81. The energies of activation for forward and reverse reactions for $A_2 + B_2 \implies 2AB$ are 180 kJ mol⁻¹ and 200 kJ mol⁻¹ respectively. The presence of catalyst lowers the activation energy of both (forward and reverse) reactions by 100 kJ mol⁻¹. The enthalpy change of the reaction $(A_2 + B_2 \implies 2AB)$ in the presence of catalyst will be (in kJ mol⁻¹)

	pH = 4.5		
	pH + pOH = 14		
	\Rightarrow pOH = 14 – 4.5 = 9.5		
	Hence (3) is correct.		
84.	Consider the reaction,		
	$2A + B \rightarrow Products$		
	When concentration of B alone was doubled, the half-life did not change. When the concentration of A		
	alone was doubled, the rate increased by two times. The unit of rate constant for this reaction is		
	(1) $L mol_{1}^{-1} s_{1}^{-1}$ (2) no unit		
	(3) mol $L^{-1} s^{-1}$ (4) s^{-1}		
Ans.	(1)		
Sol.	$2A + B \rightarrow Product$		
	When conc. of B is doubled, the half life did not change, hence reaction is of first order w.r.t. B.		
	When concentration of A is doubled, reaction rate is doubled, hence reaction is of first order w.r.t. A.		
	Hence over all order of reaction is $1 + 1 = 2$		
	So, unit of falle constant more interest.		
85	Identify the incorrect statement among the following		
00.	(1) d-Block elements show irregular and erratic chemical properties among themselves		
	(2) La and Lu have partially filled d orbitals and no other partially filled orbitals		
	(3) The chemistry of various lanthanoids is very similar		
	(4) 4f and 5f orbitals are equally shielded		
Ans.	(4)		
Sol.	4f and 5f belongs to different energy levels, hence the shielding effect is on them is not the same.		
	Shielding of 4f is more than 5f.		
	Hence (4) is correct.		
86.	Which one of the following has a square planar geometry?		
	(1) $[CoCl_4]^{2^-}$ (2) $[FeCl_4]^{2^-}$		
	(2) $[NiCL]^{2^{-}}$ (4) $[PtCL]^{2^{-}}$		
	$(3) [\operatorname{INIOI}_4] \qquad (4) [\operatorname{PIOI}_4]$		
Ans.	(4)		
501.	$_{27}C0^{-}$ - 1s ⁻ 2s ⁻ 2p ⁻ 3s ⁻ 3p ⁻ 30 ⁻ 4s ⁻		
	4s $4p$		
	As Cl ⁻ is weak field ligand so no pairing up.		
	Hence it is sp ³ hybridized giving tetrahedral geometry.		
	$Fe^{2\tau} - 1s^2 2s^2 2p^0 3s^2 3p^0 3d^0 4s^0$		
	<u> </u>		
	Due to Cl ⁻ , back pairing is not observed so it will be sp^3 hybridized giving tetrahedral geometry.		
	$Ni^{2+} - 1s^2 2s^2 2p^6 3s^2 3p^6 3d^8 4s^0$		
	3d 4s 4p		
	Descuse week linear heat activity is not choose at a it will be an ³ is total advelopments.		
	Because weak ligand, back pairing is not observed so it will be splite, tetranedral geometry.		
	thus (4) is correct		
87	Which of the following molecules is expected to rotate the plane of plane polarized light?		
07.	(1) CHO (2)		
	но—— ——н sh		
	с́н₂он		

	(3) H ₂ N	NH ₂	(4)	соон
	н)—	——(н		
	Ph	Ph	H_2N —	— —H
				H
Ans. Sol.	 (1) The plane of polarized light is rotated by optically active compound, i.e. it should be chiral. So, (1) has, chiral C-atom. So, it is optically active. In (2), (3) and (4) plane of symmetry is present. Hence, (1) is correct. 			
88.	The secondary st (1) α -helical back (3) sequence of	ructure of a protein ref kbone α-amino acids	fers to (2) hydrophobi (4) fixed config	ic interactions guration of the polypeptide backbone
Ans. Sol.	(1) Secondary struct structures are for Hence, (1) is corr	ure of proteins involves med as a result of H-b ect	s α – helical back bond a onding between different	and β – sheet structures. These t peptide groups.
89.	Which of the follo (1) $CH_3 - C \equiv CH_3$	wing reactions will yiel 1+2HBr ——→	ld 2, 2-dibromopropane? (2) $CH_3CH = C$	CHBr + HBr ───→
Ans.	$(3) CH \equiv CH + 2h$ (1)	lBr —→	(4) CH ₃ – CH =	$= CH_2 + HBr \longrightarrow$
Sol.	$CH_3-C\equivCH+H$	HBr <u>electrophilic addition of H</u> (from HBr)	$\xrightarrow{I^+} CH_3 - C = CH_2 \xrightarrow{HBr}_{Br}$	$\rightarrow CH_3 - C - CH_3$
	Hence, (1) is corr	ect.	Di	Di
90.	In the chemical re $CH_3CH_2NH_2 - CH_3CH_2NH_2 - CH_3CH_2NH_2$ (1) C_2H_5CN and (3) C_2H_5NC and	ection, ICl₃ + 3KOH→→(A) 3KCl K₅CΩ₅	$+(B)+3H_2O$, the compo (2) CH ₃ CH ₂ CC (4) C ₂ H ₂ NC an	und (A) and (B) are respectively DNH₂ and 3KCl nd 3KCl
Ans. Sol.	(4) It is example of c Hence, (4) is the	arbylamine reaction. so	o, the product will be C_2 H	l₅NC and KCl.
91.	The reaction of to (1) benzoyl chlor	luene with Cl ₂ in prese ide	ence of FeCl ₃ gives predo (2) benzyl chlo	ominantly oride
Ans.	(3) (3)	otoldene		
Sol.	Due to o- and p-	directing nature of CH ₃	₃ group. CH₂	
	CH ₃ —Cl ₂ FeC	GH ₃ Cl +	CI	
92.	Hence, (3) is corr Presence of a niti (1) activates the (2) renders the ri (3) deactivates the (4) deactivates the	ect answer. ro group in a benzene ring towards electroph ng basic ne ring towards nucleo ne ring towards electro	ring ilic substitution philic substitution	
Ans. Sol.	(4) - NO ₂ group shov	vs – M effect, so withdr	raws the electron density	r from the ring and hence deactivate

the ring towards electrophilic aromatic substitution. Hence, (4) is correct. 93. In which of the following ionization processes, the bond order has increased and the magnetic behaviour has changed?

(2) NO-

→NO⁺

(1) $C_2 \longrightarrow C_2^+$

(3)
$$O_2 \longrightarrow O_2^+$$
 (4) $N_2 \longrightarrow N_2^+$

Ans.

(2)

Sol. $\ln C_2 - C_2^+$ electron is removed from bonding molecular orbital so bond order decreases. In NO -----> NO⁺, electron is removed from anti bonding molecular orbital so bond order increases and nature changes from paramagnetic to diamagnetic.

Hence, (2) is correct.

- 94. The actinoids exhibits more number of oxidation states in general than the lanthanoids. This is because
 - (1) the 5f orbitals are more buried than the 4f orbitals
 - (2) there is a similarity between 4f and 5f orbitals in their angular part of the wave function
 - (3) the actinoids are more reactive than the lanthanoids
 - (4) the 5f orbitals extend further from the nucleus than the 4f orbitals (4)
- Ans.
- Sol. The actinoids exhibit more number of oxidation states in general than the lanthanoids. This is because the 5f orbitals extend further from the nucleus than the 4f orbitals. Hence, (4) is correct.
- 95. Equal masses of methane and oxygen are mixed in an empty container at 25°C. The fraction of the total pressure exerted by oxygen is

(1)	$\frac{2}{3}$	(2)	$\frac{1}{3} \times \frac{273}{298}$
(3)	$\frac{1}{3}$	(4)	<u>1</u> 2
(3)			

Ans.

Sol. Let the mass of methane and oxygen is w

mole fraction of oxygen =
$$\frac{\frac{W}{32}}{\frac{W}{32} + \frac{W}{16}}$$

 $\frac{1}{22}$ $\frac{1}{22}$ 1

$$=\frac{\overline{32}}{\frac{1}{32}+\frac{1}{16}}=\frac{\overline{32}}{\frac{3}{32}}=\frac{1}{3}$$

Let the total pressure be P

The pressure exerted by oxygen (partial pressure) = $X_{O_2} \times P_{total}$

$$\Rightarrow P \times \frac{1}{3}$$

Hence, (3) is correct.

A 5.25 % solution of a substance is isotonic with a 1.5% solution of urea (molar mass = 60 g mol⁻¹) in 96. the same solvent. If the densities of both the solutions are assumed to be equal to 1.0 g cm⁻³, molar mass of the substance will be

(1)	90.0 g mol ⁻¹	(2) 115.0 g mol ⁻
(3)	105.0 g mol ⁻¹	(4) 210.0 g mol ⁻

Ans.

(4)Solutions with the same osmotic pressure are isotonic Sol. Let the molar mass of the substance be M $\pi_1 = C_1 RT = C_2 RT = \pi_2$ So, $C_1 = C_2$

As density of the solutions are same So $\frac{5.25}{M} = \frac{15}{60}$

 $M = \frac{5.25 \times 60}{1.5} = 210$ Hence (4) is correct

- 97. Assuming that water vapour is an ideal gas, the internal energy (AU) when 1 mol of water is vapourised at 1 bar pressure and 100°C, (Given: Molar enthalpy of vapourization of water at 1 bar and 373 K = 41 kJ mol⁻¹ and R = 8.3 J mol⁻¹K⁻¹) will be (1) 4.100 kJ mol⁻¹ (2) 3.7904 kJ mol⁻¹ (3) 37.904 kJ mol⁻¹ (4) 41.00 kJ mol⁻¹ Ans. (3) $H_2O(\ell) \xrightarrow{vaporisation} H_2O(g)$ Sol. $\Delta n_{a} = 1 - 0 = 1$ $\Delta H = \Delta U + \Delta n_{a}RT$ $\Delta U = \Delta H - \Delta n_{_{\rm G}} R T$ $= 41 - 8.3 \times 10^{-3} \times 373$ = 37.9 kJ mol⁻¹ Hence, (3) is correct. 98. In a sautrated solution of the sparingly soluble strong electrolyte AqIO₃ (Molecular mass = 283) the
- equilibrium which sets in is

 $\textbf{AgIO}_{3(s)} \xleftarrow{} \textbf{Ag}^{\scriptscriptstyle +}_{(aq)} + \textbf{IO}^{\scriptscriptstyle -}_{3(aq)}$

(1) 28.3×10^{-2} g

(3) 1.0×10^{-7} g

If the solubility product constant K_{sp} of AgIO₃ at a given temperature is 1.0×10^{-8} , what is the mass of AgIO₃ contained in 100 ml of its saturated solution? (2) 2.83×10^{-3} g

(4) 1.0×10^{-4} g

Ans.

(2)

Sol. $AgIO_3(s) \Longrightarrow Ag^+(aq) + IO_3^-(aq)$ Let the solubility of AgIO₃ be s $K_{sp} = \left\lceil Ag^{+} \right\rceil \left\lceil IO_{3}^{-} \right\rceil$ $1.0 \times 10^{-8} = s^2$ $s = 10^{-4}$ mol/litre $= \frac{10^{-4} \times 283}{1000} \times 100$ = 283 × 10⁻⁵ = 2.83 × 10⁻³ g/ 100 ml Hence, (2) is correct.

- A radioactive element gets spilled over the floor of a room. Its half-life period is 30 days. If the initial 99. activity is ten times the permissible value, after how many days will it be safe to enter the room? (1) 1000 days (2) 300 days (3) 10 days (4) 100 days
- Ans.

Activity $\left(-\frac{dN}{dt}\right) \propto N$ Sol. $N = N_o \left(\frac{1}{2}\right)^n$ N $(1)^{n}$

(4)

$$\frac{1}{N_o} = \left(\frac{1}{2}\right)$$
$$\frac{1}{10} = \left(\frac{1}{2}\right)^n \Rightarrow 10 = 2^n$$
$$\log 10 = n \log 2$$
$$\Rightarrow n = \frac{1}{0.301} = 3.32$$

 $t = n \times t_{112}$

100. Ans. Sol.	 = 3.32 × 30 = 99.6 days Hence, (4) is correct. Which one of the following conformation of cycle (1) Twist boat (3) Chair (1) Twisted boat is chiral as it does not have plane Hence, (1) is correct. 	ohexane is chiral? (2) Rigid (4) Boat of symmetry.	
101.	Which of the following is the correct order of dec (1) $RCH_X > R_CX > R_CHX$	creasing SN ² reactivity? (2) RCH ₂ X > R ₂ CHX > R ₂ CX	
	(3) $R_3CX > R_2CHX > RCH_2X$	(4) $R_2CHX > R_3CX > RCH_2X$	
	(X = a halogen)		
Ans. Sol.	(2) More is the steric hindrance at the carbon bearing the halogen, lesser is the S_N2 reactivity. Hence, (2) is correct.		
102.	In the following sequence of reactions, $CH_3CH_2OH \xrightarrow{P+l_2} A \xrightarrow{Mg} B \xrightarrow{HCHO} C \xrightarrow{H_2C}$	D→D	
A pc	the compound 'D' is (1) butanal (3) n-propyl alcohol	(2) n-butyl alcohol(4) propanal	
Ans.	(3) $CH_{2}CH_{3}OH \xrightarrow{P+l_{2}} CH_{2}CH_{3}I \xrightarrow{Mg} CH_{2}CH_{3}CH_{3}$	Mal	
Sol.	(A) (B)		
	H-C=O		
	$\longrightarrow CH_3 - CH_2 - CH_2 OMgI \xrightarrow{120} CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3 CH_3$	$H_2CH_2OH + Mg(OH)I$	
	(C) (D) ∴ the compound D is n-propyl alcohol. Hence, (3) is correct option.		
103.	Which of the following sets of quantum numbers (1) $n = 3$, $l = 2$, $m = 1$, $s = +1/2$	represents the highest energy of an atom? (2) $n = 3$, $l = 2$, $m = 1$, $s = +1/2$	
	(3) $n = 4, l = 0, m = 0, s = +1/2$	(4) $n = 3, l = 0, m = 0, s = +1/2$	
Ans. Sol.	(2)(2) is the correct option because it has the maxHence, (2) is correct.	imum value of n + ℓ	
104.	Which of the following hydrogen bonds is the str (1) O–HN	rongest? (2)F–HF	
Ans.	(3) O–HO (2)	(4) O–HF	
Sol.	The hydrogen bond in HF is strongest, because fluorine is the most electronegative element. Thus, (2) is the correct option.		
105.	In the reaction. $2AI_{(s)} + 6HCI_{(s)} \longrightarrow 2AI^{s_+}_{(aq)} + 6$	$CF_{(aq)}^{-} + 3H_{2(g)}^{-},$	
Ans.	 6 L HCl_(aq) is consumed for every 3L H_{2(g)} produced 33.6 L H_{2(g)} is produced regardless of temperature and pressure for every mole AI that reacts 67.2 L H_{2(g)} at STP is produced for every mole AI that reacts 11.2 H_{2(g)} at STP is produced for every mole HCl_(aq) consumed (4) 		
Sol.	$2AI(s) + 6HCI(aq) \longrightarrow 2AI^{3+}(aq) + 6CI^{-}(aq) +$	$-3H_{2}(g)$	
	For each mole of HCl reacted, 0.5 mole of H ₂ ga 1 mole of an ideal gas occupies 22.4 lit at STP. Volume of H ₂ gas formed at STP per mole of HCl re Hence, (4) is correct.	as is formed at STP. acted is 22.4 × 0.5 litre	
106.	Regular use of which of the following fertilizer in (1) Potassium nitrate (3) Superphosphate of lime	creases the acidity of soil? (2) Urea (4) Ammonium sulphate	

Ans. (4)(NH₄)₂ SO₄ is a salt of strong acid and weak base, on hydrolysis it ill produce H⁺ ion. This will Sol. increase the acidity of soil. $(NH_4)_2 SO_4 \longrightarrow 2NH_4^+ + SO_4^{2-}$ $NH_4^+ + H_2O \Longrightarrow NH_4OH + H^+$ Hence, (4) is correct answer. 107. Identify the correct statement regarding a spontaneous process (1) For a spontaneous process in an isolated system, the change in entropy is positive (2) Endothermic processes are never spontaneous (3) Exothermic processes are always spontaneous (4) Lowering of energy in the reaction process is the only criterion for spontaneity Ans. (1)Sol. For a spontaneous process in an isolated system, the change in entropy is positive. Hence, (1) is correct. 108. Which of the following nuclear reactions will generate an isotope? (1) neutron particle emission (2) positron emission (3) α-particle emission (4) β-particle emission Ans. (1) $^{A}_{z}X \longrightarrow ^{A-1}_{z}X +^{1}_{0}n$ Sol. Hence, (1) is correct. 109. The equivalent conductances of two strong electrolytes at infinite dilution in H₂O (where ions move freely through a solution) at 25°C are given below: $\wedge^{\circ}_{CH_{3}COONa} = 91.0 \text{ S cm}^{2} / equiv$ $^{\circ}_{HCl} = 426.2 \text{ S cm}^2 / \text{equiv}$ What additional information/quantity one needs to calculate \wedge° of an aqueous solution of acetic acid? (1) ∧° of NaCl (2) ∧° of CH₃COOK (3) The limiting equivalent conductance of $H^+(\wedge^{\circ}_{\mu^+})$ (4) ∧° of chloroacetic acid (C/CH₂COOH) Ans. (1)From Kohlrausch's law Sol. $\Lambda_{\mathsf{CH}_3\mathsf{COOH}}^{\circ} = \Lambda_{\mathsf{CH}_3\mathsf{COONa}}^{\circ} + \Lambda_{\mathsf{HCI}}^{\circ} - \Lambda_{\mathsf{NaCI}}^{\circ}$ Hence, (1) is the correct answer. 110. Which one of the following is the strongest base in aqueous solution? (1) Trimethylamine (2) Aniline (3) Dimethylamine (4) Methylamine Ans. (3)In aqueous solution basicity order of 1°, 2° and 3° amine with methyl group is Sol. $2^{\circ} > 1^{\circ} > 3^{\circ}$ In case of aniline lone pair of nitrogen is involved in resonance, so it is weaker base than aliphatic amines. Hence, (3) is correct. 111. The compound formed as a result of oxidation of ethyl benzene by KMnO₄ is (2) acetophenone (1) benzophenone (4) benzyl alcohol (3) benzoic acid Ans. (3)Any aliphatic carbon with hydrogen attached to it, in combination with benzene ring, will be oxidized to Sol. benzoic acid by KMnO₄/H⁺. Hence, (3) is correct.

112.	The IUPAC name of	is
	\frown	
Ans. Sol.	(1) 1, 1-diethyl-2,2-dimethylpentane (3) 5, 5-diethyl-4, 4-diemthylpentane (4) 7 6 7 2 1 5 3 1	 (2) 4, 4-dimethyl-5, 5-diethylpentane (4) 3-ethyl-4, 4-dimethylheptane
	The correct answer is 3-ethyl-4, 4-dimethylhepta	ane.
113.	Which of the following species exhibits the diam (1) $O_2^{2^-}$	agnetic behaviour? (2) O ₂ ⁺
	(3) O ₂	(4) NO
Ans.	(1) The correct ention is Ω^{2-}	
501.	This species has $18 e^{-1}$ which are filled in such a	a way that all molecular orbitals are fully filled, so
	diamagnetic. σ 1s ² σ *1s ² , σ 2s ² σ *2s ² , σ 2p _z ² , π 2p _x ² = π 2p _y ² , π *2p _x ²	$= \pi^* 2p_y^2$
111	Hence, (1) is correct.	process standily in the acqueres
114.	(1) $GeX_2 \ll SiX_2 \ll SnX_2 \ll PbX_2$	(2) $SiX_2 \ll GeX_2 \ll PbX_2 \ll SnX_2$
	(3) $\operatorname{SiX}_2 \ll \operatorname{GeX}_2 \ll \operatorname{SnX}_2 \ll \operatorname{PbX}_2$	(4) $PbX_2 \ll SnX_2 \ll GeX_2 \ll SiX_2$
Ans.	(3)	
Sol.	Due to inert pair effect, the stability of +2 oxidati \therefore SiX ₂ \ll GeX ₂ \ll SnX ₂ \ll PbX ₂	on state increases as we move down this group.
115.	Hence, (3) is correct. Identify the incorrect statement among the follow (1) Ozone reacts with SO_2 to give SO_3 (2) Silicon reacts with $NaOH_{(aq)}$ in the presence	ving of air to give Na $_2$ SiO $_3$ and H $_2$ O
	(3) Cl_2 reacts with excess of NH_3 to give N_2 and (4) Br_2 reacts with hot and strong NaOH solutio	l HCl n to give NaBr. NaBrO₄ and H₂O
Ans.	(4)	
Sol.	Br_2 reacts with hot and strong NaOH to give NaB Hence (4) is incorrect statement	Br, NaBrO ₃ and H ₂ O.
116.	The charge/size ratio of a cation determines sequences represents the increasing order of cationic species, K^+ , Ca^{2+} , Mg^{2+} , Be^{2+} ?	s its polarizing power. Which one of the following the polarizinig order of the polarizing power of the
	(1) Mg ²⁺ ,Be ²⁺ ,K ⁺ ,Ca ²⁺	(2) Be ²⁺ ,K ⁺ ,Ca ²⁺ ,Mg ²⁺
	(3) K ⁺ ,Ca ²⁺ ,Mg ²⁺ ,Be ²⁺	(4) Ca ²⁺ ,Mg ²⁺ ,Be ²⁺ ,K ⁺
Ans. Sol.	(3) Higher the charge/size ratio, more is the polarizi $\therefore K^+ < Ca^{2+} < Mg^{2+} < Be^{2+}$	ng power.
	Hence, (3) is correct.	
117.	The density (in g mL ^{-1}) of a 3.60 M sulphuric mol ^{-1}) by mass will be	acid solution that is 29% H_2SO_4 (Molar mass = 98 g
	(1) 1.64	(2) 1.88
Ans.	(3) 1.22	(4) 1.45
Sol.	Let the density of solution be 'd' Molarity of solution given = 3.6 i.e. 1 litre of solution contains 3.6 moles of H_2SC or 1 litre of solution contains 3.6 × 98 gms of H_2S Since, the solution is 29% by mass.	D ₄ SO ₄

100 gm solution contains 29 gm H₂SO₄ $\frac{100}{d}$ mI solution contains 29 gm of H₂SO₄ 1000 ml solution contains 3.6×98 gm of H₂SO₄ $\therefore 3.6 \times 98 = \frac{29 \times d}{100} \times 1000$ d = 1.22Hence, (3) is correct. The first and second dissociation constants of an acid H₂A are 1.0×10^{-5} and 5.0×10^{-10} respectively. 118. The overall dissociation constant of the acid will be (1) 5.0×10^{-5} (2) 5.0×10^{15} (3) 5.0×10^{-15} (4) 0.0×10^5 Ans. (3) $H_{2}A \xrightarrow{} HA^{-} + H^{+} \qquad K_{1} = \frac{\left[HA^{-}\right]\left[H^{+}\right]}{\left[H_{2}A\right]}$ Sol. ...(1) $HA^{-} \xrightarrow{} H^{+} + A^{2-} \qquad K_{2} = \frac{\left[H^{+}\right]\left[A^{2-}\right]}{\left[HA^{-}\right]}$...(2) For the reaction $H_2A = 2H^+ + A^{2-}$ $\mathsf{K} = \frac{\left[\mathsf{H}^{+}\right]^{2} \left[\mathsf{A}^{2^{-}}\right]}{\left[\mathsf{H}_{2}\mathsf{A}\right]} = \mathsf{K}_{1} \times \mathsf{K}_{2}$ $= 1 \times 10^{-5} \times 5 \times 10^{-10}$ $= 5 \times 10^{-15}$ Hence, (3) is correct. 119. A mixture of ethyl alcohol and propyl alcohol has a vapour pressure of 290 mm at 300 K. The vapour pressure of propyl alcohol is 200 mm. If the mole fraction of ethyl alcohol is 0.6, its vapour pressure (in mm) at the same temperature will be (1) 350 (2) 300 (3) 700 (4) 360 Ans. (1)Let the vapour pressure of pure ethyl alcohol be P, Sol. According to Raoult's law 290 = 200 × 0.4 + P × 0.6 $P = \frac{290 - 80}{0.6} = 350 \text{ mm Hg}$ Hence, (1) is correct. 120. In conversion of lime-stone to lime, $CaCO_3(s) \longrightarrow CaO(s) + CO_2(g)$ the vales of ΔH° and ΔS° are +179.1 kJ mol⁻¹ and 160.2 J/K respectively at 298 K and 1 bar. Assuming that ΔH° do not change with temperature, temperature above which conversion of limestone to lime will be spontaneous is (1) 1008 K (2) 1200 (3) 845 K (4) 1118 K Ans. (4)We know, $\Delta G = \Delta H - T \Delta S$ Sol. So, lets find the equilibrium temperature, i.e. at which $\Delta G = 0$ $\Delta H = T \Delta S$ $T = \frac{179.1 \times 1000}{1000}$ 160.2 = 1118 K So, at temperature above this, the reaction will become spontaneous. Hence, (4) is correct answer.