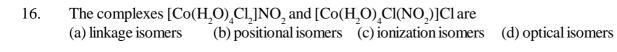
CHEMISTRY-CY

	Q.1 – Q.25 : Carry ONE mark each.				
1.			can perform at constant I		
	(a) Δ <i>H</i>	(b) ΔG	(c) ΔS	(d) ΔA	
2.	Consider the reaction:				
	$A + B \rightleftharpoons C$				
	The unit of the therr (a) mol L ⁻¹	nodynamic equilibrium c (b) L mol ⁻¹	constant for the reaction i (c) $mol^2 L^{-2}$	s (d) dimensionless	
3.	The number of IR active vibrational normal modes of CO ₂ is				
4.	The number of C_2 axes in CCl_4 is				
5.	The value of the magnetic quantum number of a p _x orbital is				
	(a) -1	(b) 0	(c) + 1	(d) undefined.	
6.	The molecular parti	The molecular partition function for a system in which the energy levels are equispaced by ϵ , is			
	(a) $\frac{1}{1+e^{\beta \epsilon}}$	(b) $\frac{1}{1-e^{\beta\epsilon}}$	(c) $\frac{1}{1+e^{-\beta \varepsilon}}$	$(d) \frac{1}{1 - e^{-\beta \varepsilon}}$	
7.	A monoatomic gas, X, adsobred on a surface, Langmuir adsorption isotherm. A plot of the fraction of surface				
	coverage, θ against the concentration of the gas [X], for very low concentration of the gas, is described by the				
	equation			[]	
	(a) $\theta = K[X]$	$(b) 1 - \theta = \frac{1}{K[X]}$	(c) $\theta = K^{1/2} [X]^{1/2}$	$(d) \theta = \frac{K[X]}{1 - K[X]}$	
8.	At a given temperature and pressure, the ratio of the average speed of hydrogen gas to that of helium gas is approximately				
9.	An example of nido-borane from the following is				
	(a) $B_4 H_{10}$	(b) $B_6 H_{10}$	(c) $B_6 H_{12}$	$(d) B_8 H_{14}$	
10.	The geometries of Ni(CO) ₄ and [NiCl ₄] ²⁻ , respectively, are (a) tetrahedral and square planar (b) square planar and tetrahedral (c) tetrahedral and tetrahedral				
11.	The number of S–S	The number of S–S bonds in $H_2S_5O_6$ is			
12.	In atomic absorption spectroscopy, the atomization process utilizes (a) flame (b) electric field (c) magnetic field (d) electron beam				
13.	At room temperature, the number of singlet resonances observed in the ¹ H NMR spectrum of Me ₃ CC(O)NMe (N N-dimethyl pivalamide) is				
14.	Amongst the following, the metal that does NOT form homoleptic polynuclear carbonyl is				



The reaction of $[Cp_{2}TaMe_{2}]I(Cp = C_{5}H_{5}^{-})$ with NaOMe yields.

(b) Fe

15.

(a) $[Cp_{2}Ta(OMe)_{2}]I^{2}$

(c) $Cp_2^T Ta(Me) = CH_2$



(b) [Cp,Ta(Me)OMe]I

(d) $Cp_2^T Ta(OMe) = CH_2$

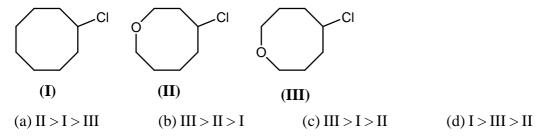
(d) Co

17. The major product of the following reaction is

18. Amongst the following, the structure of guanosine is

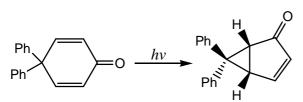
19. The correct order of IR stretching frequency of the C=C in the following olefins is

20. The correct order of the solvolysis for the following chlorides in acetic acid is

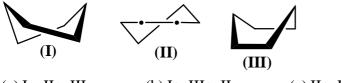




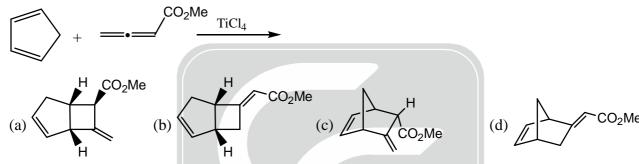
21. Formation of the product in the following photochemical reaction involves



- (a) di- π -methane rearrangement
- (b) Paterno-Buchi reaction
- (c) [2, 3]-sigmatropic rearrangement
- (d) Norrish type I reaction
- 22. The correct order of stability for the following conformations of cyclohexane is

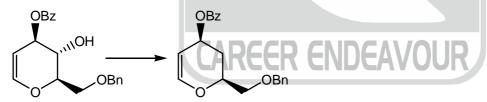


- (a) I > II > III
- (b) I > III > II
- (c) II > I > III
- (d) III > I > II
- 23. The major product formed in the following reaction is



24. The overall yield (in %) for the following reaction sequence is ____

25. The most suitable reagent combination to effect the following conversion is



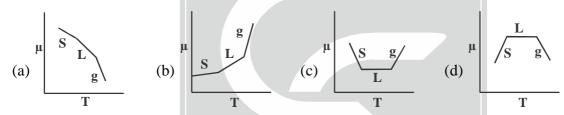
- (a) (i) NaH, CS₂, then MeI; (ii) Bu₃SnH, AlBN, C₆H₆, Reflux.
- (b) (i) I₂, PPh₂, imidazole; (ii) H₂, 10% Pd-C, AcOH, high pressure
- (c) (i) Me₂SiCl, pyridine, DMAP; (ii) Bu₂SnH, AlBN, C₆H₆, reflux
- (d) (i) MsCl, pyridine, DMAP; (ii) LiAlH, THF, reflux,

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

- 26. $\psi = N r (6-Z r) e^{-Zr/3} \cos \theta$, is a proposed hydrogenic wavefunction, where Z = atomic number, r = radial distance from the nucleus, θ = azimuthal angle, N is a constant. The **INCORRECT** statement about ψ is
 - (a) $\psi = 0$ in the xy-plane
 - (b) two radial nodes are presentt in ψ
 - (c) one angular node is present in ψ
 - (d) the size of the orbital decreases with increase in atomic number



- 27. The van der waals constant a and b of CO_2 are $3.64 L^2$ bar mol⁻² and 0.04 L mol⁻¹, respectively. The value of R is 0.083 bar dm³ mol⁻¹K⁻¹. If one mole of CO_2 is confined to a volume of 0.15L at 300K, then the pressure (in bar) exerted by the gas, is ______
- 28. A plot of osmotic pressure against concentration (gL^{-1}) of a polymer is constructed. The slope of the plot (a) increases with increase in temperature
 - (h) increases with increase in temperature
 - (b) increases with increase in molar mass of the polymer
 - (c) decreases with decrease in concentration of the polymer
 - (d) decreases with increase in temperature.
- 29. A platinum electrode is immersed in a solution containing 0.1 M Fe²⁺ and 0.1 M Fe³⁺. Its potential is found to be 0.77V against SHE. Under standard conditions and considering activity coefficients to be equal to unity, the potential of the electrode, when the concentration of Fe³⁺ is increased to 1 M, is ______
- 30. Molybdenum crystallizes in a bcc structure with unit cell dimensions of 0.314 nm. Considering the atomic mass of molybdenum to be 96, its density (in kg m⁻³) is ______
- 31. The ratio of molecules distributed between two states is 9.22×10⁶ at 300K. The difference in energy (in kJ mol⁻¹) of the two states is _____
- 32. A Carnot engine operates at 55% efficiency. If the temperature of reject steam is 105°C, then the absolute temperature of input steam is ______
- 33. Of the following plots, the correct representation of chemical potential (μ) against absolute temperature (T) for a pure substance is (S, L and g denote solid, liquid and gas phases, respectively)



- 34. The enthalpy of fusion of ice at 273K is 6.01 kJ mol⁻¹ and the enthalpy of vaporization of water at 273K is 44.83 kJ mol⁻¹. The enthalpy of sublimation (in kJ mol⁻¹) of ice at 273K, is ______
- 35. Suppose ψ_1 and ψ_2 , are two hybrid orbitals:

$$\psi_1 = 0.12 \, \psi_{3s} + 0.63 \psi_{3p_x} + 0.77 \psi_{3p_y} \text{ and } \psi_2 = 0.12 \, \psi_{3s} - 0.63 \psi_{3p_x} - 0.77 \psi_{3p_y}$$
The angle (in degrees) between there is

The angle (in degrees) between them is _____

- 36. BCl_3 and NH_4Cl were heated at $140^{\circ}C$ to give compound X, which when treated with $NaBH_4$ gave another compound Y. Compounds X and Y are
 - (a) $X = B_3 N_3 H_3 C l_3$ and $Y = B_3 N_3 H_6$
- (b) $X = B_3 N_3 H_9 C l_3$ and $Y = B_3 N_3 H_6$
- (c) $X = B_3N_3H_3Cl_3$ and $Y = B_3N_3H_{12}$
- (d) $X = B_3 N_3 C l_6$ and $Y = B_3 N_3 H_6$
- 37. The number of microstates in term ¹G is _____
- 38. The set of protons (underlined) in $CH_3CH_2CH_3CH_3$ that would exhibit different splitting patterns in high (500 MHz) and low (60 MHz) field 1H NMR, is
 - (a) CH₃CH₂CH₂OCH₃

(b) CH₃CH₂CH₂OCH₃

(c) CH₃CH₂CH₂OCH₃

- (d) CH₃CH₂CH₂OCH₃
- 39. Amongst the following, the complex ion that would show strong Jahn-Teller distortion is
 - (a) $[Cr(H_2O)_6]^+$
- (b) $[\mathrm{Ti}(\mathrm{H_2O})_6]^{3+}$
- (c) $[\mathrm{Co(H_2O)}_6]^{2+}$
- (d) $[Fe(H_2O)_6]^{2+}$
- 40. Amonst the following, the metal carbonyl species having the highest v_{CO} stretching frequency is
 - (a) $[Mn(CO)_{6}]^{+}$
- (b) $Cr(CO)_6$
- (c) $[V(CO)_{6}]^{-}$
- (d) [Fe(CO)₄]²⁻

- 41. The correct order of thermal stability for the given compounds is
 - (a) $TiMe_4 > Ti(CH_2CMe_3)_4 > TiEt_4$
- (b) $TiEt_4 > Ti(CH_2CMe_3)_4 > TiMe_4$
- (c) $TiMe_4 > TiEt_4 > Ti(CH_2CMe_3)_4$
- (d) $Ti(CH_2CMe_3)_4 > TiMe_4 > TiEt_4$
- 42. Amongst the following, the complex ion that is expected to the highest magnetic moment at room temperature

- (a) $[Ni(CN)_4]^{2-}$ (b) $[Fe(CN)_6]^{3-}$ (c) $[Cu(H_2O)_6]^{2+}$ (d) $[Co(CN)_6]^{3-}$
- 43. MnCr₂O₄ is
 - (a) normal spinel with total CFSE of –15.5 Dq
 - (b) inverse spinel with total CFSE of –15.5 Dq
 - (c) normal spinel with total CFSE of -24 Dq
 - (d) inverse spinel with total CFSE of -24 Dq
- 44. Mg²⁺ is preferred in photosynthesis by chlorphyll because
 - (a) it has strong spin-orbit coupling
- (b) it has weak spin-orbit coupling

(c) it is a heavy metal

- (d) it binds strongly with chlorophyll
- 45. In Monsanto acetic acid process shown below, the role of HI is

$$CH_3OH + CO \xrightarrow{Rh(\ell) \text{ catalyst/HI}} CH_3CO_2H$$

- (a) to convert CH₂OH to a stronger nucleophile (CH₂O⁻)
- (b) to reduce the Rh(I) catalyst to a Rh(0) species
- (c) to reduce a Rh(III) active species to a Rh(I) species in the catalytic cycle
- (d) to convert CH₂OH to CH₂I
- 46. Formation of the ketone H from the diazoketone I involves

- (a) generation of carbene and a [2, 3]-sigmatropic rearrangement
- (b) generation of carbene and an electrocyclic ring closing reaction
- (c) generation of ketene and a [2+2] cycloaddition
- (d) generation of ketene and a [3, 3] sigmatropic rearrangement
- 47. The major products X and Y formed in the following reaction sequence are

$$+ CO_2Me \xrightarrow{heat} X \xrightarrow{Raney Ni} Y$$

$$\text{(a) } X = \text{PhS} \qquad \text{CO}_2 \text{Me} \qquad Y = \text{PhS} \qquad \text{CO}_2 \text{Me}$$

$$(b) \ X = \begin{array}{c} \text{Me} \\ \text{PhS} \end{array} \begin{array}{c} \text{Me} \\ \text{PhS} \end{array} \begin{array}{c} \text{Ne} \\ \text{PhS} \end{array} \begin{array}{c} \text{OH} \\ \text{PhS} \end{array} \begin{array}{c} \text{Me} \\ \text{CO}_2 \text{Me} \end{array} \begin{array}{c} \text{Me} \\ \text{Y} = \\ \text{OH} \end{array} \begin{array}{c} \text{OH} \\ \text{OH} \end{array}$$

48. The major products X and Y formed in the following reactions are

49. The major product X and Y formed in the following reaction sequence are

$$cis-stilbene \xrightarrow{\quad h\nu \quad \quad } X \xrightarrow{\quad I_2 \quad \quad } Y$$

(a)
$$X = H$$

$$(d) X = \bigcup_{H \in H}$$

50. The product of the following reaction gave 6 line 13 C NMR spectrum with peaks at δ 175, 52, 50, 46, 37, 33 ppm. The structure of the product is

$$N_2$$
 N_2 N_2 N_2

51. The major product formed in the following reaction is

$$(c) \bigcup_{\mathsf{Ph}} \mathsf{Pr}$$

$$(d)$$
 Ph Ph

52. The major products X and Y formed in the following reaction sequence are



(a)
$$X = \bigcup_{NO_2}$$

$$Y = \bigcup_{NH_2}$$

$$(c) \ \ X =$$

$$(d) \ X = \bigcup_{i=1}^{NO_2} OH$$

$$Y =$$
 $\begin{array}{c} \text{NH}_2 \\ \text{OH} \end{array}$

53. The major products X and Y formed in the following reaction sequence are

$$\begin{array}{c|c} \hline \text{MeOH} & X & \hline \\ \hline \text{H}_2\text{SO}_4 & X & \hline \\ \hline \text{aq. NaHCO}_3 & Y \\ \hline \end{array}$$

$$\text{(a)} X = \text{NH}_2$$

$$Y = \begin{pmatrix} CO_2H \\ HN \\ Cbz \end{pmatrix}$$

$$Y = \begin{array}{c} \text{CO}_2\text{Me} \\ \text{HN} \\ \text{Cbz} \end{array}$$

$$\text{(c)} \ X = \text{HO} \qquad \text{NH}_2$$

$$Y = \begin{array}{c} \text{CO}_2\text{Me} \\ \text{NH}_2 \\ \text{CO}_2\text{Me} \end{array}$$

$$Y = \begin{array}{c} \text{CO}_2\text{Me} \\ \text{NHMe} \end{array}$$



54. Given the fact that 1, 3-butadiene has a UV absorption of 217nm, the absorption wavelength (in nm) for the conjugated system shown below is ______

(Use these absorption values for auxochromic groups:

alkyl: +5; exo-cyclic double bond: +5; every additional conjugated C = C : +30)

55. The m/z value of the detectable fragment formed by McLafferty like rearrangement of the following compound in mass spectrometer is ______



