

CY : CHEMISTRY

Duration : Three Hours

Maximum Marks :150

Read the following instructions carefully

1. This question paper contains **24** printed pages including pages for rough work. Please check all pages and report discrepancy, if any.
2. Write your registration number, your name and name of the examination centre at the specified locations on the right half of the ORS.
3. Using HB pencil, darken the appropriate bubble under each digit of your registration number and the letters corresponding to your paper code.
4. All the questions in this question paper are of objective type.
5. Questions must be answered on **Objective Response Sheet (ORS)** by darkening the appropriate bubble (marked A, B, C, D) using HB pencil against the question number on the left hand side of the ORS. **Each question has only one correct answer.** In case you wish to change an answer, erase the old answer completely. More than one answer bubbled against a question will be treated as a wrong answer.
6. Questions 1 through 20 are 1-mark questions and questions 21 through 85 are 2-mark questions.
7. Questions 71 through 73 is one set of common data questions, questions 74 and 75 is another pair of common data questions. The question pairs (76, 77), (78, 79), (80, 81), (82, 83) and (84, 85) are questions with linked answers. The answer to the second question of the above pairs will depend on the answer to the first question of the pair. If the first question in the linked pair is wrongly answered or is un-attempted, then the answer to the second question in the pair will not be evaluated.
8. Un-attempted questions will carry zero marks.
9. **NEGATIVE MARKING:** For Q.1 to Q.20, **0.25** mark will be deducted for each wrong answer. For Q.21 to Q.75, **0.5** mark will be deducted for each wrong answer. For the pairs of questions with linked answers, there will be negative marks only for wrong answer to the first question, i.e. for Q.76, Q.78, Q.80, Q.82 and Q.84, **0.5** mark will be deducted for each wrong answer. There is no negative marking for Q.77, Q.79, Q.81, Q.83 and Q.85.
10. Calculator **without data connectivity** is allowed in the examination hall.
11. Charts, graph sheets and tables are **NOT** allowed in the examination hall.
12. Rough work can be done on the question paper itself. Additional blank pages are given at the end of the question paper for rough work.

Some Useful Data

1. Physical Constants:

Universal gas constant, $R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1} = 0.082 \text{ L atm K}^{-1} \text{ mol}^{-1}$

Planck constant, $h = 6.626 \times 10^{-34} \text{ J s}$

Avogadro number, $N_A = 6.023 \times 10^{23} \text{ mol}^{-1}$

Faraday constant, $F = 96485 \text{ C mol}^{-1}$

Electron charge, $e = 1.602 \times 10^{-19} \text{ C}$

Speed of light, $c = 2.998 \times 10^8 \text{ m s}^{-1}$

Boltzmann constant, $k_B = 1.381 \times 10^{-23} \text{ J K}^{-1}$

Electron mass, $m_e = 9.109 \times 10^{-31} \text{ kg}$

2. Atomic Numbers:

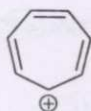
B = 5; F = 9; Al = 13; Si = 14; P = 15; S = 16; Cl = 17; Ti = 22; V = 23; Cr = 24; Mn = 25;

Fe = 26; Co = 27; Ni = 28; Cu = 29; Ru = 44; Rh = 45; Re = 75; Os = 76.

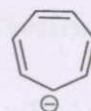
Q. 1 – Q. 20 Carry one mark each

- Q.1 The total number of isomers of $\text{Co}(\text{en})_2\text{Cl}_2$ (en = ethylenediamine) is
 (A) 4 (B) 3 (C) 6 (D) 5
- Q.2 Metal-metal quadruple bonds are well-known for the metal
 (A) Ni (B) Co (C) Fe (D) Re
- Q.3 The reaction of Al_4C_3 with water leads to the formation of
 (A) methane (B) propyne (C) propene (D) propane
- Q.4 The correct statement about C_{60} is
 (A) C_{60} is soluble in benzene
 (B) C_{60} does not react with *tert*-butyllithium
 (C) C_{60} is made up of 10 five-membered and 15 six-membered rings
 (D) Two adjacent five-membered rings share a common edge
- Q.5 The lattice parameters for a monoclinic crystal are
 (A) $a \neq b \neq c$; $\alpha = \gamma = 90^\circ$ (B) $a = b \neq c$; $\alpha \neq \beta \neq \gamma$
 (C) $a \neq b \neq c$; $\alpha \neq \beta \neq \gamma$ (D) $a = b = c$; $\alpha = \gamma = 90^\circ$
- Q.6 The magnetic moment of $[\text{Ru}(\text{H}_2\text{O})_6]^{2+}$ corresponds to the presence of
 (A) four unpaired electrons (B) three unpaired electrons
 (C) two unpaired electrons (D) zero unpaired electrons
- Q.7 The compound that is **NOT** aromatic is

(A)



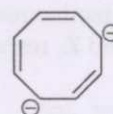
(B)



(C)



(D)



Q.8 The order of stability for the following cyclic olefins is



I



II



III



IV

(A) I < II < III < IV

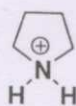
(B) II < III < IV < I

(C) II < III < I < IV

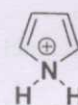
(D) IV < II < I < III

Q.9 The most acidic species is

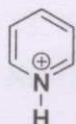
(A)



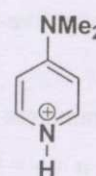
(B)



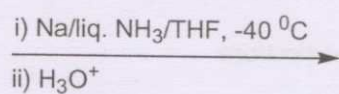
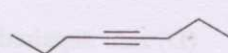
(C)



(D)



Q.10 The major product of the following reaction is



(A)



(B)



(C)



(D)



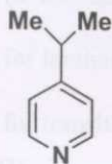
Q.11 In the carbylamine reaction, R-X is converted to R-Y via the intermediate Z. R-X, R-Y and Z, respectively, are

(A) R-NH₂, R-NC, carbene(B) R-NH₂, R-NC, nitrene(C) R-NC, R-NH₂, carbene

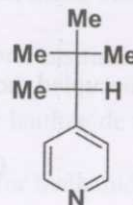
(D) R-OH, R-NC, nitrene

Q.12 The compound that is **NOT** oxidized by KMnO_4 is

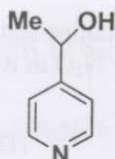
(A)



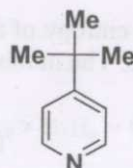
(B)



(C)



(D)



Q.13 Cyanogen bromide (CNBr) specifically hydrolyses the peptide bond formed by the C-side of

(A) methionine

(B) glycine

(C) proline

(D) serine

Q.14 The Hammett reaction constant ρ is based on

(A) the rates of alkaline hydrolysis of substituted ethyl benzoates

(B) the dissociation constants of substituted acetic acids

(C) the dissociation constants of substituted benzoic acids

(D) the dissociation constants of substituted phenols

Q.15 The lifetime of a molecule in an excited electronic state is 10^{-10} s. The uncertainty in the energy (eV) approximately is

(A) 2×10^{-5} (B) 3×10^{-6}

(C) 0

(D) 10^{-14}

Q.16 For a one component system, the maximum number of phases that can coexist at equilibrium is

(A) 3

(B) 2

(C) 1

(D) 4

Q.17 At $T = 300$ K, the thermal energy ($k_B T$) in cm^{-1} is approximately

(A) 20000

(B) 8000

(C) 5000

(D) 200

- Q.18 For the reaction $2 X_3 = 3 X_2$, the rate of formation of X_2 is
(A) $3 (-d[X_3] / dt)$ (B) $1/2 (-d[X_3] / dt)$ (C) $1/3 (-d[X_3] / dt)$ (D) $3/2 (-d[X_3] / dt)$
- Q.19 The highest occupied molecular orbital of HF is
(A) bonding (B) antibonding (C) ionic (D) nonbonding
- Q.20 The residual entropy of the asymmetric molecule N_2O in its crystalline state is $5.8 J K^{-1} mol^{-1}$ at absolute zero. The number of orientations that can be adopted by N_2O in its crystalline state is
(A) 4 (B) 3 (C) 2 (D) 1
- Q. 21 to Q.75 Carry two marks each**
- Q.21 The spectroscopic ground state symbol and the total number of electronic transitions of $[Ti(H_2O)_6]^{2+}$ are
(A) ${}^3T_{1g}$ and 2 (B) ${}^3A_{2g}$ and 3
(C) ${}^3T_{1g}$ and 3 (D) ${}^3A_{2g}$ and 2
- Q.22 The structures of the complexes $[Cu(NH_3)_4](ClO_4)_2$ and $[Cu(NH_3)_4](ClO_4)$ in solution respectively are
(A) square planar and tetrahedral (B) octahedral and square pyramidal
(C) octahedral and trigonal bipyramidal (D) tetrahedral and square planar
- Q.23 In biological systems, the metal ions involved in electron transport are
(A) Na^+ and K^+ (B) Zn^{2+} and Mg^{2+} (C) Ca^{2+} and Mg^{2+} (D) Cu^{2+} and Fe^{3+}
- Q.24 In a homogeneous catalytic reaction, 1.0M of a substrate and 1.0 μM of a catalyst yields 1.0 mM of a product in 10 seconds. The turnover frequency (TOF) of the reaction (s^{-1}) is
(A) 10^{-2} (B) 10^2 (C) 10^{-3} (D) 10^3

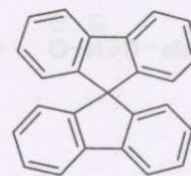
- Q.25 The expected magnetic moments of the first-row transition metal complexes and those of the lanthanide metal complexes are usually calculated using
- (A) $\mu_{s.o}$ equation (s. o. = spin only) for both lanthanide and transition metal complexes
 (B) $\mu_{s.o}$ equation for lanthanide metal complexes and μ_J equation for transition metal complexes
 (C) $\mu_{s.o}$ equation for transition metal complexes and μ_J equation for lanthanide metal complexes
 (D) μ_{L+S} equation for transition metal complexes and $\mu_{s.o}$ equation for lanthanide metal complexes
- Q.26 The Brønsted acidity of boron hydrides follows the order
- (A) $B_2H_6 > B_4H_{10} > B_5H_9 > B_{10}H_{14}$ (B) $B_2H_6 = B_4H_{10} > B_5H_9 = B_{10}H_{14}$
 (C) $B_{10}H_{14} > B_5H_9 > B_4H_{10} > B_2H_6$ (D) $B_5H_9 > B_4H_{10} > B_2H_6 > B_{10}H_{14}$
- Q.27 NaCl is crystallised by slow evaporation of its aqueous solution at room temperature. The correct statement is
- (A) The crystals will be non-stoichiometric
 (B) The crystals should have Frenkel defects
 (C) The percentage of defects in the crystals will depend on the concentration of the solution and its rate of evaporation
 (D) The nature of defects will depend upon the concentration of the solution and its rate of evaporation
- Q.28 $CaTiO_3$ has a perovskite crystal structure. The coordination number of titanium in $CaTiO_3$ is
- (A) 9 (B) 6 (C) 3 (D) 12
- Q.29 If ClF_3 were to be stereochemically rigid, its ^{19}F NMR spectrum (I for $^{19}F = \frac{1}{2}$) would be (assume that Cl is not NMR active)
- (A) a doublet and a triplet (B) a singlet
 (C) a doublet and a singlet (D) two singlets
- Q.30 The point group of NSF_3 is
- (A) D_{3d} (B) C_{3h} (C) D_{3h} (D) C_{3v}

- Q.31 When NiO is heated with a small amount of Li_2O in air at 1200°C , a non-stoichiometric compound $\text{Li}_x\text{Ni}_{(1-x)}\text{O}$ is formed. This compound is
- (A) an n-type semiconductor containing only Ni^{1+}
(B) an n-type semiconductor containing Ni^{1+} and Ni^{2+}
(C) a p-type semiconductor containing Ni^{2+} and Ni^{3+}
(D) a p-type semiconductor containing only Ni^{3+}
- Q.32 White phosphorus, P_4 , belongs to the
- (A) *closo* system (B) *nido* system
(C) *arachno* system (D) *hypho* system
- Q.33 Among the compounds Fe_3O_4 , NiFe_2O_4 and Mn_3O_4
- (A) NiFe_2O_4 and Mn_3O_4 are normal spinels
(B) Fe_3O_4 and Mn_3O_4 are normal spinels
(C) Fe_3O_4 and Mn_3O_4 are inverse spinels
(D) Fe_3O_4 and NiFe_2O_4 are inverse spinels
- Q.34 The number of M-M bonds in $\text{Ir}_4(\text{CO})_{12}$ are
- (A) four (B) six (C) eight (D) zero
- Q.35 Schrock carbenes are
- (A) triplets and nucleophilic (B) triplets and electrophilic
(C) singlets and nucleophilic (D) singlets and electrophilic
- Q.36 The **INCORRECT** statement about linear dimethylpolysiloxane, $[(\text{CH}_3)_2\text{SiO}]_n$, is
- (A) it is extremely hydrophilic
(B) it is prepared by a KOH catalysed ring-opening reaction of $[\text{Me}_2\text{SiO}]_4$
(C) it has a very low glass transition temperature
(D) it can be reinforced to give silicon elastomers

Q.37 Match the entries **a-d** with their corresponding structures **p-s**.

a bridged system

p



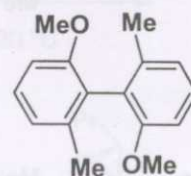
b atropisomeric system

q



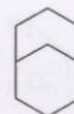
c spiro system

r



d fused system

s



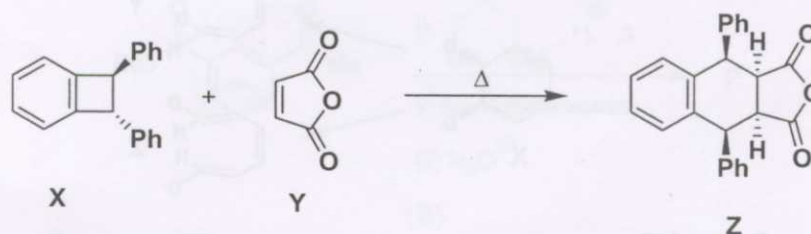
(A) a-s; b-r; c-q; d-p

(B) a-p; b-s; c-q; d-r

(C) a-q; b-p; c-s; d-r

(D) a-s; b-r; c-p; d-q

Q.38 The reaction between **X** and **Y** to give **Z** proceeds via



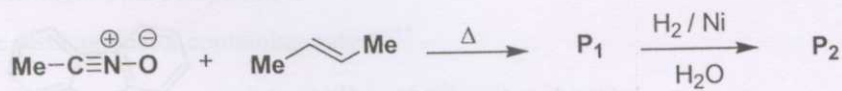
(A) 4π -conrotatory opening of **X** followed by *endo* Diels-Alder cycloaddition

(B) 4π -disrotatory opening of **X** followed by *endo* Diels-Alder cycloaddition

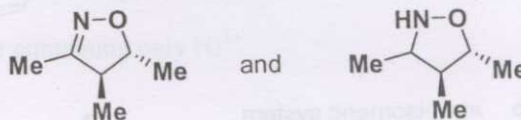
(C) 4π -conrotatory opening of **X** followed by *exo* Diels-Alder cycloaddition

(D) 4π -disrotatory opening of **X** followed by *exo* Diels-Alder cycloaddition

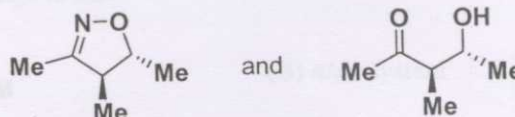
Q.39 The major products P_1 and P_2 , respectively, in the following reaction sequence are



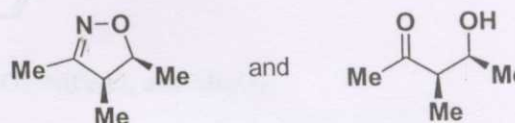
(A)



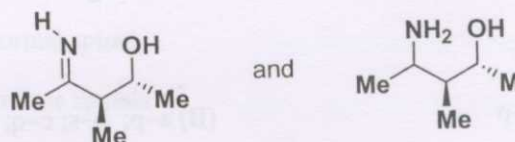
(B)



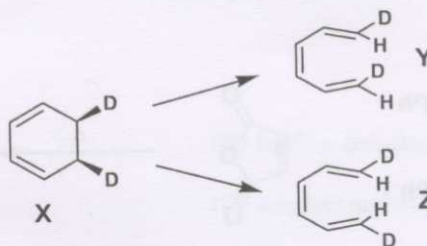
(C)



(D)



Q.40 The products Y and Z are formed, respectively, from X via

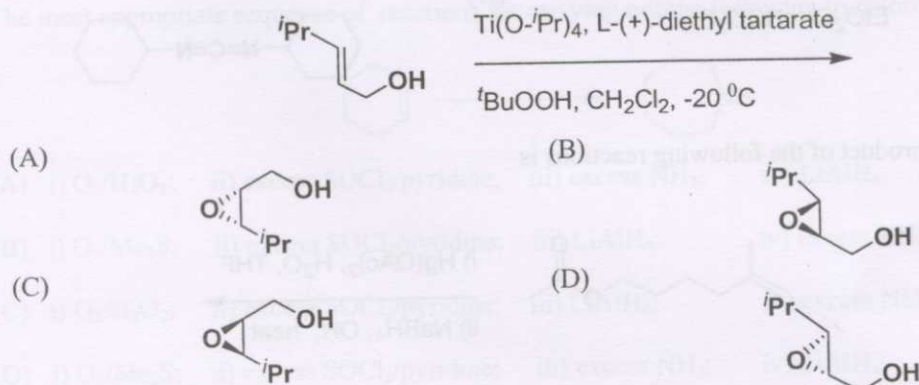


- (A) $h\nu$, conrotatory opening and Δ , disrotatory opening
 (B) $h\nu$, disrotatory opening and Δ , conrotatory opening
 (C) Δ , conrotatory opening and $h\nu$, disrotatory opening
 (D) Δ , disrotatory opening and $h\nu$, conrotatory opening

Q.41 *o*-Bromophenol is readily prepared from phenol using the following conditions

- (A) i) $(\text{CH}_3\text{CO})_2\text{O}$; ii) Br_2 ; iii) $\text{HCl-H}_2\text{O}, \Delta$
 (B) i) $\text{H}_2\text{SO}_4, 100^\circ\text{C}$ ii) Br_2 iii) $\text{H}_3\text{O}^+, 100^\circ\text{C}$
 (C) N-Bromosuccinimide, dibenzoyl peroxide, CCl_4, Δ
 (D) $\text{Br}_2/\text{FeBr}_3$

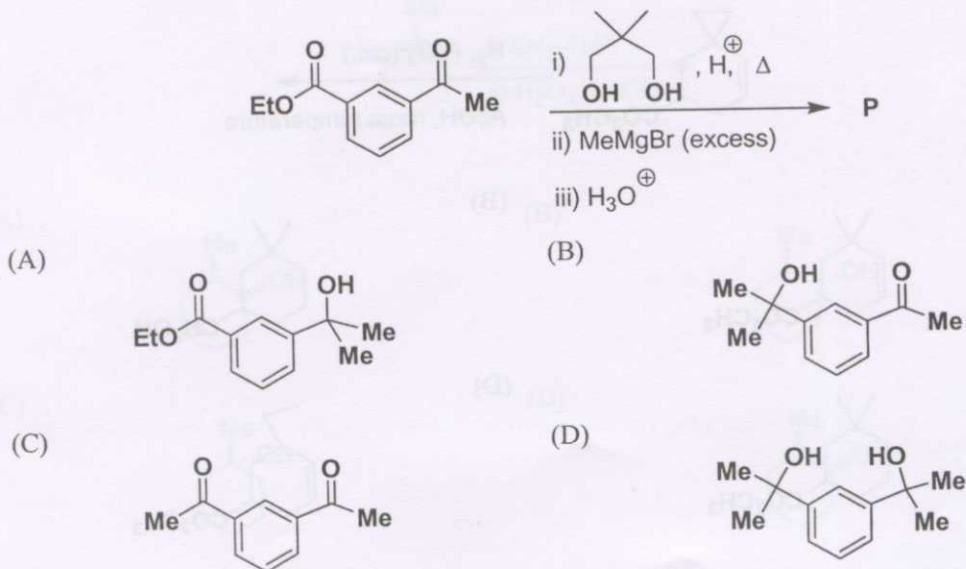
Q.42 The major product of the following reaction is



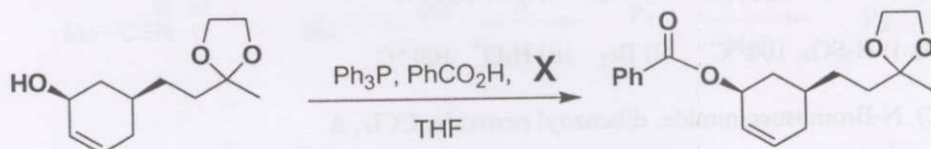
Q.43 The photochemical reaction of 2-methylpropane with F_2 gives 2-fluoro-2-methylpropane and 1-fluoro-2-methylpropane in 14:86 ratio. The corresponding ratio of the bromo products in the above reaction using Br_2 is most likely to be

- (A) 14 : 86 (B) 50 : 50 (C) 1 : 9 (D) 99 : 1

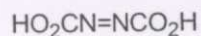
Q.44 The major product **P** of the following reactions is



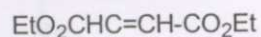
Q.45 The reagent X in the following reaction is



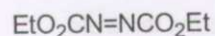
(A)



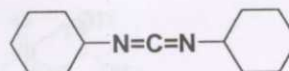
(B)



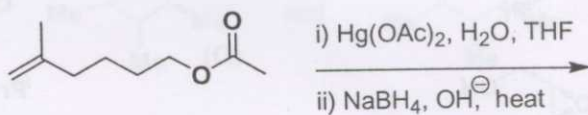
(C)



(D)



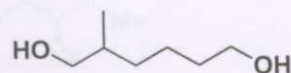
Q.46 The major product of the following reactions is



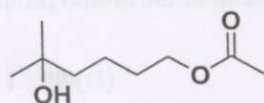
(A)



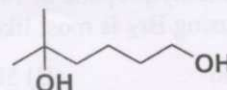
(B)



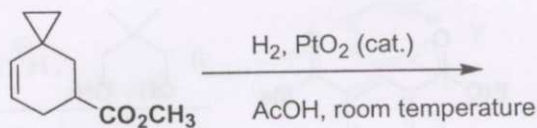
(C)



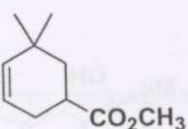
(D)



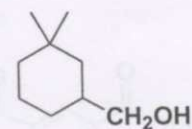
Q.47 The major product of the following reaction is



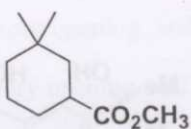
(A)



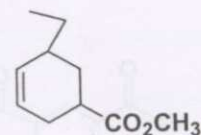
(B)



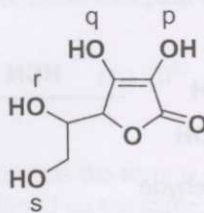
(C)



(D)



Q.48 In the following compound, the hydroxy group that is most readily methylated with CH_2N_2 is



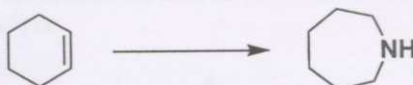
(A) p

(B) q

(C) r

(D) s

Q.49 The most appropriate sequence of reactions for carrying out the following transformation is

(A) i) $\text{O}_3/\text{H}_2\text{O}_2$; ii) excess $\text{SOCl}_2/\text{pyridine}$; iii) excess NH_3 ; iv) LiAlH_4 (B) i) $\text{O}_3/\text{Me}_2\text{S}$; ii) excess $\text{SOCl}_2/\text{pyridine}$; iii) LiAlH_4 ; iv) excess NH_3 (C) i) $\text{O}_3/\text{H}_2\text{O}_2$; ii) excess $\text{SOCl}_2/\text{pyridine}$; iii) LiAlH_4 ; iv) excess NH_3 (D) i) $\text{O}_3/\text{Me}_2\text{S}$; ii) excess $\text{SOCl}_2/\text{pyridine}$; iii) excess NH_3 ; iv) LiAlH_4

Q.50 The number of optically active stereoisomers possible for 1,3-cyclohexanediol in its chair conformation is

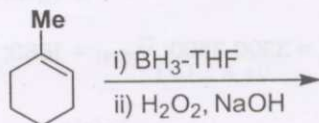
(A) 4

(B) 3

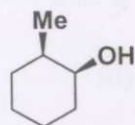
(C) 2

(D) 1

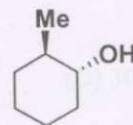
Q.51 The major product of the following reactions is



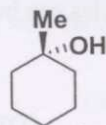
(A)



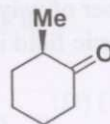
(B)



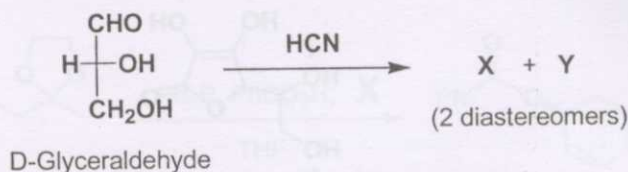
(C)



(D)

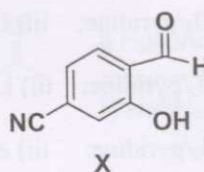


Q.52 In the following reaction,



The absolute configurations of the chiral centres in X and Y are

- (A) 2*S*, 3*R* and 2*R*, 3*R* (B) 2*R*, 3*R* and 2*R*, 3*S*
 (C) 2*S*, 3*S* and 2*R*, 3*R* (D) 2*S*, 3*R* and 2*S*, 3*R*
- Q.53 The IR stretching frequencies (cm^{-1}) for the compound X are as follows: 3300-3500 (s, br); 3000 (m); 2225 (s); 1680 (s).



The correct assignment of the absorption bands is

- (A) $\bar{\nu}_{(\text{OH})} = 3300-3500$; $\bar{\nu}_{(\text{CH})} = 3000$; $\bar{\nu}_{(\text{CN})} = 2225$; $\bar{\nu}_{(\text{CO})} = 1680$
 (B) $\bar{\nu}_{(\text{OH})} = 3000$; $\bar{\nu}_{(\text{CH})} = 3300-3500$; $\bar{\nu}_{(\text{CN})} = 2225$; $\bar{\nu}_{(\text{CO})} = 1680$
 (C) $\bar{\nu}_{(\text{OH})} = 3300-3500$; $\bar{\nu}_{(\text{CH})} = 3000$; $\bar{\nu}_{(\text{CN})} = 1680$; $\bar{\nu}_{(\text{CO})} = 2225$
 (D) $\bar{\nu}_{(\text{OH})} = 3000$; $\bar{\nu}_{(\text{CH})} = 3300-3500$; $\bar{\nu}_{(\text{CN})} = 1680$; $\bar{\nu}_{(\text{CO})} = 2225$
- Q.54 The T_d point group has 24 elements and 5 classes. Given that it has two 3-dimensional irreducible representations, the number of one dimensional irreducible representations is

- (A) 1 (B) 6 (C) 2 (D) 3

Q.55 The total number of ways in which two nonidentical spin $\frac{1}{2}$ particles can be oriented relative to a constant magnetic field is

- (A) 1 (B) 2 (C) 3 (D) 4

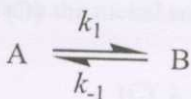
Q.56 Approximately one hydrogen atom per cubic meter is present in interstellar space. Assuming that the H-atom has a diameter of 10^{-10} m, the mean free path (m) approximately is

- (A) 10^{10} (B) 10^{19} (C) 10^{24} (D) 10^{14}

Q.57 The wavefunction of a diatomic molecule has the form $\psi = 0.89 \phi_{\text{covalent}} + 0.45 \phi_{\text{ionic}}$. The chance that both electrons of the bond will be found on the same atom in 100 inspections of the molecule approximately is

- (A) 79 (B) 20 (C) 45 (D) 60

Q.58 For the reaction given below, the relaxation time is 10^{-6} s. Given that 10% of A remains at equilibrium, the value of k_1 (s^{-1}) is



- (A) 9×10^5 (B) 10^{-5} (C) 10^5 (D) 9×10^{-5}

Q.59 The minimum number of electrons needed to form a chemical bond between two atoms is

- (A) 1 (B) 2 (C) 3 (D) 4

Q.60 The ground state electronic energy (Hartree) of a helium atom, neglecting the inter-electron repulsion, is

- (A) - 1.0 (B) - 0.5 (C) - 2.0 (D) - 4.0

Q.61 A particle is confined to a one-dimensional box of length 1 mm. If the length is changed by 10^{-9} m, the % change in the ground state energy is

- (A) 2×10^{-4} (B) 2×10^{-7} (C) 2×10^{-2} (D) 0

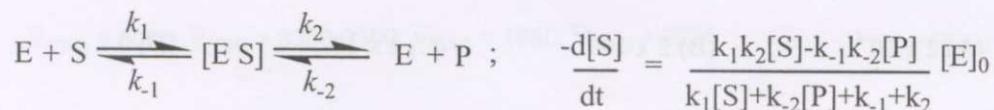
Q.62 A certain molecule can be treated as having only a doubly degenerate state lying at 360 cm^{-1} above the nondegenerate ground state. The approximate temperature (K) at which 15% of the molecules will be in the upper state is

- (A) 500 (B) 150 (C) 200 (D) 300

Q.63 A box of volume V contains one mole of an ideal gas. The probability that all N particles will be found occupying one half of the volume leaving the other half empty is

- (A) $1/2$ (B) $2/N$ (C) $(1/2)^N$ (D) $(1/2)^{6N}$

- Q.64 According to the Debye-Hückel limiting law, the mean activity coefficient of $5 \times 10^{-4} \text{ mol kg}^{-1}$ aqueous solution of CaCl_2 at 25°C is (the Debye-Hückel constant 'A' can be taken to be 0.509)
- (A) 0.63 (B) 0.72 (C) 0.80 (D) 0.91
- Q.65 The operation of the commutator $[x, d/dx]$ on a function $f(x)$ is equal to
- (A) 0 (B) $f(x)$ (C) $-f(x)$ (D) $x df/dx$
- Q.66 If a gas obeys the equation of state $P(V - nb) = nRT$, the ratio $(C_p - C_v) / (C_p - C_v)_{\text{ideal}}$ is
- (A) > 1 (B) < 1 (C) 1 (D) $(1 - b)$
- Q.67 Physisorbed particles undergo desorption at 27°C with an activation energy of $16.628 \text{ kJ mol}^{-1}$. Assuming first-order process and a frequency factor of 10^{12} Hz , the average residence time (in seconds) of the particles on the surface is
- (A) 8×10^{-10} (B) 8×10^{-11} (C) 2×10^{-9} (D) 1×10^{-12}
- Q.68 The rotational constants for CO in the ground and the first excited vibrational states are 1.9 and 1.6 cm^{-1} , respectively. The % change in the internuclear distance due to vibrational excitation is
- (A) 9 (B) 30 (C) 16 (D) 0
- Q.69 The mechanism of enzyme (E) catalysed reaction of a substrate (S) to yield product (P) is



If a small amount of S is converted to P, the maximum rate for the reaction will be observed for

- (A) $(k_1 + k_2) \gg k_1 [S]_0$ (B) $(k_1 + k_2) \ll k_1 [S]_0$
 (C) $(k_2 + k_2) = (k_1 + k_1)$ (D) $k_2 \ll k_1$
- Q.70 The lowest energy state of the $(1s)^2 (2s)^1 (3s)^1$ configuration of Be is

- (A) 1S_0 (B) 1D_2 (C) 3S_1 (D) 3P_1

Common Data Questions**Common Data for Questions 71, 72 and 73:**

An electron accelerated through a potential difference of ϕ volts impinges on a nickel surface, whose (100) planes have a spacing $d = 351.8 \times 10^{-12}$ m (351.8 pm).

Q.71 The de-Broglie wavelength of the electron is $\lambda / \text{pm} = (a/\phi)^{1/2}$. The value of 'a' in volts is

- (A) 1.5×10^{-18} (B) 1.5×10^6 (C) 6.63×10^{-5} (D) 2.5×10^{18}

Q.72 The condition for observing diffraction from the nickel surface is

- (A) $\lambda \gg 2d$ (B) $\lambda \leq 2d$ (C) $\lambda \leq ad$ (D) $\lambda \geq ad$

Q.73 The minimum value of ϕ (V) for the electron to diffract from the (100) planes is

- (A) 3000 (B) 300 (C) 30 (D) 3

Common Data for Questions 74 and 75:

An iron complex $[\text{FeL}_6]^{2+}$ (L = neutral monodentate ligand) catalyses the oxidation of $(\text{CH}_3)_2\text{S}$ by perbenzoic acid.

Q.74 The formation of the organic product in the above reaction can be monitored by

- (A) gas chromatography (B) cyclic voltammetry
(C) electron spin resonance (D) fluorescence spectroscopy

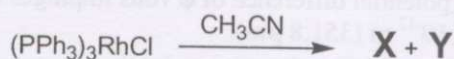
Q.75 The oxidation state of the metal ion in the catalyst can be detected by

- (A) atomic absorption spectroscopy (B) Mössbauer spectroscopy
(C) HPLC (D) gas chromatography

Linked Answer Questions: Q.76 to Q.85 carry two marks each

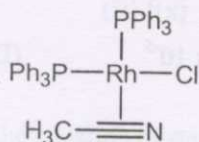
Linked Answer Questions 76 and 77:

In the reaction,

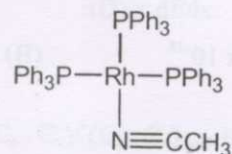


Q.76 Compound X is

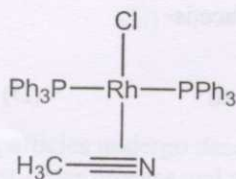
(A)



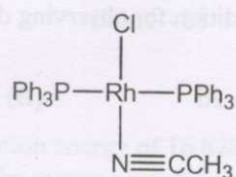
(B)



(C)

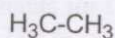


(D)

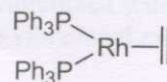


Q.77 $\text{Rh}(\text{PPh}_3)_3\text{Cl}$ reacts very fast with a gaseous mixture of H_2 and C_2H_4 to immediately give Z. The structure of Z is

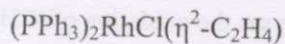
(A)



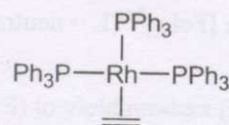
(B)



(C)



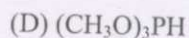
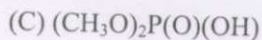
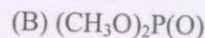
(D)



Linked Answer Questions 78 and 79:

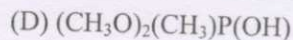
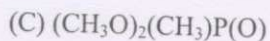
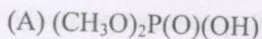
The reaction of PCl_3 with methanol in the presence of triethylamine affords compound **X**. EI mass spectrum of **X** shows a parent ion peak at $m/z = 124$. Microanalysis of **X** shows that it contains C, H, O and P. The ^1H NMR spectrum of **X** shows a doublet at 4.0 ppm. The separation between the two lines of the doublet is approximately 15 Hz (1 for ^1H and $^{31}\text{P} = 1/2$).

Q.78 Compound **X** is



Q.79 Upon heating, compound **X** is converted to **Y**, which has the same molecular formula as that of **X**. The ^1H NMR spectrum of **Y** shows two doublets centered at 3.0 ppm (separation of two lines ~ 20 Hz) and 4.0 ppm (separation of two lines ~ 15 Hz) respectively.

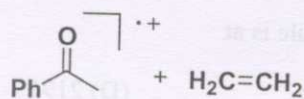
Compound **Y** is


Linked Answer Questions 80 and 81:

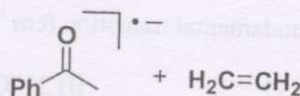
For butyrophenone ($\text{PhCOCH}_2\text{CH}_2\text{CH}_3$),

Q.80 the most probable fragmentation observed in the electron impact ionization (EI) mass spectrometry is

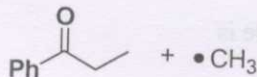
(A)



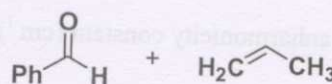
(B)



(C)

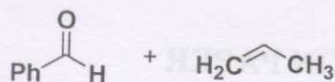


(D)

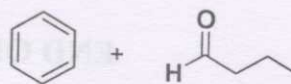


Q.81 photoirradiation leads to the following set of products.

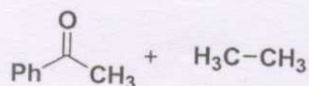
(A)



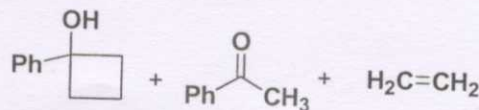
(B)



(C)

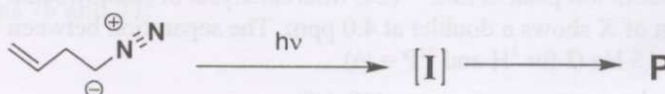


(D)



Linked Answer Questions 82 and 83:

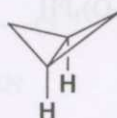
In the following reaction,



Q.82 the reactive intermediate **I** and the product **P** are

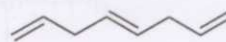
(A)

carbene and



(B)

radical and



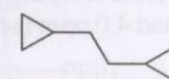
(C)

carbene and



(D)

radical and



Q.83 the product **P** shows 'm' and 'n' number of signals in ^1H and ^{13}C NMR spectra, respectively. The values of 'm' and 'n' are

(A) $m = 3$ and $n = 2$

(B) $m = 2$ and $n = 3$

(C) $m = 2$ and $n = 2$

(D) $m = 4$ and $n = 3$

Linked Answer Questions 84 and 85:

The infrared spectrum of a diatomic molecule exhibits transitions at 2144 , 4262 and 6354 cm^{-1} corresponding to excitations from the ground state to the first, second and, third vibration states respectively.

Q.84 The fundamental transition (cm^{-1}) of the diatomic molecule is at

(A) 2157

(B) 2170

(C) 2183

(D) 2196

Q.85 The anharmonicity constant (cm^{-1}) of the diatomic molecule is

(A) 0.018

(B) 0.012

(C) 0.006

(D) 0.003

END OF THE QUESTION PAPER