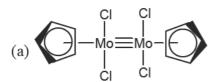
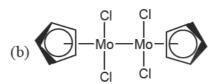
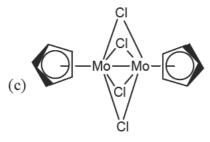
CHEMISTRY-CY

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

[CpMoCl₂]₂ obeys the 18 electron rule. The correct structure of this compound is (atomic number of Mo 1. =42)







- During oxygen transport by hemerythrin, oxygen is bound as 2.
 - (a) O₂ to one Fe(III) only
- (b) HO, to one Fe(III) only
- (c) O_2^{2-} to one Fe(II) and one Fe(III)
- (d) O₂²⁻ to two Fe(II)
- Among the following, the most stable isotope to radioactive decay is 3.
 - (a) $^{206}_{82}$ Pb
- (b) $^{210}_{92}$ Pb
- (c) 212 Pb
- At pH 7.2 and 10 Torr oxygen partial pressure, the extent of O₂ binding is 4.
 - (a) high for both haemoglobin and myoglobin
 - (b) high for haemoglobin and low for myoglobin
 - (c) high for myoglobin and low for haemoglobin
 - (d) low for both haemoglobin and myoglobin
- 5. In the first row high-spin transition metal complexes [M(H₂O)₆]Cl₂ with d⁵ and d⁷ metal ions, the d-d transitions are
 - (a) spin-forbidden for both
 - (b) spin-allowed for both
 - (c) spin-forbidden for d⁵ and spin-allowed for d⁷
 - (d) spin-allowed for d⁵ and spin-forbidden for d⁷
- Among the given boranes and heteroboranes, the example which belongs to 'closo' type is 6.
 - (a) B_5H_8
- (b) $[C_2B_0H_{11}]^{2-}$
- (c) $GeC_2B_0H_{11}$
- (d) B_6H_{10}
- The reaction of P₂O₅ with HNO₃ and HClO₄, respectively, gives 7.
 - (a) NO, and ClO,

(b) N_2O_5 and Cl_2O_6 (d) N_2O_5 and Cl_2O_7

(c) N₂O₃ and Cl₂O₇

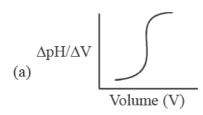
- 8. When crystals of sodium chloride are heated in the presence of sodium vapor, they turn yellow. This is due to the formation of
 - (a) Schottky defects

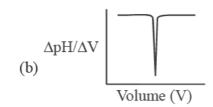
(b) Frenkel defects

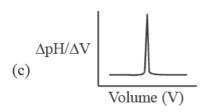
(c) F-centres

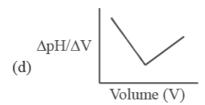
- (d) H-centres
- One mole of an ideal gas is compressed from 5L to 2L at constant temperature. The change in entropy, 9. in J K^{-1} , of the gas is . (R = 8.314 J K^{-1} mol⁻¹)

- The linear momentum of a particle described by the wavefunction e^{-ikx} is 10.
 - (a) kh
- (b) -kh
- (c) kħ
- (d) $-k\hbar$
- For an elementary bimolecular gas phase reaction, activation energy is 5.5 kJ mol⁻¹. Enthalpy of activa-11. tion, in kJ mol⁻¹, at 300K is $(R = 8.314 \text{ J K}^{-1} \text{ mol}^{-1})$
- 12. The titration of a strong acid with a strong base is represented by the plot



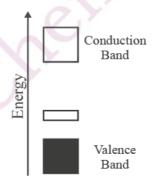






- 13. Of the following inequalities, the criteria for spontaneity of a chemical reaction is/are
 - (i) $(\Delta G)_{\tau p} < 0$
- (ii) $(\Delta U)_{S,V} > 0$ (iii) $(\Delta S)_{U,V} > 0$ (b) (ii) only (c) (i) and (ii)
- (a) (i) only

- (d) (i) and (iii)
- A protein sample consists of an equimolar mixture of ribonuclease (molar mass = 13.7 kg mol⁻¹), hemo-14. globin (molar mass = 15.5 kg mol⁻¹), and myoglobin (molar mass = 17.2 kg mol⁻¹). The statement that is true about the weight-average molar mass (\overline{M}_w) , the number-average molar mass (\overline{M}_n) , and the polydispersity index (PDI) for this sample is
 - (a) $\overline{M}_w > \overline{M}_n = 15.5~kg~mol^{-1}$ and PDI > 1 (b) $\overline{M}_w > \overline{M}_n = 15.5~kg~mol^{-1}$ and PDI < 1
 - (c) $\overline{M}_{w} = 15.5 \text{ kg mol}^{-1} > \overline{M}_{n} \text{ and } PDI > 1$ (d) $\overline{M}_{w} = 15.5 \text{ kg mol}^{-1} < \overline{M}_{n} \text{ and } PDI < 1$
- 15. The band structure given below represent a



- (a) n-type semiconductor formed by doping Si with B
- (b) *n*-type semiconductor formed by doping Si with P
- (c) p-type semiconductor formed by doping Si with P
- (d) p-type semiconductor formed by doping Si with B
- 16. The experimental ionization energies of hydrogen and helium atoms in their ground states are, respectively, 13.6 eV and 24.6 eV. The ground state energy of helium atom, in eV, is
 - (a) $-\frac{1}{2}(13.6) 24.6$ (b) -4(13.6) 24.6 (c) $-\frac{1}{4}(13.6) 24.6$ (d) -2(13.6) 24.6

17. Ring flipping of the compound in the following conformation leads to

$$(a) \bigcup_{\mathsf{Br}}^{\mathsf{Br}} \mathsf{CH}_3 \qquad (b) \bigcup_{\mathsf{Br}}^{\mathsf{CH}_3} \mathsf{CH}_3 \qquad (c) \bigcup_{\mathsf{OH}}^{\mathsf{CH}_3} \mathsf{CH}_3 \qquad (d) \bigcup_{\mathsf{H}_3\mathsf{C}}^{\mathsf{OH}} \mathsf{Br}$$

18. The total number of lines expected (due to spin-spin coupling of proton with fluorine and deuterium nuclei) in the ¹H NMR spectrum of the following compound is

19. The compound in 'R' configuration is

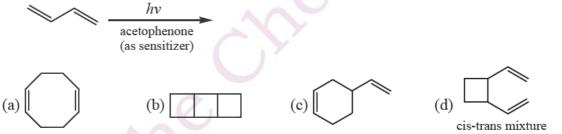
[GATE 2016]

(b)
$$H_3C$$
 H C_2H_5 H

20. The most suitable reagent for performing the following transformation, is

(a)
$$\text{LiAlH}_4$$
 (b) H_2 , Pd/C (c) PPh_3 , H_2O (d) Li , Liq . NH_3

21. The major product obtained in the following reaction, is



22. The Favourable transition state leading to the formation of the product in the following reaction, is

Chx = cyclohexyl

23. The major product of the following reaction is,

(a)
$$CH_3$$
 (b) CH_3 (c) CH_3 (d) CH_3 (d) CH_3

24. The major product obtained in the following reaction, is

H-C
$$\equiv$$
C-CH₂OH $\frac{\text{(i) LiNH}_2 \text{ (2equiv.)/liq. NH}_3}{\text{(ii) C}_2\text{H}_5\text{Br}\text{(1 equiv.)}}$

(ii) H₃O⁺

(a)
$$H_3CH_2C-C\equiv C-CH_2OH$$

(c)
$$H_3CH_2C-C\equiv C-CH_2NH_2$$

(d)
$$H-C \equiv C-CH_2NH-CH_2CH_3$$

25. The major product formed in the following reaction, is

$$N(C_2H_5)_2 \frac{(i) \text{ CH}_3\text{I}}{(ii) \text{ NaHCO}_3/\text{MeOH, warm}}$$

$$(a) \xrightarrow{H_3C} \bigvee N(C_2H_5)_2$$

$$(b) \qquad \qquad N(C_2H_5)_2$$

$$(d) \begin{array}{|c|c|} \hline \\ \hline \\ C_2H_5 \\ \hline \end{array}$$

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

- The Larmor frequency of ¹H at 1 Tesla (T) is 42.57 MHz. If the magnetogyric ratios for ¹H and ¹³C are 26.75 26. × 10⁷ rad T⁻¹ s⁻¹ and 6.72×10⁷ rad T⁻¹ s⁻¹, respectively, the Larmor frequency of ¹³C, in MHz, at 1 Tesla will
- At 1 bar and 298K, for the process $A(s) \rightarrow A(\ell)$, the ΔG is 200 J mol⁻¹ and ΔV_m is -2×10^{-6} m³ mol⁻¹. 27. The minimum pressure, in bar, at which the process becomes spontaneous at 298K is ______. (1 bar $= 10^5 \, \text{Pa}$).
- The reaction, A \B, is first order in both the directions. The forward and reverse rate constants are 28. 4.2×10^{-4} s⁻¹ and 1.04×10^{-3} s⁻¹, respectively. The relaxation time for this reaction, in seconds, in a temperature jump experiment is
- Adsorption of CO on charcoal at 273K follows Langmuir isotherm. A plot of P(kPa)/V(cm3) versus P 29. (kPa) is linear with a slope of 0.01 y-intercept of 0.5. The equilibrium constant, K (kPa⁻¹), for the adsorption is
- 30. For the following reaction,

$$A+B \xrightarrow[k_{-1}=10^5 \text{ L mol}^{-1}s^{-1}]{k_{-1}=10^4 s^{-1}} I \xrightarrow{k_2=10 s^{-1}} P$$

if steady state approximation can be applied on [I], the observed rate constant of product formation, in L mol⁻¹ s⁻¹, will be

The correct set of infra-red spectral bands (in cm⁻¹) for the ν_{CO} stretching mode of the given carbonyl 31. complex is



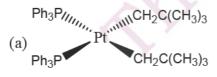
- (a) 1827, 1783, 1766
- (c) 1833, 1775, 1650

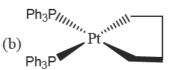
- (b) 1973, 1827, 1794
- (d) 1960, 1918
- 32. The ¹⁹F NMR spectrum of ClF₃ when measured at -60°C will be observed as a
 - (a) singlet

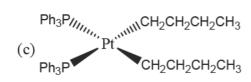
(b) doublet

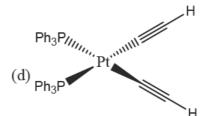
(c) doublet and triplet

- (d) doublet of doublet and a doublet of triplet
- Among the given platinum(II) complexes, the one that is thermally the most unstable is 33.









- 34. The shapes of XeF₅ and XeF₅, respectively, are
 - (a) pentagonal planar and square pyramidal
 - (b) pentagonal planar and trigonal bipyramidal
 - (c) square pyramidal and pentagonal bipyramidal
 - (d) square pyramidal and pentagonal planar
- 35. Sodium salt of pseudohalogens, X, Y and Z form colorless solutions in water. Solution of X decolorizes I_3^- solution with brisk effervescence. Solution of Y gives an intense red colour on reaction with Fe³⁺ solution. Solution of Z gives an intese blue color on reaction with a solution containing Fe³⁺ and Fe²⁺ ions. The pseudohalogens X, Y and Z respectivley are
 - (a) CN^-, N_3^- and CNS^-

(b) N_3^- , CNS⁻ and CN⁻

(c) N_3^- , CN^- , and CNS^-

- (d) N₃, CNS⁻, and CNO⁻
- 36. On reacting 1.55g of a diol with an excess of methylmagnesium iodide, 1.12L (corrected to STP) of methane gas is liberated. The molecular mass (g mol⁻) of the diol is
- 37. The structure of the compound having the following characteristics spectral data, is IR: 1690 cm^{-1} . $^{1}\text{H-NMR}: 1.30 \text{ (3H, t, J} = 7.2 \text{ Hz)}; 2.41 \text{ (2H, q, J} = 7.2 \text{ Hz)}; 2.32 \text{ (3H, s)}; 7.44 \text{ (1H, t, J} = 7.0 \text{ Hz)}; 7.57 \text{ (1H, dt, J} = 7.0, 3.0 \text{ Hz)}; 7.77 \text{ (1H, t, J} = 3.0 \text{ Hz)}; 7.90 \text{ (1H, dt, J} = 7.0, 3.0 \text{ Hz)}; EI mass: m/z 119 (100%); 57 (80%)$

$$(a) \begin{picture}(60,0) \put(0,0){\line(1,0){100}} \put(0,0){\line(1,0)$$

38. The major products X and Y formed in the following formed in the following synthetic scheme, are

$$(i) \bigvee_{\substack{N \\ \text{benzene (solvent)} \\ (ii) \bigvee_{\substack{O}} \text{ (X)}} (X) \xrightarrow{\text{(H}_3C)_2S \longrightarrow \text{CH}_2} (Y)$$

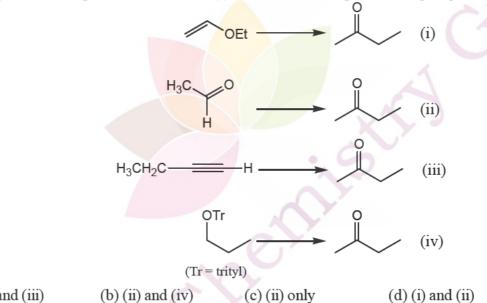
(iv) aq. NaOH, heat

This disaccharide is

39. The major product S and T formed in the following synthetic schemem, are

$$(a) S = (b) S = (c) S = (c)$$

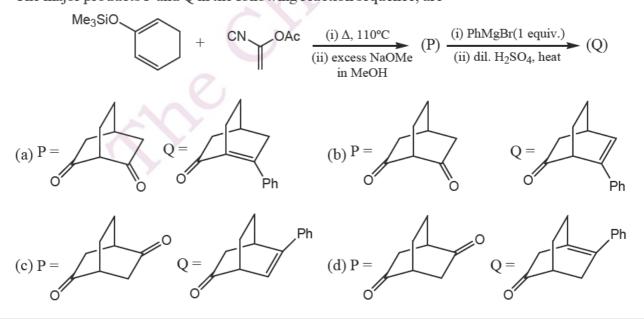
40. Among the following, the transformation(s) that can be accomplished using umpolung concept is(are)



(a) (i) and (iii)
(b) (ii) and (iv)
(c) (ii) only
(d) (i) and (ii)
A disaccharide does NOT give a positive test for Tollen's reagent. Upon acidic hydrolysis, it gives an equimolar mixture of two different monosaccharides, both of which can be oxidized by bromine water.

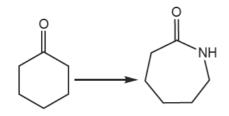
42. The major products M and N in the following reaction sequence are

43. The major products P and Q in the following reaction sequence, are



44. The major product formed in the following reaction, is

45. The following synthetic transformation can be achieved using

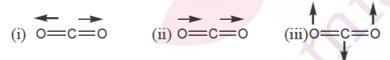


Reagents:

- (p) (i) NH₂OH/H⁺, (ii) H₂SO₄
- $(q) NH_3/H^+$
- (r) (i) NH₂OH/H⁺, (ii) NaOH
- (a) p only
- (b) p and q
- (c) q and r
- (d) r only

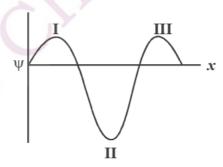
46. Consider a two-state system at thermal equilibrium with equal degeneracy where the excited state is higher in energy than the ground state by 0.1 eV. The ratio of the population of the excited state to that of the ground state, at a temperature for which $k_B T = 0.05 \text{ eV}$, is _____

47. Of the vibrational modes given below, the IR active mode(s) is(are)



- (a) (ii) only
- (b) (iii) only
- (c) (i) and (ii)
- (d) (ii) and (iii)

48. A system is described by the following real wavefunction.



The probability (P) of finding the particle in a region dx around points I, II and III in the figure obeys the trend

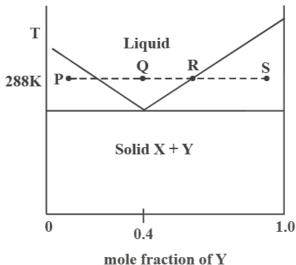
(a) P(I) > P(II) > P(III)

(b) P(II) > P(III) > P(I)

(c) P(II) > P(I) > P(III)

(d) P(III) > P(I) > P(II)

49. The temperature-composition (T-x) phase diagram of the two-component system made of X and Y is given below. At a temperature of 288K and starting at the point P, Y is added until the composition reaches S. Which of the following statements is NOTR TRUE?



- (a) At P, the solid and liquid are present in almost equal proportions
- (b) At Q, the system is all liquid
- (c) At S, the system has more solid than liquid
- (d) At R, the liquid is pure X
- 50. For a system subjected to only P-V work, entropy is given by
- $(II) \left(\frac{\partial G}{\partial P} \right)_T \qquad (III) \left(\frac{\partial A}{\partial V} \right)_T$

- (a) I and II
- (b) I and IV

- 51. According to Irving-Williams series, the number of d electrons for the first row transition metal (M) ion having the highest overall stability cosntant $(\log \beta)$ for $[M(EDTA)]^{2-}$ is
- The magnitude of the difference in the crystal field stabilization energies, in Δ_0 (irnoring pairing energy), of 52. $[Fe(H_2O)_6]^{2+}$ and $[Fe(CN)_6]^{4-}$ is
- The calculated and observed magnetic moments differ considerably for an aqua complex of a Lanthanide 53. (III) ion as a result of low lying states of high J. The ion, among the follwing, is
 - (a) Ce^{3+}

- In the electronic spectra of $\left[\text{CrF}_{6}\right]^{3-}$, absorption bands observed at 670, 440 and 290 nm are, respectively, 54. due to the transitions.
 - (a) ${}^{4}A_{2\sigma} \rightarrow {}^{4}T_{1\sigma}(P)$, ${}^{4}A_{2\sigma} \rightarrow {}^{4}T_{1\sigma}(F)$ and ${}^{4}A_{2\sigma} \rightarrow {}^{4}T_{2\sigma}$
 - (b) ${}^{4}A_{2g} \rightarrow {}^{4}T_{1g}(P)$, ${}^{4}A_{2g} \rightarrow {}^{4}T_{2g}$ and ${}^{4}A_{2g} \rightarrow {}^{4}T_{1g}(F)$
 - (c) ${}^4A_{2g} \rightarrow {}^4T_{1g}(F)$, ${}^4A_{2g} \rightarrow {}^4T_{1g}(P)$ and ${}^4A_{2g} \rightarrow {}^4T_{2g}$
 - (d) ${}^4A_{2g} \rightarrow {}^4T_{2g}$, ${}^4A_{2g} \rightarrow {}^4T_{1g}(F)$ and ${}^4A_{2g} \rightarrow {}^4T_{1g}(P)$
- Amongst the following, the group that is bound to the metal ion in coenzyme B₁₂ is 55.
 - (a) methyl
- (b) cyanide
- (c) adenosyl
- (d) hydroxyl

***** END OF THE QUESTION PAPER *****