June-2013 BOOKLET-C

Part-B

21. Which of the following pairs has the highest difference in their first ionization ene			st ionization energy?		
	(a) Xe, Cs	(b) Kr, Rb	(c) Ar, K	(d) Ne, Na	
22.	The ligand in uranoc	ene is:			
	(a) $C_8H_8^{2-}$	(b) $C_5H_5^{2-}$	(c) C_6H_6	(d) $C_4H_4^{2-}$	
23.	(a) lead to a decrease(b) change the forma(c) change the hybrid(d) increase with the	e in $C = C$ bond length l oxidation state of the lisation of the olefin ca presence of electron de	metal rbon from sp ² to sp ³ . onating substituent on		
24.	The oxidation state of	of molybdenum in $\lfloor (\eta^7) \rfloor$	$-tropylium \Big) Mo \Big(CO \Big)$	\int_3 is:	
	(a) +2	(b) +1	(c) 0	(d)-1	
25.	The reaction of [PtC	$\left[1_4\right]^{2-}$ with two equivale			
	(a) $\operatorname{cis} - \left[\operatorname{Pt} \left(\operatorname{NH}_{3} \right)_{2} \right]$	Cl_2	(b) trans $-[Pt(NH_3)]$	$_{2}$ Cl_{2}	
	(c) boths $\operatorname{cis} - \left[\operatorname{Pt}(\operatorname{NH}_3)_2 \operatorname{Cl}_2\right]$ and $\operatorname{trans} - \left[\operatorname{Pt}(\operatorname{NH}_3)_2 \operatorname{Cl}_2\right]$				
	(d) $\operatorname{cis} - \left[\operatorname{Pt} \left(\operatorname{NH}_{3} \right)_{2} \right]$	Cl_4			
26. The electronic transition responsible for the color of the transition metal ions is			metal ions is		
	(a) $d_{\pi} \rightarrow d_{\sigma}$	(b) $d_{\pi} \rightarrow d_{\sigma^*}$	$(c) d_{\pi} \rightarrow d_{\pi^*}$	$(d) d_{\sigma} \rightarrow d_{\pi^*}$	
27.	The number of metal	-metal bonds in [W ₂ (O	Ph) ₆] is:		
20	(a) 1	(b) 2	(c) 3	(d) 4	
28.	•		states arising from the		
			(c) $T_{1g} + T_{2g} + A_{2g}$		
29.			at for whipping creams		
30.	(a) N ₂ O Flame proof fabrices	(b) NO contain	(c) N_2O_3	(d) N_2O_5	
	(a) $H_2NC(O)NH_2.N$		(b) $H_2NC(S)NH_2.N$	a_2SO_4	
	(c) $H_2NC(O)NH_2.H$	I_3PO_4	(d) $H_2NC(S)NH_2.H$	$_{3}PO_{4}$	
31.	Among the compoun	ds A-D, those which h	ydrolyse easily are		
	(a) NCl ₃	(b) NF ₃	(c) BiCl ₃	(d) PCl ₃ .	
32.			the type I copper prot		
33.	(a) square planar The metal ions preser	(b) tetrahedral	(c) octahedral nitrogenase enzyme co-	(d) distorted tetrahedral	
<i>JJ</i> .	(a) Fe, Mo	(b) Fe, W	(c) Fe, Cu	(d) Fe, Ni	

34.	The reaction, $\lceil (CO) \rceil$	$_{5} \operatorname{Mn}(\operatorname{Me}) + \operatorname{CO} \rightarrow [$	$(CO)_5 Mn \{C(O)Me\}$	
	is an example for			
	(a) oxidative addition	n	(b) electrophilic subs	stitution
	(c) nucleophilic subs		(d) migratory insertion	
35.	•		ctahedral Ni(II) comple	exes is
	(a) One	(b) Two	(c) Three	(d) Zero
36.	For neutron activation	on analysis of an eleme	ent, the favourable char	acteristics of both the target and
	the product are from	the following		
	(A) high neutron cro	ss-section area of targe	et	
	(B) long half-life of t	the product		
		s-section area of target		
	(D) low half-life time	<u>-</u>		
		ristics from the above a		😾
	(a) A and B	(b) C and D	(c) B and C	(d) A and D
37.				is $1.0, 0.5, 0.33, 0.25 \text{ mol dm}^{-3}$ at
		= -	order of the reaction is	
	(a) two	(b) one	(c) zero	(d) three
38.				- a one dimensional box is 6 units
		e units, what is the diff	terence in energy levels	of $n = 3$ and $n = 2$ for the above
	system?	(1) 5		(I) 10
20	(a) 4	(b) 5	(c) 9	(d) 10
39.	The wave function \	of a certain system is	s the linear combination	1
		$\overline{1}$ $\overline{3}$		
$\Psi = \sqrt{\frac{1}{4}}\Psi_1 + \sqrt{\frac{3}{4}}\overline{\Psi}_2$				
	where ψ_1 and ψ_2 are energy eigen functions with eigen values (non-degenerate) E_1 and E_2 , respectively.			
			m energy will be obser	
	$\sqrt{3}$	3	1	$\sqrt{1}$
	(a) $\sqrt{\frac{3}{16}}$	(b) $\frac{3}{4}$	(c) $\frac{1}{4}$	(d) $\sqrt{\frac{1}{4}}$.
40.			atom with electronic of	· ·
-10.				
	(a) ${}^{2}S_{\frac{1}{2}}$	(b) ${}^{1}P_{0}$	(c) ${}^{1}S_{0}$	(d) $^{1}S_{1}$
41.	A molecules contains	s the following symme	try operations: E, 2C	, $2C_3$, C_2 , $3\sigma_d$, $3\sigma_v$. The number
		of the symmetry point		5. 2. u. v
	(a) 3, 12	(b) 5, 12	(c) 6, 12	(d) 6, 6
	(4) 3, 12	(6) 5, 12	(0) 0, 12	(4) 5, 5
42.	A triatomic molecule structure of the mole		vs two IR absorption li	nes and one IR-Raman line. The
			,B,	A
	(a) B - B - A	(b) $B - A - B$	(c) B A	$(d)_{R}$
12			_	
43.			icical g factor (g _N), th	e nuclear magneton $\left(\beta_{N}\right)$ and the
	magnetic field streng	· ·		
	(a) energy of transition	on from α to β state	(b) chemical shift	
	(c) spin-spin coupling	g constant	(d) magnetogyric rati	lo

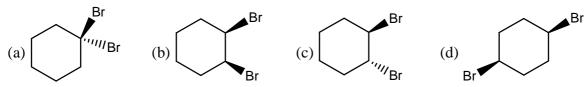
44.	An aqueous mixed solution of NaCl and I The number of components in the final m	•	lized by an aqueous NaOH solution		
	(a) 1 (b) 2	(c) 3	(d) 4		
45.	The lowest pressure at which the liquid pl	hase of a pure substa	nce can exist is known as		
	(a) critical point pressure	(b) super-incumb	_		
	(c) triple-point pressure		(d) saturation vapour pressure.		
46.	A chemical reaction involving nonlinear r	molecule + nonlinear	molecule in nonlinear activated		
	complex				
	The number of vibrational degrees of free				
477	(a) 3N-5 (b) 3N-6	(c) 3N-7	(d) 3N–8		
47.	Calculate the total number of microstates 2, 3} in three states is:	-	•		
	(a) 6 (b) 12	(c) 60	(d) 720		
48.	If the concentration (c) is increased to 4 till for strong electrolytes is (where b is Kohl	lrausch constant)			
	(a) 0 (b) $b\sqrt{c}$	(c) $2b\sqrt{c}$	(d) $4b\sqrt{c}$		
49.	In atom recombination reactions	(-)			
	(a) 0 (b) $b\sqrt{c}$ In atom recombination reactions (a) $E_a = 0$, $\Delta S^{\#} = +ve$, $\Delta H^{\#} = +ve$ (c) $E_a = +ve$, $\Delta S^{\#} = -ve$, $\Delta H^{\#} = -ve$	(b) $E_a = 0, \Delta S^{\#} =$	$=$ -ve, $\Delta H^{\#} = -ve$		
	(c) $E_a = +ve, \Delta S^{\#} = -ve, \Delta H^{\#} = -ve$	(d) $E_a = +ve$, ΔS	$^{\#}$ = +ve, Δ H $^{\#}$ = +ve		
50.	In the Lindemann mechanism of unimolec	cular reactions, the ob-	oserved order at low concentration is		
	(a) 0.5 (b) 1	(c) 1.5	(d) 2		
51.	The aggregation of surfactant molecules i				
	(a) micelles (b) clusters	(c) gel	(d) colloid		
52.	The coordinates for the atoms in a body of	The coordinates for the atoms in a body centred cubic unit cell are			
	(a) $(0, 0, 0)$ and $(1/2, 0, 0)$	(b) $(0, 0, 0)$ and $($	1/2, 1/2, 1/2)		
	(c) (0, 0, 0) and (0, 1/2, 0) The inter planar distance (Å) for a (100) pl	(d) $(0, 0, 0)$ and $($	0, 0, 1/2)		
53.					
E 1	(a) 1 (b) 2	(c) 4	(d) 8		
54.	The correlation coefficient of two parameters are	ters is found to be –0	.99. It may be concluded that the two		
	(a) strongly correlated				
	(c) connected by a cause-effect relationsh	nin (d) not connected	d by a cause-effect relationship		
55.	The IUPAC name for the compound gives	1			
	OH				
		, Ph			
		\checkmark			
	(a) (2R, 3Z)-7-phenylhept-3-en-2-ol	•	nenylhept-3-en-2-ol		
5 .	(c) (2R, 3E)-7-phenylhept-3-en-2-ol	- · · · · · - · · · ·	nenylhept-3-en-2-ol		
56.	Among the following esters, the one that	undergoes acid hydro	olysis fastest is		
	OCOCH- > OCOC	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	-		
	OCOCH ₃	,,,,,	OCOCH ₃		
	$(a) \parallel \qquad \qquad (b) \parallel \qquad \qquad $	(c)	(d) L		
	ОН	 OH	CH ₃		
57.	Rection of cyclohexyl benzyl ether with h		nce of 10% Pd/C vields		
	(a) cyclohexanol and toluene	•	and benzyl alcohol		
	(c) cyclohexane and benzyl alcohol	(d) cyclohexane a	•		

Jia Sarai,Near IIT Delhi-110016

Con: 9871547744

28A,1st Floor

58. Among the following dibromocyclohexanes, the one that reacts fastest with sodium iodide to give cyclohexene is



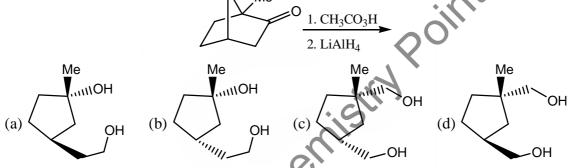
- 59. Match the following drugs with their medicinal activity
 - (A) 5-fluorouracil

(i) anti-bacterial

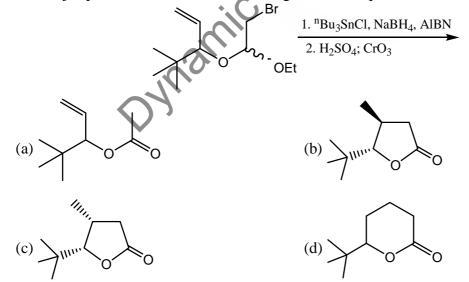
(B) amoxicillin lowering

- (ii) cholesterol
- (iii) anticancer
- (iv) anti-inflammatory

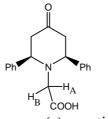
- (a) A-i, B-ii
- (b) A-iv, B-iii
- (c) A-iii, B-iv
- (d) A-iii, B-i
- 60. The major product formed in the following reaction sequence is



- 61. The biosynthetic precursor for the steroids is
 - (a) secologanin
- (b) shikimic acid
- (c) mevalonic acid
- (d) α-ketoglutaric acid
- 62. The major product formed in the following reaction sequence is

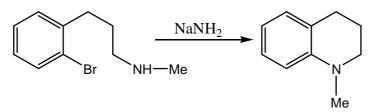


63. In the compound given below, the hydrogenes marked A and B are



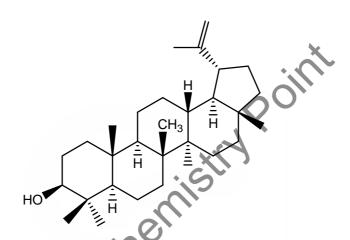
- (a) homotopic
- (b) isotopic
- (c) enantiotopic
- (d) diastereotopic

- 64. In the IR spectrum, the absorption band due to carbonyl group in phenyl acetate appears at
 - (a) 1800 cm⁻¹
- (b) 1760 cm⁻¹
- (c) 1710 cm⁻¹
- (d) 1660 cm⁻¹
- 65. The reactive intermediate involved in the following reaction is:

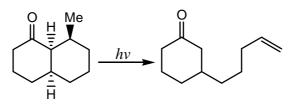


- (a) a carbocation
- (b) a carbanion
- (c) a free radical
- (d) an aryne

66. Number of isoprene units present in lupeol is



- (a) two
- (b) four
- (d) eight
- The heterocyclic ring present in the amino acid histidine is: 67.
 - (a) pyridine
- (b) tetrahydropyrrole (c) indole
- (d) imidazole
- The gauche conformation $(\varphi = 60^{\circ})$ of n-butane posseses 68.
 - (a) plane of symmetry; and is achiral
- (b) C₂-axis of symmetry; and is chiral
- (c) centre of symmetry; and is achiral
- (d) plane of symmetry; and is chiral
- The following photochemical conversion proceeds through 69.

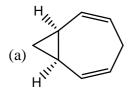


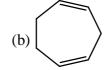
(a) Barton reaction

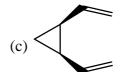
(b) Paterno-Büchi reaction

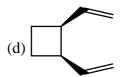
(c) Norrish type I reaction

- (d) Norrish type II reaction
- 70. Among the following dienes, the one that undergoes a degenerate Cope rearrangement is









PART-C

71.			ate of 34, 500 disintegraminutes. The $t_{1/2}$ for 41 A	ations/minute, but decay rate falls		
	(a) 90 minutes	(b) 110 minutes	(c) 180 minutes	(d) 220 minuts.		
72.	The orders of reactiv			plexes MeTiCl ₃ and (CO) ₅ Mo(thf)		
	are	, ,	3	3		
	(a) $CO > PMe_2 > NN$	Me_3 and $CO > NMe_3 >$	PMe ₂			
	(b) $PMe_2 > CO > NN$	Me_3^3 and $NMe_3 > CO >$	PMe,			
		CO^3 and $CO > PMe_3 > 1$				
		Me_3 and $PMe_3 > NMe_3$				
73.		pairs are identical in the				
			(c) XeO ₂ F ₂ , ICl ₄	(d) XeO ₄ , ClF ₂		
74.		g, those can act as Mö		4 3		
	(A) ^{129}I ,	(B) ⁵⁷ Co	(C) ⁵⁷ Fe	(D) ¹²¹ Sb		
	(a) A, B, C and D	` '	(c) A, B, and D only	(d) A. C and D only.		
75.	* / / /			n complexes, when ligands are		
	Cl- or OH-	,				
	(A) Be(II), Ba(II)	(B) Ba(II), Co(II)	(C) Co(II), Zn(II)	(D) Be (II) , $Zn(II)$		
	(a) A and B	(b) B and C	(c) C and D	(d) A and D		
76.	Silica gel contains [CoCl ₄] ²⁻ as an indicate		ca gel becomes dark blue while		
			anges to pale pink. This			
			rahedral to octahedral.			
	(b) Co(II) changes its	s oxidation state to Co	(III)			
	(c) Tetrahedral crysta	al field splitting is NO	requal to octahedral c	rystal field splitting.		
		(c) Co(II) forms kinetically labile while Co(III) forms kinetically inert complexes.				
77.		-	tement that is NOT TR	_		
	(a) there are two ion centres per active site.					
		(b) both iron centres are hexacoordinated in the active state.				
(c) one iron is hexacoordinated while the other is pentacoordinated in the active state.				d in the active state.		
	(d) it is found in marine invertebrates.					
78.	For a tetragonally dis	storted Cr(III) comple	x, zero-field splitting re	sults in the following number of		
	Kramers doublets:	\	, 1 0	8		
	(a) 1	(b) 2	(c) 3	(d) 4		
79.	• •	` /	` '	₂ Re ₂ Cl ₈ is due to the transition		
	(a) $\pi - \pi^*$		(c) $\delta - \pi^*$			
90						
80.		eduction of $Ce(SO_4)_2$,	$RMIO_4$, HNO_2 and I_2	with hydrazine in acidic medium,		
	respectively is	(b) 1 a 2 a 2 a and 4 a	(a) 2a 2a 1a and 4a	(d) 2a 4a 1a and 2a		
0.1			(c) 2e, 3e, 1e and 4e	(d) 2e, 4e, 1e and 3e		
81.		will behave as an acid	2 4	(4) 11 0		
0.2	(a) CH ₃ COOH	(b) HNO ₃	(c) HClO ₄	$(d) H_2O$		
82.				and(s) having N–N bond is/are		
02			(c) N_2O_3 and N_2O_4	(a) $N_2 O_5$ only		
83.	The treatment of PhBr with n-BuLi yields:					
	(a) $2 \text{ n-BuPh} + \text{Br}_2 +$	· L1 ₂	(b) PhPh + octane + 1	2 LiBr		
	(c) n -BuPh + LiBr		(d) $PhLi + n-BuBr$			

- 84. Though cyclobutadiene (C₄H₄) is highly unstable and readily polymerizes in its free state, its transition metal complexes could be isolated because
 - (a) it engages in long-range interaction with transition metals.
 - (b) it gains stability due to formation of $C_4H_4^{2-}$ on binding to transition metals.
 - (c) its polymerization ability reduces in presence of transition metal.
 - (d) it becomes stable in presence of transition metals due to formation of C₄H₄²⁺.
- 85. Identify the order representing increasing π -acidity of the following ligands

 C_2F_4 , NEt₃, CO and C_2H_4

- (a) $CO < C_2F_4 < C_2H_4 < NEt_3$
- (c) $C_2H_4 < NEt_2 < CO < C_2F_4$
- (b) C₂F₄ < C₂H₄ < NEt₃ < CO (d) NEt₂ < C₂H₄ < C₂F₄ < CO
- 86. The species with highest magnetic moment (spin only value) is
 - (a) VCl₆⁴⁻

- (b) $\left(\eta^5 C_5 H_5\right)_2 Cr$ (c) $\left[Co(NO_2)_6\right]^{3-}$ (d) $\left[Ni(EDTA)\right]^{2-}$
- The number of metal-metal bonds in $Ir_4(CO)_{12}$ is 87.
 - (a) 4

- Three bands in the electronic spectrum of $\left[\text{Cr} \left(\text{NH}_3 \right)_6 \right]^{3+}$ are due to the following transitions 88.
 - $\text{(A)}~^4A_{2g} \rightarrow {}^4T_{1g} \qquad \text{(B)}~^4A_{2g} \rightarrow {}^4T_{2g}$

Identify the correct statement about them

- (a) Intensity of (A) is lowest
- (b) Intensity of (C) is lowest
- (c) Intensitites of (A), (B) and (C) are similar
- (d) Intensities of (B) and (C) are similar
- 89. Identify the pairs in which the covalent radii of elements are almost similar
 - (A) Nb, Ta
- (B) Mo, W
- (C) La, Lu
- (D) Sc, Y

- (a) A and B only
- (b) A and C only
- (c) B and C only
- (d) A, B and C only

- 90. Consider following lanthanide (III) ions
 - (A) Nd(III)
- (B) Gd(III)
- (C) Dy(III)

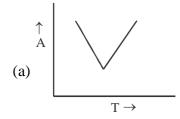
The magnetic moment closest to the spin onlyh value is(are) for

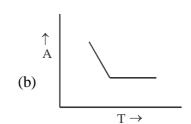
- (a) B only
- (b) A and B only (c) A and C only (d) B and C only
- 91. The Δ_{+} of the following complexes
 - (A) $\left[\text{CoCl}_{4}\right]^{2-}$
- (B) $\left[\text{CoBr}_4 \right]^{2^-}$ (C) $\left[\text{Co} \left(\text{NCS} \right)_4 \right]^{2^-}$ follows the order (b) A > B > C (c) B > A > C (d) C > B > A
- (a) C > A > B

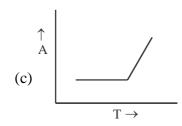
92. In complexometric titration

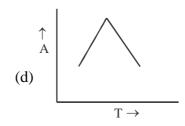
 $S(substrate) + T(titrant) \rightarrow P(product)$

The end point is estimated spectrophotometrically. If S and P have $\varepsilon = 0$, the shape of the titration curve would look like









- 93. Identify the chiral complexes from the following
 - (A) $\left\lceil \operatorname{Cr}(\operatorname{EDTA}) \right\rceil^{-}$
- (B) $\left[\text{Ru} \left(\text{bipy} \right)_{3} \right]^{3+}$ (C) $\left[\text{PtCl} \left(\text{diene} \right) \right]^{+}$

- (a) A only
- (c) A and C only
- (d) B and C only
- 94. Distribution ratio of 'A' between CHCl₂ and water is 9.0. It is extracted with several, 5 mL aliquots of CHCl₃. The number of aliquots needed to extract 99.9% of 'A' from its 5 mL aqueous solution are
- (b) 3
- (c) 4
- The correct equilibrium order for the interconversion of different forms of SiO, is 95.
 - (a) Tridymite quartz cristobalite liquid SiO
 - (b) quartz \longrightarrow Tridymite \longrightarrow cristobalite \longrightarrow liquid SiO₂
 - (c) quartz ← cristobalite ← tridymite ←
 - (d) Cristobalite \times tridymite \times quartz \tilde{\text{liquid SiO}}_2
- The rate equation for the reaction $2AB + B_2 \rightarrow 2AB_2$ is given by 96. rate = $k[AB][B_n]$

A possible mechanism consistent with this rate law is

(a)
$$2AB + B_2 \xrightarrow{\text{slow}} 2AB$$

(b)
$$A_2B_2 + B_2 \xrightarrow{\text{slow}} 2AB_2$$

$$AB + B_2 \xrightarrow{slow} AB$$

$$AB + B_2 \longrightarrow AB_3 (fast)$$

(c)
$$AB_3 + AB \xrightarrow{fast} 2AB_2$$

$$(d)^{\dagger}$$
 $AB + B_2 \Longrightarrow AB_3 (fast)$
 $(AB_3 + AB \xrightarrow{slow} 2AB_2$

- 97. Observe the following statements
 - (I) In the H₂-O₂ reaction, explosion occurs when the rate of chain branching exceeds that of chain termination.
 - (II) The order of the reaction, $nA \to \text{products}$, is 2.5. For this reaction, $t_{1/2} \propto [A]_0^{-\frac{7}{2}}$
 - (III) Unimolecular gas phase reactions are second order at low pressure but becomes first order at high pressure.

Which of the following is correct?

(a) I, II and III are correct

(b) Only II is correct

(c) Only III is correct

(d) I and II are correct.

- 98. For the particle-in-a-box problem in (0,L) an approximate wave function is given as x(L/2-x)(L-x). The average energy \overline{E} for such a state will obey
 - (a) $\frac{h^2}{2mL^2} < \overline{E} < \frac{h^2}{2mL^2}$

(b) $\overline{E} > \frac{h^2}{2mI^2}$

(c) $\frac{h^2}{4mL^2} < \overline{E} < \frac{h^2}{2mL^2}$

- (d) $0 < \overline{E} < \frac{h^2}{8mL^2}$
- 99. For two variables x and y, the following data set is given:

$$\begin{array}{c|cc}
x & y \\
\hline
-1 & 1 \\
0 & 2 \\
1 & 3
\end{array}$$

The correct statement for the covariance A and correlation coefficient B of x and y is

- (a) A=2/3, B=1(b) A=-2/3, B=1(c) A=-2/3, B=-1(d) A=0, B=0The hydrogenic orbital with the form of the radial function 100.

$$r^2(\alpha_1-r)(\alpha_2-r)exp[-\beta r]$$
, where α_1,α_2 and β

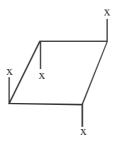
are constants, may be identified as a

- (a) 3d orbital
- (b) 4f orbital
- (d) 5f orbital
- The operator $\left[x,\left[x,p^2\right]\right]$ is identical with (a) $\left[px,\left[x,p\right]\right]$ (b) $\left[xp,\left[x,p\right]\right]$ (c) $-\left[p,\left[x^2,p\right]\right]$ (d) $\left[x,\left[x^2,p\right]\right]$ 101.

- For the particle -in-a-box problem in (0,L), the value of $\langle x^3 \rangle$ in the $n \to \infty$ limit would be 102.
- (c) $\frac{L^3}{4}$
- Identify the Mulliker notation for the following irreducible representation 103.

$$\begin{array}{|c|c|c|c|c|c|} \hline E & C_n & nC_2 & i & \sigma_h \\ \hline 1 & 1 & -1 & -1 & -1 \\ \hline \end{array}$$

- (b) A_{2u}^{-}
- (c) B'_{2u}
- (d) A_{2n}
- Identify the point group symmetry of the following molecule (all C-C bond lengths are equal) 104.



- (a) C₃
- (b) S_4
- (c) D_{2d}
- (d) D_{4d}

105.	The ground state term symbol for Nb(atomic number 41) is ⁶ D. The electronic configuration corresponding to this term symbol is			
	(a) $[Kr]4d^35s^2$	(b) $\left[Kr \right] 4d^4 5s^1$	(c) $\left[\mathrm{Kr}\right] 4\mathrm{d}^5 5\mathrm{s}^0$	$(d) \left[Kr \right] 4d^3 5s^1 5p^1$
106.	In the presence of an external magnetic field (normal Zeeman effect), the transition $^1D_2 \rightarrow ^1P_1$			ffect), the transition ${}^{1}D_{2} \rightarrow {}^{1}P_{1}$
	splits into (a) 9 lines	(b) 8 lines	(c) 7 lines	(d) 6 lines
107.	Identify the Huckel	determinant for cyclob	outadiene	
	$\begin{vmatrix} \alpha - E & \beta \\ \beta & \alpha - E \end{vmatrix}$	$\begin{bmatrix} 0 & 0 \\ \beta & 0 \end{bmatrix}$	$\begin{vmatrix} \alpha - E & \beta \\ \beta & \alpha - E \end{vmatrix}$	$\begin{bmatrix} 0 & \beta \\ \beta & 0 \end{bmatrix}$
	(a) $\begin{bmatrix} 0 & \beta \\ 0 & 0 \end{bmatrix}$	$egin{array}{ccc} 0 & 0 & \ eta & 0 & \ lpha - E & eta & \ eta & lpha - E & \ \end{array}$	(b) 0 β β β	$ \alpha - E \qquad \beta \\ 0 \qquad \alpha - E $
	$ \begin{array}{c cc} \alpha - E & \beta \\ \beta & \alpha - E \\ 0 & \beta \\ 0 & 0 \end{array} $	$ \begin{vmatrix} 0 & \beta \\ \beta & 0 \\ \alpha - E & \beta \\ \beta & \alpha - E \end{vmatrix} $	$ \begin{vmatrix} \alpha - E & \beta \\ \beta & \alpha - E \end{vmatrix} $ (d) $ \begin{vmatrix} 0 & \beta \\ 0 & \alpha \end{vmatrix} $	$\begin{bmatrix} \beta & \beta \\ \beta & 0 \\ \alpha - E & \beta \end{bmatrix}$
100				
108.				f NaOH, the pH of the solution is
	$(p\mathbf{K}_a = -\log \mathbf{K}_a)$	(b) $pK_a + 0.301$	8	
109.	A system consists of	of gaseous H_2 , O_2 , H_2O	and CO ₂ where the a	amount of CO ₂ is specified and the
	equilibrium constan	at for the reaction 2H ₂	$(g) + O_2(g) \Longrightarrow 2H$	$H_2O(g)$ is known. The number of
	degrees of freedom (a) 2	of the system is (b) 3	(c) 4	(d) 5
110.	"Colloids are therm correct pair	odynamically unstable v	with reference to bulk	but kinetically stable". Identify the
	Statements (A) thermodynamic (B) kinetically stable	•	Reasons (C) interfacial surfacial double (D) electrical double (D)	
	(a) $(A) \leftrightarrow (D)$ and	$(B) \leftrightarrow (C)$	(b) $(A) \leftrightarrow (C)$ and	$I(B) \leftrightarrow (D)$
(c) $(A) \leftrightarrow (C)$ and $(B) \leftrightarrow (C)$		(d) $(A) \leftrightarrow (D)$ and	(d) $(A) \leftrightarrow (D)$ and $(B) \leftrightarrow (D)$	
111.	An AX system gave 4 lines at 4.72, 4.6, 1.12 and 1.0 ppm away from the TMS using an nn		from the TMS using an nmr spec-	
	trometer operating	at 100 MHz. What are	the values of J_{AX} (in H	(z) and δ_{AX} (in ppm), respectively
	(a) 12 and 3.6	(b) 6 and 3.6	(c) 12 and 2.86	(d) 6 and 2.86
112.	The equilibrium pop	oulation ratio $\left(n_{j}/n_{i}\right)$	of a doubly-degenerat	e energy level (E _j) lying at energy 2
		lower non-degenerate e	energy level (E _j), assur	ming $k_B T = 1$ unit, will be
	(a) $2e^{-2}$	(b) $2e^2$	(c) e^2	(d) e^{-2}

114.	Identify, from the following, the correct ionic st a 0.01 molal solution of Na ₂ SO ₄ .	rengths for (A) 0.01 molal solution of NaCl and (B)	
	(a) $(A)0.010 \text{ mol kg}^{-1}(B)0.010 \text{ mol kg}^{-1}$ (b)	$(A)0.010 \text{ mol kg}^{-1} (B)0.030 \text{ mol kg}^{-1}$	
	(c) $(A)0.010 \text{ mol kg}^{-1}(B)0.025 \text{ mol kg}^{-1}$ (d)	$(A)0.010 \text{ mol kg}^{-1} (B) 0.015 \text{ mol kg}^{-1}$	
115.	system at equilibrium.	1 100 bosons are kept in it. Find the entropy of the	
	(a) $10^{-2} k_B$ (b) $10^2 k_B$ (c)) $460.6 k_B$ (d) $4.606 k_B$	
116.	Which is correct Nernst equation for redox rea	ction $O + ne \longrightarrow R$?	
	Which is correct Nernst equation for redox real (a) $E = E^0 - \frac{RT}{nF} \ln \frac{[O]}{[R]}$ (b) (b) $\frac{[O]}{[R]} = e^{-\frac{nF}{RT}(E-E^0)}$ (d)	$\frac{\left[O\right]}{\left[R\right]} = e^{\frac{nF}{RT}\left(E - E^{0}\right)}$	
	(c) $\frac{[O]}{[R]} = e^{-\frac{nF}{RT}(E-E^0)}$ (d	$)\frac{\left[\mathrm{O}\right]}{\left[\mathrm{R}\right]}=\mathrm{e}^{\frac{\mathrm{Rf}}{\mathrm{nF}}\left(\mathrm{E}-\mathrm{E}^{\mathrm{0}}\right)}$	
117.	A plane of spacing 'd' shows first order Bragg		
	(a) shows Bragg diffraction at 20 (b)) shows Bragg diffraction at $\frac{\theta}{2}$	
	(c) shows Bragg diffraction at $\sin^{-1}\left(\frac{\sin\theta}{2}\right)$ (d)) shows Bragg diffraction at $\sin^{-1} \left(\frac{\sin 2\theta}{2} \right)$	
118.	In the formation of H, molecules from 2H ator	ns placed at positions A and B, and separated by a	
	distance r _{AB} , a part of the spatial wave function		
	$\Phi_{A}(1)\phi_{A}(2)+\phi_{B}(1)\phi_{B}(2)$		
	(a) This is a covalent term and is important as $r_{AB} \rightarrow \infty$		
	(b) This is an ionic term and is important as $r_{AB} \rightarrow \infty$		
	(c) This is a covalent term and is important as a	$\dot{A}_{AB} \rightarrow 0$	
	(d) This is an ionic term and is important as $r_{\!_{AB}}$	$\rightarrow 0$	
119.	A 0.1 M solution of compound A shows 50% tr	ansmittance when a cell of 1 cm width is used at λ_1	

Which of the following statements is true for a cyclic process?

(b) $\oint dw = 0$

(c) Heat can be completely converted into work(d) Work can be completely converted into heat

(a) 33.3%

at λ_1 nm.

the same cell and at the same wave length?

 $[\log 20 = 1.301; \log 30 = 1.4771; \log 50 = 1.699]$

(b) 50%

113.

(a) $\oint dq = 0$

nm. Another 0.1 M solution of compound B gives the optical density value of 0.1761 using 1cm cell

What will be the transmittance of a solution that is simultaneously 0.1 M in A and 0.1 M in B using

(c) 66.7%

(d) 70%

Using standard equation for intrinsic viscosity $[\eta] = K \overline{M}_{v}^{a}$, for a solution of polymer and any infor-120. mation from the graph identify viscosity average molar mass (\overline{M}) [given that a = 0.5, $K = 5 \times 10^{-5}$ L g^{-1}]

$$\frac{\eta - \eta_0}{\eta_0} \cdot \frac{1}{c}$$

- (a) 10^3 g/mol
- (b) $10^4 \, \text{g/mol}$
- (c) 10^5 g/mol
- 121. Among the following, the correct statement for the following reaction is

O 1. MeMgBr,
$$Et_2O$$
 + (A) (B)

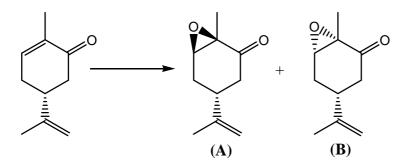
- (a) A is the major product and it will have five signals in the proton decoupled ¹³C NMR spectrum
- (b) A is the minor product and it will have eight signals in the proton decoupled ¹³C NMR spectrum
- (c) B is the major product and it will have five signals in the proton decoupled ¹³C NMR spectrum
- (d) B is the minor product and it will have five signals in the proton decoupled ¹³C NMR spectrum
- 122. For the following three step conversion of A to B, the appropriate sequence of reactions is

- (a) MnO₂; (CH₂OH)₂/p-TSA; PCC
- (b) PCC; MnO₂; (CH₂OH)₂/p-TSA;
- (c) PCC; (CH₂OH)₂/p-TSA; Jones' reagent (d) Jones' reagent; (CH₂OH)₂/p-TSA; MnO₂.
- 123. Which one of the following statements is true for the following transformation?

Ph CHO MgBr ,
$$Et_2O$$
 Ph Me (A) (B)

- (a) A is the major product and it is a Cram product.
- (b) A is the major product and it is anti-Cram product.
- (c) B is the major product and it is a Cram product.
- (d) B is the major product and it is anti-Cram product.

124. Which one of the following statements is true for the following transformation?



- (a) Suitable reagent is m-CPBA and B is the major product
- (b) Suitable reagent in m-CPBA and A is the major product.
- (c) Suitable reagent is aq. H₂O₂/NaOH and B is the major product.
- (d) Suitable reagent is aq. H₂O₂/NaOH and A is the major product.
- 125. The compound formed in the following reaction sequence is

$$\begin{array}{c|c} \hline 1. \text{ Li, liq. NH}_3, \text{ t-BuOH} \\ \hline 2. 10\% \text{ aq. H}_2\text{SO}_4 \\ \hline \\ \text{(a)} \\ \text{H}_2\text{N} \\ \hline \\ \text{(c)} \\ \hline \end{array}$$

126. Among the following compounds, the one which has highest dipole moment is

127. In the UV-visible is spectrum, a diterpenoid exhibited a λ_{max} at 275 nm. The compound, among the choices given below is

$$(c) \\ HOOC^{IIII} \\ \hline H$$

128. The major product formed in the following reaction is

129. In the broad band decoupled ¹³C NMR spectrum, the number of signals appearing for the two pyrenediols A and B

$$(A) \qquad (B) \qquad (B)$$

- (a) eight and eight (b) eight and sixteen (c) five and ten (d) five and eight.
- 130. An organic compound exhibited the following ¹H NMR spectra data:

$$\delta$$
 7.80 (2 H, d, J = 8 Hz), 6.80 (2 H, d, J = 8 Hz), 4.10 (2H, q, J = 7.2 Hz), 2.4(3H, s), 1.25(3 H, t, J = 7.2 Hz)

The compound, among the choices given below is,

131. α -Pinene on reaction with dilute alkaline KMnO₄ produces a diol, which on further oxidation with chromium trioxide gives product A, which undergoes a positive haloform test. The compound A is

$$(a) \qquad (b) \qquad (c) \qquad (c) \qquad (d) \qquad (d)$$

132. The major product formed in the reaction of guanosine with one equivalent of methyl iodide is

$$(a) \underset{H_2N}{\overset{\text{Me}}{\longrightarrow}} \underset{N}{\overset{\text{Me}}{\longrightarrow}} \underset{N}{\overset{\text{O}}{\longrightarrow}} \underset{N}{\overset{\text{Me}}{\longrightarrow}} \underset{N}{\overset{\text{O}}{\longrightarrow}} \underset{N}{\overset{\text{Ne}}{\longrightarrow}} \underset{N}{\overset{\text{O}}{\longrightarrow}} \underset{N}{\overset{\text{Ne}}{\longrightarrow}} \underset{N}{\overset{\text{O}}{\longrightarrow}} \underset{N}{\overset{\text{Ne}}{\longrightarrow}} \underset{N}{\overset{\text{O}}{\longrightarrow}} \underset{N}{\overset{\text{Ne}}{\longrightarrow}} \underset{N}{\overset{\text{O}}{\longrightarrow}} \underset{N}{\overset{\text{Ne}}{\longrightarrow}} \underset{N}{\overset{N}} \underset{N}{\overset{N}{\overset{N}} \underset{N}{\overset{N}} \underset{N}{\overset$$

133. The major product formed in the following reaction is

134. Reaction of the dipeptide, given below, with hydrogen in the presence of 10% palladium over carbon, produces a mixture of

- (a) Gly-Leu + toluene + carbon dioxide
- (b) Phe-Leu + toluene + carbon dioxide
- (c) Phe-Leu + benzyl alcohol + carbon dioxide
- (d) Gly-Leu + benzyl alcohol + carbon dioxide

135. Among the following, the most suitable reagent for carrying out resolution of racemic 3-methylcyclohexanone is

136. In the following reaction sequence, structures of the major product X and Y are

137. Consider the following reaction sequence

The overall yield for the formation of p-hydroxyacetanilide and o-hydroxyacetanilides from phenol, respectively, are approximately

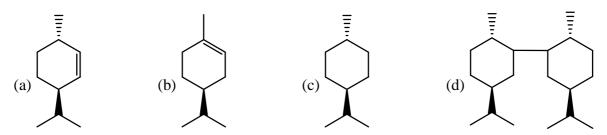
- (a) 57 and 20%
- (b) 57 and 68%
- (c) 83 and 68%
- (d) 83 and 20%
- 138. The most stable conformations of 1, 2-difluoroethane and dl-2, 3-butanediol are

139. Reaction of (S)-1, 2, 4-butanetriol with benzaldehyde in the presence of catalytic amount of p-TSA furnished the major product A. The structure of A is

$$(a) \qquad \qquad (b) \qquad (b) \qquad (c) \qquad (d) \qquad (d$$

140. The major product formed in the following reaction is

141. The major product formed in the following reaction is



142. The major product formed in the following reaction is

a)
$$N$$
 (b) N (c) N (d) N OH

143. The major product formed in the following reaction is

144. Predict the condition A and the structure of the major product B in the following sequence.

A
$$1. O_3$$
; Me_2S B $2. Ph_3P = CHCOOEt$ (excess)

(a) A is hv, B is $COOEt$ (b) A is hv, B is $COOEt$ (c) A is Δ , B is $COOEt$ (d) A is Δ , B is $COOEt$

145. The most appropriate mode of cyclisation in the following transformation is



- (a) con-rotatory in photochemical; and dis-rotatory in thermal conditions.
- (b) con-rotatory in thermal; and dis-rotatory in photochemical conditions.
- (c) con-rotatory in thermal; and con-rotatory in photochemical conditions.
- (d) dis-rotatory in photochemical; and dis-rotatory in thermal conditions.

Dynamic Chemistry Point
Oynamic

Con: 9871547744