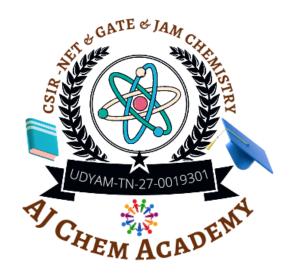
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Q.1 - Q.25 MCQ & NAT, carry ONE mark each (for each wrong

	answer: - 1/3). (** No Negative Marks for NAT)							
1.	Consider N particles at temperature-T, pressure-P, volume-V and chemical							
	potential- μ having energy-E. The parameters that are kept constant for a canonical							
	ensemble are							
	(a) N, V, T	(b) N, V, E	(c) N, P, T	(d) μ , V, T				
2.	For ortho-hydroge	n, the nuclear wavefur	nction and the rotational	l quantum number,				
	respectively, are							
	(a) antisymmetric ar	nd even	(b) symmetric and	d odd				
	(c) symmetric and e	ven	(d) antisymmetric	and odd				
3.	m_1 and m_2 are the slopes (dP/dT) of the solid-liquid equilibrium lines in the P-T							
	phase diagrams of H_2O and CO_2 , respectively. For $P < 10$ atm, the values of m_1							
	and m ₂ are							
	(a) $m_1 > 0$ and $m_2 > 0$ (b) $m_1 > 0$ and $m_2 < 0$							
	(c) $m_1 < 0$ and m_2	< 0	(d) $m_1 < 0$ a	$nd m_2 > 0$				
4.	The rate constant of a reaction is $1.25 \times 10^{-4} \text{ mol L}^{-1}\text{s}^{-1}$. If the initial							
	concentration of the reactant is 0.250 mol L ⁻¹ , the total time (in seconds) required							
	for complete conve	ersion is DYAM TN-27	7-0019301					
5.								
	of the gas is S, the partial derivative $(\partial P/\partial S)_V$ is equal to							
	(a) $(\partial T/\partial P)_S$	(b) $(\partial T/\partial V)_P$	(c) $-(\partial T/\partial V)_S$	(d) $(\partial T/\partial S)_P$				
6.	The wavelength as	sociated with a particle	e in one-dimensional box	of length L is				
			(n refers to the	e quantum number)				
	(a) 2L/n	(b) L/n	(c) nL	(d) L/2n				
7.	The dependence of	rate constant-k on ten	mperature-T (in K) of a	reaction is given by				
	the expression, $\ln k = [(-5000 \text{ K})/T] + 10$. The activation energy of the reaction							
	(in kJ mol ⁻¹) is	(up to two dec	cimal places)					
8.	The lowest energy of a quantum mechanical one-dimensional simple harmonic							
	oscillator is 300 cm ⁻¹ . The energy (in cm ⁻¹) of the next higher level is							
9.	The electronic grou	and state term for the o	chromium ion in [Cr(CN	$[)_6]^{4-}$ is				
	(a) ${}^{3}F$	(b) $^{3}\mathrm{H}$	(c) 3G	(d) 5D				
10.	The VO_4^{3-} , CrO_4^{2-}	and MnO ₄ ions exhib	it intense ligand to me	tal charge transfer				
	transition. The wavelengths of this transition follow the order							







(a)
$$CrO_4^{2-} < VO_4^{3-} < MnO_4^{-}$$
 (b) $MnO_4^{-} < VO_4^{3-} < CrO_4^{2-}$ (c) $VO_4^{3-} < CrO_4^{2-} < MnO_4^{-}$ (d) $CrO_4^{2-} < MnO_4^{-} < VO_4^{3-}$

(b)
$$MnO_4^- < VO_4^{3-} < CrO_4^{2-}$$

(c)
$$VO_4^{3-} < CrO_4^{2-} < MnO_4^{-}$$

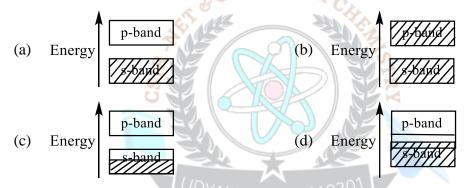
(d)
$$CrO_4^{2-} < MnO_4^{-} < VO_4^{3-}$$

11. The lanthanide ion that exhibits color in aqueous solution is

- (d) Lu(III)
- 12. The hapticity of cycloheptatriene, (C_7H_8) , in $Mo(C_7H_8)(CO)_3$ is _____
- 13. The v_{0-0} resonance Raman stretching frequency (in cm⁻¹) of the O_2 coordinated to iron centre in oxyhemoglobin is nearly

14. The energy band diagram for magnesium is

(The hatched and unhatched regions in the figure correspond to filled and unfilled regions of the band, respectively)



15. P, F and I represent primitive, face-centered and body-centered lattices, respectively. The lattice types of NaCl and CsCl, respectively, are

- (d) P and I
- 16. The characteristic feature of an electron spin resonance (ESR) spectrum of frozen aqueous solution of CuSO₄. 5H₂O at 77 K is

(a)
$$g_{\parallel} > g_{\perp}$$

(b)
$$g_{\parallel} < g_{\perp}$$

(c)
$$g_{\parallel} = g_{\perp}$$

(d)
$$g_x \neq g_y \neq g_z$$

17. The most suitable reagent for the following transformation is

(b)
$$PtO_2/H_2$$
 (c) LiAlH₄ (d) B₂

(a) Li/Liq. NH₃

- (d) B_2H_6
- 18. The major products M and N formed in the following reactions are







(a)
$$\mathbf{M} = \begin{pmatrix} \mathbf{H}_3 \mathbf{C} \\ \mathbf{N} \end{pmatrix} OH$$
 ; $\mathbf{N} = \begin{pmatrix} \mathbf{N} \\ \mathbf{N} \end{pmatrix} OCH_3$

(c)
$$\mathbf{M} = \bigvee_{\substack{\mathsf{CH}_3}} \mathsf{O}$$
 ; $\mathbf{N} = \bigvee_{\mathsf{N}} \mathsf{OCH}_3$

(d)
$$\mathbf{M} = \begin{pmatrix} \mathbf{N} \\ \mathbf{C} \\ \mathbf{H}_3 \end{pmatrix}$$
 ; $\mathbf{N} = \begin{pmatrix} \mathbf{H}_3 \\ \mathbf{N} \\ \mathbf{N} \end{pmatrix}$ OH

19. The ¹³C-NMR spectrum of acetone-d⁶ has a signal at 30 ppm as a septet in the intensity ratio

(a) 1:6:15:20:15:6:1

(b) 1:3:6:7:6:3:1

(c) 1:2:3:5:3:2:1

(d) 1:3:7:10:7:3:1

20. The major product formed in the following reaction is

21. The major product obtained in the following reaction is





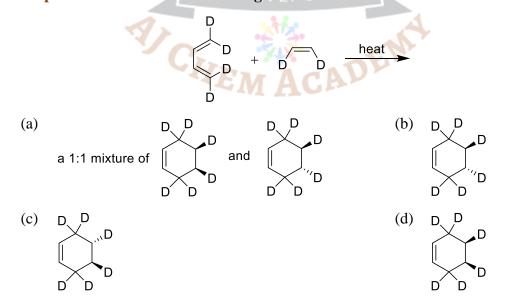


22. In the two step reaction sequence given below, the starting bis-sulfone acts as

- (a) a dienophile and synthetic equivalent of acetylene
- (b) a dienophile and synthetic equivalent of ethylene
- (c) a dipolarophile and synthetic equivalent of acetylene
- (d) a dipolarophile and synthetic equivalent of ethylene

23. The major product formed in the following photochemical reaction is

24. The product formed in the following reaction is



25. The number of possible stereoisomers for cyclononene is _____

Q.26 - Q.55 MCQ & NAT, carry TWO marks each (for each wrong answer: - 2/3). (** No Negative Marks for NAT)

26. The mobility of a univalent ion in aqueous solution is 6.00×10^{-8} m² s⁻¹ V⁻¹ at





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300 K. Its diffusion coefficient at 300 K is $X \times 10^{-9}$ m² s⁻¹. The value of X is _____ (up to two decimal places)

27. For the following consecutive first order reactions

$$X \xrightarrow{k_1 = 2.0 \text{ s}^{-1}} Y \xrightarrow{k_2 = 0.1 \text{ s}^{-1}} Z$$

the time (in seconds) required for Y to reach its maximum concentration (assuming only X is present at time t = 0) is _____ (up to two decimal places)

- 28. Under physiological conditions, the conversion of CO_2 to bicarbonate ion by carbonic anhydrase enzyme (MW = 30,000 g mol⁻¹) has a turnover number of 4.00×10^5 s⁻¹. The minimum amount of enzyme (in μ g) required to convert 0.44 g of CO_2 to bicarbonate ions in 100 seconds is _____ (up to two decimal places)
- 29. Assume 1,3,5-hexatriene to be a linear molecule and model the π electrons as particles in a one-dimensional box of length 0.70 nm. The wavelength (in nm) corresponding to the transition from the ground-state to the first excited-state is ____
- 30. The standard Gibbs free energy change of the reaction shown below is -2.7 kJ mol^{-1} . $Sn_{(s)} + Pb^{2+} \rightleftharpoons Sn^{2+} + Pb_{(s)}$ $\{E^0(Pb^{2+}/Pb) = -0.126 \text{ V}\}$ the value of $E^0(Sn^{2+}/Sn)$ in V is _____ (up to two decimal places)
- 31. The dissociative chemisorption of $X_{2(g)}$ on a metal surface follows langmuir adsorption isotherm. The ratio of the rate constants of the adsorption and desorption processes is 4.0 atm^{-1} . The fractional surface coverage of X (adsorbed) at 1.0 atm pressure is _____ (up to two decimal places)
- 32. The ionic activity coefficients of Ca^{2+} and F^{-} are 0.72 and 0.28, respectively. The mean activity coefficient of CaF_2 is _____ (up to two decimal places)
- 33. The angle of orientation (in degrees) of the angular momentum vector with respect to z-axis for $\ell = 2$ and $m_{\ell} = +2$ state of H-atom is _____(up to two decimal places).
- 34. The Gibbs free energy of mixing is denoted as ΔG_{mix} . 1.0 mole of He, 3.0 moles of Ne and 2.0 moles of Ar are mixed at the same pressure and temperature. Assuming ideal gas behavior, the value of $\Delta G_{mix}/RT$ is ______. (up to two decimal places)
- 35. $\Psi = \left[c\varphi_1 \left(\frac{1}{\sqrt{3}}\right)\varphi_2\right]$ represents a normalized molecular orbital constructed from two different atomic orbitals φ_1 and φ_2 that form an orthonormal set. The value of |c| is ______. (up to two decimal places)
- 36. In cyclophosphazenes, $(NPX_2)_3$ (X = F, Cl, Br and Me), the strength of P–N π -bond varies with X in the order







(a)
$$F > Cl > Br > Me$$

(b) Me > F > Cl

(c)
$$Br > Cl > F > Me$$

(d) Me > Br > Cl >

- 37. The structure type and shape of the polyhedral (skeletal) framework of the carborane $Me_2C_2B_{10}H_{10}$, respectively, are
 - (a) nido and dodecahedron

(b) closo and icosahedron

(c) nido and icosahedron

(d) closo and dodecahedron

38. If Δ_0 is the octahedral splitting energy and P is the electron pairing energy, then the crystal-field stabilization energy (CFSE) of $[\text{Co}(\text{NH}_3)_6]^{2+}$ is

(a)
$$-0.8\Delta_0 + 2P$$

(b)
$$-0.8\Delta_0 + 1P$$

$$(c) -0.8\Delta_0$$

(d)
$$-1.8\Delta_0 + 3P$$

The rates of substitution for the following reaction vary with L in the order **39.**

 $L = CH_3^-, Cl_-, Ph_and H_a$

(a)
$$CH_3^- > Cl^- > Ph^- > H^-$$
 (b) $Cl^- > Ph^- > H^- > CH_3^-$

(b)
$$Cl^- > Ph^- > H^- > CH_3^-$$

(c)
$$Ph^{-} > CH_{3}^{-} > H^{-} > Cl^{-}$$
 (d) $H^{-} > CH_{3}^{-} > Ph^{-} >$

(d)
$$H^- > CH_3^- > Ph^- > Cl^-$$

- 40. The product formed in the reaction of MeMn(CO)₅ with ¹³CO is
 - (a) $(Me^{13}CO)Mn(CO)_5$

(c)
$$(MeCO)Mn(CO)_4(^{13}CO)$$

- UDYAM-TN-27-0019 (d) (Me¹³CO)Mn(CO)₄(¹³CO)
- 41. For the following three alkenes I, II and III, the rates of hydrogenation using Wilkinson's catalyst at 25 °C vary in the order



II



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(a) I > III > II

III < II < I (d)

(c) II > I > III

- (d) II > III > I
- 42. 210 Bi undergoes β^- decay to $\frac{1}{8}$ of its initial amount in 15 days. The time required for its decay to $\frac{1}{4}$ of its initial amount is _____ days. (up to two decimal places)
- 43. The metal ion and the macrocyclic skeleton present in the green pigment of plants, respectively, are
 - (a) Mg(II) and chlorin

(b) Mg(II) and corrin

(c) Mn(II) and chlorin

- (d) Mg(II) and porphine
- 44. The spinel structure of MgAl₂O₄ has cubic close packed arrangement of oxide ions.

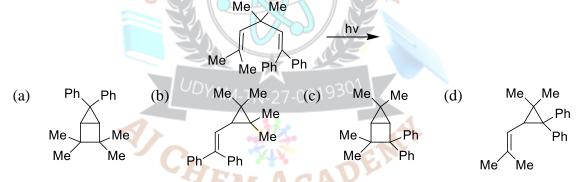
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The fractions of the octahedral and tetrahedral sites occupied by cations, respectively, are

- (a) $\frac{1}{8}$ and $\frac{1}{2}$

- (b) $\frac{1}{4}$ and $\frac{1}{2}$ (c) $\frac{1}{2}$ and $\frac{1}{4}$ (d) $\frac{1}{2}$ and $\frac{1}{8}$
- 45. The diffusion limiting current (I_d) at a dropping mercury electrode for an aqueous Mg(II) solution of concentration 'c' (mol L⁻¹) is 300 μA. If 'c' is increased by 0.1 mol L^{-1} , I_d increases to 900 μ A. The value of 'c' (in mol L^{-1}) is _____ (up to two decimal places)
- The major product formed in the following reaction is

47. The product formed in the following photochemical reaction is



48. Among the following decahydroquinoline toluenesulfonates (Ts), the one that yields 9-methylamino-E-non-5- enal as a major product upon aqueous solvolysis is

$$(a) \quad \mathsf{TSO} \\ \vdots \\ \mathsf{H} \\ \mathsf{CH}_3 \\ (b) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ (c) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ (d) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ \mathsf{CH}_3 \\ (d) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ \mathsf{CH}_3 \\ (d) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ \mathsf{CH}_3 \\ (d) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ \mathsf{CH}_3 \\ (d) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ \mathsf{CH}_3 \\ (d) \quad \mathsf{TSO} \\ \mathsf{H} \\ \mathsf{CH}_3 \\ \mathsf{$$

49. The product obtained in the following solvolysis reaction is

(a) a racemic mixture of trans 1,2-diacetoxycyclohexane





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(b) enantiomerically pure trans 1,2-diacetoxycyclohexane

(c) racemic cis 1,2-diacetoxycyclohexane

(d) a mixture of cis and trans 1,2-diacetoxycyclohexane

50. The spectroscopic data for an organic compound with molecular formula $C_{10}H_{12}O_2$ are given below, the compound is

IR band : $\approx 1750 \text{ cm}^{-1}$

¹H NMR : δ 7.3 (m, 5H), 5.85 (q, 1H, J = 7.2 Hz), 2.05 (s, 3H),

1.5 (d, 3H, J = 7.2 Hz) ppm

(a) methyl 2-phenylpropionate

(b) 1-(phenylethyl) acetate

(c) 2-(phenylethyl) acetate

(d) methyl 3-phenylpropionate

51. The structures of the intermediate [P] and major product Q formed in the following reaction sequence are

(a)
$$P = Me$$

$$Me$$

$$Me$$

$$Me$$

$$Me$$

(b)
$$P = Me$$
; $Q = Me$

(c)
$$P = Me$$
 ; $Q = Me$

(d)
$$P = Me$$
 ; $Q = Me$

52. Hydration of fumaric acid gives malic acid as shown below. Assume that addition of water takes place specifically from A face or B face. The correct statement





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pertaining to stereochemistry of malic acid formed is

- (a) addition specifically from A face gives S isomer of malic acid
- (b) addition specifically from B face gives S isomer of malic acid
- (c) addition specifically from A face gives R isomer of malic acid
- (d) addition specifically from B face gives a racemic mixture of malic acid
- 53. Hydroboration of 2-butyne with $(C_6H_{11})_2BH$ yields the intermediate U, which on treatment with I_2 and NaOMe at $-78\,^{\circ}$ C, gives product V. The structures of U and V are

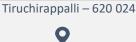
(a)
$$U = H_3C \xrightarrow{B(C_6H_{11})_2} ; V = H_3C \xrightarrow{CH_3} C_6H_{11}$$

(c)
$$U = H_3C \xrightarrow{B(C_6H_{11})_2} ; V = H_3C \xrightarrow{C_6H_{11}} CH_3$$

(d)
$$U = H_3C CH_3 ; V = H_3C C_6H_{11}$$

54. The structures of the major products W and X in the following synthetic scheme are

(a)
$$\mathbf{W} = \bigvee_{\mathbf{N}}^{\mathbf{OH}} ; \mathbf{X} = \bigvee_{\mathbf{N}}^{\mathbf{O}}$$







(b)
$$W = \bigvee_{N}^{OH} ; X = \bigvee_{N}^{O}$$

(c)
$$\mathbf{W} = \begin{pmatrix} 1 & 1 & 1 \\ 1 & 1 & 1 \\ 1 & 1 & 1 \end{pmatrix}$$
 ; $\mathbf{X} = \begin{pmatrix} 1 & 1 & 1 \\ 1 & 1 & 1 \\ 1 & 1 & 1 \end{pmatrix}$

(d)
$$\mathbf{W} = \begin{pmatrix} \mathbf{V} \\ \mathbf{N} \\ \mathbf{V} \end{pmatrix}$$
; $\mathbf{X} = \begin{pmatrix} \mathbf{V} \\ \mathbf{V} \\ \mathbf{V} \end{pmatrix}$

55. The major products Y and Z in the following reaction sequence are

(a)
$$Y = NCO$$
; $Z = NCO$

(b)
$$\mathbf{Y} = N_3$$

$$\mathbf{Z} = N_3$$

$$\mathbf{Z} = N_3$$

(d)
$$Y = \bigvee_{N_3}^{O} : Z = \bigvee_{Ph O}^{O} \bigvee_{N_3}^{O}$$

Answer Key

Q.No	Ans	Q.No	Ans	Q.No	Ans
1.	a	21.	b	41.	b
2.	b	22.	a	42.	9.90 to 10.10
3.	d	23.	b	43.	a
4.	1999 to 2001	24.	d	44.	d
5.	c	25.	3	45.	0.04 to 0.05
6.	a	26.	1.50 to 1.60	46.	c
7.	41.50 to 41.60	27.	1.56 to 1.60	47.	d







8.	899 to 901	28.	7.40 to 7.60	48.	c
9.	b	29.	225 to 240	49.	a
10.	c	30.	-0.15 to -0.13	50.	b
11.	b	31.	0.65 to 0.70	51.	b
12.	6	32.	0.36 to 0.40	52.	a
13.	a	33.	33.30 to 36.90	53.	d
14.	d	34.	-6.10 to -6.05	54.	d
15.	b	35.	0.80 to 0.84	55.	a
16.	a	36.	a		
17.	a	37.	b		
18.	c	38.	c		
19.	b	39.	d		
20.	d	40.	c		

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