Electronic Spectra of Transition Metal Complexes

| 1. | Electronic spectra of transition metal complex observed in region. | | | | |
|----|--|---------|---|--|--|
| | a) 1R | b) | UV | | |
| | c) visible | d) | UV-visible | | |
| 2. | Electronic spectra of transition metal comple | exer gi | ve the information of of transition | | |
| | metal complexes. | | | | |
| | a) colour | b) | magnetic properties | | |
| | c) Tendency to form soluble complex | d) | All above | | |
| 3. | Metal complex absorb the light in the visible | e regio | on | | |
| | a) 100-200 nm | b) | 200-250 nm | | |
| | c) 400-700 nm | d) | 800-1200 nm | | |
| 4. | The spectra of coloured solution of transition | on met | al complexes is measured by using | | |
| | a) Spectrophotometer | b) | Conductometer | | |
| | c) pH meter | d) | lonometer | | |
| 5. | Spectrophotometer is based on the principle | e of | | | |
| | a) Ostwald's law | b) | Lambert's - Beer's law | | |
| | c) Ostribution law | d) | None of the above | | |
| 6. | Which of the following is the type of electro | onic tr | ansition | | |
| | a) d-d transition | b) | LMCT | | |
| | c) MLCT | d) | All of the above | | |
| 7. | Purple colour of $\left[Ti (H_2 O)_6 \right]^{3+}$, Blue colou | ur of [| $\operatorname{Cu}(\operatorname{NH}_3)_4^{2+}$ and pink colour of $[\operatorname{CoF}_6^{3-}]^{3-}$ | | |
| | is / are dueto | | | | |
| | a) d-d transition | b) | LMCT | | |
| | c) MLCT | d) | interligand transition | | |
| 8. | Intense red colour of $\left[\text{Fe}(\text{SCN})_6 \right]^{3-1}$ | | | | |
| | Yellow colour of $K_2 CrO_4$ | | | | |
| | Yellow brown colour of $[Fe(H_2O)_5 OH]^{3+}$ | | | | |
| | Dark purple colour of KHnO ₄ , | | | | |
| | Intense colour of [V(Ben30 hydroxidic acid | ł)], | | | |
| | Intense colour of [Fe (N-phenyl hydroximic | c acid) |], | | |
| | orange colour of $K_2 Cr_2 O_7$ | | | | |
| | is/are due to | | | | |

| | a) | d-d transition | b) | LMCT |
|-----|------|--|------------------|---|
| | c) | MLCT | d) | interligand transition |
| 9. | Bri | ght yellow colour of $[Cu(Phen)_2]$, colour o | fme | tal carbonyl such as $Ni(CO)_4$, $Fe_2(CO)_9$ and |
| | [Fe | $e(dipy)_3]^{2+}$ is / are due to | | |
| | a) | d-d transition | b) | LMCT |
| | c) | MLCT | d) | interligand transition |
| 10. | Du | ring the transition between the energy leve | ls, ai | h electron does not change its spin i.e. $\Delta S =$ |
| | 0 is | s the statement of | | |
| | a) | spin selection rule | b) | orbital selection rule (Laporte rule) |
| | c) | Both a & b | d) | None of the above |
| 11. | Wł | nich of the following is Laporte selection r | ule. | |
| | a) | $\Delta 1 = \pm 1$ | b) | $\Delta 1 = 0$ |
| | c) | Both a and b | d) | None of thes above |
| 12. | Spe | ectroscopic ground state term symbol of d | ¹ cor | nfiguration is |
| | a) | 2 D | b) | 2 F ₂ |
| |) | $-\frac{3}{2}$ | -, | <u>5</u> 2 |
| | c) | 2 D ₂ | d) | ${}^{2}F_{2}$ |
| | | <u></u> | | <u>-</u> 3 |
| 13. | Spe | etroscopic ground state term symbol of | ••••• | configuration. |
| | | (for oxygen) $d^2 = {}^4F_{\frac{3}{2}}$ | | |
| | | $d^2 = {}^3F_2$ | | |
| | | $d^8 = {}^3F_4$ | | |
| | | oxygen = ${}^{3}P_{2}$ | | |
| | | $\mathbf{V}^{+}=\ {}^{4}F_{\underline{3}}$ | | |
| 14. | Org | gel energy level digram is widely used for | | complexes |
| | a) | Weak ligand | b) | strong ligand |
| | c) | Both a and b | d) | None of the above |
| 15. | Но | le formulation concept observed in | | |
| | | | | |

- a) $\left[ri(H_2O)_6 \right]^{3+}$ b) $\left[Cu(H_2O)_6 \right]^{2+}$
- c) Both a & b d) none of the above

| 16. | 16. The electronic septra of $\left[\operatorname{Ti}(\operatorname{H}_{2}\operatorname{O})_{6}\right]^{3+}$ show only one band with a peak at | | | | |
|-----|--|-------|-----------------------------|--|--|
| | a) 20300 cm ⁻¹ | b) | 10150 cm^{-1} | | |
| | c) 25600 cm^{-1} | d) | 30600 cm^{-1} | | |
| 17. | $\left[Ti (H_2 O)_6 \right]^{3+}$ ion absorb and abea | r pur | ple. | | |
| | a) Red visible light | b) | Green-yellow visible light | | |
| | c) Violet visible light | d) | Blue visible light | | |
| 18. | Ground state term symbol for He (16 ²) is | ••••• | | | |
| | a) ${}^{1}S_{\frac{1}{2}}$ | b) | ${}^{1}S_{0}$ | | |
| | c) ${}^{0}S_{\frac{1}{2}}$ | d) | none of these | | |
| 19. | For $L = 0$ and $S = 1$ full spectroscopic term i | s | | | |
| | a) ${}^{5}D_{1}$ | b) | ¹ S ₁ | | |
| | c) ${}^{3}S_{1}$ | d) | ¹ P ₃ | | |
| 20. | Ground state for 2P ³ is | | | | |
| | a) ${}^{4}S_{3}$ | b) | ${}^{3}F_{4}$ | | |
| | c) ${}^{4}S_{\frac{5}{2}}$ | d) | ${}^{2}P_{1}$ | | |
| 21. | Which of the following corruspoun to at | sorp | tion beak of maximum wave n | | |
| | $\left[Cr(H_2O)_6 \right]^{3+}$ complex 100? | | | | |

- a) ${}^{4}A_{2g} \longrightarrow {}^{4}T_{2g}(F)$ b) ${}^{4}A_{2g} \longrightarrow {}^{4}T_{1g}(P)$
- c) ${}^{4}A_{2g} \longrightarrow {}^{4}T_{1g}(F)$ d) ${}^{4}A_{2g}(F) \longrightarrow {}^{4}T_{1g}(F)$

22. Solution of $[COCl_4]^{2-}$ are deep blue because.....

- a) it is tetrahedral ion b) it has d⁷ configuration
- c) its transition in blue spectrum **d**) its transition is ${}^{4}A_{2}(F) \longrightarrow {}^{4}T_{2}(P)$

number in

Coordination Theory (Part - Second)

| 1. | Valence bond theory of coordination compound was put by the scientist | | | | | | | | |
|--|---|--|--------|--------------------------------|--|--|--|--|--|
| | a) | Pauling and slater | b) | G.M. Lewis | | | | | |
| | c) | Van vleck | d) | Pearson | | | | | |
| 2. | Val | lence bond theory of coordination explain | ••••• | | | | | | |
| | a) | Formation of coordination compound | b) | Stereosomerism | | | | | |
| | c) | Magnetic properties | d) | All of the above | | | | | |
| 3. | Ac | According to valence bond theory of coordination compound the nd orbital involved in | | | | | | | |
| | hył | nybridisation maybe | | | | | | | |
| | a) | Only inner orbitals | b) | Only outer orbitals | | | | | |
| | c) | Inner orbitals or outer orbitals | d) | None of the above | | | | | |
| 4. | Wł | nich of the following pair is correct? | | | | | | | |
| | a) | Coordination no. 4 - dsp^2 | b) | coordination No. 6 - d^2sp^2 | | | | | |
| | c) | Coordination no. 7 - $d^3sp^{3 }$ | d) | All above pairs are correct | | | | | |
| 5. | d ² s | sp^3 and sp^3d^2 type of complexes are also k | nowr | n as | | | | | |
| | a) | inner orbital complexes | b) | Outer orbital complexes | | | | | |
| | c) | Both inner and outer orbital complex | es | d) None of the above | | | | | |
| 6. | Inn | ner orbital complexes are also known as lo | ow sp | in complexes because | | | | | |
| | a) | Complex having less No. of unpaired | rons | | | | | | |
| | b) | Complex having maximum No. of unpai | d eleo | ctrons | | | | | |
| | c) | Both a & b | d) | Can't say | | | | | |
| 7. | F | $e(CN)_6^{-3}$ is an example of | | | | | | | |
| | a) | inner orbital complexes | b) | Outer orbital complexes | | | | | |
| | c) | Sandwitch complexes | d) | Crown ether | | | | | |
| 8. | Hy | bridisation in $\left[\text{Fe}(\text{CN})_6 \right]^{-3}$ is | | | | | | | |
| | a) | d ² sp ³ | b) | dsp ² | | | | | |
| | c) | dsp ³ | d) | d ³ sp ³ | | | | | |
| 9. | Ma | agnetic nature of $\left[Fe(CN)_6 \right]^{-3}$ is | | | | | | | |
| | a) | Paramagnetic | b) | Dimagnetic | | | | | |
| | c) | Ferromagnetic | d) | Antiferromagnetic | | | | | |
| 10. | Ou | ter orbital complexes are also known as h | igh s | pin complexes because | | | | | |
| a) Complex having less No. of unpaired electrons | | | | | | | | | |

b) Complex having maximum No. of unpaid electrons

| | c) Both a & b | d) | None of the above |
|-----|--|------------|---------------------------------|
| 11. | $\left[\operatorname{Fe} F_{6}\right]^{-3}$ is an example of | | |
| | a) Inner orbital complexes | b) | Outer orbital complexes |
| | c) Sandwitch complexes | d) | Crown ether complexes |
| 12. | Hybridisation in $[FeF_6]^{-3}$ is | | |
| | a) sp ³ d ² | b) | dsp ² |
| | c) dsp^3 | d) | d ³ sp ³ |
| 13. | Magnetic nature of $[FeF_6]^{-3}$ is | | |
| | a) Paramagnetic | b) | Dimagnetic |
| | c) Ferromagnetic | d) | Antiferromagnetic |
| 14. | Inner and outer orbital complexes respect | tively kn | own as |
| | a) d^2sp^3 and sp^3d^2 complexes | b) | Lowspin and high spin complexes |
| | c) Spin paired and spin free complexes | d) | All above are correct |
| 15. | Limitahon of VBT (Valence bond theory) |) of coor | dination compound is/are |
| | a) Fail to explain splitting of d orbitals | | |
| | b) Fail to interpret spectra of complex | | |
| | c) Fail to predict the relative energies of | f comple | exes |
| | d) All of the above | | |
| 16. | eg orbital is a set of | | |
| | a) <i>dxy</i> , <i>dxz</i> , <i>dyz</i> orbitals | b) | dx^2y^2 & dz^2 orbitals |
| | c) $dxy \& dz^2$ orbitals | d) | $dyz \& dx^2 - y^2$ orbitals |
| 17. | t ₂ g orbital is a set of | | |
| | a) <i>dxy</i> , <i>dxz</i> , <i>dyz</i> orbitals | b) | dx^2y^2 & dz^2 orbitals |
| | c) $dxy \& dz^2$ orbitals | d) | $dyz \& dx^2 - y^2$ orbitals |
| 18. | t ₂ g orbitals | | |
| | a) lie along with the axis | b) | lie inbetween the axis |
| | c) Both a & b | d) | None of the above |
| 19. | eg orbitals | | |
| | a) lie along with axis | b) | lie in between the axis |
| | c) Both a and b | d) | None of the above |
| 20. | Crystal field theory (CFT) was put by the | e scientis | t |
| | a) Huggins | b) | Pauling |
| | c) Shter | d) | Bethe & Vanvleck |
| 21. | The correct statements of CFT is / are | ••••• | |

a) Ligands are point charges

| | b) | b) Attraction is purely electrostatic | | | | | |
|-----|----|--|--------|----------------------|--|--|--|
| | c) | Degeneracy of a orbital is destroyed when comples form | | | | | |
| | d) | All above are correct | | | | | |
| 22. | Ax | ial orbitals of d-orbitals are | | | | | |
| | a) | dx^2 - y^2 & dz^2 | b) | dxy, dxz & dyz | | | |
| | c) | $dxy \& dz^2$ | d) | dx^2y^2 & dyz | | | |
| 23. | Ac | cording to CFT the bond between metal & | k liga | and is | | | |
| | a) | purely electrostatic | b) | Covalent | | | |
| | c) | Metalic | d) | Coordinate bond | | | |
| 24. | Un | der the influence of ligands d orbitals of n | netal | ion split into | | | |
| | a) | $dxy \& dz^2$ | b) | eg & t _{2g} | | | |
| | c) | eg & dx^2-y^2 | d) | $t_{2g} \& dxy$ | | | |
| 25. | Th | e splitting of degenerate d orbitals into tw | o set | s is known as | | | |
| | a) | crystal field splitting | b) | ionisation energy | | | |
| | c) | sublimation energy | d) | excitation energy | | | |
| 26. | Cr | ystal field splitting energy is denoted by | ••••• | | | | |
| | a) | 10 Dq | b) | 1.0 Δ_{o} | | | |
| | c) | Both a and b | d) | None of the above | | | |
| 27. | In | octahedral complexes the energy of orbita | l is | | | | |
| | a) | $eg > t_{2g}$ | b) | $t_{2g} > eg$ | | | |
| | c) | $t_{2g} = eg$ | d) | cant say | | | |
| 28. | Wł | nich CFSE is correct in octahedral cmplex | es | | | | |
| | a) | $d^2 = -4 Da$ | b) | $d^2 = -8 Da$ | | | |
| | c) | $d^3 = -12 \text{ Da}$ | d) | All are correct | | | |
| 29. | CF | SE of weak and strong field octaheral cor | npley | kes respectively is | | | |
| | | $d^4 = -6 Dq and -16 Dq + P$ | | | | | |
| | | $d^5 = -0 Dq and -20 Dq + 2P$ | | | | | |
| | | $d^6 = -4 Dq + P and -24 Dq + 3P$ | | | | | |
| | | $d^7 = -8 Dq + 2P and -18 Dq + 3P$ | | | | | |
| | | $d^8 = -12 Dq + 3P and -12 Dq + 3P$ | | | | | |
| | | $d^9 = -6 Dq + 4P and -6 Dq + 4P$ | | | | | |
| | | $d^{10} = 0 + 5P$ Da and 0 Dq + 5P | | | | | |
| 30. | In | tetrahedral complexes the energy of orbita | ıl is | | | | |
| | a) | $eg > t_{2g}$ | b) | $t_{2g} > eg$ | | | |
| | c) | $t_{2g} = eg$ | d) | can't say | | | |

31. The comparison of energy of tetrahedral complexes with octahedral complexes is shown by equation....

| | a) $\Delta t = -\frac{4}{9} \Delta_0$ | | b) | $\Delta t = \frac{4}{9} \Delta_0$ |
|-----|--|-------------------|-------|--|
| | c) $\Delta_0 = \frac{-4}{9} \Delta t$ | | d) | $\Delta_0 = \frac{5}{9}\Delta t$ |
| 32. | Low spin complexs of tetrah | edral geometry | have | not yet been obtain, the reason is / are |
| | a) Not a single d orbital is | along with axis | b) | Less no. of ligaod |
| | c) a correct | | d) | Both a & b correct |
| 33. | Only high spin complexes ar | e observed in tet | trahe | dral complexes the reason is/are |
| | a) Not a single d orbital is | along with axis | b) | Less no. of ligands |
| | c) Only a corred | | d) | Both a and b are correct |
| 34. | The energy of tetrahedral co | omplexes is less | than | octahedral complexes i.e. $\Delta t = \frac{-4}{9} \Delta_0$ the |
| | reason is | | | |
| | a) Not a single d orbital is | along with axis | b) | Less no. of ligands |
| | c) Only a corred | | d) | Both a and b are correct |
| 35. | CFSE of tetrahedral complex | xes is as follows | ••••• | |
| | Configuration | C.F.S.E. value | | |
| | d ¹ | – 6 Dq | | |
| | d^2 | – 12 Dq | | |
| | d ³ | – 8 Dq | | |
| | d^4 | – 4 Dq | | |
| | d ⁵ | 0 Dq | | |
| | d^6 | - 6 Dq + P | | |
| | d^7 | - 12 Dq + 2P | | |
| | d^8 | - 8 Dq + 3P | | |
| | d ⁹ | -4 Dq $+4$ P | | |
| | d^{10} | 0 Dq + 5P | | |
| 36. | Splitting of d orbitals in tetra | agonal and squar | e bla | nar complexes arises from |
| | a) Regular tetrahedral com | plexes | b) | Regular square planar complexes |
| | c) Regular octahedral cm | plexes | d) | None of these |
| 37. | In tegragonal (elongated oct | ahedral) comple | xes e | energy of orbital is |
| | a) $dx^2 - y^2$, $dxy > dz^2$, dxy , | dyz. | b) | $dx^2 - y^2, dxy < dz^2, dxy, dyz$ |
| | c) $dxy dxz dyz > dz^2$ | | d) | $dx^2 - y^2, dxy = dz^2, dxy, dyz$ |
| 38. | In square planar complexes of | energy of orbital | is | |
| | a) $dx^2 - y^2$, $dxy > dz^2$, dxy , | dyz. | b) | $dx^2 - y^2, dxy < dz^2, dxy, dyz$ |
| | c) dx^2y^2 , $dxz = dz^2$, dxy , dy | 27. | d) | None of the above |
| | | | | |

| 39. | Which of the following factors affecting the 10 Dg (magnitude of crystal field splitting) | | | | | |
|-----|--|--|--|--|--|--|
| | a) Nature of ligand | b) Oxidation state of metalion | | | | |
| | c) Size of d orbitals | d) Geometry of complexes | | | | |
| | e) All are above correct | | | | | |
| 40. | According to spectrochemical series which | h one of the following correct increasing order of | | | | |
| | strength of ligaods. | | | | | |
| | a) $Br^- < H_2O < CN^- < CO$ | b) $CO < CN^{-} < H_2O < Br^{-}$ | | | | |
| | c) $Br^{-} < CO < H_{2}O < CN^{-}$ | d) $CN^{-} < Br^{-} < H_2O < CO$ | | | | |
| 41. | Which one of the following equation is con- | rrect about the crystal field splitting of cobalt ion. | | | | |
| | a) Co (II) < Co (III) | b) Co (III) = Co(II) | | | | |
| | c) Co (II) > Co (III) | d) All above are correct | | | | |
| 42. | Which one of the following is correct about | at crystal field splitting | | | | |
| | a) $3d < 4d < 5d$ | b) $3d > 4d > 5d$ | | | | |
| | c) $3d = 5d > 4d$ | d) $3d > 5d > 4d$ | | | | |
| 13 | According to CET $\begin{bmatrix} Ti(H O) \end{bmatrix}^{+3}$ complete | y ion show colour | | | | |
| 45. | According to $CI + [\Pi(\Pi_2 O)_6]$ complete | | | | | |
| | a) Purple | b) Green | | | | |
| | c) Blue | d) | | | | |
| 44. | The colour of coordination compound is a | rises due to transition of electrons | | | | |
| | a) $S \longrightarrow P$ | b) $P \longrightarrow d$ | | | | |
| | c) $\mathbf{d} \longrightarrow \mathbf{d}$ | d) $d \longrightarrow F$ | | | | |
| 45. | The electronic configuration of $[Ti(H_2O)]$ | 1 ³⁺ is | | | | |
| | | 6 _ | | | | |
| | a) $t_{2g}^0 e_g^0$ | b) $t_{2g}^1 e_g^0$ | | | | |
| | $x^{2} = x^{0}$ | $1 + \frac{3}{2} = 0$ | | | | |
| | c) $l_{2g} e_g$ | a) $l_{2g} e_g$ | | | | |
| 46. | $\left[\operatorname{Ti}(\operatorname{H}_2\operatorname{O})_6\right]^{3+}$ absorb radiation and | nd show purple colour | | | | |
| | a) I.R. | b) U.V. | | | | |
| | c) Visible | d) Radio | | | | |
| 47. | When $\left[Ti (H_2 O)_6 \right]^{3+}$ absorb visible light r | resulting excitation of electron takes place from | | | | |
| | a) $t_{2g}^1 \longrightarrow e_g^0$ | b) $t_{2g}^0 \longrightarrow e_g^1$ | | | | |
| | c) $t_{2g}^{1} \longrightarrow e_{g}^{1}$ | d) $e_g^1 \longrightarrow t_{2g}^1$ | | | | |
| | | | | | | |

| 48. | According to CFT, how much unpaired electrons are present in $[CoF_6]^{3-}$ | | | | |
|-----|---|--|--------|---|--|
| | a) | 1 | b) | 2 | |
| | c) | 3 | d) | 4 | |
| 49. | Ace | cording to CFT, how much unpaired electr | rons | are present in $\left[Co(NH_3)_6 \right]^{3+}$ | |
| | a) | 0 | b) | 1 | |
| | c) | 2 | d) | 4 | |
| 50. | In o | octahedral complexes t ₂₉ orbitals are stabil | ized | by | |
| | a) | -4 Dq OR -0.4 Δ_{o} | b) | +6 Dq OR +0.6 Δ_{o} | |
| | c) | 10 Dq OR 1.0 Δ_{0} | d) | none of these | |
| 51. | In o | octahedral complese eg orbitals are destab | ilizec | 1 by | |
| | a) | - 4 Dq | b) | +6 Dq | |
| | c) | Both | d) | None of these | |
| 52. | In t | etrahedral complexes eg orbital stabilised | by | | |
| | a) | -6 Dq | b) | + 4 Dq | |
| | c) | 10 Dq | d) | None of these | |
| 53. | In t | etrahedral complexes $t_{2\sigma}$ orbital destabiliz | ed by | <i>J</i> | |
| | a) | -6 Dq | b) | +4 Dq | |
| | c) | 10 Dq | d) | None of these | |
| 54. | "Aı | ny non linear moleculer system in a deger | nerat | e electronic state will be unstable and will | |
| | unc stat | lergo some sort of distortion to lower it tement of | s syr | netry and remove the degeneracy." is the | |
| | a) | Trans effect | b) | Bhol's effect | |
| | c) | John-Teller effect | d) | None of these | |
| 55. | The | e complex ion show the highest crystal fiel | d spl | litting energy is | |
| | a) | $\left[\mathrm{CO}(\mathrm{NH}_3)_6\right]^{2+}$ | b) | $\left[\text{CO}(\text{NH}_3)_6 \right]^{3+}$ | |
| | c) | $\left[Rh(NH_3)_6 \right]^{3+}$ | d) | $\left[\operatorname{Ir}\left(\mathrm{NH}_{3}\right)_{6}\right]^{3+}$ | |
| 56. | Co | lour of complex is explained by | | | |
| | a) | Werner's theory | b) | VBT | |
| | c) | CFT | d) | Ligand field theory | |
| 57. | Wh | ich of the following ligand having largest | cryst | al field splitting? | |
| | a) | F- | b) | H ₂ O | |
| | c) | NH ₃ | d) | CN ⁻ | |

| 58. | Wł | nich of the following has maximum numbe | rofu | unpaired electrons? |
|-----|----|---|--------|---|
| | a) | d ⁶ (Tetrahedral) | b) | d ⁹ (octahedral) |
| | c) | d ⁷ (octahedral, High spin) | d) | d ⁴ (octahderal low spin) |
| 59. | Wł | nich of the following has smallest crystal f | ield s | plitting? |
| | a) | $\left[\text{CO}(\text{NH}_3)_6 \right]^{2+}$ | b) | $\left[\text{CO}(\text{NH}_3)_6 \right]^{3+}$ |
| | c) | $\left[Rh(NH_3)_6 \right]^{3+}$ | d) | $\left[\operatorname{Ir}\left(\mathrm{NH}_{3}\right)_{6}\right]^{3+}$ |
| 60. | Th | e number of unpaired electrons, in $NiCl_4^{2-}$ | (tet | rahedral) is / are |
| | a) | 2 | b) | 0 |
| | c) | 1 | d) | 4 |
| 61. | Wł | nich of the following ligands having maxin | num | crystal field splitting? |
| | a) | NH ₃ | b) | F [−] |
| | c) | Oxalate ^{2–} | d) | CO |
| 62. | Th | e number of unpaired electron in a d ⁷ tetra | ahedr | al configuration is |
| | a) | 3 | b) | 2 |
| | c) | 1 | d) | 7 |
| 63. | Wł | nich of the following has no CFSE in octal | hedra | l field? |
| | a) | Fe ³⁺ (High spin) | b) | CO ³⁺ (Low spin) |
| | c) | Fe ³⁺ (Low spin) | d) | Cr ³⁺ (High spin) |
| 64. | Wł | nich of the following is correct? | | |
| | a) | Hydrated $CuSO_4$ is colourless | | |
| | b) | Anhydrated CuSO ₄ in solid form is w | hite | |
| | c) | Hydrated CuSO_4 in solid form is white | | |

d) $CuSO_4$ solution is yellow

Metal Cluster

| 1. | Hydride of boron is commonly known as | • | |
|-----|---|--------------------|--------------------------------|
| | a) Boron | b) | Boranes |
| | c) Borazine | d) | All above |
| 2. | The boranes which baring the general form | ıla B _n | H _{n+4} is said to be |
| | a) Nido boranes | b) | Closo boranes |
| | c) Arachnoboranes | d) | All these |
| 3. | The boranes which having the general form | ula B _r | H_{n+6} is said to be |
| | a) Nido boranes | b) | Closo boranes |
| | c) Arachnoboranes | d) | All these |
| 4. | The borones which having the formula B_nH | n^{-2} is | said to be |
| | a) Nido boranes | b) | Closo boranes |
| | c) Arachnoboranes | d) | All these |
| 5. | B ₂ H ₆ | | |
| | a) Diborane (6) | b) | Diborane (2) |
| | c) Diborane (8) | d) | none of these |
| 6. | $B_4 H_{10}$ | | |
| | a) Tetraborane (10) | b) | Tetraborane (14) |
| | c) Tetraborane (14) | d) | Tetraborane (0) |
| 7. | $B_{10}H_{14}$ | | |
| | a) Decarbone (14) | b) | Decaborane (10) |
| | c) Decarbonae (24) | d) | Decaborone |
| 8. | Diborane is prepared by | | |
| | a) $BCl_3 + H_2$ | b) | $BCl_3 + LiAlH_4$ |
| | c) BCl ₃ + NaH | d) | All these |
| 9. | Diborane is | | |
| | a) rolatile gas | b) | nonvolatile gas |
| | c) liquid | d) | solid |
| 10. | Diborane on heating at 100-250°C, gires | • | |
| | a) Boron | b) | Hydrogen |
| | c) Boron & Hydrogen | d) | Mixture of hydrides |
| 11. | Diborane on hydrolysis, gires | | |
| | a) mixture of boron | b) | Boric acid |
| | c) a and b | d) | none |

| 12. | Diborane is | | |
|-----|---|--------|--------------------------------------|
| | a) Electron rich compound | b) | Electron deficient compound |
| | c) both a and b | d) | None of these |
| 13. | Diborane show | | |
| | a) Hydrogen bridge structure | b) | cage like structure |
| | c) puckted 7ng structure | d) | All these |
| 14. | In diborane bond length between boron and | bridg | ing hydrogenation is |
| | a) 1.33 A° | b) | 1.19 A° |
| | c) 1.77 A° | d) | 1.00 A° |
| 15. | In diborane bona length between boron and | termi | nal hydrogen atom is |
| | a) 1.33 A° | b) | 1.19 A° |
| | c) 1.77 A° | d) | 1.00 A° |
| 16. | In diborane bond length between two boron | atom | n is |
| | a) 1.33 A° | b) | 1.19 A° |
| | c) 1.77 A° | d) | 1.00 A° |
| 17. | In diborane the bond angle between one bon | on at | om and two bridging hydrogen atom is |
| | a) 95° | b) | 120° |
| | c) 180° | d) | 45° |
| 18. | In diborane the bond angle between one bon | on an | d two terminal hydrogen atom is |
| | a) 95° | b) | 120 ° |
| | c) 180° | d) | 45° |
| 19. | Experimental evidences of hydrogen bndge | struct | ure of diborones is / are |
| | a) NMR sturcutre | b) | Methylation |
| | c) Specific heat | d) | All these |
| 20. | Three-centred electron pair bond is observe | d in | |
| | a) diborane | b) | borazine |
| | c) silicates | d) | fullerenes |
| 21. | How much three centied electron pair bond | prese | ent in diboranes |
| | a) Two | b) | Three |
| | c) Four | d) | One |
| 22. | Banana type bond observed in | | |
| | a) diborane | b) | borazene |
| | c) silicone | d) | silicates |
| 23. | In diborane each boron atom undergoes | | |
| | a) Sphybridisation | b) | Sp ² hybridisation |
| | c) Sp ³ hybridisation | d) | dsp ² hybridisation |

| 24. | B-] | H-B bridge in B_2H_6 is formed by the over | laping | g of | | | | |
|-----|--|---|--------|-----------------------------|--|--|--|--|
| | a) | two centred one electrons | b) | three centred two electrons | | | | |
| | c) | four centred one electrons | d) | two centred two electrons | | | | |
| 25. | Mi ske | Mixed hydride of carbon and boron having both carbon and boron atoms in an electron deficient skeletal framework known as | | | | | | |
| | a) | boranes | b) | carboranes | | | | |
| | c) | hydrocarborane | d) | none of these | | | | |
| 26. | Wł | Which of the following is the type of carboranes | | | | | | |
| | a) | Closo | b) | Nido | | | | |
| | c) | Arachno | d) | All these | | | | |
| 27. | Clo | oso means | | | | | | |
| | a) | closed structure | b) | nest like structure | | | | |
| | c) | open structure | d) | none | | | | |
| 28. | Nic | do means | | | | | | |
| | a) | Nest like | b) | Cup like | | | | |
| | c) | open | d) | All these | | | | |
| 29. | Ca | rboranes are derived from | | | | | | |
| | a) | hydrocarbon | b) | beranes | | | | |
| | c) | both a and b | d) | none of these | | | | |
| 30. | Th | The formula of dicarbacloso dodecacaborane is | | | | | | |
| | a) | $C_2B_{10}H_{12}$ | b) | $C_2B_{12}H_{10}$ | | | | |
| | c) | $C_2B_2H_6$ | d) | $C_2B_8H_{12}$ | | | | |
| 31. | Th | The structure of dicarbaclosododecaborane is | | | | | | |
| | a) | icosahedral | b) | Tetrahedral | | | | |
| | c) | Square planar | d) | Octahedral | | | | |
| 32. | $C_2 B_{10} H_{12}$ occure in isomeric form | | | | | | | |
| | a) | Two | b) | Three | | | | |
| | c) | Four | d) | Five | | | | |
| 33. | Isometric form of $C_2 B_{10} H_{12}$ are | | | | | | | |
| | a) | Ortho | b) | Para | | | | |
| | c) | Meta | d) | All these | | | | |
| 34. | In meta $C_2 B_{10} H_{12}$ carbon occuples the position | | | | | | | |
| | a) | 1, 2 | b) | 1, 7 | | | | |
| | c) | 1, 12 | d) | None of the above | | | | |
| 35. | In para $C_2 B_{10} H_{12}$ carbon occupies the position | | | | | | | |
| | a) | 1, 2 | b) | 1, 7 | | | | |
| | c) | 1, 12 | d) | None of the above | | | | |

| 36. | Ortho $C_2B_{10}H_{12}$ is conrected into meta $C_2B_{10}H_{12}$ at | | | | |
|-----|---|----------------------------|------------|-----------------------|--|
| | a) 100°C | | b) | 470°C | |
| | c) 700°C | | d) | 1000° C | |
| 37. | Meta $C_2B_{10}H_{12}$ is converted into para $C_2B_{10}H_{12}a^+$ | | | | |
| | a) 100°C | | b) | 470°C | |
| | c) 700°C | | d) | 1000° C | |
| 38. | (m + 1) electron | pairs are brosent in the | structure | of | |
| | a) closo carbo | rane | b) | Nido carborane | |
| | c) Arachno car | borane | d) | none of these | |
| 39. | (m + 2) electron | pairs are present in the | structure | of | |
| | a) closo carbor | ane | b) | Nido carborane | |
| | c) Arachno car | borane | d) | none of these | |
| 40. | (m + 3) electron | pairs are present in the | structure | of | |
| | a) closo carbor | ane | b) | Nido carborane | |
| | c) Arachno ca | rborane | d) | none of these | |
| 41. | $C_2 B_{10} H_{12}$ is isoe | lectronic with | | | |
| | a) B ₁₂ H ₁₂ ²⁻ | | b) | $B_{12}H_{12}$ | |
| | c) $B_{12}H_{10}^{-2}$ | | d) | $B_{12}H_{10}$ | |
| 42. | $C_2 B_{10} H_{12}$ & their | r isomers are | | | |
| | a) White cryst | talline solid | b) | White amorphous solid | |
| | c) colourless lie | quid | d) | colourless gas | |
| 43. | $C_2B_{10}H_{12}$ resist t | to attack | | | |
| | a) heat | | b) | chemical | |
| | c) both a and | b | d) | none of these | |
| 44. | $C_2 B_{10} H_{12}$ resistte | o attack | | | |
| | a) oxidising ag | ent | b) | reducing agent | |
| | c) both a and | b | d) | none of these | |
| 45. | $C_2B_{10}H_{12}$ involv | ed in reaction | | | |
| | a) Lithiation | | b) | bromination | |
| | c) chlorination | | d) | iodination | |
| 46. | $C_2 B_{10} H_{12}$ involv | ed in the preparation of. | ••••• | | |
| | a) Griganard | derivate | b) | Amide deriradire | |
| | c) halogen deri | ivation | d) | Acid derivative | |
| 47. | Which of the fol | lowing is the types of the | e boranes | | |
| | a) Nido | | b) | Arachno | |
| | c) Closo | | d) | All of the above | |

| 48. | ST | YX number of diborane is | | | | | | |
|-----|---|---|-------|----------------------------------|--|--|--|--|
| | a) | 2002 | b) | 4012 | | | | |
| | c) | 4020 | d) | 3203 | | | | |
| 49. | Dil | Diborane, on treatment with aqueous KOH gives | | | | | | |
| | a) | orthoborate | b) | Metaborate | | | | |
| | c) | Both | d) | None | | | | |
| 50. | 0. Diborane, on action with ammonia at low temp gives | | | | | | | |
| | a) | Diammoniate of diborade | b) | Boron nitride | | | | |
| | c) | Borazine | d) | Carborane | | | | |
| 51. | Dil | borane, on action with ammonia at high te | mp. g | gives | | | | |
| | a) | Diammoniat of diborade | b) | Boron nitride | | | | |
| | c) | Borozine | d) | Borazole | | | | |
| 52. | Dil | borane, on treatment with ammonia, at 20 | 0°C ያ | gives | | | | |
| | a) | Diammoniate of diborane | b) | Boron nitrite | | | | |
| | c) | Borozine | d) | None | | | | |
| 53. | Me | etaloborane are | | | | | | |
| | a) | Polyhedral borane anion | b) | Polyhedral borane cation | | | | |
| | c) | Both a and b | d) | none of the above | | | | |
| 54. | Me | etaloborane having the general formula | • | | | | | |
| | a) | $B_n H_{n+4}$ | b) | B_nH_{n+6} | | | | |
| | c) | $B_n H_n^{-2}$ | d) | None of the above | | | | |
| 55. | Metaloborane is | | | | | | | |
| | a) Stable, aromatic, electron deficient compound | | | | | | | |
| | b) | o) Unstable, aromatic, electron deficient compound | | | | | | |
| | c) |) Stable, aliphatic, electron deficient compound | | | | | | |
| | d) | d) Unstable, aliphatic, electron deficient compount | | | | | | |
| 56. | Me | Metaloborane is prepared by | | | | | | |
| | a) | Action of decaborane with triethylamine | | | | | | |
| | b) | b) Action of sodium boron hydride with diborane | | | | | | |
| | c) | Action of diborane with triethyl amine | | | | | | |
| | d) | All above | | | | | | |
| 57. | B ₁₂ | $_{2}$ H ₁₂ ²⁻ show | | | | | | |
| | a) | Tetrabedral structure | b) | Octahedral distorted | | | | |
| | c) | Icosahedral structure | d) | Prism structure | | | | |
| 58. | B ₁₂ | ${}_{2}H_{12}^{2-}$ having | | | | | | |
| | a) | Two centred electron pair bond | b) | Three centred electron pair bond | | | | |
| | c) | Both a and b | d) | None of these | | | | |
| | | | | | | | | |

59. The number of electron in cage or icosahedral structure of metaloborane can be calculated by..... a) Wade's rule b) EAN rules c) Hunds rule Thumbs rule d) Borazine react with excess bromine and gives 60. Tribromobenzene a) Benzene b) c) both None of these d) Borazine react with water and gives 61. a) Boric acid b) diborane c) Benzene d) All these 62. On pyrolysis borazine gives a) diphenyl like product Napthalene like protect b) c) both d) None of these Planar hexagonal ring structure is shown by 63. a) Borazine b) Methanol d) none of these c) both In boranzine the bond length between boron and nitrogen is 64. a) 0.144 nm b) 0.100 nm c) 0.200 nm 0.215 nm d) 65. In borazine boron and nitrogen atom undergoes b) SP² hybridisation a) SP hybridisation c) SP³ hybridisation d) dsp² hybridisation According to Wade's rule the number of electrons present in framework of $B_{12}H_{12}^{2-}$ are.... 66. a) 24 b) 25 c) 26 27 d) 67. The compound of carborane with some metal ion called..... a) Metalocarborane Metalobarane b) c) Carborane None of the above d) The trival name of $C_2 B_9 H_{11}^{2-}$ is..... 68. Dicabollide a) Carbollide b) None fo the above c) Carbinide d) 69. Dicarbollide word derived from spanish word..... a) Olla b) Bolla c) Colla None of these d) Which of the following compound form sandwith compound..... 70. a) Borane b) Carborane d) Metallocarborane c) Metalloborane

Bio-Inorganic Chemistry

| 1. | Which of the following elements are bulk element | | | | | |
|-----|--|---|----|----------------------------|--|--|
| | a) | С | b) | Н | | |
| | c) | 0 | d) | All of these | | |
| 2. | Which of the following are trace elements | | | | | |
| | a) | Fe | b) | Cu | | |
| | c) | Zn | d) | All of these | | |
| 3. | Wł | nich of the following are ultratrace elemen | ts | | | |
| | a) | F | b) | Se | | |
| | c) | Si | d) | All of these | | |
| 4. | Na | ⁺ and K ⁺ are | | | | |
| | a) | Charge carrier | b) | Maintaine osmotic pressure | | |
| | c) | Maintain sensityvity of nerve cells | d) | All of these | | |
| 5. | •••• | essential codetituent of chlorophyll. | | | | |
| | a) | Mg | b) | Na | | |
| | c) | Mn | d) | All | | |
| 6. | Calcium is important for | | | | | |
| | a) | Bone | b) | Blood cloating | | |
| | c) | Rhythm of heart beat | d) | All of these | | |
| 7. | Essential constituent of haemoglobin is | | | | | |
| | a) | Mg | b) | Fe | | |
| | c) | Ca | d) | F | | |
| 8. | Wi | lson disease is caused by | | | | |
| | a) | Exess of Fe | b) | Excess of Cr | | |
| | c) | Excess of Cu | d) | Excess of Au | | |
| 9. | Co | balt is essential constituent of | | | | |
| | a) | Vitamin A | b) | Vitamin C | | |
| | c) | Vitamin K | d) | Vitamin B ₁₂ | | |
| 10. | Which of the following synthesize the insulin in pancres | | | | | |
| | a) | Co ⁺⁺ | b) | Zn ⁺⁺ | | |
| | c) | Ni ⁺⁺ | d) | K^+ | | |
| 11. | Which of the following causes the toxicity loss of hairs | | | | | |
| | a) | Se | b) | Te | | |
| | c) | Ро | d) | K | | |

| 12. Which of the following has glucose tolerance factor | | or | | |
|---|------|--|-------|-------------------------------------|
| | a) | Cu ⁺⁺ | b) | CO ⁺⁺⁺ |
| | c) | er*** | d) | Ca ⁺⁺ |
| 13. | itai | -itai disease caused by the toxicity of | | |
| | a) | Cd++ | b) | Ag^+ |
| | c) | Au ⁺⁺ | d) | Pt ⁺⁺ |
| 14. | Mi | namata disease caused by the toxicity of | ••• | |
| | a) | Hg ⁺⁺ | b) | Cl⁻ |
| | c) | Br- | d) | I [_] |
| 15. | De | ficiency of causes Goiter. | | |
| | a) | F^- | b) | Cl⁻ |
| | c) | Br [_] | d) | I⁻ |
| 16. | •••• | maintain the balance between inside a | and c | outside the fluid of all |
| | a) | Na ⁺ / K ⁺ pump | b) | Ni ⁺⁺ / Co ⁺⁺ |
| | c) | Cr+++ / Fe++ | d) | Sn / Al |
| 17. | •••• | essential for coagulation of blood as | nd ry | thum of heart heat |
| | a) | Mg | b) | Ca |
| | c) | Sr | d) | Bg |
| 18. | In | photosynthesis reaction play an imp | porta | nt role |
| | a) | Mg | b) | Ca |
| | c) | Br | d) | Ba |
| 19. | Po | rphin molecule is a ligand of ring | syste | em |
| | a) | Tetrapyrol | b) | Tetra furon |
| | c) | Tetra thiophine | d) | None of these |
| 20. | Th | e porphyrin ring with metal ion called | | |
| | a) | Carborane | b) | Metals carborane |
| | c) | Metaloborane | d) | metaloporphyrine |
| 21. | Me | etaloporphyrin of iron is / are | | |
| | a) | Chlorophyll | b) | Vitamin B ₁₂ |
| | c) | Haemoglobin & myoglobin | d) | none of these |
| 22. | Me | etaloporphirin of magnesium is | | |
| | a) | Chlorophyll | b) | Myoglobin |
| | c) | Cobalamine | d) | Haemoglobin |
| 23. | Me | etaloporphyrin of iron is | | |
| | a) | Haemoglobin | b) | Myoglobin |
| | c) | Cytochrome | d) | |

| 24. | Me | etaloporphitio of cobalt is | | | | | | | |
|------------------|--|--|-------------|--|--|--|--|--|--|
| | a) | Haemogiobin | b) | Myoglobin | | | | | |
| | c) | Cytochrome | d) | Cobalamine or B ₁₂ vitamin | | | | | |
| 25. | Haemoglobin transport & to the different | | | | | | | | |
| | a) | O ₂ & CO ₂ | b) | $O_2 \& NH_3$ | | | | | |
| | c) | $CO_2 \& SO_2$ | d) | None of these | | | | | |
| 26. | In | In heamoglobin heme groups are attached on one globin molecule | | | | | | | |
| | a) | 01 | b) | 02 | | | | | |
| | c) | 03 | d) | 04 | | | | | |
| 27. | Haemoglobin when attached to oxygen is called | | | | | | | | |
| | a) | Oxyhaemoglobin | b) | Deoxy haemoglobin | | | | | |
| | c) | Both | d) | None of these | | | | | |
| 28. | The binding of oxygen by haemoglobin is depends on | | | | | | | | |
| | a) | pH of blood | b) | Colour of blood | | | | | |
| | c) | Smell of blood | d) | Complex nature of blood | | | | | |
| 29. | At | low pH and high concentration of CO_2 of | bloo | d the affinity for oxygen is | | | | | |
| | a) | increases | b) | decreases | | | | | |
| • | c) | Cant say | d) | None of these | | | | | |
| 30. | The | e effect of pH & con ¹¹ of CO_2 on binding and | d rele | casing of oxygen by haemoglobin is called | | | | | |
| | a) | Bohr's effect | b) | Common ion effect | | | | | |
| 01 | c) | Steric effect | d) | None of these | | | | | |
| 31. | In o | deoxyhaemoglobin Fe ⁺⁺ is | 1.) | H , 1 , 6 , 6 , 7 , 7 , 7 , 7 , 7 , 7 , 7 , 7 | | | | | |
| | a) | low spin & parammagnetic | D) | High spin & paramagnetic | | | | | |
| 20 | C) | low spin & dimagnetic | a) | High spin & paramagnetic | | | | | |
| 32. | In (| Lowanin & dimognatic | h) | Louisin & nonomegnetic | | | | | |
| | a) | Poth a % h | d) | Lowspin & paramagnetic | | | | | |
| 22 | с) Мъ | | u) | None of these | | | | | |
| 55. | NIY D) | Mussolo nigmont | b) | ostalvat | | | | | |
| | a) | Roth a & b | d) | None of these | | | | | |
| 3/ | U) In 1 | myoglobin baeme group attached | u) on gl | obin molecule | | | | | |
| J 4 . | a) 111 1 | | b) | | | | | | |
| | a) | 03 | d) | 04 | | | | | |
| 35 | C) Mv | voglobin store in mussele | u) | | | | | | |
| | a) | 0 . | b) | CO. | | | | | |
| | с) | \sim_2 Both a and b | d) | None of these | | | | | |
| | 2) | | 4) | | | | | | |

| 36. | ••••• | actas oxygen reserviour within the musscle | | | |
|-----|-------|---|---------|---|--|
| | a) | Haemoglobin | b) | Mycglobin | |
| | c) | Cytocbrome | d) | Cobalamine | |
| 37. | Mo | elecular weight of Haemoglobin is | | | |
| | a) | 10,000 | b) | 20,000 | |
| | c) | 64,000 | d) | 100,000 | |
| 38. | Mo | elecular weight of myoglobin is | | | |
| | a) | 5,000 | b) | 10,000 | |
| | c) | 15,000 | d) | 17,000 | |
| 39. | | molecule consist of only one polyper | otide | chain of protein and one haeme group | |
| | a) | Mroglobin | b) | Haemoglobin | |
| | c) | Cytochrome | d) | none of these | |
| 40. | Atr | nospheric nitrogen is | | | |
| | a) | Very active | b) | inert | |
| | c) | Both these | d) | None of these | |
| 41. | Co | nversion of atmospheric nitrogen into usef | ùl nit | trogeneous empound by natural or artificial | |
| | me | thod called | | | |
| | a) | Nitration | b) | Nitrogenfixation | |
| | c) | Both | d) | None of these | |
| 42. | Wh | hich of the following way for the fixation of | of niti | rogen is used | |
| | a) | Lightening | b) | Blue green algae & bacteria | |
| | c) | Both a & b | d) | None of these | |
| 43. | The | e enzyme that is responsible for nitrogen f | ixatic | on in microorganism is | |
| | a) | Nitrogenase | b) | Cytochrome-c | |
| | c) | Both a & b | d) | None of these | |
| 44. | Nit | rogenase is made up of with protein | • | | |
| | a) | one | b) | two | |
| | c) | three | d) | four | |
| 45. | Ree | duction of molecular nitrogen occur at site | e of | | |
| | a) | Fe | b) | S | |
| | c) | Мо | d) | None of these | |
| 46. | Но | w much electrons required for conversion | of n | itrogen to ammonia by nitrogenase are | |
| | a) | 2 | b) | 4 | |
| | c) | 6 | d) | | |
| 47. | Sm | aller brotein of nitrogenase contain one | ••• | | |
| | a) | Fe ₄ S ₄ | b) | Fe ₂ S ₂ | |
| | c) | FeS | d) | Fe ₃ S ₃ | |

- Larger protein or nitrogenase contain...... 48.
 - a) 2 MO, 30 Fe, 30S b) 1 MO, 15 Fe, 15S
 - c) 2 MO, 50Fe, 50S 2 MO, 100Fe, 1000S d)

Nitrogenase convert atmospheric nitrogen to..... 49.

- a) Nitrate b) Ammonia
- c) Both
- d) None of these