B. Ethyl chloride

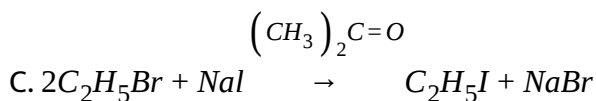
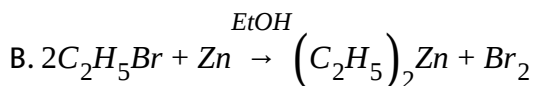
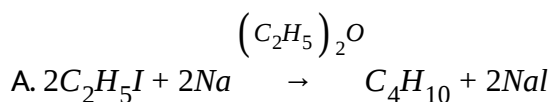
C. Isobutyl chloride

D. tert-Butyl chloride

**Answer: d**

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**88.** Choose the incorrect reaction.



D. All are correct reactions.

**Answer: b**

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89. In order to convert aniline into chlorobenzene the reagents needed are

A.  $CuCl$

B.  $NaNO_2/HCl$  and  $CuCl$

C.  $Cl_2/CCl_4$

D.  $Cl_2/AlCl_3$

Answer: b



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90. The reaction between chloral and chlorobenzene in  $H_2SO_4$  yields:

A. Chloretone

B. *p, p* - Dichlorodiphenyl trichloroethane

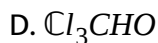
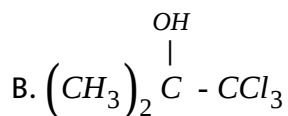
C. *o* - Chlorobenzaldichloride

D. Chloralphenylchloride

**Answer: b**

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91. Under the influence of air and light chloroform decomposes into



**Answer: c**

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92. When iodoform is heated with silver powder, the gaseous product formed is:

- A. Ethhene
- B. Ethyne
- C. Ethane
- D. Silver iodate

**Answer: *b***

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**93.** The reaction involving the treatment of benzene diazonium chloride with copper powder and HCl is treated as

- A. Sandmeyer's reaction
- B. Gattermann's reaction
- C. Ullmann's reaction
- D. Kolbe's reaction

**Answer: *b***



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94.  $C_6H_5Cl \xrightarrow{NaOH(aq.)} \xrightarrow{625\text{ K } 300\text{ atm...}}$  The product can be:

- A. Benzal
- B. Sodium benzote
- C. Benzol
- D. Sodium phenate

**Answer: d**

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95.  $(CH_3)_2CHI \xrightarrow{KOH} EtOHA \xrightarrow{SO_2Cl_2} 475KB$

The compound B in the sequence is

- A. Dimethyl sulphate
- B. 1, 2 - - Dichloro ethane

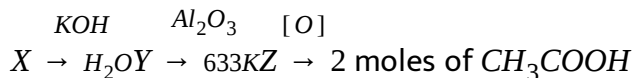
C. 3 - Chloro propene

D. 1 - Chloro-2 - iosopropane

**Answer: c**

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**96.** In given sequence of reaction predict X



A.  $CH_3CH_2CH(I)CH_3$

B.  $CH_3CH_2CH_2CH_2I$

C.  $CH_3CH(I)CH(I)CH_3$

D.  $CH_3CH(I)CH_2CH_2I$

**Answer: a**

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97. Iodoform can be prepared from:

- A. Isoamyl alcohol
- B.  $\alpha$  - Phenyl ethanol
- C. Isobutyl alcohol
- D.  $\beta$  - ethanol

**Answer: b**

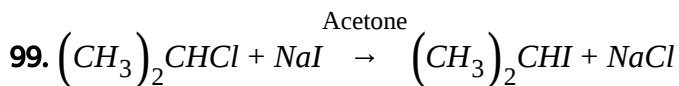
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98. When bromoethane is treated with potassium sulphide, the main product formed is

- A. Ethanethiol
- B. Ethanol
- C. Mustard gas
- D. Thioethyl ethane

**Answer: d**

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The above reaction is known as :

- A. Perkin's reaction
- B. Finkelstein's reaction
- C. Fitting reaction
- D. Sabatier and Senderan's reaction

**Answer: b**

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100. Isopropyl alcohol is heated with a suspension of bleaching powder  $(CaOCl_2)$  with water. The products are :

A. Etane and propane

B. Etyne and ethene

C. Trichloromethane and sodium acetate

D. Carbon tetrachloride

**Answer:** c

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**101.** Out of monochloro, monobromo and monoiodo derivatives of ethane, the least reactive compound towards nucleophilic substitutions will be :

A.  $C_2H_5Br$

B.  $C_2H_5Cl$

C.  $C_2H_5I$

D. All are equally reactive

**Answer: c**

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**102.** Which of the following reacting substances will not liberate ethyne gas ?

A.  $CH_3Cl$  and  $Ag$

B.  $CaC_2$  and  $H_2O$

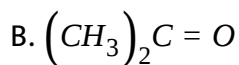
C.  $CHI_3$  and  $Ag$

D.  $CHCl_3$  and  $Ag$

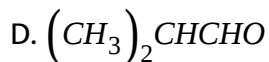
**Answer: a**

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**103.** Which of the following on reaction with chloroform will give chloretone ?



C. Chloral



**Answer: b**

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**104.** The product formed by the reaction between 2,2,2-trichloroethanal (chloral) and chlorobenzene in  $H_2SO_4$  is

A. Chloretone

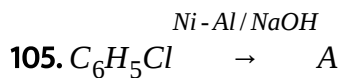
B. *DDT*

C. Chlorobenzaldichloride

D. Benzene sulphonic acid

**Answer: b**

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In this reaction A is :

- A. Phenol
- B. Sodium phenoxide
- C. Benzol
- D. Benzene

**Answer: d**

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106. A sample of chloroform being used as anaesthetic is tested by .

- A.  $AgNO_3(aq. )$
- B. Fehling's solution



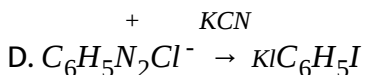
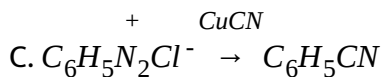
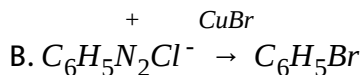
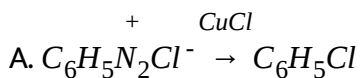
C.  $AgNO_3(ag.)$  after boiling with  $KOH$

D. Any of the above

**Answer: a**

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**107.** Which of the following is not an example of Sandmeyer's reaction?



**Answer: d**

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108. In order to prepare fluorobenzene from benzene diazonium chloride which of the following reagent is used

- A. Flouing
- B.  $HF$
- C. Hydrofluoroilic acid
- D. Fluroboric acid

**Answer: d**



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109. Ulimann reaction involes the use of the follwing reacants?

- A. Iodobenzene and sodium
- B. Benzene and copper
- C. Isodonenzene and copper powder
- D. Benzene diazonium chloride and  $Cu/HCl$

**Answer: c**

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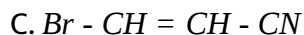
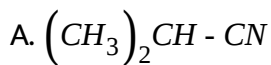
**110.** The yield of chlorobenzene obtained by reaction of phenols with  $PCl_5$  is less due to formation of

- A. *o* - Chlorophenol
- B. *p*- Chlorophenol
- C. Phosphorous oxychloride
- D. Triphenyl phosphate

**Answer: d**

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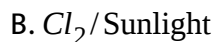
**111.** Identify C in the following series  $C_3H_7I \xrightarrow{KOH} AlcA \xrightarrow{NBS} \Delta B \xrightarrow{KCN} AlcC$ .



**Answer: b**

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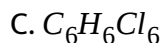
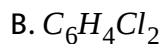
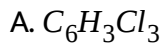
**112.** Lindane can be obtained by the reaction of benzene with



**Answer: b**

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113. IUPAC name of Gammexane is



D. Diphenylrichlorocethane

Answer: c



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114. Which of the following haldies has the least dipole moment?

A. 1, 2 - Dichlorobenzane

B. Dichloromethane4

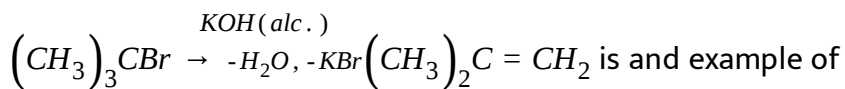
C. Trichloromethane

D. Ethyl chloride

Answer: *c*

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115. The chemical reaction :



A. Nucleophilic substitution

B. Electrophilic substitution

C. Free radical substitution

D.  $\beta$  - Elimination

Answer: *d*

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116. An ethereal solution of 4-Nitrochlorobenzene is treated with metallic sodium. The product formed is :

- A. Aminobenzene
- B. 4, 4 - Dinitrodiphenyl
- C. *p* - Chloroaniline
- D. Benzene diazonium chloride

**Answer: b**



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117. Ethanol  $\xrightarrow{P_4/I_2}$  *X*  $\xrightarrow{(i) KOH(alc)}$  *Y*  $\xrightarrow{(ii) HBr}$  *Z*

In this sequence of reactions *Y* is :

- A. Ethene
- B. Bromethane
- C. Ethanol

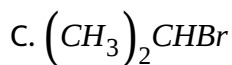
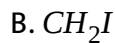
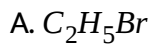
D. None

**Answer: b**



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**118.** Which of the following halides can yield ethane and also methane in a single step?



D. None

**Answer: b**



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119. A halide with formula  $C_6H_{13}Br$  gave two isomeric alkenes A and B with formula  $C_6H_{12}$ . On reductive ozonolysis of mixture of A and B following compounds were obtained  $CH_3COCH_3$ ,  $CH_3CHO$ ,  $CH_3CH_2CHO$  and  $(CH_3)_2CHCHO$ . The halide is

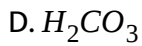
- A. 2 - Bromohexane
- B. 3 - Bromo -2 - methylpentane
- C. 2, 2 - Dimethyl -1 bromohexane
- D. Unpredictable

**Answer: b**

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120. Fire extinguisher is:

- A.  $CO_2$
- B.  $Cl_4$

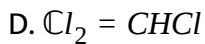
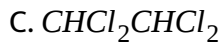


**Answer: b**



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**121.** Which of the following is called Westron ?



**Answer: c**



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122. Which of the following represents Freen?

- A. Acetylene tetrachloride
- B. Trichlorethylene
- C. Dichlordifluoromethane
- D. Ethylene dichloride

Answer: c



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123. Which of the following represents Westrosol?

- A.  $CHCl_3$
- B.  $CH_2Cl_2$
- C.  $CHCl_2CH_2Cl$
- D.  $Cl_2C = CHCl$

**Answer: d**



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**124.** Optically active isomers but not mirror images are called

- A. Enantiomers
- B. Mesomers
- C. Diastereomers
- D. Metamers

**Answer: c**



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**125.** The process of converging one enantiomer of an optically active compound into a racemic mixture is called:

- A. Resolution
- B. Inversion
- C. Epimerisation
- D. Racemisation

**Answer: d**

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**126.** The formation of an optically active compound from a chiral molecule is called:

- A. Asymmetric synthesis
- B. Resolution
- C. Walden inversion
- D. Epimerisation

**Answer: a**

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127. Which of the following reagents will be able to distinguish between allyl bromide and *n* - propyl bromide?

A. Aqueous  $AgNO_3$

B.  $NaOH$ ,  $AgNO_3$

C. *Alk.*  $KMnO_4$

D. Tollens reagent

**Answer: c**

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128. Rasching's process is employed for the commercial preparation of:

A. Ethyl chloride

B. Grignard's reagent

C. Chlorobenzene

D. Ethanol

**Answer: c**



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129. Vinyl chloride and ethyl chloride can be distinguished by

A. Lucas reagent

B.  $KOH, AgNO_3$

C.  $AgCl$

D.  $HCl/AgCl$

**Answer: b**



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130. The distillation of bleaching powder and acetone gives:

A.  $\text{CHCl}_3$

B. Chloral

C.  $\text{CH}_3\text{Cl}$

D.  $\text{Cl}_4$

**Answer: a**



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131. Chlorination of methane proceeds by

A. Electrophilic substitution

B. Nucleophilic substitution

C. Free radical mechanism

D. None of these



**Answer: c**



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**132.** Aryl halides are less reactive towards nucleophilic substitution reaction as compared to alkyl halides due to

- A. The formation of less stable carbonium ion
- B. Resonance stabilisation
- C. Larger-carbon-halogen bond
- D. Inductive effect

**Answer: b**



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**133.** How many chiral carbon atoms are present in 2,3,4-trichloropentane?

A. 3

B. 2

C. 1

D. 4

**Answer: b**

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**134.** Carbylamine test is performed in alc .  $KOH$  by heating a mixture of :

A. Chloroform and silver powder

B. Trihalogenated methane and a primary amine

C. An alkyl halide and a primary amine

D. An alkyl cyanide and a primary amine

**Answer: b**

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135. What happens when  $CCl_4$  is treated with  $AgNO_3$  solution ?

- A.  $NO_2$  is evolved
- B. A white precipitate of  $AgCl$  is formed
- C.  $CCl_4$  will dissolve in  $AgNO_3$  solution
- D. No reaction

**Answer: d**



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136. Among the following compounds the one that is most reactive towards electrophilic nitration is

- A. Toluene
- B. Benzoic acid
- C. Nitrobenzene

D. Benzene

**Answer: a**

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**137.** Slow oxidation of chloroform in air leads to

A. Formly chloride

B. Formic acid

C.  $\text{COCl}_2$

D. Trichloro acetic acid

**Answer: c**

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**138.** Reaction of *t* - butyl bromide with sodium methoxide produces

- A. Isobutane
- B. Isobutylene
- C. *t*- Butyl methyl ether
- D. Sodium tert-butoxide

**Answer: b**

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**139.** Which is most reactive towards  $Br_2$  in the presence of  $FeBr_3$ ?

- A. Anisole
- B. Benzene
- C. Bromobenzene
- D. Nitrobenzene

**Answer: a**

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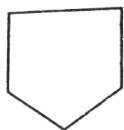
140. Which of following is most easily cleaved by  $HBr$ .

a.



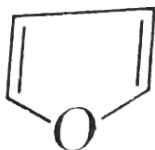
A.

b.



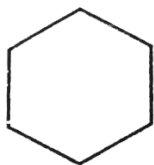
B.

c.



C.

d.



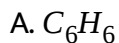
D.

Answer: a



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141. Which will react faster with *NBS*?



C. Toluene

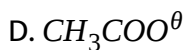
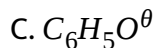
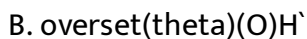
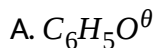
D. Cyclopropane

Answer: c



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142. Which is the most effective ion in an  $SN^2$  displacement on methyl bromide?



**Answer: a**



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**143.** Of the following four groups are *m* - directing when present on a benzene ring. The one which is not meta-directing is:

A.  $-COOH$

B.  $-NO_2$

C.  $-CHO$

D.  $-NH_2$

**Answer: d**



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**144.** The chemistry of benzene is characterised by which fo the following types of reaction?



A. Additomn

B. Elimination

C. Polymerisation

D. Substituion

**Answer: d**

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**145.** A reaction involving an aromatic nucleus is usually initiated by:

A. Free radicals

B. Molecules possessing a lone pair of electrons

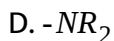
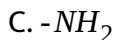
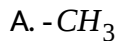
C. Nucliphiles

D. Eletrophiles

**Answer: d**

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146. Which of the following deactivates the aromatic nucleus?



Answer: *b*



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147. The experimental determined rate equation for the alkaline hydrolysis of  $RBr$  is given by:

$$\text{Rate} = k[RBr]^{1/2}[OH^-]$$

Which of the following statements is inconsistent with these observations?

A. The reaction is first order with respect to  $RBr$

- B. The reaction is second order overall
- C. The reaction process is false.
- D. The rate-determining step is bimolecular.

**Answer:** c

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**148.** Which of the following undergoes nitration most readily?

- A. Toluene
- B. Styrene
- C. Chlorobenzene
- D. phenol

**Answer:** c

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149. Which of the following reactions is not stereospecific?

A.  $SN^2$

B. Addition of  $Br_2$  to ethylene in  $CCl_4$

C.

D.

Answer: c



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150. Which of the following on reaction with acetylene ( $C\equiv CH$ ) produce gas(es) ?

A. (I), (II), and (III)

B. (I), (II), and (IV)

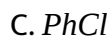
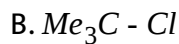
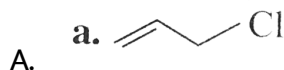
C. (II), (III), and (IV)

D. (I), (III), and (IV)

Answer: a

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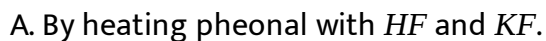
151. Which is least reactive towards nucleophilic substitution ( $S_N2$ )



Answer: c

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152. Fluorobenzene ( $C_6H_5F$ ) can be synthesized in the laboratory .



B. From aniline by diazotisation followed by heating the diazonium salt

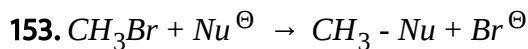
with  $BF_3$ .

C. By direct fluorination of benzene with  $F_2$  gas.

D. By reacting  $PhBr$  with  $NaF$  solution

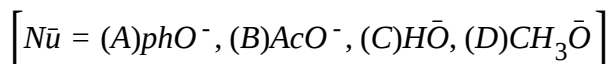
**Answer: b**

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The decreasing order of the rate of the above reaction with nucleophiles

$Nu^{\ominus}$  A to D is-



A.  $(D) > (C) > (A) > (B)$

B.  $(D) > (C) > (B) > (A)$

C.  $(C) > (D) > (A) > (B)$

D.  $(B) > (D) > (C) > (A)$

**Answer: c**

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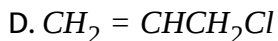
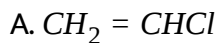
**154.** *HBr* reacts with  $CH_2 = CH - OCH_3$  under anhydrous conditions at room temperature to give:

- A.  $CH_3CHO$  and  $CH_3Br$
- B.  $BrCH_2CHO$  and  $CH_3OH$
- C.  $BrCH_2CH_2OCH_3$
- D.  $CH_3 - CH(Br) - OCH_3$

**Answer: d**

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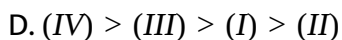
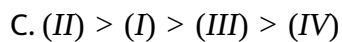
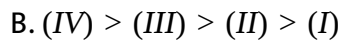
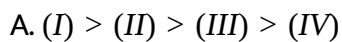
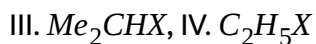
**155.** The halogen compound which most readily undergoes nucleophilic substitution is



**Answer: d**

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**156.** The correct decreasing order of  $SN^1$  reactivity of the following is:



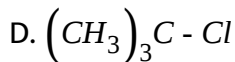
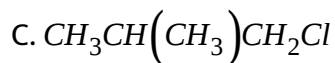
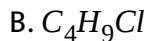
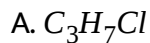


Answer: a



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157. Which of the following has the highest boiling point?



Answer: b



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158. Which of the following sequence of reaction (reagents) can be used for conversion of  $C_6H_5CH_2CH_3$  into  $C_6H_5CH = CH_2$  ?

A.  $\text{SOCl}_2, \text{H}_2\text{O}$

B.  $\text{SO}_2\text{Cl}_2, \text{alc. KOH}$

C.  $\text{Cl}_2/h\nu, \text{H}_2\text{O}$

D.  $\text{SOCl}_2, \text{alc. KOH}$

**Answer: b**

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**159.** The decreasing order of dipole moment of the following is:

I.  $\text{CH}_3\text{Cl}$ , II.  $\text{CH}_3\text{Br}$ , III.  $\text{CH}_3\text{F}$

A. (I) > (II) > (III)

B. (I) > (III) > (II)

C. (II) > (I) > (III)

D. (II) > (III) > (I)

**Answer: b**

160. Bottles containing  $C_6H_5I$  and  $C_6H_5 - CH_2I$  lost their original labels. They were labelled A and B for testing. A and B were separately taken in a test tube and boiled with  $NaOH$  solution. The end solution in each tube was made acidic with dilute  $HNO_3$  and then some  $AgNO_3$  solution was added. Substance B gave a yellow precipitate. Which one of the following statements is true for this experiment.

- A. Addition of  $HNO_3$  was unnecessary
- B. (A) was  $PhI$
- C. (A) was  $PhCH_2I$
- D. (B) was  $PhI$

**Answer: b**

161. One mole of 1,2-dibromopropane on treatment with X moles of  $\text{NaNH}_2$  followed by treatment with ethyl bromide gave a pentyne. The value of X is:

A. 1

B. 2

C. 3

D. 4

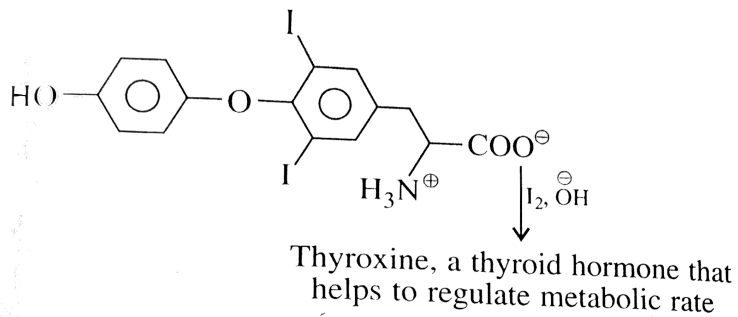
**Answer: c**



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Exercises Single Correct Answer Type

1. In the reaction is:



Thyroxine is:

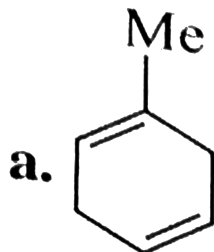
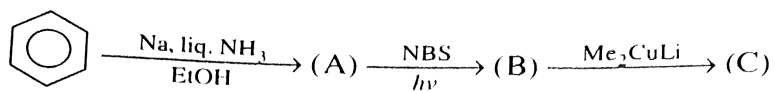
- A.
- B.
- C.
- D.

Answer: *b*

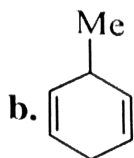


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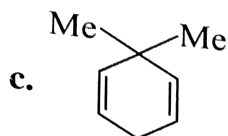
2. The final product (C) in the following reactions is:



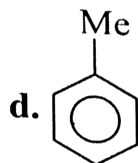
A.



B.



C.

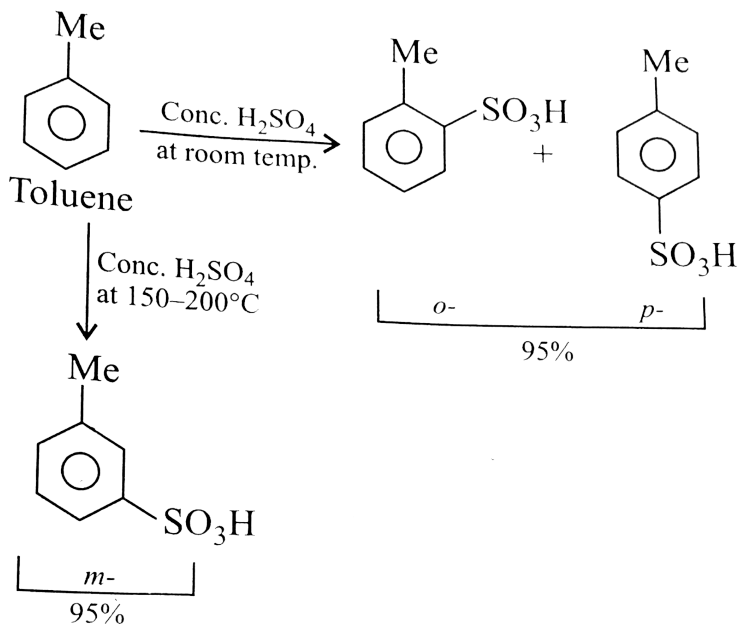


D.

Answer: *b*



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3.

Which of the following statements is wrong about the reactions?

- A. At lower temperature, the reaction is kinetically controlled and *o/p* directive effects of the (*Me*) group operate.
- B. At a higher temperature, the reaction is thermo-dynamically controlled, and longer reaction times are employed for equilibrium to be reached. The most stable form of *m*-toluene sulphonic acid is obtained.
- C. (*Me* - ) group is activated by +*I* effect, and *o* - , *p* - directing.

D. ( $Me^-$ ) group is deactivating by hyperconjugation and is  $m$ -directing.

**Answer: d**

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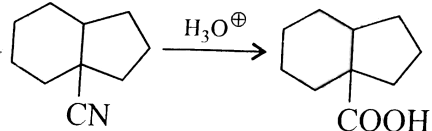
4. In the following reaction, the final product can be prepared by two paths (I) and (II).

Which of the following statements is correct?

Path (I)



Path (II)  $\downarrow$  NaCN



A. Path (I) is feasible.

B. Path (II) is feasible.

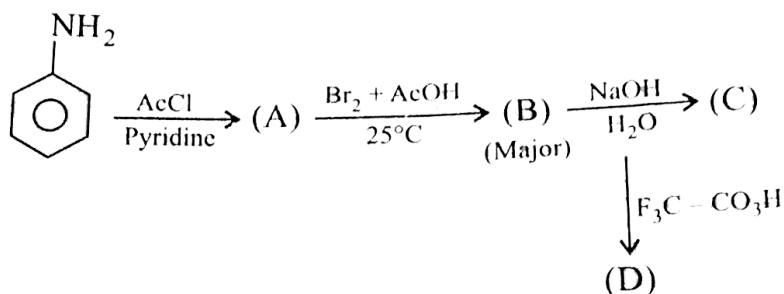


C. Both paths are feasible.

D. Neither of the two paths is feasible

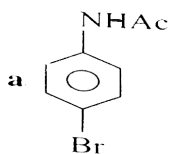
Answer: a

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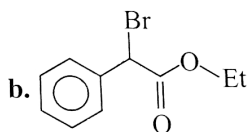


5.

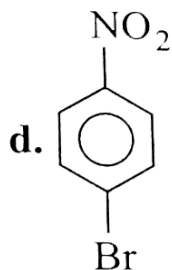
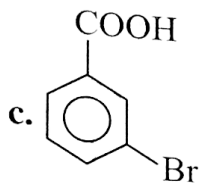
(D) would be:



A.

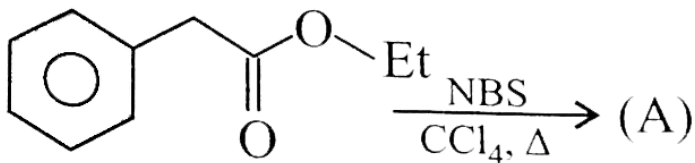


B.



Answer: *d*

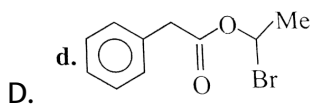
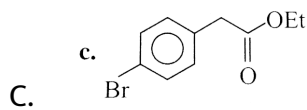
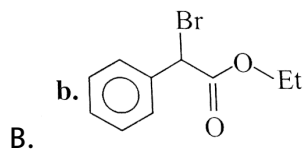
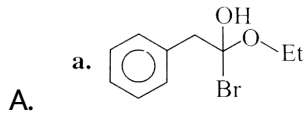
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(A) would be:

6.

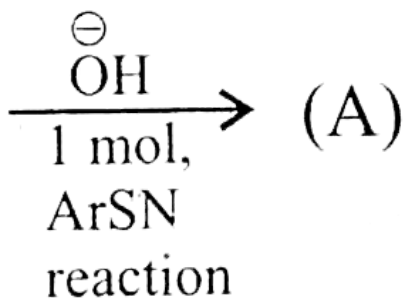
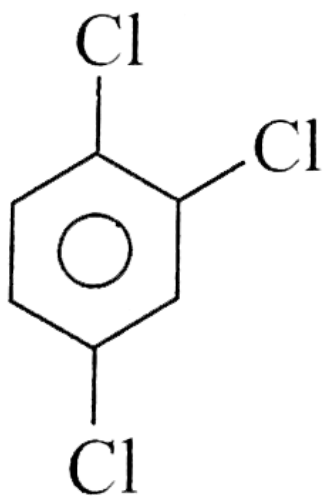
(A) would be:



**Answer: b**

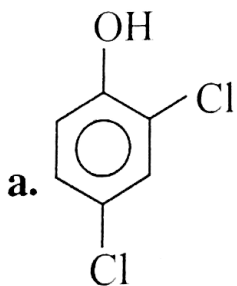


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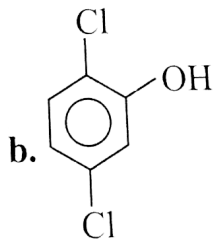


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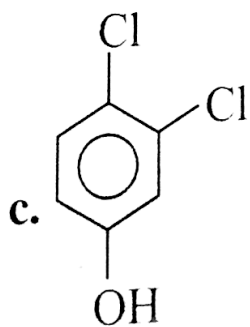
(A) would be:



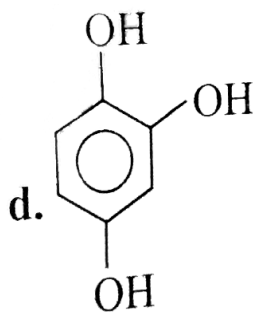
A.



B.



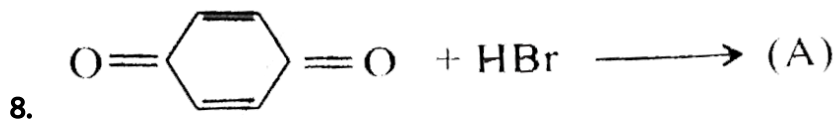
c.



D.

Answer: a

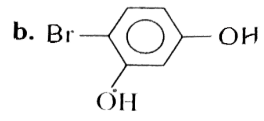
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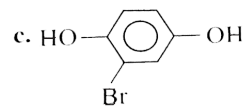
(A) would be:



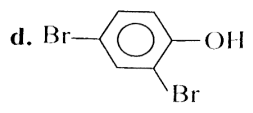
A.



B.



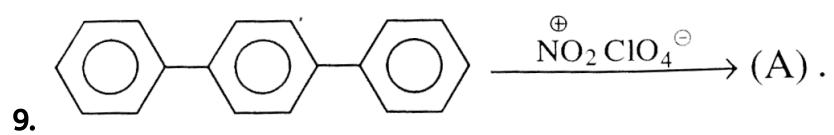
C.



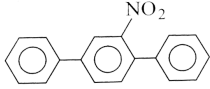
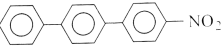
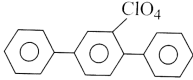
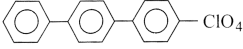
D.

Answer: c

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(A) would be:

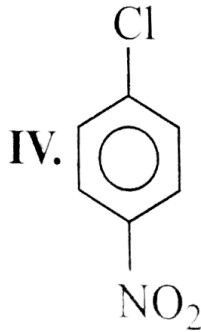
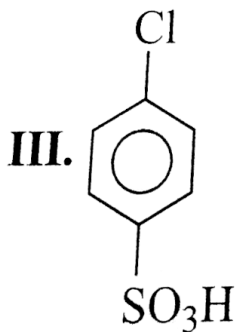
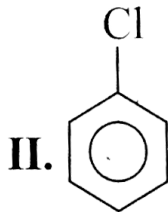
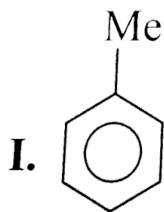
- A. a.  c1ccc(cc1)-c2cc(ccc2)[N+](=O)[O-]
- B. b.  c1ccc(cc1)-c2cc(ccc2)[N+](=O)[O-]
- C. c.  c1ccc(cc1)-c2cc(ccc2)Cl([O-])[O+]
- D. d.  c1ccc(cc1)-c2cc(ccc2)Cl([O-])[O+]

**Answer: a**



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10. The decreasing order of  $ArSN$  reaction with  $C_2H_5O^- / EtOH$  is:



A. (I) > (II) > (III) > (IV)

B. (IV) > (III) > (II) > (I)

C. (III) > (IV) > (II) > (I)

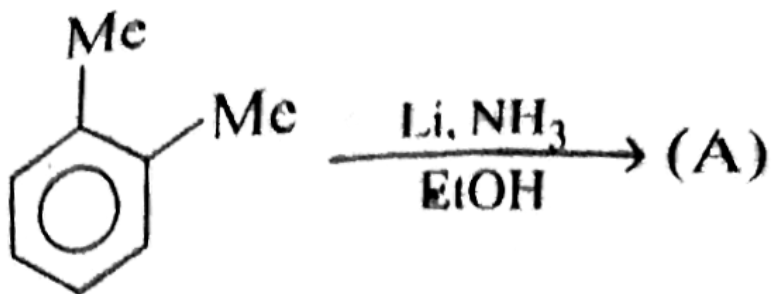
D. (IV) > (III) > (I) > (II)

Answer: c



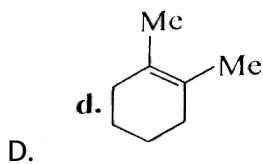
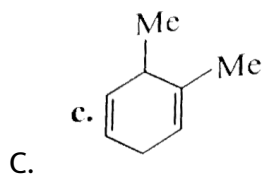
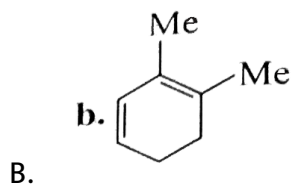
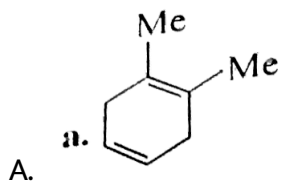
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11.

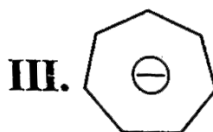
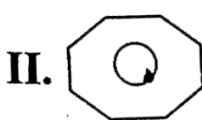
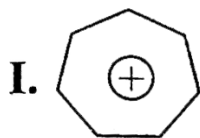
(A) would be:



Answer: a

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12. Which of the following is the correct order of stability of the given compounds?



A. (I) > (II) > (III)

B. (III) > (II) > (I)

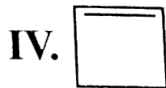
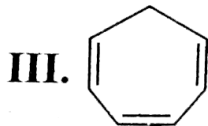
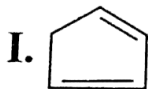
C. (II) > (I) > (III)

D. (II) > (III) > (I)

Answer: a

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13. Give the decreasing order of  $K_a$  value of the following compounds.



A. (I) > (IV) > (III) > (II)

B. (II) > (III) > (IV) > (I)

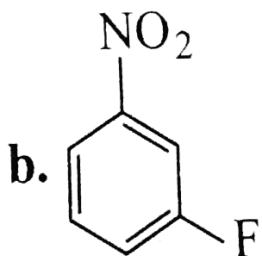
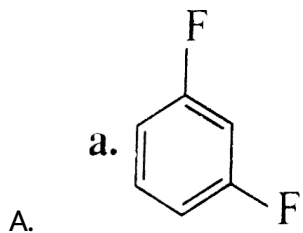
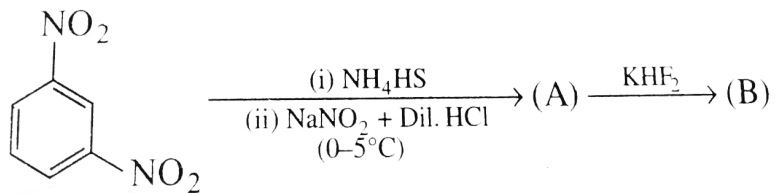
C. (I) > (III) > (II) > (IV)

D. (I) > (IV) > (II) > (III)

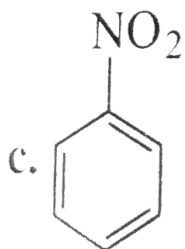
Answer: a

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14. What is the end product of the reaction?

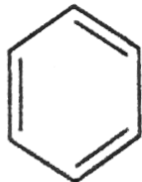


C.



C.

d.

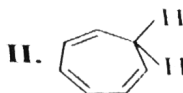
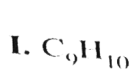


D.

Answer: *b*

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15. Give the increasing order of  $pK_a$  value of the following compounds?



A. (I) < (III) < (II)

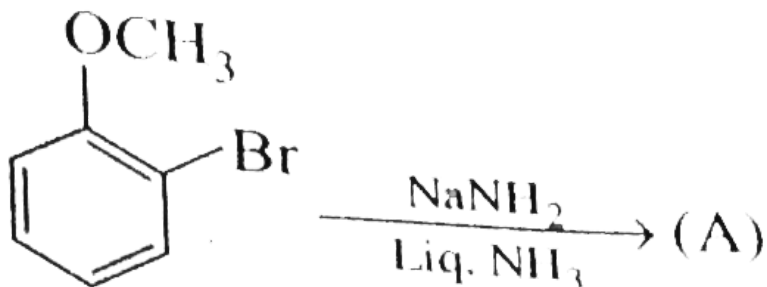
B. (I) > (II) > (III)

C. (II) < (III) < (I)





D. (III) > (II) > (I)

Answer: *a*

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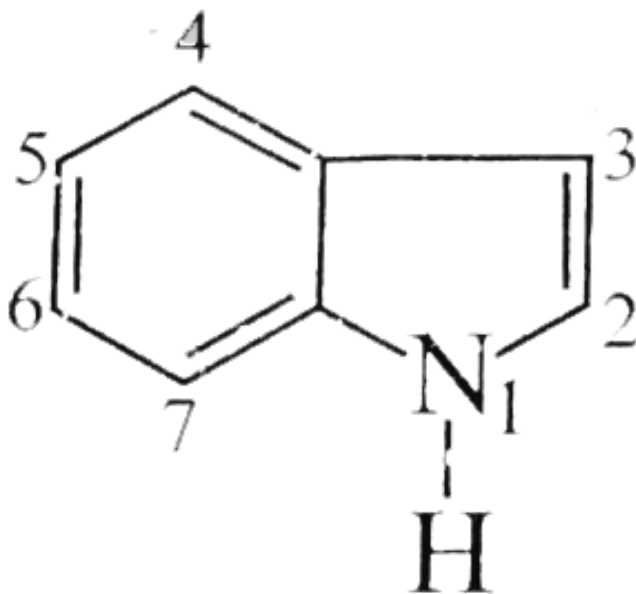
The major product (A) and reaction (R) are:

- A.  COc1cccc(N)c1 *Cine substitution*
- B.  COc1cccc(N)c1 *Addition-elimination reaction*
- C.  COc1cccc(N)c1 *SN<sup>1</sup> reaction*
- D.  COc1cccc(N)c1 *SN<sup>2</sup> reaction*

**Answer: a**

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17. Which of the following mentioned positions in the given compound is more reactive towards electrophilic substitutions?



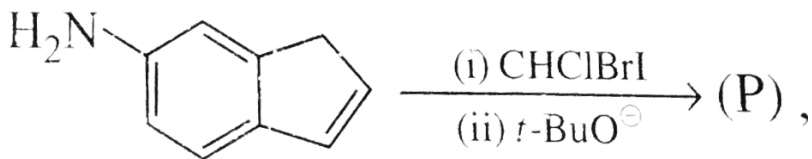
- A. 3
- B. 2
- C. 5
- D. 6

**Answer: a**

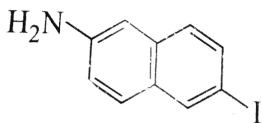


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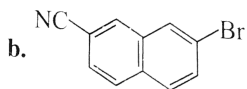
18. In the reaction is



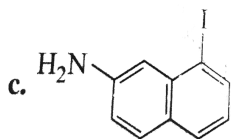
Product (P) will be:



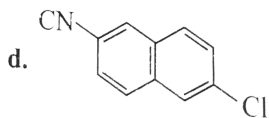
A.



B.



C.

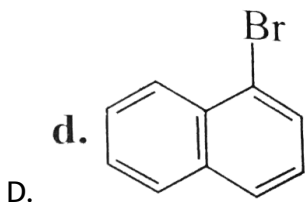
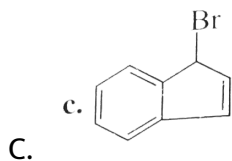
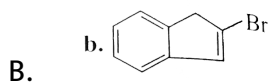
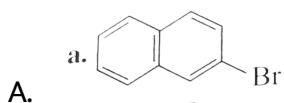
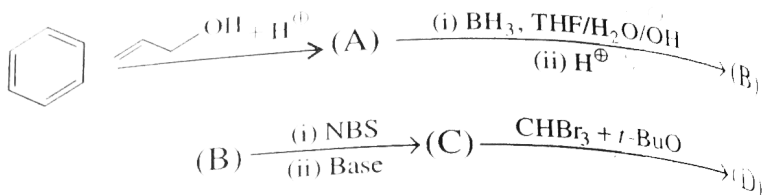


D.

Answer: d

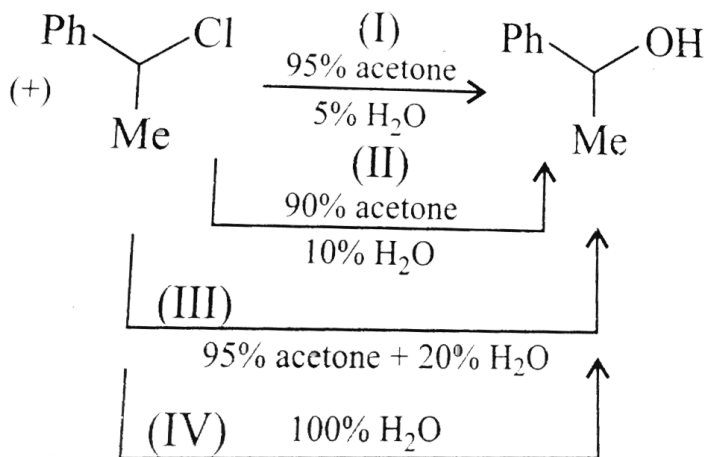


19. What is the end product (D) of the following reaction?



Answer: a

20. Consider the following reactions:



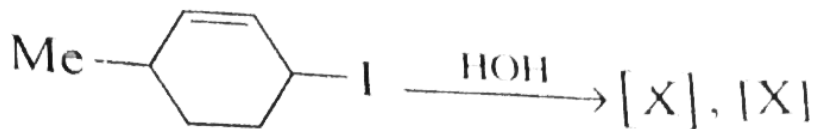
Arrange the following reactions in the decreasing order of greater proportion of inverted product and select the correct answer.

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

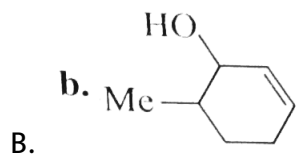
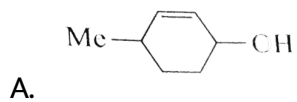
Answer: d



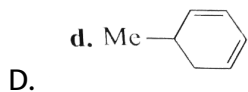
21. In the reaction



will be:



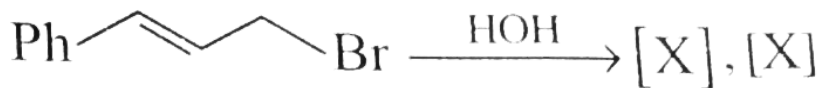
C. Both (a) and (b)



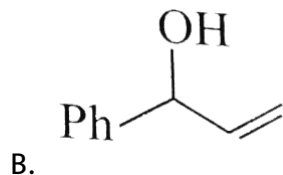
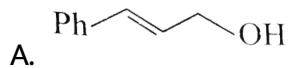
Answer: a

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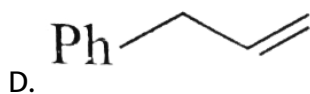
22. In the reaction



will be:



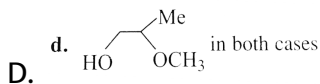
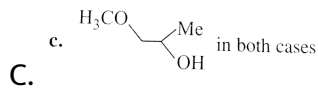
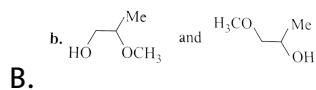
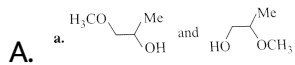
C. Equimolar mixture of (a) and (b)



Answer: c

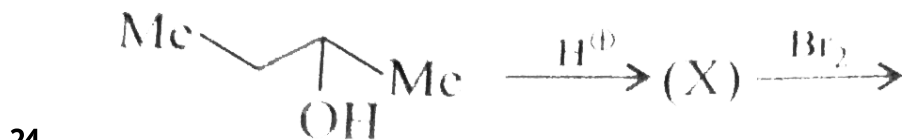
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23. [Y]  $\xleftarrow[\text{CH}_3\text{ONa}]{\text{CH}_3\text{OH}}$    $\xrightarrow[\text{H}^+]{\text{CH}_3\text{OH}}$  [X], [X] are:



Answer: *b*

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five

compounds with formula  $C_4H_8Br_2$ . How many structures of (X) are possible?

A. 2

B. 3

C. 4

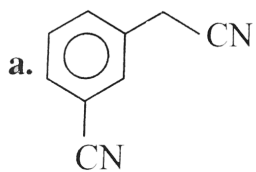
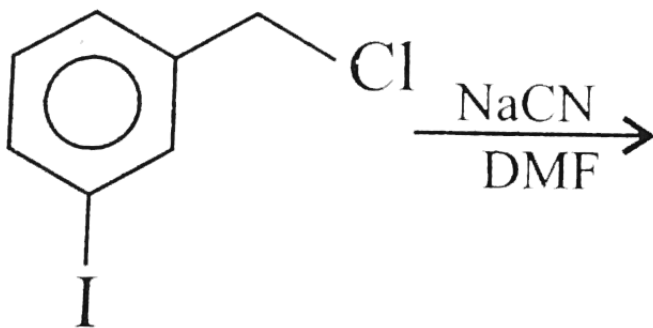
D. 5

**Answer: b**

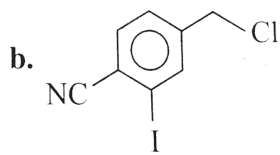
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25. The structure of the major product formed in the following reaction is:

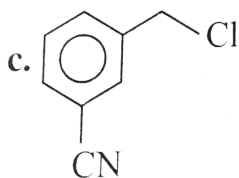
is:



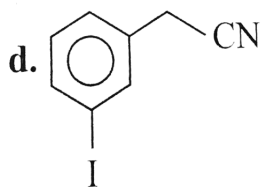
A.



B.



C.



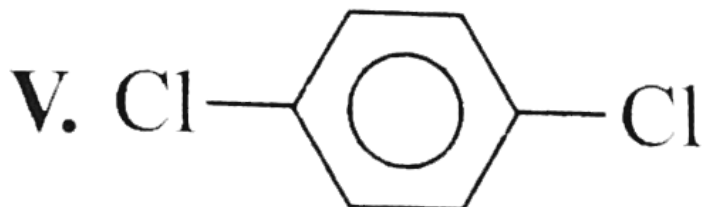
D.

**Answer: d**

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**26.** Consider the following halogen containing compounds:

I.  $CHCl_3$ , II.  $Cl_4$ , III.  $CH_2Cl_2$



IV.  $CH_3Cl$ , V.

A. (II) < (V)

B. (II)

C. (III) < (IV)

D. (I), (IV)

Answer: a

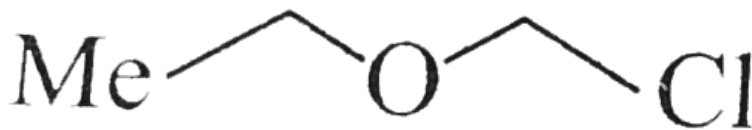
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## Exercises Assertion-Reasoning Type

1.

Assertion

(A):



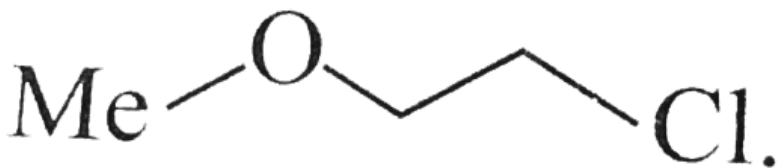
with

$H_2O$

reacts faster

than



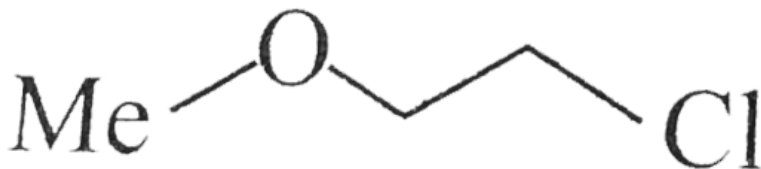


Reason (R): The carbocation of



is more than

the carbocation of



A. Both (A) and (R) are true and (R) is the correct explanation of (A).

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

(A).

C. (A) is true but (R) is false.

D. (A) is false but (R) is true.

**Answer: d**



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## Exercises Assertion-Reasoning

1. Assertion (A):  $SN^1$  reaction is carried out in the presence of a polar protic solvent.

Reason (R): A polar protic solvent increases the stability of carbocation due to solvation.

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

**Answer: d**



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2. Assertion (A):  $\text{SN}^2$  reaction is carried out in the presence of polar aprotic solvent. Reason (R): Polar aprotic solvents do not contain acidic hydrogen.

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

**Answer: b**

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3. Assertion (A): Reaction between  $(\text{Me}_3\text{CONa})$  (sodium ter-butoxide) and ethyl iodide  $(\text{C}_2\text{H}_5\text{I})$  does not produce an ether.

Reason (R): Sodium ter-butoxide is a very strong base but is not a nucleophile.

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

**Answer: a**

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4. Assertion (A): *tert*-Butyl bromide ( $\text{Me}_3\text{C}-\text{Br}$ ) and sodium ethoxide ( $\text{NaOEt}$ ) will react to form only ether.

Reason (R): Ethers are prepared from sodium alkoxide and alkyl halide.

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

C. (A) is true but (R) is false.

D. (A) is false but (R) is true.

**Answer: b**

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5. Assertion: Benzyl bromide when kept in acetone water produces benzyl alcohol.

Reason: The reaction follows  $S_N2$  mechanism.

A. Both (A) and (R) are true and (R) is the correct explanation of (A).

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

C. (A) is true but (R) is false.

D. (A) is false but (R) is true.

**Answer: b**



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6. Assertion: Hydroxyketones are not directly used in Grignard reaction.

Reason : Grignard reagents react with hydroxyl group.

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

Answer: a



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7. Assertion (A): 2 - Bromobutane on reaction with sodium ethoxide in ethanol gives 1 - butene as a major product.

Reason (R): *i*-Butene is more stable than 2-butene.

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

**Answer: d**

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8. Assertion (A): The presence of nitro group facilitates nucleophilic substitution reaction in aryl halide.

Reason (R): The intermediate carbonion is stabilised due to the presence of the nitro group.

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

C.  $(A)$  is true but  $(R)$  is false.

D.  $(A)$  is false but  $(R)$  is true.

**Answer: a**

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9. Assertion  $(A)$ :  $MeMgBr$  should be prepared under perfectly anhydrous conditions.

Reason  $(R)$ : Grignard reagent reacts with water.

A. Both  $(A)$  and  $(R)$  are true and  $(R)$  is the correct explanation of  $(A)$ .

B. Both  $(A)$  and  $(R)$  are true but  $(R)$  is not the correct explanation of  $(A)$ .

C.  $(A)$  is true but  $(R)$  is false.

D.  $(A)$  is false but  $(R)$  is true.

**Answer: a**





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10. Assertion (A): IN comparison to  $C_2H_5Br$ , it is difficult to carry out  $SN$  reaction o vinyl bromide.

Reason (R): Vinyl group is electron donating.

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

**Answer: c**



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11. Assertion (A):  $PhBr$  is less reactive than  $C_2H_5Br$  towards  $SN$  reactions.

Reason (R): The difference in reactivity between  $RX$  and  $H_2O$

$m_o \leq \underline{c}$ esare weaker than those present between the  $m_o \leq \underline{c}$ es of  $RX'$  and water molecules separate, u.

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

**Answer: b**



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12. Assertion : Reaction of tert - butyl chloride with  $Na$  gives 2, 2, 3, 3 - tetramethyl butane.

Reason : Tert - butyl chloride on Wurtz reaction give alkene.

- A. Both (A) and (R) are true and (R) is the correct explanation of (A).

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

**Answer: d**

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## Archives Single Correct

1. Among the following compounds the one that is most reactive towards electrophilic nitration is
- A. Toluene
- B. Benzene
- C. Benzoic acid
- D. Nitrobenzene

**Answer: a**



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2. The reaction of toluene with chlorine in the presence of ferric chloride gives mainly

- A. Benzoyl chloride
- B. *m* - Chlorotoluene
- C. Benzyl chloride
- D. *o* - and *p* - Chlorotoluene

**Answer: d**



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3. In the reaction of *p*-chlorotoluene with  $KNH_2$  in liquid  $NH_3$  the major product is .

A. *o* - Toluidine

B. *m* - Toluidine

C. *p* - Toluidine

D. *p* - Chloronailine

**Answer: b**

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4. Benzylchloride ( $C_6H_5CH_2Cl$ ) can be prepared from toluene by chlorination with :

A.  $SO_2Cl_2$

B.  $SOCl_2$

C.  $Cl_2$

D.  $NaOCl$

**Answer: c**



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5. A solution of (+)-2-chloro-2-phenyl ethane in toluene racemises slowly in the presence of small amount of  $SbCl_5$ , due to the formation of:

- A. Carbanion
- B. Carbene
- C. Free radical
- D. Carbocation

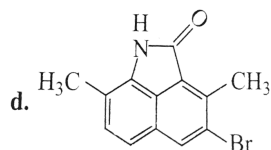
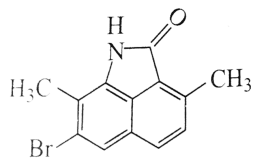
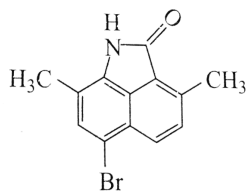
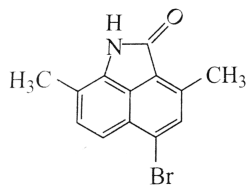
**Answer: d**



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6. 

product on monobromination of this compound is



D.

**Answer: b**



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7. Which of the following is soluble in water?

A.  $CS_2$

B.  $C_2H_5OH$

C.  $CCl_4$

D.  $CHCl_3$

**Answer: b**

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8. When phenol is treated with excess of bromine water, it gives

A. *m* - Bromophenol

B. *o* - and *p* - bromophenol

C. 2, 4 - Dibromophenol

D. 2, 4, 6 - Tribromophenol

**Answer: d**

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9. Chlorobenzene can be prepared by reacting aniline with

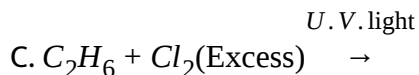
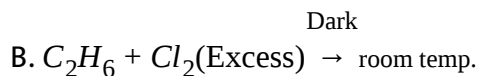
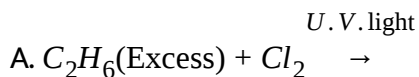
- A. Hydrochloric acid
- B. Cuprous chloride
- C. Chlorine in the presence of anhydrous aluminium chloride
- D. Nitrous acid followed by heating with cuprous chloride

Answer: d

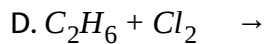


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10. The reaction conditions leading to the best yield of  $C_2H_5Cl$  are



U. V. light



**Answer: a**



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11. n-Propyl bromide on treatment with ethanolic potassium hydroxide produces .

A. Propane

B. Propene

C. Propyne

D. Propanol

**Answer: b**



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12. Phenol reacts with bromine in carbon disulphide at low temperature to give

- A. *m* - Bromophenol
- B. *o* - and *p* - bromophenol
- C. *p* - Bromophenol
- D. 2, 4, 6 - Tribromophenol

**Answer: b**



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13. Chlorination of toluene in the presence of light and heat followed by treatment with aqueous  $NaOH$  gives

- A. *o* - Cresol
- B. *p* - Cresol
- C. 2,4- Dihydroxy toluene

D. Benzoic acid

**Answer: d**

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14. 1 - Chlorobutane on reaction with alcoholic potash gives:

A. 1 - Butene

B. 1 - Butanol

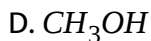
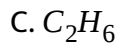
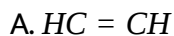
C. 2 - Butene

D. 2 - Butanol

**Answer: a**

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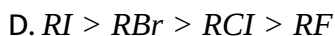
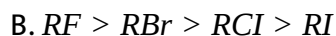
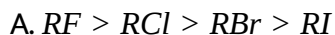
15. Among the following compounds , the strongest acid is:



**Answer: d**

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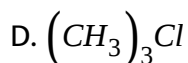
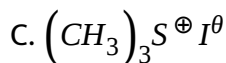
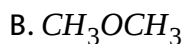
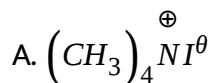
**16.** The order of reactivities of the following alky halides for an  $S_N2$  reaction is .



**Answer: d**

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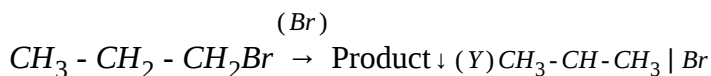
17. The compound that will react most readily with  $\text{NaOH}$  to form methanol is



Answer: a

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18. Identify the set of reagents/reaction conditions (X) and (Y) in the following set of transformation:



A. (X) = Dilute aqueous  $\text{NaOH}$ ,  $20^\circ \text{C}$

(Y) =  $\text{HBr}$ /acetic acid,  $20^\circ \text{C}$

B. (X) = Concentrated alcoholic  $\text{NaOH}$ ,  $80^\circ \text{C}$

(Y) =  $\text{HBr}$ /acetic acid,  $20^\circ \text{C}$

C. (X) = Dilute aqueous  $\text{NaOH}$ ,  $20^\circ \text{C}$

(Y) =  $\text{Br}_2/\text{CHCl}_3$ ,  $0^\circ \text{C}$

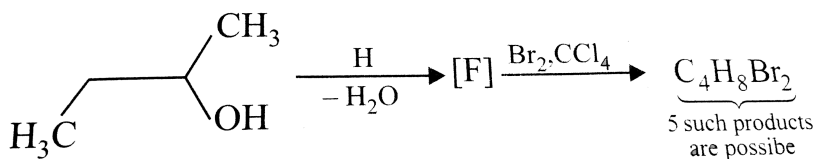
D. (X) = Concentrated alcoholic  $\text{NaOH}$ ,  $80^\circ \text{C}$

(Y) =  $\text{Br}_2/\text{CHCl}_3$ ,  $0^\circ \text{C}$

Answer: b

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19. How many structures for  $F$  are possible



A. 2

B. 5

C. 6

D. 3

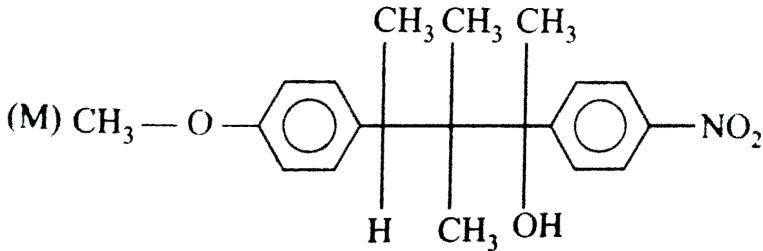
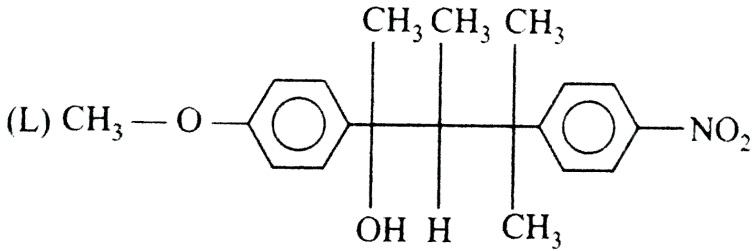
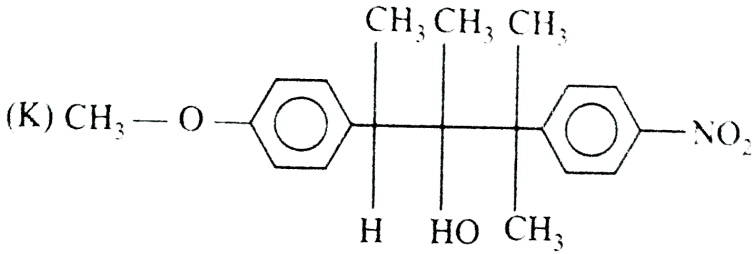
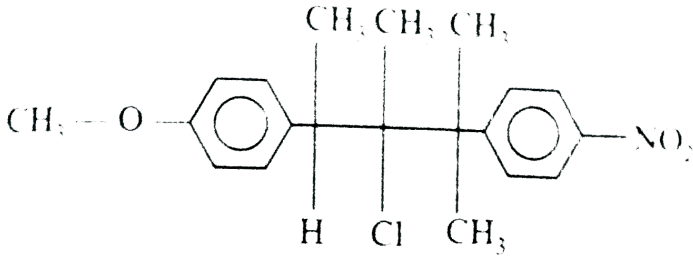
**Answer: *d***



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20. The following compound on hydrolysis in aqueous acetone will give .



A. Mixture of (K) and (L)

B. Mixture of (K) and (M)

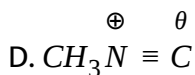
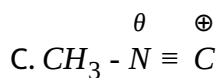
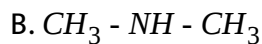
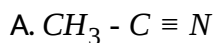
C. Only (M)

D. Only (K)

Answer: a

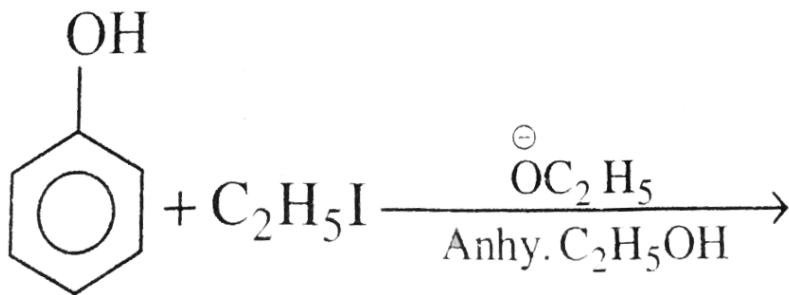
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21.  $CH_3NH_2 + CHCl_3 + KOH \rightarrow$  nitrogen containing compound  
 $+ KCl + H_2O$ . Nitrogen containing compound is

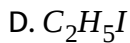
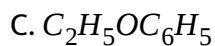
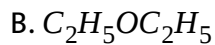
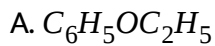


Answer: d

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1.

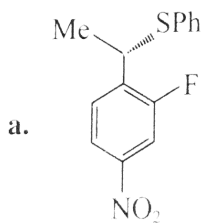
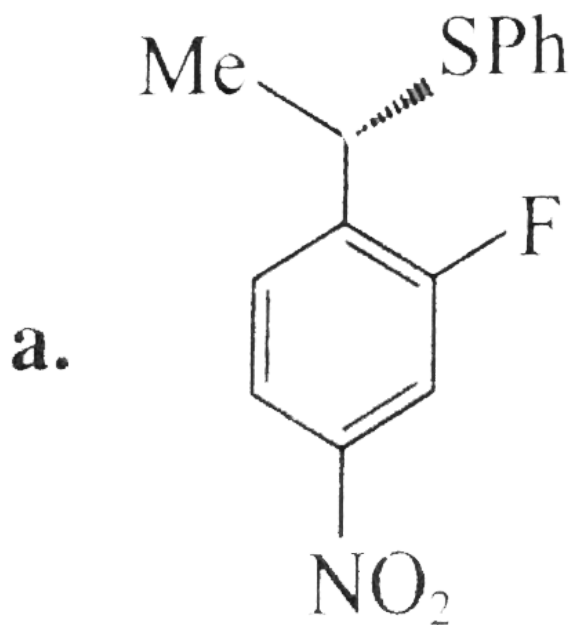


**Answer: b**

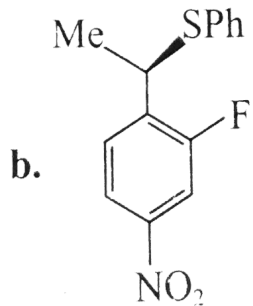


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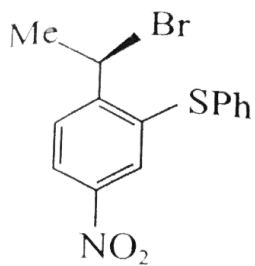
2. The major product of the following reaction is"



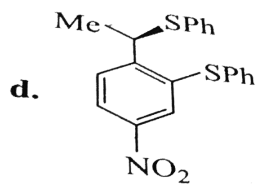
**A.**



**B.**



**C.**



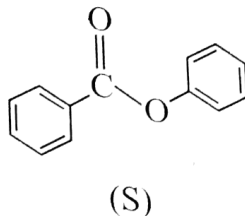
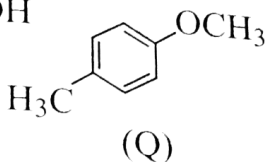
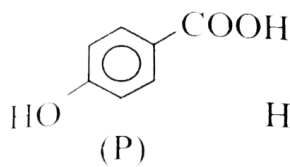
**D.**

**Answer: a**

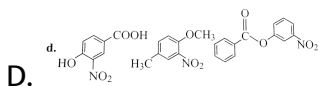
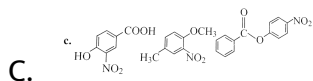
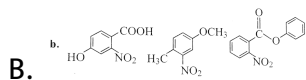
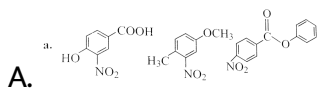


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### 3. Compounds (P), (Q) and (S)



were separately subjected to nitration using  $HNO_3/H_2SO_4$  mixture. The major product formed in each case, respectively is:



Answer: c



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1. An aromatic molecule will

- A. Have  $4n\pi$  - electrons
- B. Have  $(4n + 2)\pi$  - electrons
- C. Be planar
- D. Be cyclic

Answer: (b, c, d)



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2. Toluene, when treated with  $Br_2/Fe$  gives p-bromotoluene as the major product, because the  $CH_3$  group:

- A. is para-directing
- B. is meta-directing
- C. activates the ring by hyperconjugation
- D. deactivates the ring

**Answer: (a, c)**

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3. Aryl halides are less reactive towards nucleophilic substitution reaction as compared to alkyl halides due to

- A. The formation of less stable carbonium ion
- B. Resonance stabilisation
- C. Longer carbon halogen bond
- D.  $sp^2$  - Hybridised carbon attached to the halogen.

**Answer: (b, d)**

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4. The compounds used as refract are:



A.  $NH_3$

B.  $CCl_4$

C.  $CF_4$

D.  $CF_2Cl_2$

**Answer: (a, d)**



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5. The products of reaction of alcoholic silver nitrate with ethyl bromide are

A. Ethane

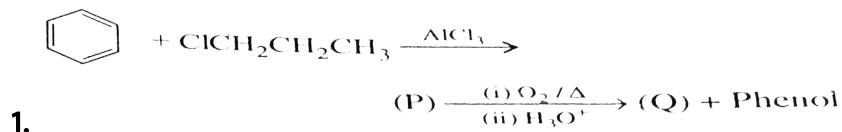
B. Ethene

C. Nitroethane

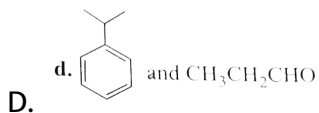
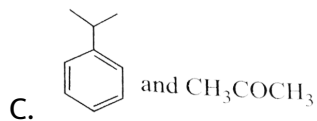
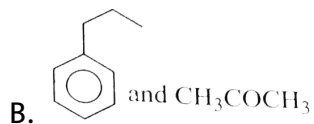
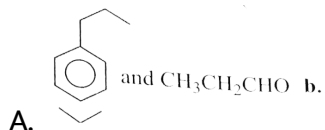
D. Ethyl nitrite

**Answer: (c,d)**

Archives Multiple Correct Answers Type



The major products (P) and (Q) are:



Answer: c

## Archives Fill In Theblanks

1. The structure of the intermediate prouduct formed by the oxidation of toluene with  $CrO_3$  and acetic anhydride ,whose hydrolysis gives benzaldehyde is.....

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2. The halogen which is most reactions , in the halogenation of alkanes under sunlight is .....

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3. The starting material for the manufacture of polyvinyl chloride is obtained by reacting HCl with .....

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4. Formation of phenol from chlorobenzene is an example of  
.... Aromatic substitution.

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5. In benzene, carbon uses all the three p-orbitals for hybridisation .

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6. True or False?

An electron donating substituent in benzene orients the incoming electrophilic group to the meta-position.

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7. Carbon tetrachloride burns in air when lighted to give phosgene .

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8. The yield of ketone when a secondary alcohol is oxidised is more than the yield of aldehyde when a primary alcohol is oxidised.

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9. Carbon tetrachloride is inflammable.

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10. Sodium ethoxide is prepared by reacting ethanol with aqueous sodium hydroxide.

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11. The reaction of vinyl chloride with hydrogen iodide to give 1-chloro-1-iodoethane is an example of anti-Markownikoff's rule.



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12. Photobromination of 2-methylpropane gives a mixture of 1-bromo-2-methyl propane in the ratio 9 : 1.



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## Archives Assertion-Reasoning

1. Assertion: Aryl halides undergo nucleophilic substitution reactions with ease.

Reason: The carbon halogen bond in aryl halides has partial double bonds character.

A. Statement 1: is true Statement 2: is true, Statement 2 is the correct explanation of Statement 1

B. Statement 1 is true, Statement 2 is true, Statement 2 is not the correct explanation fo Statement 1.

C. Statement 1 is true, Statement 2 is false.

D. Statement 1 is false, Statement 2 is true.

**Answer: D**

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2. Statement I Benzonitrile is prepared by the reaction of chlorobenzene with potassium cyanide.

Statement II Cyanide ( $CN^-$ ) is a strong nucleoohile.

A. Statement 1: is true Statement 2: is true, Statement 2 is the correct explanation of Statement 1

B. Statement 1 is true, Statement 2 is true, Statement 2 is not the correct explainion fo Statement 1.

C. Statement 1 is true, Statement 2 is false.

D. Statement 1 is false, Statement 2 is true.

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3. Assertion: Bromobenzene upon reaction with  $Br_2/Fe$  gives 1,4-dibromobenzene as the major product

Reason In bromobenzene the inductive effect of the bromo group is more dominant than the mesomeric effect in directing the incoming electrophile .

A. Statement 1: is true Statement 2: is true, Statement 2 is the correct explanation of Statement 1

B. Statement 1 is true, Statement 2 is true, Statement 2 is not the correct explanation fo Statement 1.

C. Statement 1 is true, Statement 2 is false.



D. Statement 1 is false, Statement 2 is true.

**Answer: C**

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## Archives Subjective

1. Give reason for the following in one or two sentences:

'Although benzene is highly unsaturated, normally it does not undergo addition reaction'.

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2. Show with balanced equation what happens when the 'p-xylene is reacted with concentrated sulphuric acid and the resultant product is fused with KOH'.

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3. How many sigma and pi bonds are present in a benzene molecule?

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4. Write down the main product of the following reaction



Benzene      →

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5. Answer the following with suitable equation wherever necessary

(i) How can you prepare benzene from lime?

(ii) How will you convert toluene to m-nitrobenzoic acid?

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6. Arrange the following in increasing order of reactivity towards sulphonation with fuming sulphuric acid.

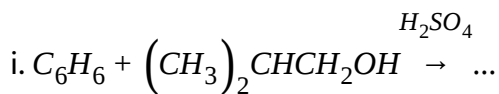
Benzene, toluene, methoxy benzene, chlorobenzene.

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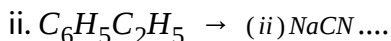
7. Give reasons in two or three sentences only for the following: Phenyl group is known to exert negative inductive effect, but each phenyl ring in biphenyl ( $C_6H_5 - C_6H_5$ ) is more reactive than benzene towards electrophilic substitution.

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8. Product the major product in the following reactions:



(i)  $Br_2$  heat light



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9. Predict the structure of the intermediates/products in the following reaction sequences



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10. Toluene reacts with bromine in the presence of the light to give benzyl bromide, while in presence of  $FeBr_3$  it gives p-bromotoluene. Give explanation for the above observation.

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11. Give reasons for the following in one or two sentences "Nitrobenzene does not undergo Friedel-Craft's alkylation."

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12. Give reasons for the following:

(i) tert-butylbenzene does not give benzoic acid on treatment with acidic  $KMnO_4$ .

(ii) Normally, benzene gives electrophilic substitution reaction rather than electrophilic addition reaction although it has double bond.

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13. 7-bromo-1,3,5-cycloheptatriene exists as ionic species in aqueous solution while 5-bromo-1,3-cyclopentadiene doesn't ionise even in presence of  $Ag^+(aq)$ , Explain.

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14. An organic liquid 'A' containing C, H and O with boiling point  $78^\circ C$ , possessing a rather pleasant odour. On heating with concentrated sulphuric acid gives a gaseous product 'B' with the empirical formula  $CH_2$ . 'B' decolourises bromine water as well as alkaline permanganate solution.

and takes up one mole of  $H_2$  (per mole of B) in presence of finely divided nickel at high temperature. Identify the substances A and B.

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15. show by chemical equations only how would you prepare the following from the indicated starting materials. Specify the reagents in each step of synthesis of:

i. Chloroform from carbon disulphide.

ii. Hexachloroethane ( $C_2Cl_6$ ) from calcium carbide.

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16. Chloroform is stored in dark colored bottles. Explain in not more than two sentences.

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17. A compound ( $X$ ) containing  $C$ ,  $H$ , and  $O$  is unreactive towards sodium. It does not add with bromine. It also and excess of  $Hl$ , ( $X$ ) yields only one organic product ( $Y$ ), ( $Y$ ) on hydrolysis yields a new compound ( $Z$ ) which can be converted to ( $Y$ ) by reaction with red phosphorus and permanganate gives a carboxylic acid. The equivalent weight of this acid is 60. What are the compounds ( $X$ ), the conversion of ( $X$ ) to ( $Y$ ).

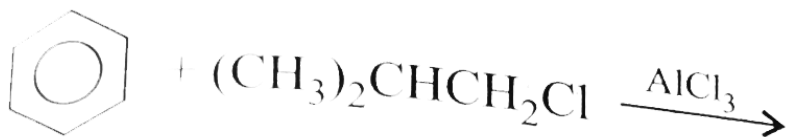
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18. State the conditions under which the following preparations are carried out. Give the necessary equations which need not be balanced.

- i. Ethanol from acetylene.
- ii. Lead tetraethyl from sodium lead alloy.
- iii. Methyl chloride from aluminium carbide.

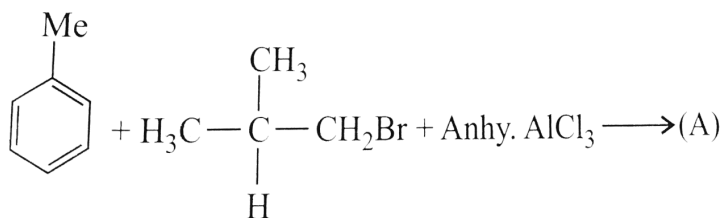
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1. Write the structure of the major organic product expected from the following reaction:



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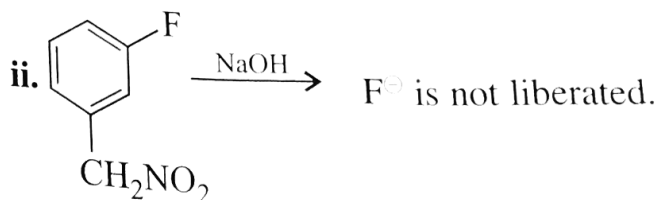
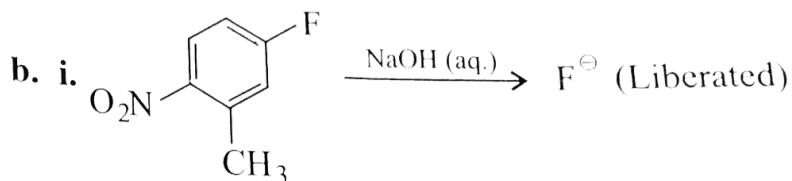
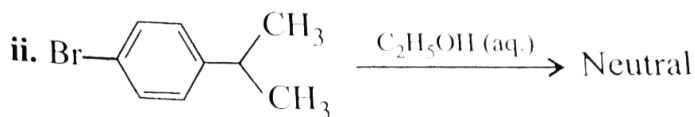
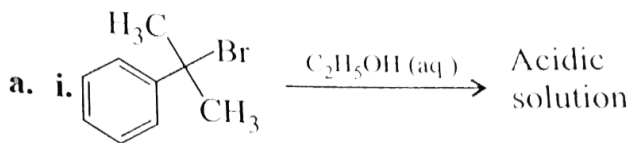
2. Complete the following giving structures of the principle organic products.



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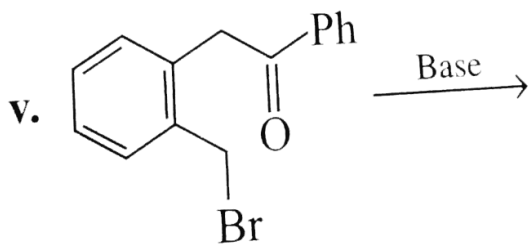
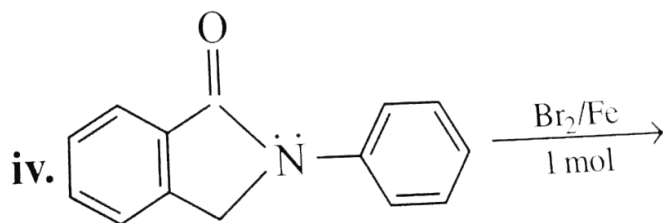
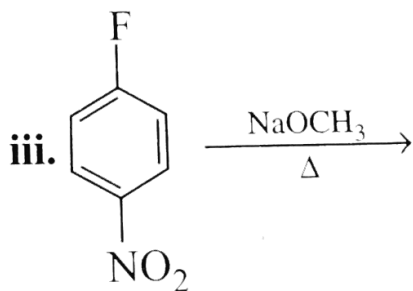
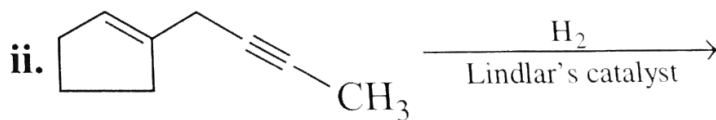
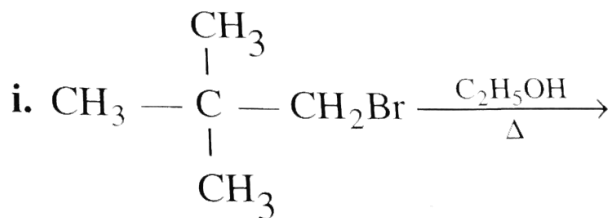


3. Give reasons for the following:



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4. What would be the major product in each of the following reactions?



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