Q.	1 -	Q.	25	carry	one	mark	each.
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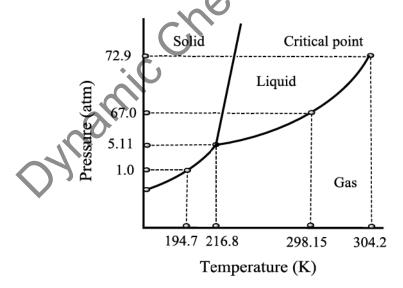
- Q.1 The **INCORRECT** statement about the solid-state structure of CsCl and CaF<sub>2</sub> is:
  - (A) Cations in both solids exhibit coordination number 8.
  - (B) CsCl has bcc type structure and CaF<sub>2</sub> has cubic close pack structure.
  - (C) Radius ratio for Cs/Cl and Ca/F is 0.93 and 0.73, respectively.
  - (D) Both exhibit close pack structure.
- Q.2 The INCORRECT statement about the interhalogen compound ICl<sub>3</sub> is:
  - (A) It exists as a dimer.
  - (B) Geometry around the iodine is tetrahedral in solid-state.
  - (C) It decomposes as ICl and Cl2 in gas-phase.
  - (D) Liquid ICl3 conducts electricity.
- Q.3 Among the following carbon allotropes, the one with discrete molecular structure is
  - (A) Diamond
- (B) α-Graphite
- (C) β-Graphite
- (D) Fullerene
- Q.4 The **INCORRECT** statement about the silicones is:
  - (A) They are thermally unstable because of the Si–C bond.
  - (B) They are insoluble in water.
  - (C) They are organosilicon polymers.
  - (D) They have stable silica-like skeleton (-Si-O-Si-O-Si-).
- Q.5 The  $\Delta_o$  value of  $[Ni(H_2O)_6]^{2+}$  is 8500 cm<sup>-1</sup>. The  $\Delta_o$  values for  $[NiCl_6]^{4-}$  and  $[Ni(NH_3)_6]^{2+}$  compared to  $[Ni(H_2O)_6]^{2+}$  are
  - (A) higher and lower, respectively.
- (B) lower and higher, respectively.
- (C) higher in both complex ions.
- (D) lower in both complex ions.

- Q.6 In Freundlich isotherm, a linear relationship is obtained in the plot of
  - ( $\theta$  = surface coverage and p = partial pressure of the gas)
  - (A)  $\theta$  vs p.

(B)  $\ln(\theta)$  vs  $\ln(p)$ .

(C)  $\ln(\theta)$  vs p.

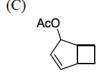
- (D)  $\theta$  vs  $\ln(p)$ .
- Q.7 Micelle formation is accompanied by the
  - (A) decrease in overall entropy due to ordering.
  - (B) increase in overall entropy mostly due to increase in solvent entropy.
  - (C) increase in overall entropy mostly due to increase in solute entropy.
  - (D) increase in overall entropy and decrease in enthalpy
- Q.8 Consider the following phase diagram of CO<sub>2</sub> (not to scale). At equilibrium, the **INCORRECT** statement is:

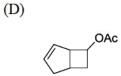


- (A) At 200 K, on increasing the pressure from 1 to 50 atm, CO2 gas condenses to liquid.
- (B) It is not possible to obtain liquid CO<sub>2</sub> from gaseous CO<sub>2</sub> below 5.11 atm.
- (C) Both liquid and gas phase of  $CO_2$  coexist at 298.15 K and 67 atm.
- (D) With increasing pressure, the melting point of solid  ${\rm CO}_2$  increases.

Q.9 The major product formed in the following reaction is







Q.10 The Woodward-Hoffmann condition to bring out the following transformation is

(A)  $\Delta$ , conrotatory

(B)  $\Delta$ , disrotatory

(C) hv, disrotatory

D) hv, conrotatory

Q.11 The major product formed in the following reaction is

- (A)
- OH O
- (B) OH O
- (C) Ph O OH OH
- Cl<sub>3</sub>C Ph O OH

Q.12 In the following reaction, the stereochemistry of the major product is predicted by the

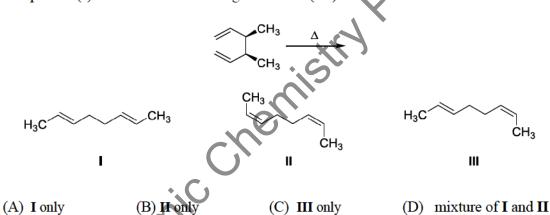
(A) Cram's model

(B) Cram's chelation model

(C) Felkin model

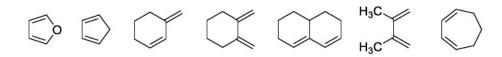
(D) Felkin-Anh model

Q.13 The product(s) formed in the following reaction is (are)



Q.14 Among the following compounds, the number of compounds that **DO NOT** exhibit optical activity at room temperature is \_\_\_\_\_.

Q.15 The number of following diene(s) that undergo Diels-Alder reaction with methyl acrylate is



Q.16 The number of <sup>1</sup>H NMR signals observed for the following compound is .

Q.17 The number of CO stretching bands in IR spectrum of trigonal bipyramidal *cis*-M(CO)<sub>3</sub>L<sub>2</sub> is \_\_\_\_\_\_.

(M = metal and L = monodentate ligand)

Q.18 On heating a sample of 25 mg hydrated compound (molecular weight = 250 g/mol) in thermogravimetric analysis, 16 mg of dehydrated compound remains. The number of water molecules lost per molecule of hydrated compound is \_\_\_\_\_\_.

(Molecular weight of water = 18 g/mol)

Q.19 The total number of  $\alpha$  and  $\beta$  particles emitted in the following radioactive decay is \_\_\_\_\_.

$$^{238}_{92}\mathrm{U} \longrightarrow ^{210}_{82}\mathrm{Pb}$$

Q.20 An ideal gas occupies an unknown volume V liters (L) at a pressure of 12 atm. The gas is expanded isothermally against a constant external pressure of 2 atm so that its final volume becomes 3 L. The work involved for this expansion process is \_\_\_\_\_\_cal. (Round off to two decimal places)

(Gas constant R = 0.082 L atm mol<sup>-1</sup>  $K^{-1} = 2$  cal mol<sup>-1</sup>  $K^{-1}$ )

Q.21 The entropy change for the melting of 'x' moles of ice (heat of fusion is 80 cal g<sup>-1</sup>) at 273 K and 1 atm pressure is 28.80 cal K<sup>-1</sup>. The value of 'x' is \_\_\_\_\_. (Round off to two decimal places)

(Molecular weight of water =18 g/mol)

Q.22 Consider a two-state system at thermal equilibrium having energies 0 and 2k<sub>B</sub>T for which the degeneracies are 1 and 2, respectively. The value of the partition function at the same absolute temperature T is \_\_\_\_\_\_. (Round off to two decimal places)

(k<sub>B</sub> is the Boltzmann constant)

- Q.23 Consider a system of three identical and distinguishable non-interacting particles and three available nondegenerate single particle energy levels having energies 0,  $\epsilon$  and  $2\epsilon$ . The system is in contact with a heat bath of temperature T K. A total energy of  $2\epsilon$  is shared by these three particles. The number of ways the particles can be distributed is \_\_\_\_\_.
- Q.24 In a 400 MHz <sup>1</sup>H NMR spectrometer, a proton resonates at 1560 Hz higher than that of tetramethylsilane. The chemical shift value of this proton is \_\_\_\_\_\_ ppm. (Round off to one decimal place)

(Chemical shift of tetramethylsilane is fixed at zero ppm)

Q.25 Gas phase bond length and dipole moment of a compound (MX) is 3 Å and 10.8 D, respectively. The ionic character in gas phase MX is \_\_\_\_\_\_\_%. (Round off to one decimal place)

 $(1D = 3.336 \times 10^{-30} \text{ C m})$ 

## Q. 26 - Q. 55 carry two marks each.

Q.26	The	experimentally	observed	magnetic	moment	values,	which	match	well	with	the
	spin-	only values for t	he pair of	aqueous io	ns is						

(Atomic number: Cr = 24, Co = 27, Gd = 64, Tb = 65, Dy = 66 and Lu = 71)

(A) Cr(III) and Gd(III)

(B) Co(II) and Gd (III)

(C) Cr(III) and Dy(III)

- (D) Lu(III) and Tb(III)
- Q.27 Among the following compounds, a normal spinel is
  - (A) MgFe<sub>2</sub>O<sub>4</sub>

(B) ZnFe<sub>2</sub>O<sub>4</sub>

(C) CoFe<sub>2</sub>O<sub>4</sub>

- D) CuFe<sub>2</sub>O<sub>4</sub>
- Q.28 Following are the examples of silicate minerals

Zircon, ZrSiO<sub>4</sub>

Beryl, Be3Al2Si6O1

Pyrophyllite, Al<sub>2</sub>(OH)<sub>2</sub>[(Si<sub>2</sub>O<sub>5</sub>)<sub>2</sub>]

I

II

Ш

The correct structural description of the minerals is

- (A) I Ortho silicate, II Cyclic silicate and III Sheet silicate
- (B) **I** − Ortho silicate, **II** − Sheet silicate and **III** − Cyclic silicate
- (C) I Cyclic silicate II Sheet silicate and III Ortho silicate
- (D) I Sheet silicate, H Ortho silicate and HI Cyclic silicate
- Q.29 In the EPR spectrum of a methyl radical, the number of lines and their relative intensities, respectively, are
  - (A) 1 and 1
- (B) 3 and 1:2:1
- (C) 4 and 1:2:2:1
- (D) 4 and 1:3:3:1
- Q.30 The product obtained in the reaction of Mn<sub>2</sub>(CO)<sub>10</sub> with Br<sub>2</sub> is
  - (A) Mn(CO)<sub>5</sub>Br
- (B)  $Mn_2(CO)_8Br_2$
- (C)  $Mn(CO)_4Br_2$
- (D) Mn<sub>2</sub>(CO)<sub>9</sub>Br

Q.31 The correct molecular representation of W(Cp)<sub>2</sub>(CO)<sub>2</sub> is

(Cp = cyclopentadienyl)

- (A)  $[W(\eta^1 Cp)(\eta^3 Cp)(CO)_2]$
- (B)  $[W(\eta^1-Cp)(\eta^5-Cp)(CO)_2]$
- (C)  $[W(\eta^3-Cp)(\eta^5-Cp)(CO)_2]$
- (D)  $[W(\eta^5-Cp)_2(CO)_2]$
- Q.32 Match the metalloproteins with their respective functions.

P	Ferritin	I	Electron transfer
Q	Rubredoxin	II	Acid-base catalysis
R	Cobalamin	III	Metal storage
S	Carbonic anhydrase	IV	Methyl transfer

- (A) P III; Q II; R I; S IV
- (B) P III; Q I; R IV; S II
- (C) P IV; Q I; R III; S II
- (D) P IV; Q II; R I; S III
- Q.33 Suppose the wave function of a one dimensional system is

$$\psi = \sin(kx) \exp(3ikx)$$

In an experiment measuring the momentum of the system, one of the expected outcomes is

- (A) 0
- (B) ħk
- (C)  $2 \hbar k$
- (D) 3 ħk

Q.34 The major product formed in the following reaction is

(D)

(B)

(AIBN = azobisisobutyronitrile)

$$\begin{array}{c} \text{O} \\ \\ \\ \text{COOCH}_3 \end{array}$$

Q.35 The major product formed in the following reaction is

HOOC 
$$COOCH_3$$
  $COOCH_3$   $COOCH_3$   $COOCH_3$ 

Q.36 The major product formed in the following reaction is

(A) 
$$OOEt$$
 + HCHO  $OEt$  EtOH,  $A$ 

Q.37 The major product formed in the following reaction is

(C) 
$$CI$$
  $CH_3$ 

## Q.38 In the following reaction sequence, the products $\mathbf{P}$ and $\mathbf{Q}$ are

(A)
$$P = \begin{array}{c} CN \\ Ts \end{array}$$

$$COCH_3$$
(B)
$$P = \begin{array}{c} CN \\ Ts \end{array}$$

$$COCH_3$$

$$COCH_3$$
(C)
$$CN \\ Ts \end{array}$$

$$COCH_3$$

$$COCH_3$$
(D)
$$CN \\ COCH_3$$

(D) 
$$Q = \begin{pmatrix} CN & CH_3 \\ NH & Ts \end{pmatrix}$$

## Q.39 The major product formed in the following reaction is

(PCC = pyridinium chlorochromate)

(A) 
$$CH_3$$
  $CH_3$   $CH_$ 

Q.40 In the following reactions, the major products  $\mathbf{P}$  and  $\mathbf{Q}$  are

(A)

$$PhCO_3H$$
 $phCO_3H$ 
 $phcO$ 

Q.41 In the following reaction sequence, the products  $\mathbf{P}$  and  $\mathbf{Q}$  are

Q.42 The major product formed in the following reaction is

(B)

(D)

Q.43 The rate of the following redox reaction is slowest when X is

$$[\text{Co}^{\text{II}}(\text{NH}_3)_5\textbf{X}]^{3+/2+} \ + \ [\text{Cr}^{\text{II}}(\text{H}_2\text{O})_6]^{2+} \ \rightarrow \ [\text{Co}^{\text{II}}(\text{NH}_3)_5(\text{H}_2\text{O})]^{2+} \ + \ [\text{Cr}^{\text{III}}(\text{H}_2\text{O})_5\textbf{X}]^{3+/2+}$$

- (A) H<sub>2</sub>O
- (B) NH<sub>3</sub>
- (C) Cl
- (D)  $N_3^-$

Q.44 A complex is composed of one chromium ion, three bromides and six water molecules. Upon addition of excess AgNO<sub>3</sub>, 1.0 g aqueous solution of the complex gave 0.94 g of AgBr. The molecular formula of the complex is

(Atomic weight: Cr = 52, Br = 80, Ag = 108, O = 16 and H = 1)

(A)  $[Cr(H_2O)_6]Br_3$ 

(B)  $[Cr(H_2O)_5Br]Br_2 \bullet H_2O$ 

(C)  $[Cr(H_2O)_4Br_2]Br•2H_2O$ 

(D)  $[Cr(H_2O)_3Br_3] \cdot 3H_2O$ 

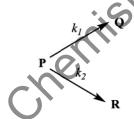
Q.45 The number of possible optically active isomer(s) for the following complex is

$$\begin{bmatrix}
O_2 \\
(en)_2 Co & \\
N \\
H_2
\end{bmatrix}$$
Co(en)<sub>2</sub>

en = ethylenediamine

Q.46 The specific rotation of optically pure (R)-2-bromobutane is -112.00. A given sample of 2-bromobutane exhibited a specific rotation of -82.88. The percentage of (S)-(+)-enantiomer present in this sample is \_\_\_\_\_\_.

Q.47 Consider the following two parallel irreversible first order reactions at temperature T,



where  $k_1$  and  $k_2$  are the rate constants and their values are  $5 \times 10^{-2}$  and  $15 \times 10^{-2}$  min<sup>-1</sup>, respectively, at temperature T. If the initial concentration of the reactant 'P' is 4 mol L<sup>-1</sup>, then the concentration of product 'R' after 10 min of reaction is \_\_\_\_\_ mol L<sup>-1</sup>. (Round off to two decimal places)

(Assume only P is present at the beginning of the reaction.)

Q.48 Consider the following equilibrium

$$SO_2(g) + \frac{1}{2}O_2 \hookrightarrow SO_3(g)$$

At 298 K, the standard molar Gibbs energies of formation,  $\Delta_f G^0$ , of SO<sub>2</sub> (g) and SO<sub>3</sub> (g) are -300 and -371 kJ mol<sup>-1</sup>, respectively. The value of the equilibrium constant,  $K_P$ , at this temperature is \_\_\_\_\_ × 10<sup>10</sup>. (Round off to the nearest integer)

(Gas constant  $R = 8.31 \text{ J mol}^{-1} \text{ K}^{-1}$ )

## Q.49 Consider the electrochemical cell

$$M(s)|MI_2(s)|MI_2(aq)|M(s)$$

where 'M' is a metal. At 298 K, the standard reduction potentials are

$$E_{\mathrm{M}^{2+}(\mathrm{aq})/\mathrm{M(s)}}^{0} = -0.12 \ \mathrm{V}, \quad E_{\mathrm{MI}_{2}(\mathrm{s})/\mathrm{M(s)}}^{0} = -0.36 \ \mathrm{V}$$
 and the temperature coefficient is  $\left(\frac{\partial E_{\mathrm{cell}}^{0}}{\partial T}\right)_{\mathrm{p}} = 1.5 \times 10^{-4} \ \mathrm{V} \ \mathrm{K}^{-1}$ . At this temperature the standard enthalpy change for the overall cell reaction,  $\Delta_{\mathrm{r}} \mathrm{H}^{0}$ , is \_\_\_\_\_kJ mol<sup>-1</sup>. (Round off to two decimal places)

(Faraday constant  $F = 96500 \text{ C mol}^{-1}$ )

Q.50 The normal boiling point of a compound (X) is 350 K heat of vaporization,  $\Delta_{\text{vap}}H$ , = 30 kJ mol<sup>-1</sup>). The pressure required to boil 'X' at 300 K is \_\_\_\_\_ Torr. (Round off to two decimal places)

(Ignore the temperature variation of  $\Delta_{\text{vap}}H$ ; Gas constant  $R=8.31~\text{J}~\text{mol}^{-1}~\text{K}^{-1}$  and 1~atm=760~Torr)

Q.51 For a bimolecular gas phase reaction  $P+Q\to R$ , the pre-exponential factor is  $1\times 10^{13}~dm^3~mol^{-1}~s^{-1}$ . The standard entropy of activation at 25 °C is \_\_\_\_\_\_ J K^{-1} mol^{-1}. (Round off to two decimal points)

(The standard concentration  $c^o=1~\text{mol}~dm^{-3};$  Planck constant  $h=6.62\times 10^{-34}~\text{J}~\text{s};$  Boltzmann constant  $k_B=1.38\times 10^{-23}~\text{J}~\text{K}^{-1};$  Gas constant  $R=8.31~\text{J}~\text{mol}^{-1}~\text{K}^{-1})$ 

Q.52 Character table of point group D<sub>8</sub> is given below.

D <sub>8</sub>	Е	2C <sub>8</sub>	2C <sub>4</sub>	$2C_8^3$	$C_2$	4C <sub>2</sub> ′	4C <sub>2</sub> "
$A_1$	a	1	1	1	1	1	1
$A_2$	b	1	1	1	1	h	i
$B_1$	С	-1	1	-1	1	1	j
$\mathbf{B}_2$	d	-1	1	-1	1	-1	1
$E_1$	e	$\sqrt{2}$	0	$-\sqrt{2}$	-2	0	0
$E_2$	f	0	-2	0	k	0	0
$E_3$	g	$-\sqrt{2}$	0	$\sqrt{2}$	-2	0	0

Value of (a+b+c+d+e+f+g+h+i+j+k) is equal to \_\_\_\_\_.

- Q.53 If  $\langle \alpha | \hat{S}_x \hat{S}_y \hat{S}_y \hat{S}_x | \alpha \rangle = i\hbar^2 a$ , where  $\hat{S}_x$  and  $\hat{S}_y$  are spin angular momentum operators and  $|\alpha\rangle$  is spin up eigen function, then the value of 'a' is \_\_\_\_\_. (Round off to one decimal place)
- Q.54 A particle in one dimensional box of length 2a with potential energy

$$V = \begin{cases} 0 & |x| < a \\ \infty & |x| > a \end{cases}$$

is perturbed by the potential V' = cx eV, where c is a constant. The 1<sup>st</sup> order correction to the 1<sup>st</sup> excited state of the system is \_\_\_\_\_× c eV.

Q.55 Consider a two dimensional harmonic oscillator with angular frequency  $\omega_x = 2\omega_y = 6.5 \times 10^{14} \text{ rad s}^{-1}$ . The wavelength of x polarized light required for the excitation of a particle from its ground state to the next allowed excited state is \_\_\_\_\_  $\times 10^{-6}$  m. (Round off to one decimal place)

(Speed of light  $c = 3.0 \times 10^8 \text{ m s}^{-1}$ )

END OF THE QUESTION PAPER