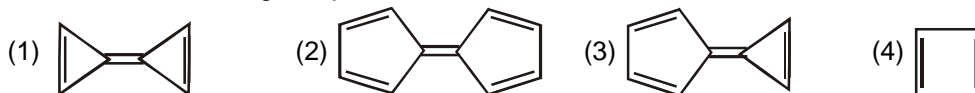


Self Practice Paper (SPP)

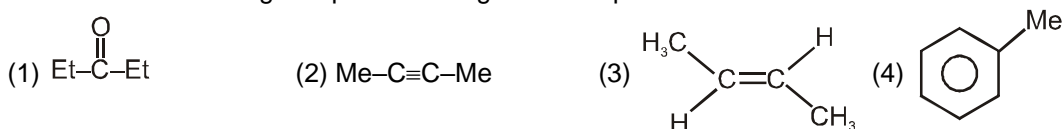
1. Which one of following compounds is aromatic in nature ?




2. Ease of ionization to produce carbocation & bromide ion under the treatment of Ag^+ will be maximum in ?



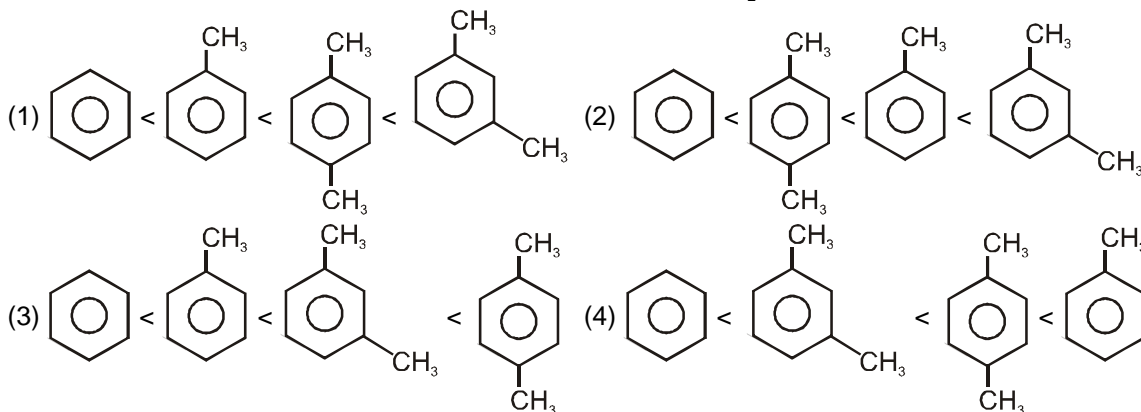
3. Which one of following compounds can give electrophilic substitution reaction ?



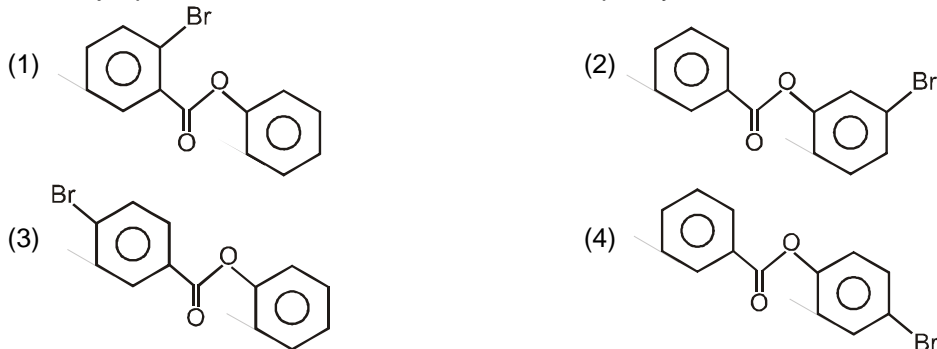
4.  $\xrightarrow{\text{E}^+}$ Name of intermediate produced during given reaction will be

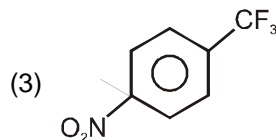
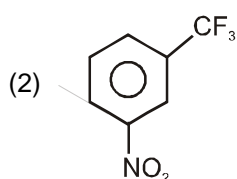
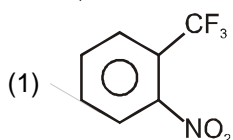
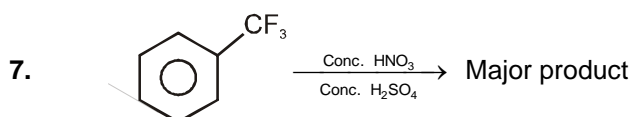
- (1) Arenium ion (2) σ complex/Wheland intermediate
(3) Non aromatic cyclohexa-dienyl carbocation (4) All of these

5. Which of following is correct order of reactivity towards the ArS_2E ?



6. The major product formed on monobromination of phenylbenzoate is :





(4) 1 & 3 both

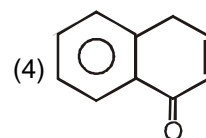
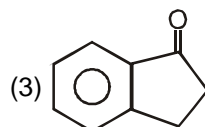
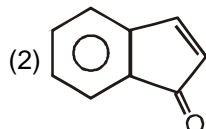
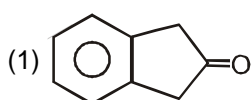
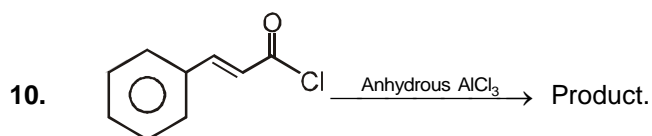
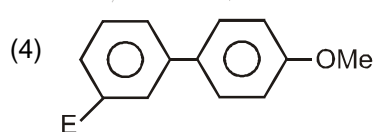
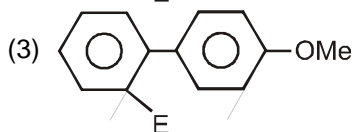
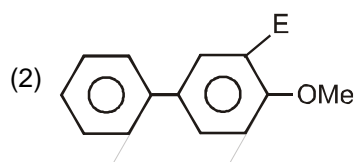
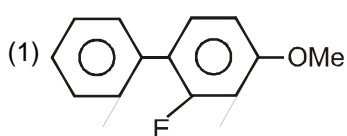
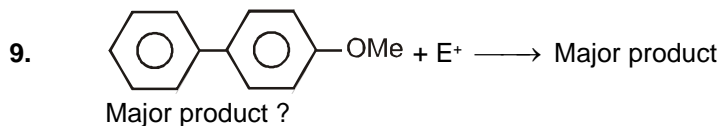
8. In the sulphonation of benzene the active electrophilic species is

(1) SO_2

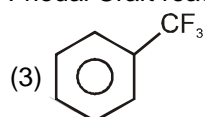
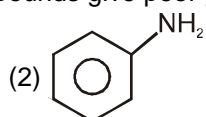
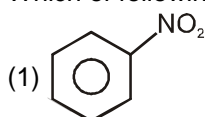
(2) SO_3

(3) SO_4^{2-}

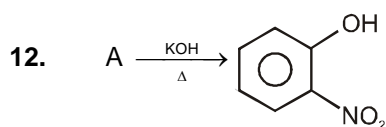
(4) HSO_4^-



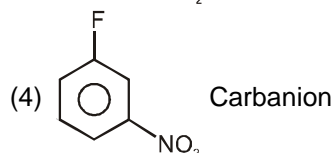
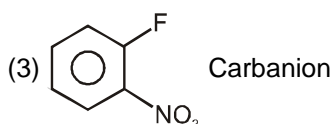
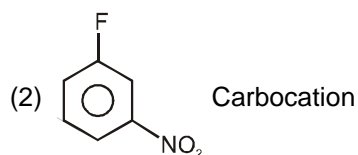
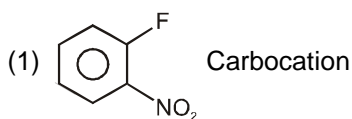
11. Which of following compounds give poor yield in Friedel Craft reaction.



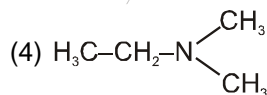
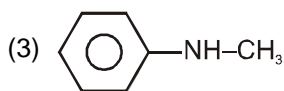
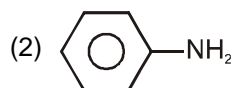
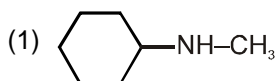
(4) All of these



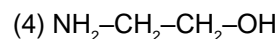
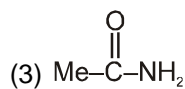
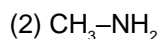
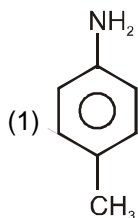
Reactant (1) & intermediate is



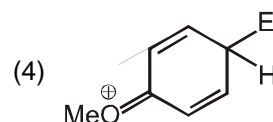
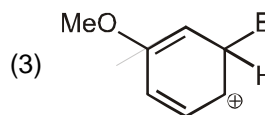
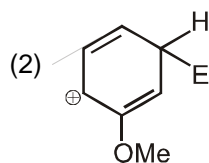
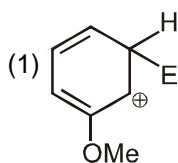
13. Which of following compound gives carbyl amine reaction



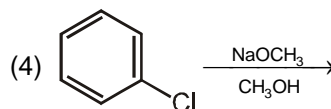
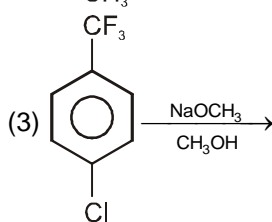
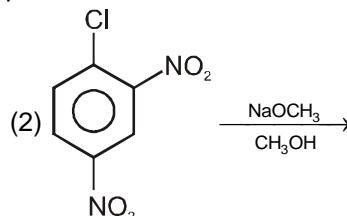
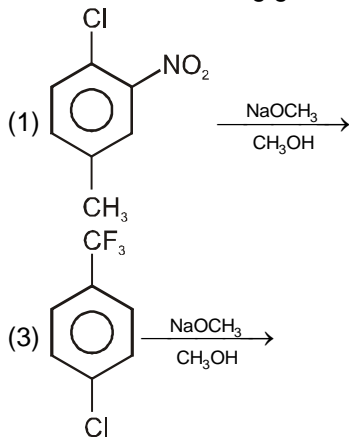
14. The compound which does not give foul smell when heated with CHCl_3 & KOH is



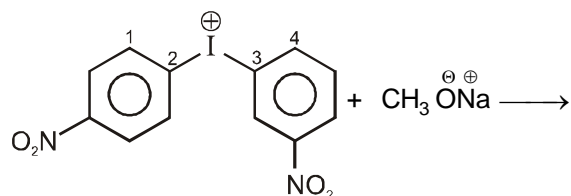
15. Which one of following species is the most stable arenium ion ?



16. Which of the following gives faster rate of nucleophilic substitution reaction ?



17.



Reagent attack at ___ site.

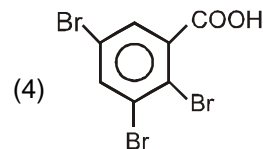
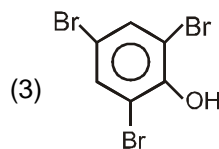
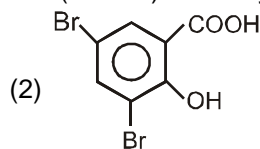
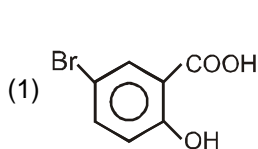
(1) 1

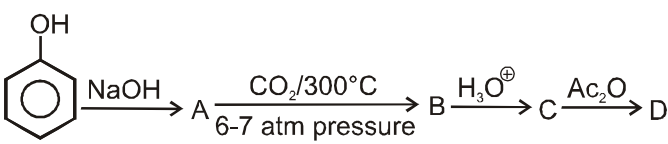
(2) 2

(3) 3

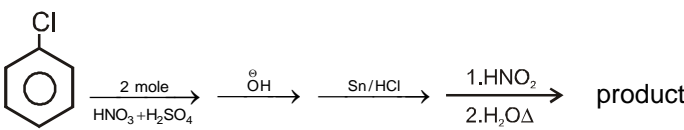
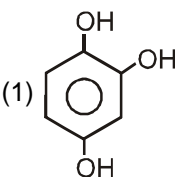
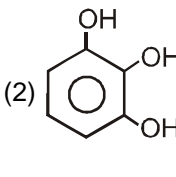
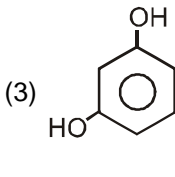
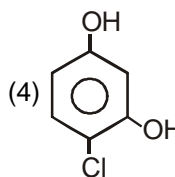
(4) 4

18. The action of bromine water (excess) on salicylic acid results in the formation of :



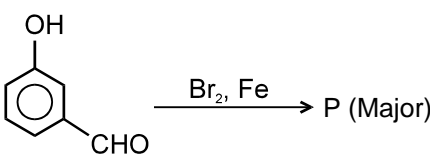
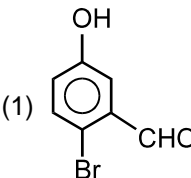
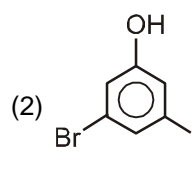
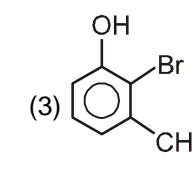
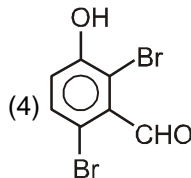
19. 
 'D' is :
 (1) Aspirin (2) Valine (3) Cumene (4) Salicylic acid

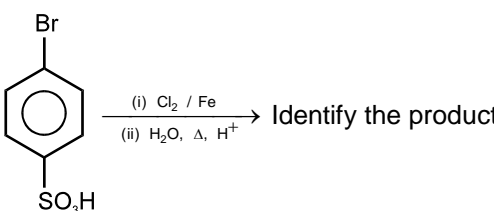
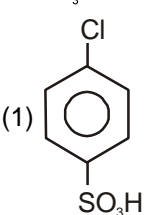
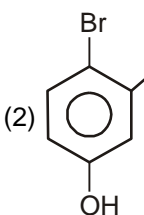
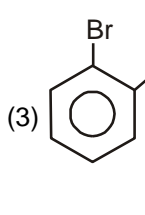
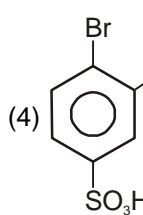
20. In the following reaction
 $X \xrightarrow{\text{Bromination}} Y \xrightarrow[\text{HCl}]{\text{NaNO}_2} Z \xrightarrow[\text{C}_2\text{H}_5\text{OH}]{\text{Boiling}} \text{Tribromobenzene}$, X is
 (1) benzoic acid (2) salicylic acid (3) phenol (4) aniline

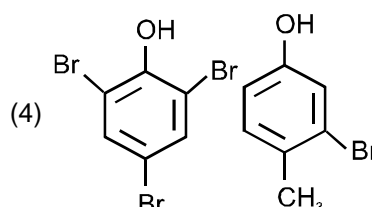
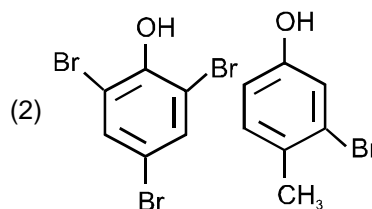
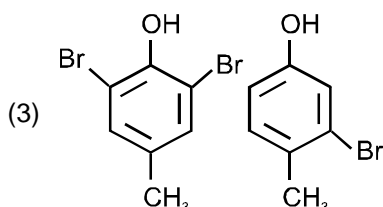
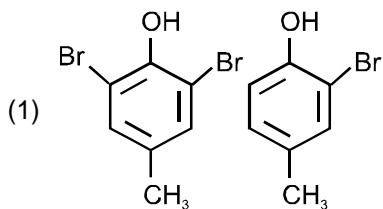
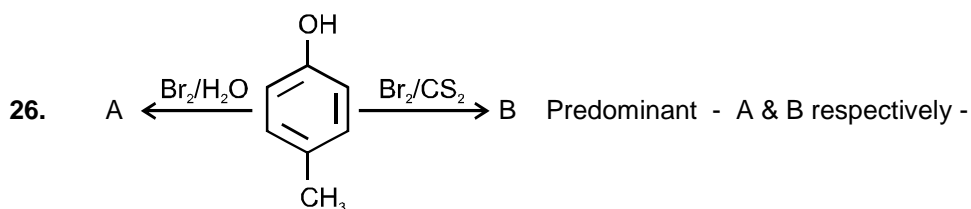
21. 
 (1)  (2)  (3)  (4) 

22. $\text{C}_6\text{H}_6 + (\text{P}) \xrightarrow{\text{AlCl}_3} \text{C}_6\text{H}_5\text{CONH}_2$
 (P) in the above reaction is :
 (1) NH_2CONH_2 (2) ClCONH_2 (3) CH_3CONH_2 (4) $\text{CH}_2(\text{Cl})\text{CONH}_2$

23. Nitrobenzene and CH_3Cl in presence of anhydrous AlCl_3 gives :
 (1) o-nitrotoluene (2) p-nitrotoluene (3) m-nitrotoluene (4) Reaction will not occur

24. 
 (1)  (2)  (3)  (4) 

25. 
 (1)  (2)  (3)  (4) 



27. An organic compound A on reduction gives compound B which on reaction with chloroform and potassium hydroxide forms C. The compound C on catalytic reduction gives N-methylaniline. The compound A is

(1) nitrobenzene (2) nitromethane (3) methylamine (4) aniline

28. Primary amine reacts with carbon disulphide and HgCl_2 to produce alkyl isothiocyanate. This reaction is

(1) Carbylamine reaction (2) Hofmann bromide reaction
(3) Perkin reaction (4) Hofmann mustard oil reaction

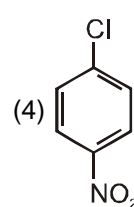
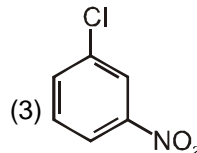
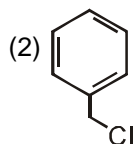
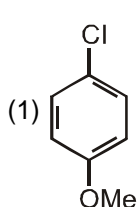
29. When aniline reacts with HNO_2 ($\text{NaNO}_2 + \text{HCl}$) diazonium chloride is formed which on reaction with H_3PO_2 gives :

(1) $\text{C}_6\text{H}_5\text{OH}$ (2) C_6H_6 (3) $\text{Ph}-\text{CH}_3$ (4) CH_3-CH_3

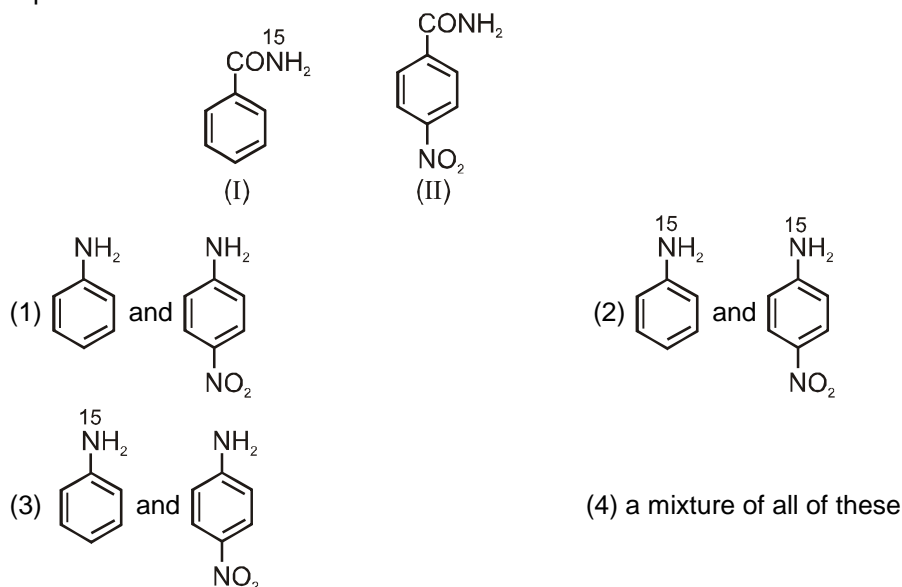
30. Aniline reacts with to yield as the final product

(1) Aqueous bromine, 2-bromoaniline (2) Aqueous bromine, 2, 4, 6-tribromoaniline
(3) chloroform/KOH, phenyl cyanide (4) acetyl chloride, benzanilide

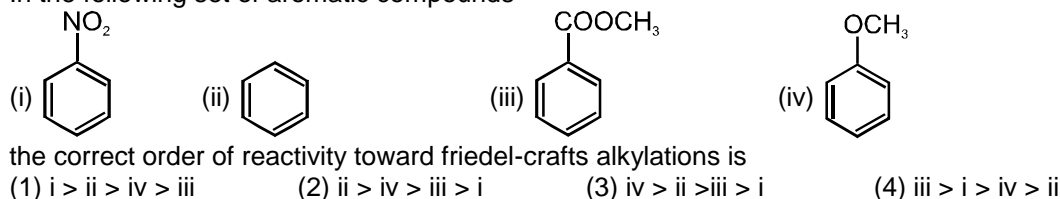
31. Compound which undergoes nucleophilic substitution reactions most readily is



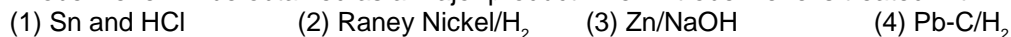
32. The products formed when a mixture of the following two amides (I and II) are treated with bromine and aqueous KOH are



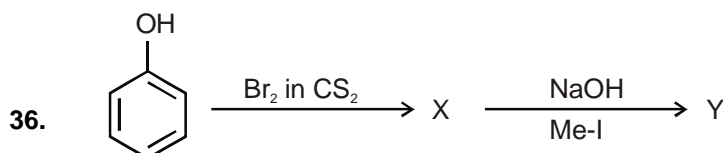
33. In the following set of aromatic compounds



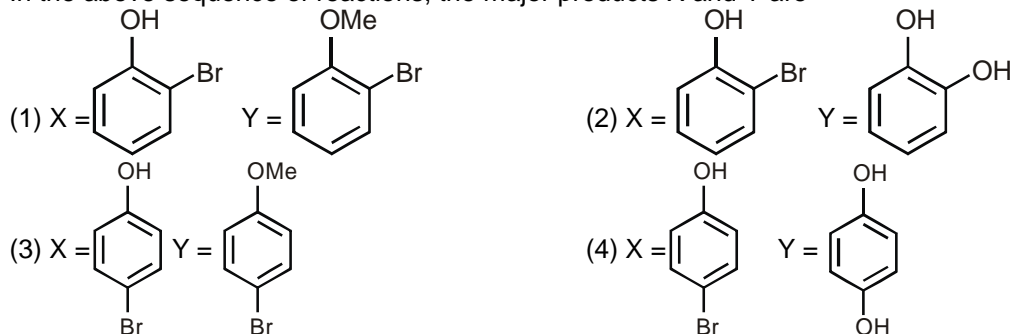
34. Azobenzene will be obtained as a major product when nitrobenzene is treated with:



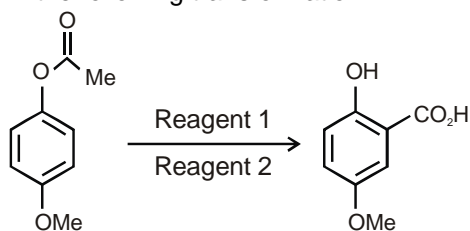
35. Among the following



In the above sequence of reactions, the major products X and Y are



37. In the following transformation



reagents 1 and 2 are :

(1) H_2SO_4 ; alkaline KMnO_4

(3) H_3PO_4 ; CHCl_3/KOH

(2) AlCl_3 ; I_2/NaOH

(4) $\text{KOH}; \text{CHCl}_3 / \text{KOH}$

38. Friedel–Crafts acylation is :

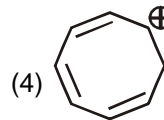
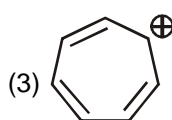
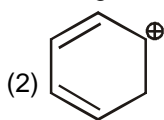
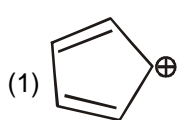
(1) α -acylation of a carbonyl compound

(3) acylation of aliphatic olefins

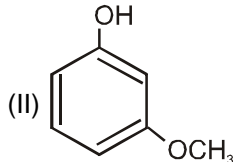
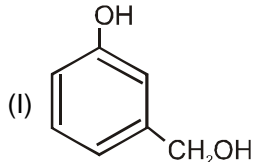
(2) acylation of phenols to generate esters

(4) acylation of aromatic nucleus

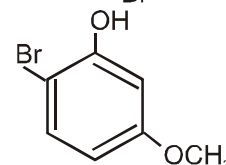
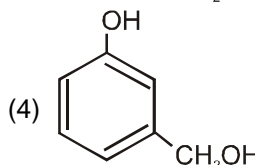
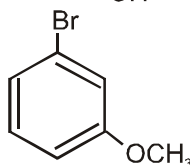
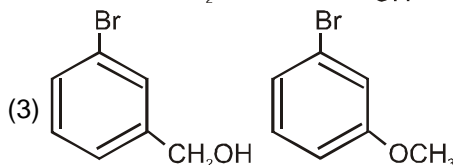
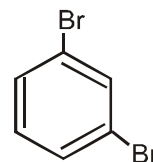
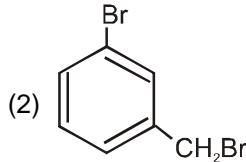
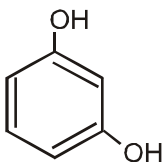
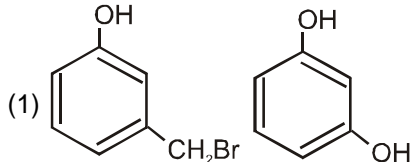
39. The aromatic carbocation among the following is :



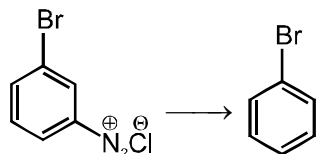
40. Two isomeric compounds I and II are heated with HBr.



The products obtained are



41. For the transformation



the reagent used is :

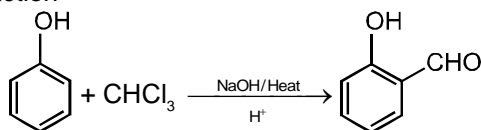
(1) LiAlH_4

(2) H_3PO_2

(3) H_3O^+

(4) H_2/Pt

42. The reaction



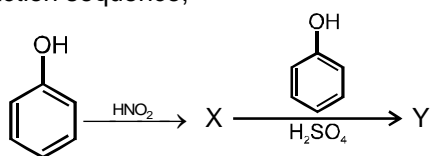
is known as :

(1) Perkin reaction

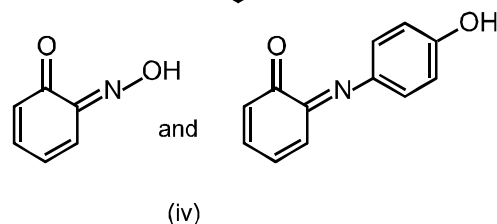
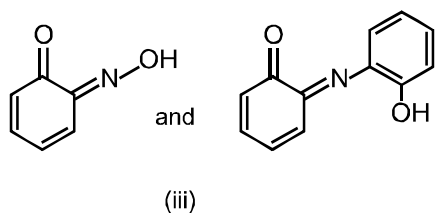
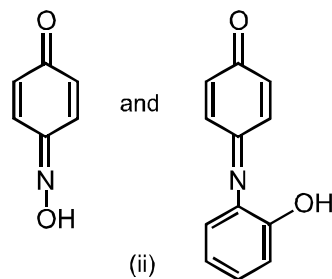
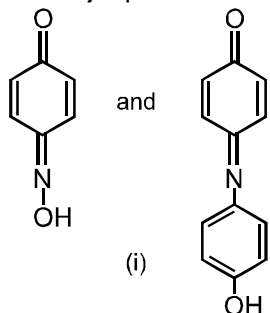
(2) Sandmeyer reaction

43. (3) Reimer-Tiemann reaction
The reaction sequence,

- (4) Cannizzaro reaction



the major products X and Y, respectively, are :



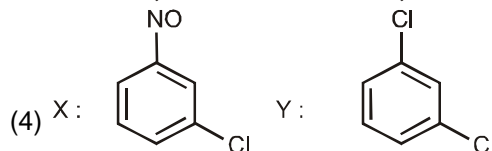
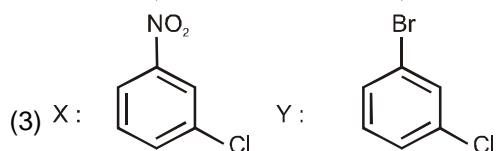
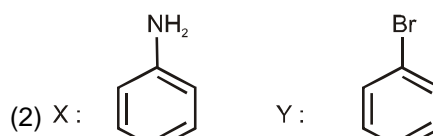
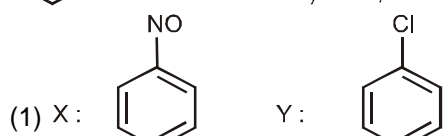
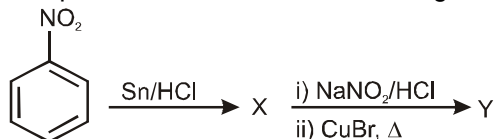
(1) i

(2) ii

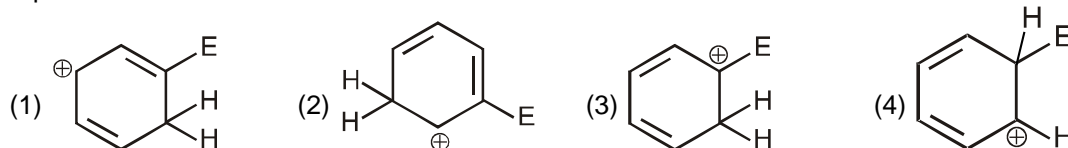
(3) iii

(4) iv

44. The products X and Y in the following reaction sequence are :



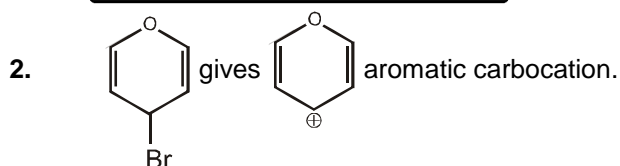
45. In the reaction of benzene with an electrophile E^+ , the structure of the intermediate σ -complex can be represented as :



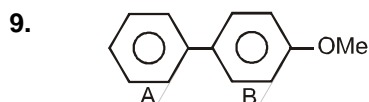
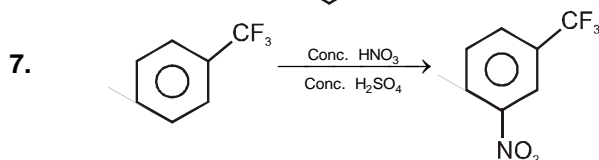
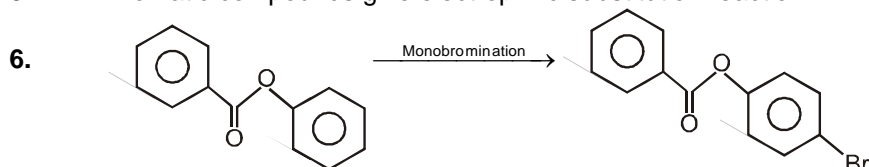
SPP Answers

1.	(3)	2.	(3)	3.	(4)	4.	(4)	5.	(1)	6.	(4)	7.	(2)
8.	(2)	9.	(2)	10.	(2)	11.	(4)	12.	(3)	13.	(2)	14.	(3)
15.	(4)	16.	(2)	17.	(2)	18.	(3)	19.	(1)	20.	(4)	21.	(1)
22.	(2)	23.	(4)	24.	(1)	25.	(3)	26.	(1)	27.	(1)	28.	(4)
29.	(2)	30.	(2)	31.	(2)	32.	(3)	33.	(3)	34.	(3)	35.	(4)
36.	(3)	37.	(2)	38.	(4)	39.	(3)	40.	(1)	41.	(2)	42.	(3)
43.	(1)	44.	(2)	45.	(4)								

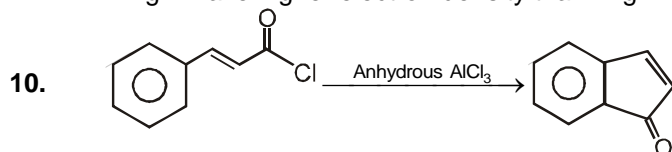
SPP Solutions



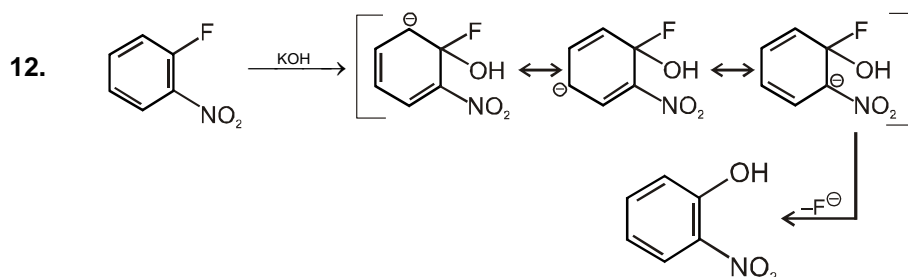
3. Aromatic compounds give electrophilic substitution reaction.



Ring B have higher electron density than ring A. So electrophile attack on ring B.



11. Highly deactivated ring gives poor yield in Friedel Craft reaction.

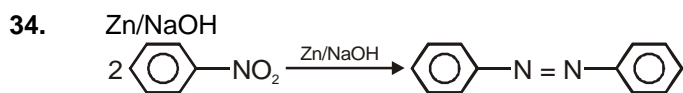


13. 1° amine gives carbyl amine test.

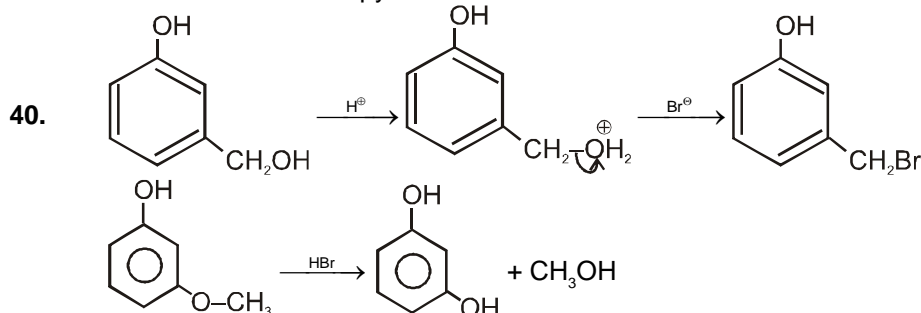
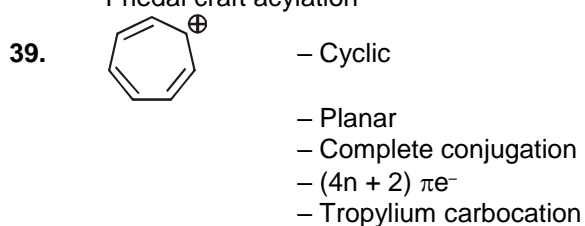
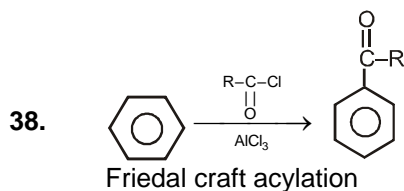
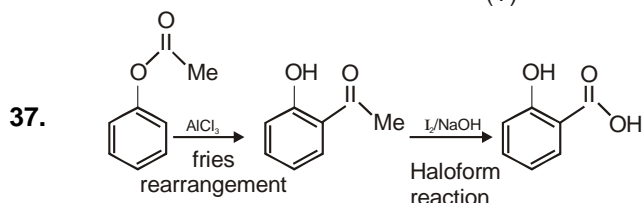
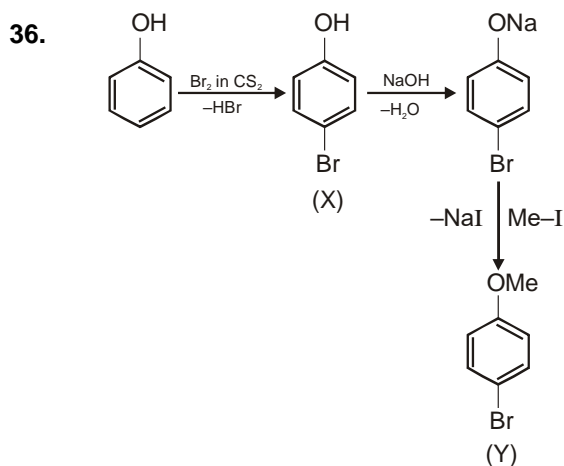
14. Only 1° amine give carbyl amine test. 1° Amide does not gives carbyl amine test.

15. Structure have complete octate of each atom.
- 18.
- 19.
- 20.
- 21.
22.
$$\text{C}_6\text{H}_6 + \text{ClCONH}_2 \xrightarrow{\text{AlCl}_3} \text{C}_6\text{H}_5\text{CONH}_2 + \text{HCl}$$
23. All -m group do not show friedel craft reaction.
24. Orientation decided by more activating -OH group [+ M effect].
- 25.
- 26.
27.
$$\text{Ph-NH}_2 + \text{CHCl}_3 + \text{KOH} \longrightarrow \text{Ph-NC} \xrightarrow{\text{Reduction}} \text{Ph-NH-CH}_3$$
29.
$$\text{PhN}_2^+\text{Cl}^- \xrightarrow{\text{H}_3\text{PO}_2, \Delta} \text{C}_6\text{H}_6$$
- 30.

33. Reativity towards Friedel-Crafts alkylation is proportional to electron density in the benzene ring.



35. (II) & (IV) Because both have close system of conjugated double bond and follow Huckel's $(4n+2) \pi e^-$ rule.



41. H_3PO_2 a mild reducing agent is used to remove the diazonium group with H.

43. Due to bulkyness reaction is favored at para position.

