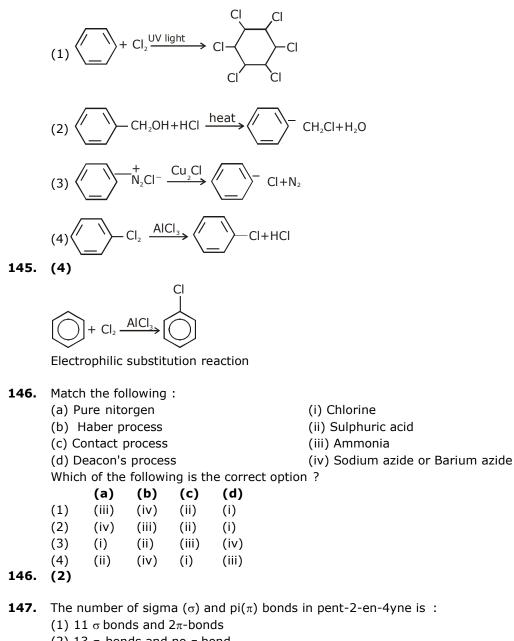
136. In which case change in entropy is negative? (1) Sublimation of solid to gas (2) 2H(g) \rightarrow H₂(g) (3) Evaporation of water (4) Expansion of a gas at constant temperature 136. (2) $2H(g) \longrightarrow H_2(g)$ No. of particle decreases from reactant to product side **137.** For the chemical reaction $N_2(g) + 3H_2(g) \square 2NH_3(g)$ the correct option is : $(1) -\frac{d[N_2]}{dt} = \frac{1 d[NH_3]}{2 dt}$ (2) $3\frac{d[H_2]}{dt} = 2\frac{d[NH_3]}{dt}$ $(3) -\frac{1}{3}\frac{d[H_2]}{dt} = -\frac{1}{2}\frac{d[NH_3]}{dt}$ $(4) -\frac{d[N_2]}{dt} = 2\frac{d[NH_3]}{dt}$ 137. (1) $N_2(g) + 3H_2(g) \Box 2NH_3(g)$ $r = -\frac{1}{2} \frac{d\left[\left[N_{2}\right]\right]}{dt} \frac{1}{3} \frac{d\left[\left[H_{2}\right]\right]}{dt} \frac{1}{2} \frac{d\left[\left[NH_{3}\right]\right]}{dt}$ $\therefore \frac{-d\left[\left\lfloor N_{2}\right.\right]\right]}{dt} \frac{1}{2} \frac{d}{dt} \left[\left\lfloor NH_{3}\right.\right]\right]}{dt}$ **138.** Which of the folloewing diatomic molecular species has only π bonds according to molecular Orbital Theory ? (1) C₂ (2) Be₂ (3) 0₂ (4) N₂ 138. (1) $12e^{-} \sigma 1s^{2} \sigma^{*}1s^{2} \sigma 2s^{2} \sigma^{*}2s^{2} \pi 2p_{x}^{2} = \pi 2p_{4}^{2}$ C_2 Bo = $\frac{8-4}{2}$ = 2 (where last 4 e⁻ present in $\pi 2px^2 = \pi 2p_4^2$) **139.** Which of the following is incorrect statement? (1) $GeX_4(x = F, CI, Br, I)$ is more stable than GeX_2 (2) SnF₄ is ionic in nature (3) PbF₄ is covalent in nature (4) SiCI₄ is easily hydrolysed 139. (3) Order of stability of OS IV A due to inert pair effect due to preudo inert gas configuration Ge4+ Ge Ge²⁺ < $\mathbf{\Lambda}$ Sn²⁺ < Sn4+ Sn ^ > Pb Pb²⁺ Pb⁴

140. Under isothermal condition a gas at 300 K expands from 0.1 L to 0.25 L against a constant external pressure of 2 bar. The work done by the gas is : [Given that 1 L bar = 100J] (1) 25 J (2) 30 J (3) -30 J (4) 5KJ 140. (3) $= -p_{ext} (V_2 - V_1)$ = -2(0.25 - 0.1) irreversible w isothermal = -2(0.15)expansion = -0.3 lt - bar $= -0.3 \times 100 \text{ J}$ = -30.1**141.** The compound that is most difficult to protonate is : (1) 0 H₃C (2) Ph H (4) H₃C (3) H H 141. (2) $\mathcal{O}_{\mathsf{Pb}}$ Lone pair one O-atom is delocalised in benzene ring 142. Which of the following is an amphoteric hydroxide ? (1) Mg(OH)₂ (2) Be(OH)₂ (4) $Ca(OH)_{2}^{-}$ (3) Sr(OH)₂ 142. (2) Be(OH), 143. The correct structure of tribromooctaoxide is : $\begin{array}{c} 0 & 0^{-} & 0 \\ 0 & 0^{-} & 0 \\ 0 & - & 0^{-} \\ 0 & 0^{-} & 0^{-} \end{array}$ $(3) \begin{array}{c} 0 & 0 & 0 \\ 0 = Br - Br - Br - Br = 0 \\ 0 \neq 1 & \infty \end{array}$ 143. (3) **144.** The biodegradable polymer is : (3) nylon-6,6 (4) nylon 2-nylon 6 (1) nylon-6 (2) Buna-S 144. (4) Nylon-2-Nylon - 6 is a biodegradable polymer

145. Among the following, the reaction that proceeds through an electrophilic substitution is :

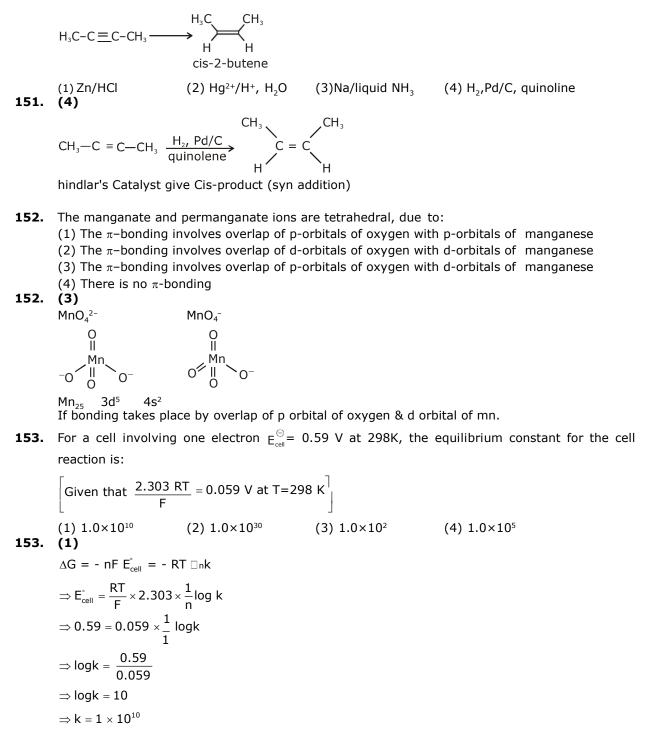


(2) 13 σ bonds and no π bond

- (3) 10 σ bonds and 3 π bonds
- (4) 8 σ bonds and 5 π bonds

147.	(3)
	σ/H
	$H \stackrel{\sigma}{=} C \stackrel{\cong}{=} C \stackrel{\sigma}{=} C \stackrel{\sigma}{=} C \stackrel{\sigma}{=} C \stackrel{\sigma}{=} C \stackrel{\sigma}{=} H$ $H \stackrel{\sigma}{=} H \stackrel{\sigma}{=} H$
	10σ and 3π – bonds Note : Given IUPAC Name is Incorrect
148.	Enzymes that utilize ATP in phosphate transfer require an alkaline earth metal (M) as the cofactor M is :
	(1) Ca (2) Sr (3) Be (4) Mg
148.	(4) Mg is required in enzymes that utilize ATP in phosphate transfer.
149.	, j j
	 (1) Axial P–Cl bonds are longer than equatorial P–Cl bonds (2) PCl₅ molecule is non-reactive
	(3) Three equatorial P-Cl bonds make an angle of 120° with each other (4) Two axial P-Cl bonds make an angle of 180° with each other
149.	
150.	If the rate constant for a first order reaction is k, the time (t) required for the completion of 99%
	of the reaction is given by :
150.	(1) $t=4.606/k$ (2) $t=2.303/k$ (3) $t=0.693/k$ (4) $t=6.909/k$ (1)
	for 1st order reaction -
	$\Box_{n} \frac{C_{0}}{C_{t}} = kt \qquad \qquad C_{t} = \frac{1}{100} C_{t}$
	$\Rightarrow \Box_n \frac{c_o}{\left(\frac{1C_o}{100}\right)} = kt$ as 99% of reactant is consumed
	$\left(\frac{1C_0}{100}\right)$ as 99% of reactant is consumed
	$\Rightarrow \Box n100 = kt$
	$\Rightarrow t = \frac{1}{\kappa} \times 2.303 \text{ log } 10^2$
	$\Rightarrow t = \frac{1}{K} \times 2.303 \times 2 \times \log 10$
	$\Rightarrow t = \frac{4.606}{K}$
	K

151. The most suitable reagent for the following conversion, is :

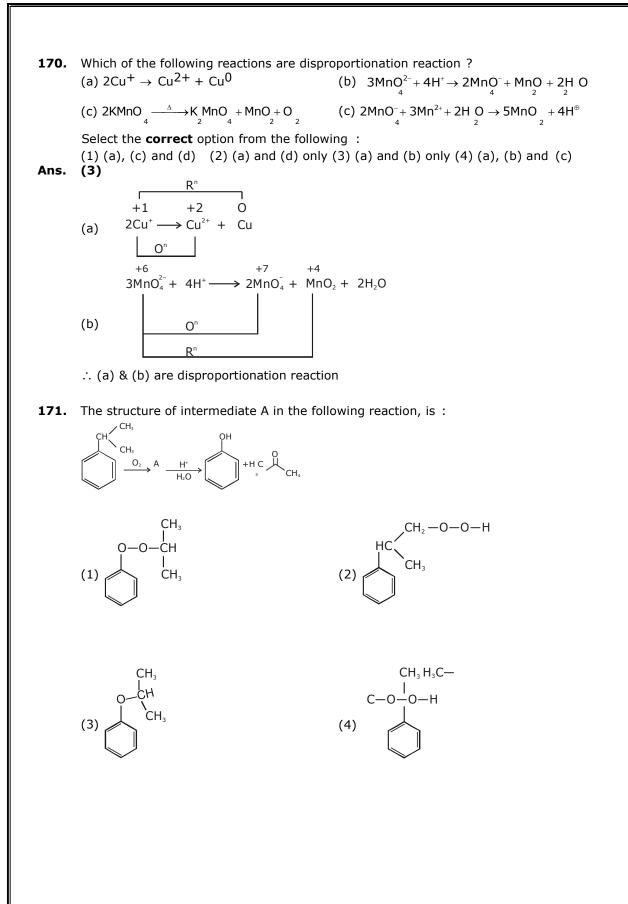


154. pH of a saturated solution of $Ca(OH)_2$ is 9. The solubility product (K_{sp}) of $Ca(OH)_2$ is : (1) 0.125×10^{-15} (2) 0.5×10^{-10} (3) 0.5×10^{-15} (4) 0.25×10^{-15} Ans. (3) pH = 9∴ pOH = 14 - 9 = 5 $[OH^{-}] = 10^{-5}$ $Ca(OH)_{2}(s) \square Ca^{2+}(aq) + 2OH^{-}(aq)$ 2S = 10⁻⁵ S : $ksp = [Ca^{2+}] [OH^{-}]^{2}$ $= S \times (2S)^{2}$ $=\frac{10^{-5}}{2}\times(10^{-5})^2$ $= 0.5 \times 10^{-15}$ **155.** For an ideal solution the correct option is : (2) Δ_{mix} G =0 at constant T and P (1) Δ_{mix} H =0 at constant T and P (3) $\Delta_{mix} S = 0$ at constant T and P (4) $\Delta_{mix} V \neq 0$ at constant T and P Ans. (1) factual 156. A gas at 350 K and 15 bar has molar volume 20 percent smaller than that for an ideal gas under the same conditions. The correct option about the gas and its compressibility factror (Z) is: (1) Z < 1 and attractive forces are dominant (2) Z < 1 and repulsive forces are dominant (3) Z > 1 and attractive forces are dominant (4) Z > 1 and repulsive forces are dominant Ans. (1) $z = \frac{(PV)_{real}}{(PV)_{ideal}}$ as real volume is lesser than an ideal gas volume ∴ z < 1 and for z < 1, attractive forces dominant

157. The correct order of the basic strength of methyl substituted amines in aqueous solution is : (1) $(CH_3)_3N > (CH_3)_2NH > CH_3NH_2$ (2) $CH_3NH_2 > (CH_3)_2NH > (CH_3)_3N$ (3) $(CH_3)_2NH > CH_3NH_2 > (CH_3)_3N$ (4) $(CH_3)_3N > CH_3NH_2 > (CH_3)_2NH$ Ans. (3) Correct order of basic strength of methyl substituted amines is $(CH_3)_2NH > CH_3NH_2 > (CH_3)_3N (2^\circ > 1^\circ > 3^\circ)$ **158.** For the second period elements the correct increasing order of first ionisation enthalpy is: (1) Li < B < Be < C < N < O < F < Ne (2) Li < Be< B < C < O < N < F < Ne (3) Li < Be < B < C < N < O < F < Ne (4) Li < B < Be < C < O < N < F < Ne Ans. (4) **159.** Which mixture of the solutions will lead to the formation of negatively charged colloidal [AgI]I⁻ Ans? (1) 50 mL of 2 M AgNO₃ + 50 mL of 1.5 M KI (2) 50 mL of 0.12 M AgNO₃ + 50mL of 0.1 M KI (3) 50 mL of 1 M AgNO₃ + 50 mL of 1.5 M KI (4) 50 mL of 1 M AgNO₃ + 50 mL of 2M KI Ans. (3,4) If in KI solution AgNO₃ is added than it will form –vely charged $[AgI]I^-$ colloid. If in AgNO₃ solution KI is added it will form AgI/Ag⁺ (positively charged) colloid. **160.** For the cell reaction $2Fe^{3+}(aq) + 2I^{-}(aq) \longrightarrow 2Fe^{2+}(aq) + I_2(aq)$ E_{coll}^{Θ} 0.24V at 298K. The standard Gibbs energy ($\Delta_{r} G^{\Theta}$) of the cell reaction is: (Given that Faraday constant $F = 96500 \text{ C} \text{ mol}^{-1}$) (1) 46.32 kJ mol⁻¹ (2) 23.16 kJ mol⁻¹ (3) -46.32 kJ mol⁻¹ (4) -23.16 kJ mol⁻¹ Ans. (3) $\Delta G^{o}_{cell} = -nFE^{o}_{cell}$ = -2 × 96500 × 0.24 = -46320 J/Mol = -46.32 J/Mol **161.** Which is the correct thermal stability order for H_2E (E = 0, S, Se, Te and Po)? (1) $H_2Po < H_2Te < H_2Se < H_2S < H_2O$ (2) $H_2Se < H_2Te < H_2Po < H_2O < H_2S$ (3) $H_2S < H_2O < H_2Se < H_2Te < H_2Po$ (4) $H_2O < H_2S < H_2Se < H_2Te < H_2Po$ Ans. (1)

162.		s of hydrogen molecu	les required to produc	ce 20 moles of ammonia through
A =c	Haber's process is: (1) 30	(2) 40	(3) 10	(4) 20
Ans.	(1) $N_2(g) + 3H_2(g) \longrightarrow$ $\frac{n_{H_2}}{3} = \frac{n_{NH_3}}{2}$ $\Rightarrow n_{H_2} = \frac{3}{2} \times$	$2NH_3(g)$ $20 \Rightarrow n_{H_2} = 30 \text{ moles}$		
163.	Which of the followi region?	ng series of transition	s in the spectrum of h	ydrogen atom falls in visible
Ans.	-	(2) Brackett series	(3) Lyman series	(4) Balmer series
164.				m hexagonal close packed (hcp) mula of the compound is:
Ans.	(1) C_3A_4 (1) (c) OV :	(2) C ₄ A ₃ hcp(A)	(3) C ₂ A ₃	(4) C ₃ A ₂
	$6 \times \frac{75}{100}$:	6		
	$\begin{array}{c} \frac{3}{4} & \vdots \\ 3 & \vdots \end{array}$	1 4		
165. Ans.	The non-essential and (1) alanine (1)	mino acid among the t (2) lysine	following is: (3) valine	(4) leucine
166.	Addition of HCl to a	lkene "A" gives "B" as	the major product. T	ne and ethanal in equimolar ratio. The structure of product "B" is:
	CH₃ I (1) H₃C - CH₂ - C I CI	CH₃	CH₃ I (2) H₃C – CH – CH I CI CH₃	
	(3) CI - CH ₂ - CH ₂ -	CH ₃ I CH CH CH ₃	CH ₂ (4) H ₃ C - CH ₂ - CH -	Cl - CH₃

(1)										
$A \xrightarrow{(i) O_3} + CH_3CHO \\ O \\$										
So, (A) shou	ıld be	\succ							
	、+ HC	1	→)® }_							
(1) [cies is not s SiCl ₆] ²⁻		SiF ₆] ²⁻		(4) [(GeCl ₆] ²⁻	
						SiF ₆] ²⁻		(4) [(GeCl ₆] ²⁻	
(1) [3 (2) Matcl	Sn(OH) n the X	6 ^{]2-}	(2) [SiCl ₆] ²⁻	(3) [GeCl ₆] ²⁻ n-II and assi	ın the
(1) [3 (2) Matcl code Colu (a) X (b) X	Sn(OH) n the Xi : mn –I eF ₄ eF ₆	6 ^{]2-}	(2) [GiCl ₆] ²⁻ ds in Colum Column (i) Pyram (ii) squar	(3) [: nn–I with i – II nidal re planar	ts struc				jn the
(1) [3 (2) Matcl code Colu (a) X (b) X (c) X (d) X	Sn(OH) n the Xi : mn –I eF ₄ eF ₆ eOF ₄ eO ₃	6 ^{]2-}	(2) [SiCl ₆] ²⁻ ds in Colum Column (i) Pyram (ii) squar (ii) Disto	(3) [: nn–I with i – II nidal	ts struc edral				յn the
(1) [3 (2) Matcl code (a) X (b) X (c) X (d) X Code (1) (3)	Sn(OH) in the Xi imm –I eF4 eF6 eOF4 eO3 :	₆] ^{2–} enon co	(2) [9	SiCl ₆] ²⁻ ds in Colum Column (i) Pyram (ii) squar (ii) Disto	(3) [: nn–I with i – II nidal re planar orted octah nre Pyramic	ts struc edral dal	ture in	Columr	n-II and assi	jn the
(1) [3 (2) Matcl code Colu (a) X (b) X (c) X (d) X Code (1) (3) (4)	Sn(OH) n the Xi : mn – I eF ₄ eOF ₄ eO ₃ : (a) (ii) (i)	6 ^{]2-} enon co (b) (iii) (ii)	(2) [9 mpoun (c) (i) (iii)	GiCl ₆] ^{2–} ds in Colum (i) Pyram (ii) squar (iii) Distc (iv) Squa (d) (iv)	(3) [: nn–I with in – II nidal re planar orted octah are Pyramic (2) (4)	edral dal (a) (iii) (ii)	ture in (b) (iv) (iii)	Columr (c) (i) (iv)	n-II and assid (d) (ii)	յn the



Ans. (4)

