## CHEMISTRY-CY

## Q. 1 - Q. 25 : Carry ONE mark each.

1. The Lewis acidity of $\mathrm{BF}_{3}$ is less than $\mathrm{BCl}_{3}$ even though fluorine is more electronegative than chlorine. It is due to
(a) stronger $2 \mathrm{p}(\mathrm{B})-2 \mathrm{p}(\mathrm{F}) \sigma$-bonding
(b) stronger $2 \mathrm{p}(\mathrm{B})-2 \mathrm{p}(\mathrm{F}) \pi$-bonding
(c) stronger $2 p(B)-2 p(C l) \sigma$-bonding
(d) stronger $2 \mathrm{p}(\mathrm{B})-2 \mathrm{p}(\mathrm{Cl}) \pi$-bonding
2. Pyrozenes are a class of silicate minerals, which exhibit a polymeric chain structure, as shown below


Its simplest repeat unit is
(a) $\left[\mathrm{SiO}_{4}\right]^{4-}$
(b) $\left[\mathrm{SiO}_{3}\right]^{2-}$
(c) $\left[\mathrm{Si}_{2} \mathrm{O}_{7}\right]^{6}$
(d) $\left[\mathrm{Si}_{4} \mathrm{O}_{11}\right]^{6-}$
3. Among the following pentachlorides the one which does not exist due to the 'inertpair effect' is
(a) $\mathrm{PCl}_{5}$
(b) $\mathrm{BiCl}_{5}$
(c) $\mathrm{SbCl}_{5}$
(d) $\mathrm{AsCl}_{5}$
4. Band theory predicts that magnesium is an insulator. However, in practice it acts as a conductor due to
(a) presence of filled 3 s orbital
(b) overlap of filled 2 p and filled 3 s orbital
(c) overlap of filled 3s and emtpy 3p orbital
(d) presence of unfilled 3p orbital
5. The number of 'framework electron pairs' present in the borane cluster $\left[\mathrm{B}_{12} \mathrm{H}_{12}\right]^{2-}$ is:
(a) 10
(b) 11
(c) 12
(d) 13
6. The reaction between $\left\{\mathrm{PdCl}_{4}\right]^{2-}$ and $\mathrm{C}_{2} \mathrm{H}_{4}$ produces a new compound. Compare to free $\mathrm{C}_{2} \mathrm{H}_{4}$, the $\mathrm{C}-\mathrm{C}$ bond order of the product is:
(a) between 1 and 2
(b) less than 1
(c) unaltered
(d) greater than 2
7. Among the following pair of metal ions present in Nature, the first one functions as an electrontransfer agent and the second one catalyzes the hydrolysis reactions. The correct pair is
(a) Fe and Zn
(b) Mg and Fe
(c) Co and Mo
(d) Ca and Cu
8. Structurally nickellocene is similar to ferrocene. Nickellocene attains stability due to the formation of
(a) a monocation
(b) a dication
(c) a monoanion
(d) a dianion
9. The absolute configurations for compounds X and Y , respectively are


X


Y
(a) R, S
(b) $\mathrm{S}, \mathrm{R}$
(c) $\mathrm{R}, \mathrm{R}$
(d) $\mathrm{S}, \mathrm{S}$
10. In the reaction,

the major product $[\mathrm{X}]$ is
(a)

(b)

(c)

(d)

11. Among the following, a pair of resolvable configurational enantiomers is given by
(a) cis-1, 2-dimethylcyclohexane
(b) cis-1, 3-dimethylcyclohexane
(c) cis-1, 4-dimethylcyclohexane
(d) trans-1, 3-dimethylcyclohexane
12. In the reaction,

the major product $[\mathrm{X}]$ is:
(a)

(b)

(c)

(d)

13. The decreasing order of isoelectric point for the following $\alpha$-amino acids is
Lysine
(I)
Alanine
(II)
Glutamic acid
(III)
(a) I $>$ II $>$ III
(b) II $>$ I $>$ III
(c) III $>$ I $>$ II
(d) I $>$ III $>$ II
14. The decreasing order of the reactivity of the following compounds towards electrophiles is


I


II


III
(a) II $>$ I $>$ III
(b) II $>$ III $>$ I
(c) III $>$ I $>$ II
(d) I > II > III
15. In the reaction,

the major product $[\mathrm{X}]$ is
(a)

(b)

(c)

(d)

16. The decreasing order of acidity of marked $\mathbf{H}$ of the following molecules is

(I)

(II)

(a) I $>$ II $>$ III
(b) III $>$ I $>$ II
(c) III $>$ II $>$ I
(d) II $>$ I $>$ III
17. The decreasing order of nucleophilicity for the following anions is $\mathrm{CH}_{3} \mathrm{CO}_{2}^{-}, \mathrm{CH}_{3} \mathrm{O}^{-}, \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}^{-}, \mathrm{NO}_{3}^{-}$
(a) $\mathrm{CH}_{3} \mathrm{CO}_{2}^{-}>\mathrm{CH}_{3} \mathrm{O}^{-}>\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}^{-}>\mathrm{NO}_{3}^{-}$
(b) $\mathrm{CH}_{3} \mathrm{O}^{-}>\mathrm{NO}_{3}^{-}>\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}^{-}>\mathrm{CH}_{3} \mathrm{CO}_{2}^{-}$
(c) $\mathrm{CH}_{3} \mathrm{O}^{-}>\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}^{-}>\mathrm{CH}_{3} \mathrm{CO}_{2}^{-}>\mathrm{NO}_{3}^{-}$
(d) $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{O}^{-}>\mathrm{CH}_{3} \mathrm{O}^{-}>\mathrm{NO}_{3}^{-}>\mathrm{CH}_{3} \mathrm{CO}_{2}^{-}$
18. The molar entropy of crystalline CO at absolute zero is
(a) Zero
(b) $-R \ln 2$
(c) $R \ln 2$
(d) $2 \mathrm{R} \ln 2$
19. For an ideal gas
(a) $\left(\frac{\partial \mathrm{P}}{\partial \mathrm{T}}\right)_{\mathrm{V}}\left(\frac{\partial \mathrm{T}}{\partial \mathrm{V}}\right)_{\mathrm{P}}\left(\frac{\partial \mathrm{V}}{\partial \mathrm{P}}\right)_{\mathrm{T}}=0$
(b) $\left(\frac{\partial \mathrm{P}}{\partial \mathrm{T}}\right)_{\mathrm{V}}\left(\frac{\partial \mathrm{T}}{\partial \mathrm{V}}\right)_{\mathrm{P}}\left(\frac{\partial \mathrm{V}}{\partial \mathrm{P}}\right)_{\mathrm{T}}=-1$
(c) $\left(\frac{\partial \mathrm{P}}{\partial \mathrm{T}}\right)_{\mathrm{V}}\left(\frac{\partial \mathrm{T}}{\partial \mathrm{V}}\right)_{\mathrm{P}}\left(\frac{\partial \mathrm{V}}{\partial \mathrm{P}}\right)_{\mathrm{T}}=+1$
(d) $\left(\frac{\partial \mathrm{P}}{\partial \mathrm{T}}\right)_{\mathrm{V}}\left(\frac{\partial \mathrm{T}}{\partial \mathrm{V}}\right)_{\mathrm{P}}\left(\frac{\partial \mathrm{V}}{\partial \mathrm{P}}\right)_{\mathrm{T}}=+2$
20. Among W (work), Q (heat), U (internal energy) and S (entropy)
(a) W and U are path functions but Q and S are state functions.
(b) W and S are path functions but Q and U are state functions.
(c) $S$ and $U$ are path functions but $Q$ and $W$ are state functions.
(d) W and Q are path functions but U and S are state functions.
21. For eigen functions $\psi_{1}=\sqrt{\frac{1}{\mathrm{~b}}} \sin \left(\frac{\pi \mathrm{x}}{\mathrm{b}}\right)$ and $\psi_{2}=\sqrt{\frac{2}{\mathrm{~b}}} \sin \left(\frac{2 \pi \mathrm{x}}{\mathrm{b}}\right)$ of particle in a 1-D box of length $\mathrm{b}(0 \leq \mathrm{x} \leq \mathrm{b})$
(a) $\psi_{1}$ is normalized but not orthogonal to $\psi_{2}$
(b) $\psi_{1}$ is normalized but not orthogonal to $\psi_{2}$
(c) $\psi_{2}$ is normalized and orthogonal to $\psi_{1}$
(d) $\Psi_{2}$ is neither normalized nor orthogonal to $\psi_{1}$
22. The bond order of $\mathrm{C}_{2}$ molecule is
(a) 0
(b) 1
(c) 2
(d) 3
23. Sulfur can exist in four phases. The possible number of triple points is
(a) 1
(b) 2
(c) 3
(d) 4
24. The standard reduction potentials at 298 K for single electrodes are given below:

## Electrode Electrode Potential (volt)

| $\mathrm{Mg}^{2+} / \mathrm{Mg}$ | -2.34 |
| :--- | :--- |
| $\mathrm{Zn}^{2} / \mathrm{Zn}$ | -0.76 |
| $\mathrm{Fe}^{2} / \mathrm{Fe}$ | -0.44 |

From this we can infer that
(a) Zn can reduce both $\mathrm{Mg}^{2+}$ and $\mathrm{Fe}^{2+}$
(b) Fe can reduce both $\mathrm{Mg}^{2+}$ and $\mathrm{Zn}^{2+}$
(c) Mg can reduce both $\mathrm{Zn}^{2+}$ and $\mathrm{Fe}^{2+}$
(d) Mg can reduce $\mathrm{Zn}^{2+}$ but not $\mathrm{Fe}^{2+}$.
25. For the pair of reactions given below
(i) $\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})$
(ii) $\frac{1}{2} \mathrm{~N}_{2}(\mathrm{~g})+\frac{3}{2} \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{NH}_{3}(\mathrm{~g})$

If at a particular temperature, $\mathrm{K}_{\mathrm{P} 1}$ and $\mathrm{K}_{\mathrm{P} 2}$ are the equilibrium constants for reactions (i) and (ii) respectively, then
(a) $\mathrm{K}_{\mathrm{P} 1}=2 \mathrm{~K}_{\mathrm{P} 2}$
(b) $\mathrm{K}_{\mathrm{P} 1}=2 \mathrm{~K}_{\mathrm{P} 2}^{2}$
(c) $\mathrm{K}_{\mathrm{P} 1}=\mathrm{K}_{\mathrm{P} 2}^{2}$
(d) $\mathrm{K}_{\mathrm{P} 1}^{2}=\mathrm{K}_{\mathrm{P} 2}$

## Q. 26 - Q. 55 : Carry TWO marks each.

26. According to VSEPR model, the shape of $\left[\mathrm{XeOF}_{5}\right]^{-}$is
(a) Octahedral
(b) Trigonal bipyramidal
(c) square pyramidal
(d) pentagonal monopyramidal
27. The number of unpaired electron(s) present in the species $\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{5}(\mathrm{NO})\right]^{2+}$ which is formed during 'brown ring test' is:
(a) 2
(b) 3
(c) 4
(d) 5
28. $\mathrm{Fe}_{3} \mathrm{O}_{4}$ and $\mathrm{Co}_{3} \mathrm{O}_{4}$ are metal oxides having spinel structure. Considering their CFSEs, the correct statment regarding their structure is
(a) both have normal spinel structure
(b) both have inverse spinel structure
(c) $\mathrm{Fe}_{3} \mathrm{O}_{4}$ has normal and $\mathrm{Co}_{3} \mathrm{O}_{4}$ has inverse spinel structure
(d) $\mathrm{Fe}_{3} \mathrm{O}_{4}$ has inverse and $\mathrm{Co}_{3} \mathrm{O}_{4}$ has normal spinel structure.
29. The mechanism of the reaction between $\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{4}$ and $\left[\mathrm{Fe}(\text { bpy })_{3}\right]^{3+}$ (bpy $=2,2^{\text {' }}$-bipyridine) is
(a) outer-sphere electron-transfer
(b) inner-sphere electron-transfer
(c) self-exchange reaction
(d) ligand-exchange followed by electron transfer.
30. The d-d absorption band of $\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}{ }^{2+}\right.$ is split due to
(a) presence of octahedral geometry
(b) static Jahn-Teller distortion
(c) dynamic Jahn-Teller distortion
(d) presence of trigonal bipyramidal geometry.
31. The crystal-field symbol for the ground-state of $\left[\mathrm{Mn}(\mathrm{CN})_{6}\right]^{4}$ is
(a) ${ }^{2} \mathrm{~T}_{2 \mathrm{~g}}$
(b) ${ }^{1} \mathrm{~A}_{1 g}$
(c) ${ }^{5} \mathrm{E}_{\mathrm{g}}$
(d) ${ }^{6} \mathrm{~A}_{1 \mathrm{~g}}$
32. In the following reactions:

the reagent/conditions X and Y are
(a) $\mathrm{X}=\mathrm{BF}_{3} ; \mathrm{Y}=$ heating at $1250^{\circ} \mathrm{C}$
(b) $\mathrm{X}=\mathrm{NaF} ; \mathrm{Y}=$ heating at $25^{\circ} \mathrm{C}$
(c) $\mathrm{X}=\mathrm{NH}_{4} \mathrm{~F} ; \mathrm{Y}=\mathrm{HCl}$
(d) $\mathrm{X}=\mathrm{CF}_{3} \mathrm{SO}_{3} \mathrm{H} ; \mathrm{Y}=\mathrm{H}_{2} \mathrm{SO}_{4}$
33. $\left[\mathrm{CoCl}_{4}\right]^{2-}$ is a blue coloured complex. Controlled-treatment of this complex with water generates two isomeric light pink coloure complexes of composition $\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4} \mathrm{Cl}_{2}\right]$.

Identify the correct point groups for $\left[\mathrm{CoCl}_{4}\right]^{2-}$ and two isomeric complexes
$\left[\mathrm{Co}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4} \mathrm{Cl}_{2}\right]$.
(d) $\mathrm{D}_{2 \mathrm{~h}}$ and $\left(\mathrm{C}_{2 \mathrm{v}}\right.$ and $\mathrm{C}_{2 \mathrm{~h}}$ )
(b) $\mathrm{T}_{\mathrm{d}}$ and $\left(\mathrm{C}_{2 \mathrm{v}}\right.$ and $\left.\mathrm{D}_{4 \mathrm{~h}}\right)$
(c) $\mathrm{D}_{4 \mathrm{~h}}$ and $\left(\mathrm{C}_{2 \mathrm{v}}\right.$ and $\left.\mathrm{D}_{4 \mathrm{n}}\right)$
(d) $\mathrm{T}_{\mathrm{d}}$ and $\left(\mathrm{C}_{2 \mathrm{v}}\right.$ and $\left.\mathrm{C}_{4 \mathrm{v}}\right)$
34. In the reaction

the major product $[\mathrm{X}]$ is:
(a)

(b)

(c)

(d)

35. In the reaction,

the major product $[\mathrm{X}]$ is:
(a)

(b)

(c)

(d)

36. In the following reaction sequence

the major product $[\mathrm{X}]$ is
(a)

(b)

(c)

(d)

37. In the reaction,

the major products, $[\mathrm{X}]$ and $[\mathrm{Y}]$, respectively, are
(a)


(b)


(c)

and

(d)


38. In the reaction,

the major product $[\mathrm{X}]$ is
(a)

(b)

(c)

(d)

39. In the reaction sequence

the major products, $[\mathrm{X}]$ and $[\mathrm{Y}]$ respectively, are
(a)
 and

(b)
 and

 and

(d)


40. The change in entropy when two moles of Argon gas are heated at constant volume from 300 K to 500 K is:
(a) $-12.74 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mole}^{-1}$
(b) $-6.37 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mole}^{-1}$
(c) $6.37 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mole}^{-1}$
(d) $12.74 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mole}^{-1}$
41. At any temperature $T$, the fugacity coefficient $(\gamma)$ is given by

$$
\operatorname{In} \gamma=\int_{0}^{\mathrm{P}} \frac{\mathrm{Z}-1}{\mathrm{P}^{\prime}} \mathrm{dP}^{\prime}
$$

where Z is the compressiblity factor. The fugacity coefficient of a real gas governed by equations of state, $\mathrm{P}(\mathrm{V}-\mathrm{b})=\mathrm{RT}$ with 'b' a constant is given by
(a) $\frac{\mathrm{RT}}{\mathrm{bP}}$
(b) $e^{\frac{R T}{b P}}$
(c) $\frac{\mathrm{bP}}{\mathrm{RT}}$
(d) $e^{\frac{b P}{R T}}$
42. The specific rate constant of decomposition of a compound is represented by

Ink $=5.0-\frac{12000}{T}$
The activation energy of decomposition for this compound at 300 K is
(a) $24 \mathrm{kcal} / \mathrm{mole}$
(b) $12 \mathrm{kcal} / \mathrm{mole}$
(c) $24 \mathrm{cal} / \mathrm{mole}$
(d) $12 \mathrm{cal} / \mathrm{mole}$
43. The commutator $\left[x^{3}, p_{x}\right]$ is equal to
(a) $-\frac{3 h x^{2}}{2 \pi i}$
(b) $\frac{h x}{2 \pi i}$
(c) $\frac{h x^{2}}{2 \pi i}$
(d) $\frac{3 h x^{2}}{2 \pi i}$
44. An electron of mass ' $m$ ' is confined to a one dimensional box of length ' $b$ '. If it makes a radiative transition from second excited state to the ground state, the frequency of the photon emitted is
(a) $\frac{9 \mathrm{~h}}{8 \mathrm{mb}^{2}}$
(b) $\frac{3 \mathrm{~h}}{8 \mathrm{mb}^{2}}$
(c) $\frac{\mathrm{h}}{\mathrm{mb}^{2}}$
(d) $\frac{2 \mathrm{~h}}{\mathrm{mb}^{2}}$
45. The point group of $\mathrm{CIF}_{3}$ molecule and its corresponding number of irreducible representation are respectively.
(a) $\mathrm{C}_{3 \mathrm{v}}$ and 4
(b) $\mathrm{C}_{2 \mathrm{v}}$ and 4
(c) $\mathrm{C}_{3 \mathrm{v}}$ and 3
(d) $\mathrm{C}_{2 \mathrm{v}}$ and 3
46. The most populated rotational state for $\mathrm{HCl}\left(\mathrm{B}=8.5 \mathrm{~cm}^{-1}\right)$ at 300 K is:
(a) 2
(b) 3
(c) 5
(d) 7
47. The ratio of life times of two states that gives rise to line widths of $1.0 \mathrm{~cm}^{-1}$ and $0.2 \mathrm{~cm}^{-1}$ respectively is:
(a) $1: 2$
(b) $1: 5$
(c) $2: 1$
(d) $5: 1$

## Commond Data Questions:

## Commond data for Q. 48 and Q.49:

A six-coordinate transition-metal complex is ESR and Mossbauer active. The effective magnetic moment of this complex is -5.9 B.M.
48. The metal-ion along with its oxidation state and the number of unpaired electron present are
(a) Fe (II) and 4
(b) Mn (II) and 5
(c) Fe (III) and 1
(d) Fe (III) and 5
49. The complex is
(a) $\left[\mathrm{Mn}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$
(b) $\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{3-}$
(c) $\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$
(d) $\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{3+}$

## Common data for Q. 50 and Q.51:

An organic compound $[\mathrm{X}]\left(\mathrm{C}_{12} \mathrm{H}_{16} \mathrm{O}_{3}\right)$ exhibits the following spectral data IR: $-1720 \mathrm{~cm}^{-1}$
${ }^{1} \mathrm{H}$ NMR: $2.35(\mathrm{~s}, 6 \mathrm{H}), 3.30(\mathrm{~s}, 3 \mathrm{H}), 3.83(\mathrm{t}, 2 \mathrm{H}), 4.42(\mathrm{t}, 2 \mathrm{H}), 7.07(\mathrm{~s}, 1 \mathrm{H}), 7.58(\mathrm{~s}, 2 \mathrm{H})$
The compound $[\mathrm{X}]$ with an excess of MeMgBr gives a $1: 1$ mixture of compounds [ Y$]$ and $[\mathrm{Z}]$. The compound [Z] exhibits the following ${ }^{1} \mathrm{H}$ NMR data: $2.0(\mathrm{bs}, 1 \mathrm{H}), 3.30(\mathrm{~s}, 3 \mathrm{H}), 3.56(\mathrm{t}, 2 \mathrm{H}), 3.70(\mathrm{t}, 2 \mathrm{H})$
50. The compound $[\mathrm{X}]$ is:
(a)

(b)

(c)

(d)

51. The compound [ $\mathrm{Y}[$ is:
(a)
 and

(b)
 and

(c)
 and

(d)
 and


## Linked Answer Q. 52 and Q.53:

In the reaction sequence

52. The compound $[\mathrm{X}]$ is:
(a)

(b)

(c)

(d)

53. The compound $[Z]$ is:
(a)

(b)

(c)

(d)


Statement for Linked Answer Q. 54 and Q.55:
In the m vs T diagram for different phases of the same substance at one atmospheric pressure, the lines $\mathrm{A}, \mathrm{B}$ and C compound to

54. Based on the above diagram.
(a) A represents the change in chemical potential as a function of temperature for the solid phase, B for the liquid and C for the gas.
(b) A represents the change in chemical potential as a function of temperature for the gas phase, B for the gas and C for the solid
(c) A represents the change in chemical potential as a function of temperaturefor the liquid phase, B for the gas and C for the solid
(d) A represents the change in chemical potential as a function of temperaturefor the gas phase, B for the gas and C for the liquid.
55. From the same diagram
(a) D represents boiling point, E sublimation point and F melting point.
(b) E represents boiling point, D sublimation point and F melting point.
(c) E represents melting point, F sublimation point and D boiling point.
(d) D represents melting point, F boiling point and E sublimation point.

