## BSC\_SEM\_V

NC-33 Physical Chem+ Inorganic Chemistry -XIII (CGPA) Time: One Hour Max. Marks: 40 Instructions • Attempt all 40 questions 1 Heteronuclear metal carbonyl is-(A)[Cr(CO)5] (B)[Mn<sub>2</sub>(CO)<sub>10</sub>] (C)[MnRe(CO)<sub>10</sub>] (D)[Co<sub>2</sub>(CO) 8] 2 Ni in Ni(CO)<sub>4</sub> undergoes-(B)SP<sup>2</sup> hybridisation (C)SP<sup>3</sup> hybridisation (D)dSP<sup>2</sup> hybridisation (A)SP hybridisation

(D)4

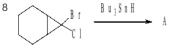
3 Metal-Metal bonds present in Fe<sub>3</sub>(CO)<sub>12</sub> is-(A)1 (C)3

(A)1 bridge and 7 terminals (B)2 bridge and 6 terminals (C)3 bridge and 5 terminals (D)4 bridge and 4 terminals 5 Ionic organometallic compound is-

(A)NaC<sub>6</sub>H<sub>5</sub> (B)(CH<sub>3</sub>)<sub>4</sub>Si (C)Al<sub>2</sub>Me<sub>6</sub> (D)None of the above 6 Compounds formed by less electropositive metal is -

(B)Covalent organometallic (C)Electron deficient organometallic (D)Transition metal organometallic (A)Ionic organometallic 7 Organolithium compounds react with alkylnitriles gives-

(A)Aldehydes (C)Alcohols (D)Acids (B)Ketones



In above reaction 'A' is-



9 Transition metal organometallic compounds formed due to-(A)Interaction between P-orbital of (B)Symmetry property of d-orbital (C)Both A and B (D)None of the above

Organic ligand and p or d-orbital of metal

10 ----- is used in the production of linear higher olefins.

4 The carbonyl groups present in bridge structure of Co2(CO)8 is-

(A)Trialkylaluminium (B)Trialkyltin halide (C)Alkyllithium (D)Dialkylmercury

11 Association of R<sub>3</sub>SnX molecule exist due to-

(A)Halogen bridges (C)Tin bridges (D)All of the above

 $RX + Hg \xrightarrow{A} R_2 Hg_+ HgX_2$ 

In the above reaction 'A' is -

(A)Na/Hg (C)Zn/Hq (D)HgCl<sub>2</sub>

 $^{3}$  LiPh +  $\bigcirc$   $\bigcirc$   $\bigcirc$   $\bigcirc$   $\bigcirc$   $\bigcirc$  Et<sub>2</sub>O  $\bigcirc$ 

In above reaction 'A' is -



14 The selection rule for diatomic rigid rotator is-

$$(A)_{\Delta J} = 0 \qquad (B)_{\Delta J} = \pm 1 \qquad (C)_{\Delta J} = \pm 2 \qquad (D)_{\Delta J} = \pm 3$$

15 By using Distribution law we can determine -

(A)Solubility of solute (B)Association of solute (C)Dissociation of solute (D)All of the above

16 For the study of the distribution law the two solvents should be -

(A)Volatile (C)<sub>Miscible</sub> (D)<sub>Non-miscible</sub> (B)Reacting with each other

 $^{17}$  KD =  $C_1/C_2$  (1-x) is used for -

(A)Dissociation of solute (B)Association of solute (C)Energy of solute (D)None of these

18 If solute 'A' is dimerized in one of the solvent then following expression is applicable-

 $(D)_{KD} = C_1 / C_2^2$  $(B)_{KD} = C_1 / {}^2 \sqrt{C_2}$  $(C)_{KD} = C_1 / \sqrt{3} \sqrt{C_2}$  $(A)_{KD} = C_1 / C_2$ 

(B) <sub>Three molecules changes their concentration during reaction.</sub>	(C) <sub>Both A and B</sub> .	(D) <sub>None</sub> of the above.
also reacts to give back the reactants is ca	illed-	
(B)Opposing reaction.	(C)Parallel reaction.	(D) <sub>Reaction mechanis</sub>
gid Rotor is-		
$(B)_{I}=\mu^{2}r.$	$(C)_{I} = \mu r.$	$(D)_{\bar{I}}=\mu^2r^2.$
(B) <sub>Gamma ray</sub> .	(C) <sub>Microwave region</sub> .	$(D)_{IR}$ region.
or Association of solute in one of the solv	ent is -	
$(B)_{K_D} = C_1/C_2$	$(C)_{K_D} = C_1/C_2 \times (1-x)$	(D) $_{\mbox{\footnotesize Both B}}$ and C
$(B)_{C} = K.P$	$(C)_{P=C \times K}$	$(D)_{\mathrm{C}} = \mathrm{K} \sqrt{\mathrm{P}}$
nethod is-		
(B) <sub>Multiple</sub> extraction.	(C)Double extraction.	(D) <sub>None of these</sub>
e having-		
(B)Lone pair of electron.	$(C)_{\sigma-electrons}$ .	(D)Both A and B.
n molecule is exposed to-		
(B) <sub>IR radiation</sub> .	(C) <sub>UV</sub> and visible radiation.	$(D)_{\gamma}$ - $rays$
force constant is -		
$(B)_{B.E} \propto 1/K.$	(C) <sub>0</sub>	(D) <sub>B.E</sub> $\propto 1/\sqrt{K}$
(B)During electronic transition bond length not changes appreciably	(C)Both A and B.	(D) <sub>None of these.</sub>
horter frequency) than that of incident lig	ht are called-	
(B) <sub>Antistoke's lines.</sub>	(C) <sub>Rayleigh lines</sub> .	$(D)_{X-Rays.}$
frequency of scattered light then Raman	shift (Δυ) is-	
$(B)_{\Delta\upsilon} = \upsilon_{i} + \upsilon_{S}$	$(C)_{\!\Delta\upsilon}=\upsilon_i^{}/\upsilon_{\!_S}$	$(D)_{\!\Delta\upsilon}=\upsilon_{s}\!/\upsilon_{i}$
(B) <sub>Antistokes lines</sub> .	(C) <sub>Rayleigh lines</sub> .	(D)Both A and B.
(B) <sub>Low</sub> Concentration.	(C)Dissolved solute remains in its molecular state.	(D) <sub>All of above</sub> .
cillator is -		
$(B)_{\Delta V} = \pm 2$	$(C)_{\Delta V} = 0$ ,±1	$(D)_{\Delta V} = 0$ , $\pm 2$
e as a Rigid rotator in cm <sup>-1</sup> is-	,	,
•	$(C)\varepsilon_{I} = BJ(J+1)^{2}$	$(D)\epsilon_J = \mathrm{BJ}(J + n)$
, , ,	3	-3 - (- )
•	(C) <sub>3B</sub>	(D) <sub>4B</sub>
(B) <sub>Mol 1</sub> -1 S-1	$(C)_{Mol}^2 1^{-2} S^{-1}$	$(D)_{Mol}^{-2} 1^2 S^{-1}$
		, ,
•	(C)Chain reaction	(D)Both B and C.
note chamban reaction.	. , Chan I would the	, , , , , , , , , , , , , , , , , , ,
		or (D)None of these.
	*	
	concentration during reaction. also reacts to give back the reactants is cat (B)Opposing reaction. gid Rotor is- gid Rotor is- (B) $I = \mu^2 r$ .  (B)Gamma ray.  Association of solute in one of the solve (B) $K_D = C_1/C_2$ (B) $C = K.P$ method is- (B)Multiple extraction.  An implementation and is exposed to- (B)IR radiation.  force constant is - (B)B.E $\propto 1/K$ .  (B)During electronic transition bond length not changes appreciably morter frequency) than that of incident light (B)Antistoke's lines.  frequency of scattered light then Raman (B) $\Delta v = v_1 + v_8$ (B)Antistokes lines.  (B)Low Concentration.  Cillator is - (B) $E_J = BJ^2(J+1)$ spectra of rigid rotor is- (B) $E_J = BJ^2(J+1)$ spectra of rigid rotor is- (B)Photo chemical reaction.  (B)The final product reacts to give back the reactants.	concentration during reaction.  also reacts to give back the reactants is called- (B)Opposing reaction.  (C)Parallel reaction.  gid Rotor is- (B)I = $\mu$ 7r.  (C)I = $\mu$ r.  (B)Gamma ray.  (C)Microwave region.  or Association of solute in one of the solvent is - (B)KD = C1/C2  (C)KD = C1/C2 X (1-x)  (B)C = K.P  (C)P= C X K  method is- (B)Multiple extraction.  (C)Double extraction.  (B)Multiple extraction.  (C)Touble extraction.  (D)Touble extraction.  (D)Touble extraction.  (D)Touble extraction.  (D)Touble extraction.  (E)Touble extraction.  (D)Touble extraction.  (D)Touble extraction.  (E)Touble extraction.  (D)Touble extraction.  (D)Touble extraction.  (D)Touble extraction.  (E)Touble extraction.  (D)Touble extraction.  (E)Touble extraction.  (C)Touble extraction.  (C)Touble extraction.  (D)Touble extraction.  (E)Touble extraction.  (C)Touble extraction.  (C)To

 $(\mathsf{C})_{B'} = B$ 

(D)Both B and C

 $(\mathsf{B})_B > B'$ 

 $(A)_{B'} > B$