

CHEMISTRY

1. Assuming $2s-2p$ mixing is **NOT** operative, the paramagnetic species among the following is

[JEE (Adv)-2014 (Paper-2)]

- (A) Be_2 (B) B_2
(C) C_2 (D) N_2

2. The standard state Gibbs free energies of formation of C(graphite) and C(diamond) at $T = 298 \text{ K}$ are

$$\Delta_f G^\circ[\text{C}(\text{graphite})] = 0 \text{ kJ mol}^{-1}$$

$$\Delta_f G^\circ[\text{C}(\text{diamond})] = 2.9 \text{ kJ mol}^{-1}$$

The standard state means that the pressure should be 1 bar, and substance should be pure at a given temperature. The conversion of graphite [C(graphite)] to diamond [C(diamond)] reduces its volume by $2 \times 10^{-6} \text{ m}^3 \text{ mol}^{-1}$. If C(graphite) is converted to C(diamond) isothermally at $T = 298 \text{ K}$, the pressure at which C(graphite) is in equilibrium with C(diamond), is

[Useful information : $1 \text{ J} = 1 \text{ kg m}^2 \text{ s}^{-2}$; $1 \text{ Pa} = 1 \text{ kg m}^{-1} \text{ s}^{-2}$; $1 \text{ bar} = 10^5 \text{ Pa}$]

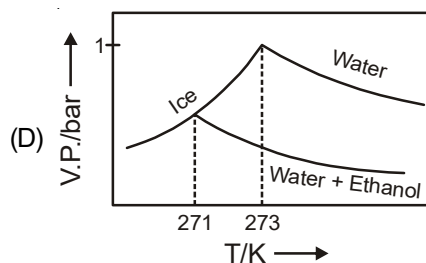
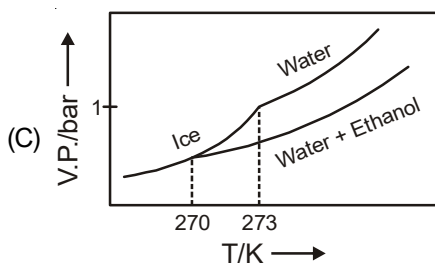
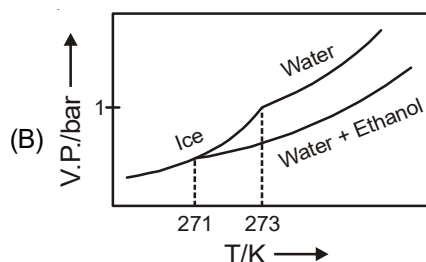
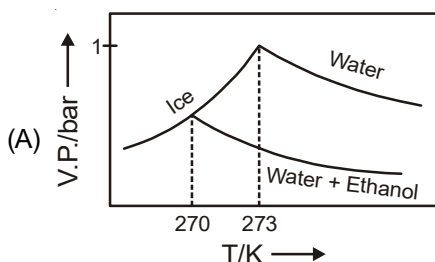
[JEE (Adv)-2017 (Paper-2)]

- (A) 14501 bar (B) 29001 bar
(C) 1450 bar (D) 58001 bar

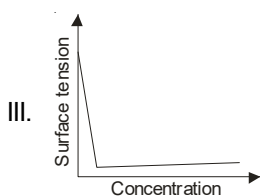
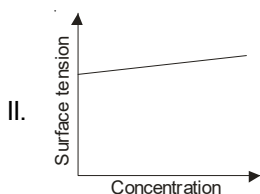
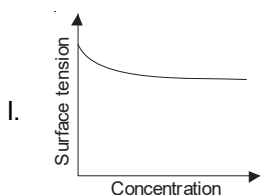
3. Pure water freezes at 273 K and 1 bar . The addition of 34.5 g of ethanol to 500 g of water changes the freezing point of the solution. Use the freezing point depression constant of water as 2 K kg mol^{-1} . The figures shown below represent plots of vapour pressure (V.P.) versus temperature (T). [molecular weight of ethanol is 46 g mol^{-1}]

Among the following, the option representing change in the freezing point is

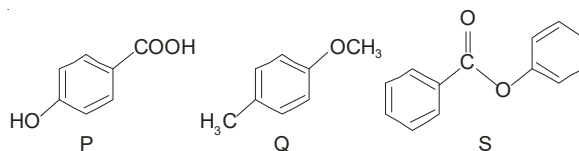
[JEE (Adv)-2017 (Paper-2)]



4. The qualitative sketches I, II and III given below show the variation of surface tension with molar concentration of three different aqueous solutions of KCl, CH_3OH and $\text{CH}_3(\text{CH}_2)_{11}\text{OSO}_3^-\text{Na}^+$ at room temperature. The correct assignment of the sketches is
[JEE (Adv)-2016 (Paper-2)]

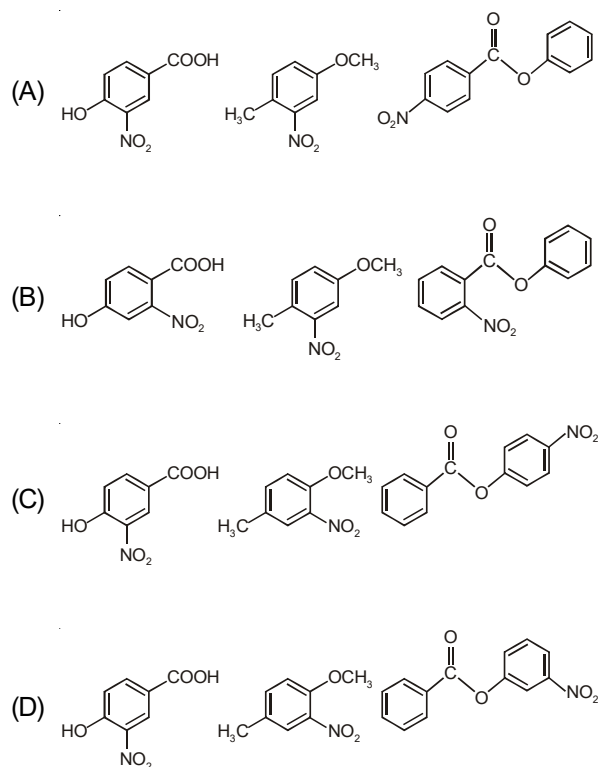


- (A) I : KCl
II : CH_3OH
III : $\text{CH}_3(\text{CH}_2)_{11}\text{OSO}_3^-\text{Na}^+$
- (B) I : $\text{CH}_3(\text{CH}_2)_{11}\text{OSO}_3^-\text{Na}^+$
II : CH_3OH
III : KCl
- (C) I : KCl
II : $\text{CH}_3(\text{CH}_2)_{11}\text{OSO}_3^-\text{Na}^+$
III : CH_3OH
- (D) I : CH_3OH
II : KCl
III : $\text{CH}_3(\text{CH}_2)_{11}\text{OSO}_3^-\text{Na}^+$
5. Among the following complexes (K – P), $\text{K}_3[\text{Fe}(\text{CN})_6](\text{K})$, $[\text{Co}(\text{NH}_3)_6]\text{Cl}_3(\text{L})$, $\text{Na}_3[\text{Co}(\text{oxalate})_3]^{-3}(\text{M})$, $[\text{Ni}(\text{H}_2\text{O})_6]\text{Cl}_2(\text{N})$, $\text{K}_2[\text{Pt}(\text{CN})_4](\text{O})$ and $[\text{Zn}(\text{H}_2\text{O})_6](\text{NO}_3)_2(\text{P})$, the diamagnetic complexes are
[IIT-JEE-2011 (Paper-2)]
- (A) K, L, M, N (B) K, M, O, P
(C) L, M, O, P (D) L, M, N, O
6. The compounds P, Q and S



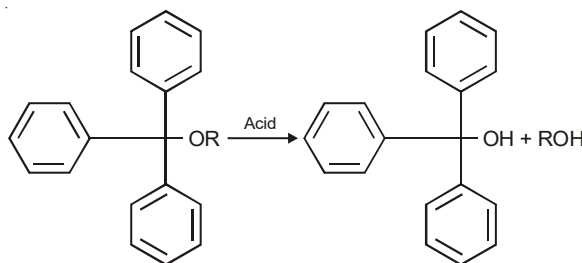
were separately subjected to nitration using $\text{HNO}_3/\text{H}_2\text{SO}_4$ mixture. The major product formed in each case respectively, is

[IIT-JEE-2010 (Paper-2)]

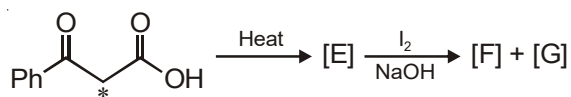


7. The acidic hydrolysis of ether (X) shown below is fastest when

[JEE (Adv)-2014 (Paper-2)]

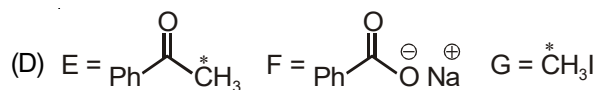
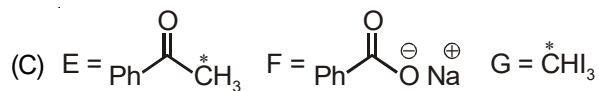
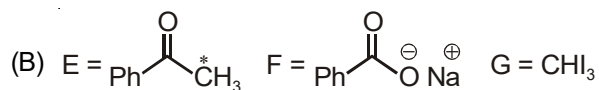
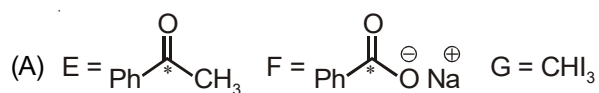


- (A) One phenyl group is replaced by a methyl group
 (B) One phenyl group is replaced by a *para*-methoxyphenyl group
 (C) Two phenyl groups are replaced by two *para*-methoxyphenyl groups
 (D) No structural change is made to X
8. In the following reaction sequence, the correct structures of **E**, **F** and **G** are

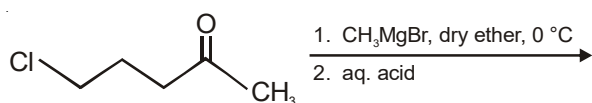


(* implies ^{13}C labelled carbon)

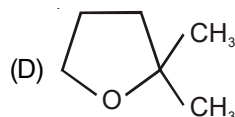
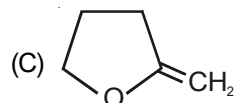
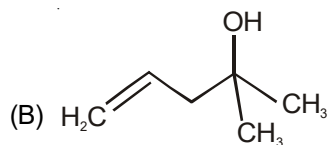
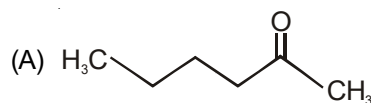
[IIT-JEE-2008 (Paper-2)]



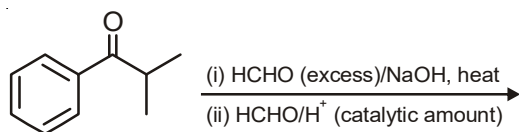
9. The major product in the following reaction is



[JEE (Adv)-2014 (Paper-2)]



10. The major product of the following reaction sequence is



[JEE (Adv)-2016 (Paper-2)]

