Section A: Q. 1 - Q. 10 Carry ONE mark each.
Q. 1
Q. 2 (2)
(S. 3


Q.5 \begin{tabular}{|l|l|}

\hline (A) \& | Which one of the following options is best suited for effecting the |
| :--- |
| transformation? | \\

(B) \& $\mathrm{DMSO}_{2}$ \\
(D) \& $\mathrm{Ag}_{2}(\mathrm{Oi}-\mathrm{Pr})_{3}$
\end{tabular}



| Q. 7 | Among the following, the compound that forms the strongest hydrogen bond is |
| ---: | :--- |
| (A) | HF |
| (B) | HCl |
| (C) | HBr |
| (D) | HI |
| Q.8 | Among the following, the biomolecule with a direct metal-carbon bond is |
| (A) | coenzyme $\mathrm{B}_{12}$ |
| (B) | nitrogenase |
| (C) | chlorophyll |



Section A: Q. 11 - Q. 30 Carry TWO marks each.



| Q.13 | The major product formed in the reaction of $(2 S, 3 R)$-2-chloro-3-phenylbutane <br> with NaOEt in EtOH is |
| ---: | :--- |
| (A) | $($ (E)-2-phenyl-but-2-ene |
| (C) | (Z)-2-phenyl-but-2-ene |
| (D) | $(2 R, 3 R)$-2-ethoxy-3-phenylbutane |
|  |  |

Q. 14


| Q.17 | The structure types of $\mathrm{B}_{10} \mathrm{H}_{10}{ }^{2-}$ and $\mathrm{B}_{10} \mathrm{H}_{14}$, respectively, are |
| ---: | :--- |
| (A) | closo and nido |
| (B) | nido and arachno |
| (D) | nido and closo |
| Q.18 and arachno |  |
| The ground state and the maximum number of spin-allowed electronic |  |
| (D) | ${ }^{4} \mathrm{~T}_{1}$ and 3 |
| (B) | ${ }^{4} \mathrm{~A}_{2}$ and 3 |
| (C) | ${ }^{4} \mathrm{~A}_{2}$ and 2 |


| Q. 19 | The correct statement about the geometries of $\mathrm{BH}_{2}{ }^{+}$and $\mathrm{NH}_{2}{ }^{+}$based on valence shell electron pair repulsion (VSEPR) theory is |
| :---: | :---: |
| (A) | both $\mathrm{BH}_{2}{ }^{+}$and $\mathrm{NH}_{2}{ }^{+}$are trigonal planar |
| (B) | $\mathrm{BH}_{2}^{+}$is linear and $\mathrm{NH}_{2}^{+}$is trigonal planar |
| (C) | $\mathrm{BH}_{2}{ }^{+}$is trigonal planar and $\mathrm{NH}_{2}{ }^{+}$is linear |
| (D) | both $\mathrm{BH}_{2}{ }^{+}$and $\mathrm{NH}_{2}{ }^{+}$are linear |
| Q. 20 | The order of increasing CO stretching frequencies in $\left[\mathrm{Co}(\mathrm{CO})_{4}\right]^{-},\left[\mathrm{Cu}(\mathrm{CO})_{4}\right]^{+}$, $\left[\mathrm{Fe}(\mathrm{CO})_{4}\right]^{2-}$ and $\left[\mathrm{Ni}(\mathrm{CO})_{4}\right]$ is |
| (A) | $\left[\mathrm{Cu}(\mathrm{CO})_{4}\right]^{+}<\left[\mathrm{Ni}(\mathrm{CO})_{4}\right]<\left[\mathrm{Co}(\mathrm{CO})_{4}\right]^{-}<\left[\mathrm{Fe}(\mathrm{CO})_{4}\right]^{2-}$ |
| (B) | $\left[\mathrm{Fe}(\mathrm{CO})_{4}\right]^{2-}<\left[\mathrm{Co}(\mathrm{CO})_{4}\right]^{-}<\left[\mathrm{Ni}(\mathrm{CO})_{4}\right]<\left[\mathrm{Cu}(\mathrm{CO})_{4}\right]^{+}$ |
| (C) | $\left[\mathrm{Co}(\mathrm{CO})_{4}\right]<\left[\mathrm{Fe}(\mathrm{CO})_{4}\right]^{2-}<\left[\mathrm{Cu}(\mathrm{CO})_{4}\right]^{+}<\left[\mathrm{Ni}(\mathrm{CO})_{4}\right]$ |
| (D) | $\left[\mathrm{Ni}(\mathrm{CO})_{4}\right]<\left[\mathrm{Cu}(\mathrm{CO})_{4}\right]^{+}<\left[\mathrm{Co}(\mathrm{CO})_{4}\right]^{-}<\left[\mathrm{Fe}(\mathrm{CO})_{4}\right]^{2-}$ |
|  |  |

Q.21

| Q.22 | The stability of adducts $\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{PF}_{3}, \mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{NMe}_{3}, \mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{CO}, \mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{OMe}_{2}$ follows the <br> order |
| ---: | :--- |
| (A) | $\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{OMe}_{2}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{CO}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{PF}_{3}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{NMe}_{3}$ |
| (C) | $\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{PF}_{3}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{CO}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{NMe}_{3}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{OMe}_{2}$ |
| (D) | $\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{PF}_{3}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{CO}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{PF}_{3}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{NMe}_{3}<\mathrm{OMe}_{2}<\mathrm{H}_{3} \mathrm{~B} \cdot \mathrm{OMe}_{2}$ |

Q. $23 \quad$| The spacing between successive rotational energy levels of a diatomic |
| :--- |
| molecule $X Y$ and its heavier isotopic analogue $X^{\prime} Y^{\prime}$ varies with the rotational |
| quantum number, $J$, as |

| Q.24 | The ratio of the $2 \mathrm{p} \rightarrow 1$ s transition energy in $\mathrm{He}^{+}$to that in the H atom <br> is closest to |
| ---: | :--- | :--- |
| (A) | 1 |
| (B) | 2 |
| (D) | 8 |
|  |  |



| Q.26 | Capillary $W$ contains water and capillary $M$ contains mercury. The contact <br> angles between the capillary wall and the edge of the meniscus at the air-liquid <br> interface in $W$ and $M$ are $\theta_{W}$ and $\theta_{M}$, <br> The contact angles satisfy the conditions |
| ---: | :--- |
| (A) | $\theta_{W}>90^{\circ}$ and $\theta_{M}>90^{\circ}$ |$\theta_{\text {(B) }}$| (D) | $\theta_{W}<90^{\circ}$ and $\theta_{M}<90^{\circ}$ |
| :--- | :--- | :--- |



Q.29 $\quad$ The number of CO stretching bands in the infrared spectrum of $\mathrm{Fe}(\mathrm{CO})_{5}$ is

| Q.30 | The standard Gibbs free energy change for the reaction <br> $\mathrm{H}_{2} \mathrm{O}(\mathrm{g}) \rightarrow \mathrm{H}_{2}(\mathrm{~g})+\frac{1}{2} \mathrm{O}_{2}(\mathrm{~g})$ <br> at 2500 K is $+118 \mathrm{~kJ} \mathrm{~mol}^{-1}$. <br> The equilibrium constant for the reaction is <br> [Given: $R=8.314 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$ ] |
| :--- | :--- |
| (A) | 0.994 |
| (C) | $3.42 \times 10^{-3}$ |
| (D) | 292.12 |

Section B: Q. 31 - Q. 40 Carry TWO marks each.

Q.31 | Among the following, the reaction(s) that favor(s) the formation of the products |
| :--- |
| at $25^{\circ} \mathrm{C}$ is/are |
| (B) |
| (C) |

| Q.32 | Among the following, the correct statement(s) is/are: |
| ---: | :--- |
| (A) | The first $\mathrm{p} K_{\mathrm{a}}$ of malonic acid is lower than the $\mathrm{p} K_{\mathrm{a}}$ of acetic acid while its <br> second $\mathrm{p} K_{\mathrm{a}}$ is higher than the $\mathrm{p} K_{\mathrm{a}}$ of acetic acid. <br> (C) <br> Both the first and the second $\mathrm{p} K_{\mathrm{a}} \mathrm{s}$ of malonic acid are lower than the $\mathrm{p} K_{\mathrm{a}}$ of of malonic acid is higher than the $\mathrm{p} K_{\mathrm{a}}$ of acetic acid while its <br> acetic acid. <br> (D) <br> Both the first and the second $\mathrm{p} K_{\mathrm{a}} \mathrm{s}$ of malonic acid are higher than the $\mathrm{p} K_{\mathrm{a}}$ of <br> acetic acid. |

Q. 33 (A) | The compound(s) that participate(s) in Diels-Alder reaction with |
| :--- |
| maleic anhydride is/are |
| (B) |

| Q.34 | Among the following, the suitable route(s) for the conversion of benzaldehyde <br> to acetophenone is/are |
| ---: | :--- |
| (A) | $\mathrm{CH}_{3} \mathrm{COCl}$, anhydrous $\mathrm{AlCl}_{3}$ |
| (B) | (i) $\mathrm{HS}\left(\mathrm{CH}_{2}\right)_{3} \mathrm{SH}, \mathrm{F}_{3} \mathrm{~B} \cdot \mathrm{OEt}_{2}$; (ii) n-BuLi; (iii) MeI; (iv) $\mathrm{HgCl}_{2}, \mathrm{CdCO}_{3}, \mathrm{H}_{2} \mathrm{O}$ |$|$| (DaNH, MeI |
| :--- |
| (i) MeMgBr ; (ii) aq. acid; (iii) pyridinium chlorochromate (PCC) |
|  |

Q.35

| Q.36 | The reason(s) for the lower stability of $\mathrm{Si}_{2} \mathrm{H}_{6}$ compared to $\mathrm{C}_{2} \mathrm{H}_{6}$ is/are |
| ---: | :--- |
| (A) | silicon is more electronegative than hydrogen |
| (C) | Si-Si bond is weaker than $\mathrm{C}-\mathrm{C}$ bond |
| (D) | the presence of low-lying d-orbitals in silicon |
|  |  |


| Q.37 | For an $N$-atom nonlinear polyatomic gas, the constant volume molar heat <br> capacity $C_{v, m}$ has the expected value of $3(N-1) R$, based on the principle of <br> equipartition of energy. The correct statement(s) about the measured value of <br> $C_{v, m}$ is/are |
| ---: | :--- |
| (A) | The measured $C_{v, m}$ is independent of temperature. |
| (B) | The measured $C_{v, m}$ is dependent on temperature. |
| (C) | The measured $C_{v, m}$ is typically lower than the expected value. |
| (D) | The measured $C_{v, m}$ is typically higher than the expected value. |
| (D) | urease |
| hydrogenase |  |
| (A) | carboxypeptidase |
| carbonic anhydrase |  |



Q.43


## Section C: Q. 51 - Q. 60 Carry TWO marks each.

| Q. 51 | The specific rotation of optically pure $(R)-1$-phenylethylamine is +40 (neat, $20^{\circ} \mathrm{C}$ ). A synthetic sample of the same compound is shown to contain 4:1 mixture of $(S)$ - and $(R)$-enantiomers. The specific rotation of the neat sample at $20^{\circ} \mathrm{C}$ is $\qquad$ (round off to the nearest integer) |
| :---: | :---: |
|  |  |
|  |  |
| Q. 52 | The number of $\beta$ particles emitted in the nuclear reaction ${ }_{92}^{238} U \rightarrow{ }_{82}^{206} \mathrm{~Pb}$ is $\qquad$ . |
|  | ¿uras |
| Q. 53 | Iron is extracted from its ore via the reaction $\mathrm{Fe}_{2} \mathrm{O}_{3}+3 \mathrm{CO} \rightarrow 2 \mathrm{Fe}+3 \mathrm{CO}_{2}$ <br> The volume of CO (at STP) required to produce 1 kg of iron is $\qquad$ liters. <br> (round off to the nearest integer) <br> [Given: Atomic wt. of $\mathrm{Fe}=56$; assume STP to be $0^{\circ} \mathrm{C}$ and 1 atm ] |
|  |  |
|  |  |




