Exercise-1

Marked Questions may have for Revision Questions.

Section (A) : Introduction to Bonding, Octet rule and Ionic bond

OBJECTIVE QUESTIONS

A-1.	 An ionic bond A⁺ B⁻ is most likely to be formed when : (1) The ionization energy of A is high and the electron affinity of B is low (2[*]) The ionization energy of A is low and the electron affinity of B is high (3) The ionization energy of A and the electron affinity of B is high (4) The ionization energy of A and the electron affinity of B is low 				
A-2.	Which forms a crystal of NaCl ? (1) NaCl molecules (2*) Na⁺ and Cl⁻ ions (3) Na and Cl atoms (4) None of these				
A-3.	Which of the following pair of elements form a compound with maximum ionic character ?(1) Na and F(2*) Cs and F(3) Na and Br(4) Cs and I				
A-4.	Two element have elec (1*) ionic	ctronegativity of 1.2 and (2) polar covalent	3.0. Bond formed betwee(3) co-ordinate	en them would be : (4) metallic	
A-5.	Which one of the follov (1) B and Cl ₂	ving pairs of elements is (2*) K and O ₂	most likely to form an ior (3) O2 and Cl2	nic compound? (4) Al and I ₂	
A-6.			ease of formation of ioni (3) Mg ²⁺ > Al ³⁺ > Na ⁺	-	
A-7.	Which of the following (1) RbF	shows the highest lattice (2) CsF	energy ? (3*) NaF	(4) KF	
A-8.	Which of the following (1) Cs – F	have low lattice energy 3 (2) Cs – Cl	? (3) Cs – Br	(4*) Cs – I	
Sect	ion (B) : Fajan's rule	9			
D 4	According to Fajan's rule covalent character is favoured by :(1) large cation and small anion(2*) small cation and large anion(3) large cation and large anion(4) small cation and small anion				
B-1.	(1) large cation and sm	nall anion	(2*) small cation and la	•	
в-1. В-2.	(1) large cation and sm(3) large cation and lar	nall anion ge anion	(2*) small cation and la	nall anion	
	 (1) large cation and sm (3) large cation and lar Which one of the follow (1) Pb²⁺, Br⁻ 	nall anion ge anion ving combination of ion v (2*) Pb ⁴⁺ , Br⁻ ent character of alkaline e aCl₂ < SrCl₂	 (2*) small cation and la (4) small cation and sm vill have highest polarisat (3) Fe²⁺, Br⁻ 	nall anion tion ? (4) Fe ³⁺ , Br⁻ Cl₂ < MgCl₂	
B-2.	(1) large cation and sm (3) large cation and lar Which one of the follow (1) Pb^{2+} , Br^- Correct order of covale (1) $BeCl_2 < MgCl_2 < Cat (3*) BeCl_2 > MgCl_2 > CWhich of the following(1) CCl_4 < BeCl_2 < BC$	nall anion ge anion ving combination of ion v (2*) Pb ⁴⁺ , Br ⁻ ent character of alkaline e aCl ₂ < SrCl ₂ aCl ₂ > SrCl ₂ is in order of increasing e l ₃ < LiCl	 (2*) small cation and la (4) small cation and sm vill have highest polarisat (3) Fe²⁺, Br⁻ earth metal chloride in (2) BeCl₂ < CaCl₂ < Sr((4) SrCl₂ > BeCl₂ > CaCl₂ covalent character ? (2) LiCl < CCl₄ < BeCl₂ 	nall anion tion ? (4) Fe ³⁺ , Br− Cl₂ < MgCl₂ Cl₂ > MgCl₂ ₂ < BCl ₃	
B-2. B-3.	 (1) large cation and sm (3) large cation and lar Which one of the follow (1) Pb²⁺, Br⁻ Correct order of covale (1) BeCl₂ < MgCl₂ < Ca (3*) BeCl₂ > MgCl₂ > C Which of the following 	nall anion ge anion ving combination of ion v (2*) Pb ⁴⁺ , Br ⁻ ent character of alkaline e aCl ₂ < SrCl ₂ aCl ₂ > SrCl ₂ is in order of increasing e l ₃ < LiCl	 (2*) small cation and la (4) small cation and sm vill have highest polarisat (3) Fe²⁺, Br⁻ earth metal chloride in (2) BeCl₂ < CaCl₂ < Sr((4) SrCl₂ > BeCl₂ > CaC covalent character ? 	nall anion tion ? (4) Fe ³⁺ , Br− Cl₂ < MgCl₂ Cl₂ > MgCl₂ ₂ < BCl ₃	
B-2. B-3.	(1) large cation and sm (3) large cation and lar Which one of the follow (1) Pb^{2+} , Br^- Correct order of covale (1) $BeCl_2 < MgCl_2 < Cat (3^*) BeCl_2 > MgCl_2 > CWhich of the following(1) CCl_4 < BeCl_2 < BCl(3*) LiCl < BeCl_2 < BCl$	nall anion ge anion ving combination of ion v (2*) Pb ⁴⁺ , Br ⁻ ent character of alkaline e aCl ₂ < SrCl ₂ aCl ₂ > SrCl ₂ is in order of increasing e l ₃ < LiCl	(2*) small cation and la (4) small cation and sm vill have highest polarisat (3) Fe^{2+} , Br^- earth metal chloride in (2) $BeCl_2 < CaCl_2 < Sr((4) SrCl_2 > BeCl_2 > CaCl_2(4) SrCl_2 > BeCl_2 > CaCl_2(2) LiCl < CCl_4 < BeCl_2(4) LiCl < BeCl_2 < CCl_4$	nall anion tion ? (4) Fe ³⁺ , Br− Cl₂ < MgCl₂ Cl₂ > MgCl₂ ₂ < BCl ₃	

СНЕ	MISTRY FOR JEE			CHEMICAL BONDING		
•	(1*) AIF ₃	(2) Al ₂ O ₃	(3) AIN	(4) Al₄C₃		
B-7.	Which compound amor (1) AICI₃	ng the following has leas (2^*) AlI $_3$	st ionic character? (3) MgI₂	(4) CsI		
B-8.	In which of the followin (1) NaCl	g compound the cation h (2) AlCl₃	nas pseudo inert gas con (3*) Cเ	-		
Secti	ection (C) : Covalent Bond and Coordinate Bond					
C-1.	1st and 2nd period) : (1) the number of unpa (2) the number of paire (3) the number of unpa	ired p-electrons d d-electrons		al charge) is equal to (excluding		
C-2.	Which of the following ((1) MgCl ₂	contains both electrovale (2) H₂O	ent and covalent bonds ? (3*) NH₄Cl	(4) none		
C-3.	The types of bond pres (1) only covalent	ent in N₂O₅ are : (2) only ionic	(3) ionic and covalent	(4*) covalent & coordinate		
C-4.	Example of super octet (1) SF ₆	t molecule is : (2) PCl₅	(3) IF7	(4*) All of these		
C-5.	The number of electror (1) 2	ns involved in the bond fo (2) 4	ormation in N₂ molecule i (3) 10	s : (4*) 6		
C-6.	The octet rule is not ob (1) CO ₂	eyed in : (2) BCl₃	(3) PCI₅	(4*) (2) and (3) both		
C-7.	For the formation of cor (1) 1.7	valent bond the differend (2) More than 1.7	ce in the value of electror (3) 1.7 or more	negativity should be : (4*) equal to or less than 1.7		
C-8.	The covalency of nitrog (1) 0	gen in HNO₃ is : (2) 3	(3*) 4	(4) 5		
C-9.	Which one of the follow (1*) NH₄Cl	/ing molecules has a co- (2) AICl₃	ordinate as well as coval (3) NaCl	lent bond ? (4) Cl ₂		
C-10.	Which of the following $1. \text{ ClO}_{4^-}$, $2. \text{ BF}_{3^-}$	species are hypervalent , 3. SO4 ²⁻ ,				
	(1) 1, 2, 3	(2*) 1, 3	(3) 3, 4	(4) 1, 2		

Section (D) : Formal charge and Resonance

D-1. What are the formal charges on central sulphur and each terminal oxygen atoms	in SO ₂ ?
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D-2.	Resonating structures have different :	
	(1) atomic arrangements	(2*) electronic arrangements
	(3) functional groups	(4) alkyl groups

D-3. Which among the following resonance structures of N₃⁻ satisfies the octet rule but is ruled out as a resonance structure?

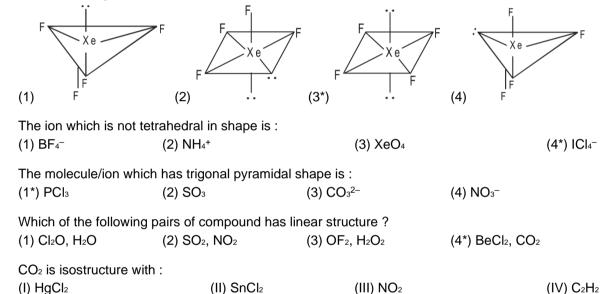
				•
	(I) [:N = N = N:] ⁻ (1*) IV only	(II) [:Ņ́ – N ≡ N:] [–] (2) I and IV only	(III) [:N ≡ N – ┆┆:]⁻ (3) I only	$(IV) \begin{bmatrix} N = N \\ N \\ N \end{bmatrix}^{-}$ (4) II and III only
D-4.		owing is not the resonatin (2) O = C = O	•	(4) ⁺ O ≡ C − O ⁻
D-5	•	of C-C bond in C6H6 is ca/k dk vkSlr ca/k Øe gS % (2) 2	(3*) 1.5	(4) 1.33
D-6.		which has the weakest c		
		H₃COO⁻ (3) (*) CO ₃ ²⁻
D-7				´ ce forms for N₂O (diamagnetic)? [Schaum's series]
	(A) $\dot{:}$ \ddot{N} = N = \ddot{O} : (B) (1) A and C	$N \equiv N - \dot{\Omega}$: (2) C , E and D	$\dot{N} - N \equiv O;$ (E (3*) D and E	$\dot{N} = O = \ddot{N}; (E)$ $\dot{N} = N = \dot{O};$ (4) C and D.
Secti	on (E) : Valence b	ond theory		
E-1.	VBT is given by : (1*) Heitler & london	(2) Pauling & Slater	(3) Hund & Mulike	n (4) Huckel & Hund
E-2.	Number and type of t (1) one sigma (σ) and (3) one σ and one an	• • •	on atoms in CaC₂ are (2*) one σ and two (4) one σ bond	
E-3.	The total number of c	σ and π bonds in C ₂ (CN)	4 are :	
	(1*) 9 σ and 9 π σ and 18π	(2) 9 σ and 1	8π (3	3) 18 σ and 9 π (4) 18
E-4.	Acetylene consists of (1*) both sigma and p (3) pi bond only		(2) sigma bond on (4) none of these	ly
E-5.	Number of bonds in §	SO2 are :	(1)	
	(1) two σ and two π		(2) two σ and one	π
	(3*) two σ, two π and	one ione pair	(4) none of these	
E-6	Effective overlapping			
	(1) ⊕⊖+⊕⊝	(2) + + + + + + + + + + + + + + + + + + +	(3*) ⊕⊖+(Э⊕ (4) All the above
E-7	(1) A sigma bond is s(2*) p-orbitals always(3) s-orbitals never for	atement according to Va tronger then π - bond have only sidewise ove rm π - bonds v one sigma bond betwee	erlapping	

Section (F) : VSEPR Theory

F-1. In which of the following molecules number of lone pairs and bond pairs on central atom are not equal ?

CHEMISTRY FOR JEE CHEMICAL BONDING (2*) I₃-(1) H₂O (3) O₂F₂ (4) SCI₂ F-2. Which one of the following species is not linear? (1) CO₂ (2*) CIO 2 (3) I₃-(4) NO₂₊ F-3. Match list I with List II and select the correct answer using the codes given below the lists. List I List II (Compound) (Shape) (a) CS_2 1. Bent SO₂ (b) 2. Linear BF₃ 3. Trigonal planer (C) (d) NH₃ 4. Tetrahedral 5. Trigonal pyramidal

F-4. Which is the right structure of XeF₄?



F-8.	CO ₂ is isostructure with :				
	(I) HgCl₂	(II) SnCl₂	(III) NO2	(IV	
	The correct answer is :				
	(1*) I, IV	(2) I, II	(3) II, III	(4) , ,	

F-9. The pair having similar geometry is : (1) BF₃, NH₃ (2*) BF₃, AIF₃ (4) BCl₃, PCl₃

Section (G) : Hybridisation and Bond angle

G-1.	Which of the following has been arranged in increasing order of % p-character?				
	$(1^*) sp < sp^2 < sp^3$	(2) $sp^3 < sp^2 < sp$	(3) sp ² < sp ³ < sp	(4) $sp^2 < sp < sp^3$	
G-2.	Choose the molecules	in which hybridisation oc	curs in the ground state	?	
	(a) BCl₃	(b) NH₃	(c) PCl₃	(d) BeF ₂	
G-3.	In C—C bond C ₂ H ₆ und	lergoes heterolytic fissior	n, the hybridisation of two	o resulting carbon atoms is :	
	(1) sp ² both	(2) sp ³ both	(3*) sp², sp³	(4) sp, sp ²	
G-4.	The hybridization in PF	₃ is :			
	(1*) sp ³	(2) sp ²	(3) dsp ³	(4) d ² sp ³	

(3) BeF₂, H₂O

F-5.

F-6.

F-7.

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• G-5.	Which of the following com (1) CH ₄ (2)	pounds have bond an) CO2	gle as nearly 90º ? (3) H₂O	(4*) SF ₆
G-6	sp ² – hybridisation is show (1) BeCl ₂ (2*	/n by : *)) BF₃	(3) NH₃	(4) XeF ₂
G-7.	The hybridisation of carbon (1*) sp ³ , sp ² , sp (2)		and acetylene is (respective) (3) sp ² , sp ³ , sp	ctively) – (4) sp, sp³, sp²
G-8.	The hybridization of the cer (1) dsp ² (2)	ntral atom in $\frac{ICI_2^+}{IS}$ is -	(3) sp²	(4*) sp ³
G-9.	Each carbon in carbon sub (1) sp² - hybridized (3*) sp-hybridized	oxide is :	(2) sp ³ -hybridized (4) sp ² -hybridized but lir	nked with one co-ordinate bond
G-10.	In which of the following pa $(1) \text{ CIF}_3$, CIF $_3\text{O}$ (2)	•	e central atom is differen (3*) [CIF₂O]⁺, [CIF₄O]⁻	
G-11.	OF ₂ is : (1) Linear molecule and sp (3*) Bent molecule and sp ³	•	ahedral molecule and sp e of these	³ hybridised
G-12.	Which has the smallest bor(1*) OSF2(2)	nd angle (X – S – X) ir) OSCl₂	the given molecules? (3) OSBr ₂	(4) OSI ₂ .
G-13.	Consider the following iodic	des : PI₃ 102°	AsI₃ 100.2°	SbI₃ 99°
	The bond angle is maximur (1) due to small size of pho (3) due to less electronegat	osphorus	(2*) due to more bp–bp(4) none of these	repulsion in PI₃
G-14.	Arrange the following in the (1) $P_4 < PH_3 < H_2O$ (2)	-		-
G-15.	The bond angle H – N – H (1) $NH_2^- > NH_3 > NH_4^+$ (2*			
G-16.	The angle between two cov (1) H ₂ O (2*	valent bond is maximu *) CO₂	ım in : (3) NH₃	(4) CH4
G-17.	00	nd angle :) PBr₃	(3) PF₃	(4*) Pl ₃
Secti	on (H) : Electron defici	ient bond and Ba	ck bonding	
H-1	Electron deficient molecule (1) I ₂ Cl ₆ (2*	e among the following *) B ₂ H ₆	is : (3) Al₂Cl₀	[Made by BKS Sir_2015] (4) All of these
H-2	For B ₂ H ₆ S ₁ : Each boron is sp ³ hybri S ₂ : from terminal 'H' & two hydrogen in different plane S ₃ : It has 4 σ bond & 2 brid	'B' atom are in same	plane but two bridge	

•				
	S ₄ : 8 σ bonds are pres (1) TTFF	sent in it (2*) T T T F	(3) F F T F	(4) FTFT
H-3	Identify the species con (1*) (BeH ₂) _n	ntaining Banana bonds (2) BF₃	(3) (AICI ₃) ₂	(4) (BeCl ₂) _n
H-4	Which is not true abour (1) Both 'B' atoms are (3) Two hydrogens occ	sp ³ hybridised	(2*) Boron atom is in ground state (4) There are two, three centre two electron bonds	
H-5	 For BF₃ molecule which of the following is true ? (1) B-atom is sp² hybridised. (2) There is a pπ–pπ back bonding in this molecule. (3) Observed B–F bond length is found to be less than the expected bond length. (4*) All of these 			
H-6	Respective order of strength of back-bonding in boron trihalides is :(1) $BF_3 < BCI_3 < BBr_3$ (2) $BF_3 > BBr_3 > BCI_3$ (3*) $BF_3 > BCI_3 > BBr_3$ (4) $BF_3 < BBr_3 < BCI_3$			
H-7	 For BF₃ molecule which of the following will not be true (1) It has less bond length than BF₄⁻ (2) It has less bond length than the compound [NH₃ → BF₃] (3*) It's bond strength is increased because of pπ–dπ back bonding (4) It forms BF₄⁻ when hydrolysed in water. 			
H-8.	Which is the true statement about (SiH ₃) ₃ N ? (1*) It is trigonal planar. (2) It is trigonal pyramidal. (3) It is stronger lewis base than that of (CH ₃) ₃ N. (4) It has a total of 9 sigma bonds.			
Secti	on (I) : Molecular O	rbital Theory (MOT))	
I-1.	When two atomic orbita (1*) two moleculear orb (3) three molecular orb		(2) one molecular orbit (4) four molecular orbit	
I-2.	During the formation of density is :	f a molecular orbital from	atomic orbitals of the sa	ame atom, probability of electron
	(1) non zero in the nod(3*) zero in the nodal p		(2) maximum in the no (4) zero on the surface	•
I-3.	Which one of the follow (1) H_2^+	ving can not exist on the (2) He ₂ +	basis of molecular orbita (3) C2	l theory ? (4*) He₂
I-4.	Which of the following (1) B ₂	has fractional bond orde (2) O ₂ ²⁻	r ? (3) F ₂	(4*) H₂ [−]
I-5.		g set, the values of bond I (2) CN, NO²+, CN⁻, F₂	d orders will be 2.5 ? (3) O ₂ + , NO ²⁺ , O ₂ ²⁺ , CN	N [−] (4) O ₂ ^{2−} , O ₂ [−] , O ₂ ⁺ , O ₂
I-6.	Among the following sp (1) B ₂	becies, which has the mir (2*) C₂	nimum bond length ? (3) F ₂	(4) O ₂ -
I-7.	Number of antibonding (1*) 4	electrons in N₂ is : (2) 10	(3) 12	(4) 14

I-8.	Pick out the incorrect statem (1) N ₂ has greater dissociation (3*) Bond length in N ₂ + is les	on energy than N_{2^+} (2)	O ₂ has lower dissocia Bond length in NO ⁺ is	•••	
I-9.	Which of the following pairs $(1^*) N_{2^+}$ and O_{2^+} (2) F			(4) C_2 and N_2	
I-10.	A simplified application of M((1) 2 (2*)			uld give its bond (4) 0.5	order as :
I-11.	Which of the following specie (1*) NO ⁻ (2) (CN ⁻	(4) CO	
I-12.	The following molecules/spe the correct order. (I) O_2 ; (II) O_2^- ; (III) $O_2^{2^-}$; (1*) I I I < I I < I V (2) I	(IV) O ₂ +		ncreasing bond o (4) I I < I I I I < I <	
I-13.		-		me magnetic pro (4) (I I) and (I I I)	-
I-14.	Negative bond order means (1*) molecule is unstable (3) molecule is neutral		(2) molecule is s none of them	table	
Sect	ion (J) : Dipole Moment				
J-1.	Which hydrogen is most pola (1) LiH (2) () HF	(4) HI	
J-2.	The most polar bond is :(1) C - H(2) I	N – H (3)	S – H	(4*) O – H	
J-2. J-3.			S-H	(4*) O – H	
	(1) C – H (2) M		7	(4*) O – H (4)	
	(1) C – H (2) M Which has maximum dipole	moment ?	hanent dipole moment	(4) ←→	[CB]
J-3.	(1) $C - H$ (2) M Which has maximum dipole (1*) (2) Of the following molecules, t (1) SiF ₄ (2) E Which of the following has th	moment ? (3) he one, which has perm BF_3 (3*)	hanent dipole moment) PF ₃	(4)	[CB] (4) NH₃
J-3. J-4.	(1) $C - H$ (2) M Which has maximum dipole (1*) (2) Of the following molecules, t (1) SiF ₄ (2) E Which of the following has th	moment ? (3) the one, which has perm BF_3 (3*) the least dipole moment 3 SO_3 unds the one that is pola	hanent dipole moment) PF ₃ ? (3) XeO ₃ ar and has central ator	(4)	(4) NH₃
J-3. J-4. J-5.	(1) $C - H$ (2) M Which has maximum dipole (1*) (2) Of the following molecules, t (1) SiF ₄ (2) E Which of the following has th (1) NF ₃ (2*) Among the following compose (1) H ₂ CO ₃ (2) S	moment ? (3) the one, which has perm BF_3 (3*) the least dipole moment 3 SO_3 unds the one that is pola SiF_4 (3) wen molecules are such	hanent dipole moment) PF ₃ ? (3) XeO ₃ ar and has central ator BF ₃	(4)	(4) NH₃ lisation is :

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	(1*) CH4 < NF3 < NH3 < H2O (3) NH3 < NF3 < CH4 < H2O		(2) $NF_3 < CH_4 < NH_3 < H_2O$ (4) $H_2O < NH_3 < NF_3 < CH_4$	
J-9.	Carbon tetrachloride has no dipole moment be (1) its planar structure (3) similar sizes of carbon and chlorine	(2*) its regular tetrahe	dral structure inities of carbon and chlorine	
J-10.	The molecules which has zero dipole moment (1) CIO_2 (2) PCI_3	is : (3*) XeF₄	(4) CHCl₃	
J-11.	The dipole moment of HCl is 1.03 D. If H–Cl character in the H–Cl bond - (1) 60% (2) 39%	bond distance is 1.275 Å (3) 29%	A, what is the percentage of ionic (4*) 17%	
Sect	ion (K) : Hydrogen bonding			
K-1.	Pure phosphoric acid is very viscous because (1) it is a strong acid (2) it is tribasi (4*) it has PO ₄ ³⁻ groups which are bonded by r	ic acid (3) it	is hygroscopic	
K-2.	Which of the following is least volatile ? (1*) HF (2) HCl	(3) HBr	(4) HI	
K-3.	Which one of the following does not have inter (1) H ₂ O (2*) o-nitro phenol	rmolecular H-bonding ? (3) HF	(4) CH₃COOH	
K-4.	Which of the following exhibits H-bonding ? (1) CH_4 (2) H_2Se	(3*) N ₂ H ₄	(4) H ₂ S	
K-5.	H–bonding is not present in : (1) NH ₃ (2) H ₂ O	(3*) H ₂ S	(4) HF	
K-6.	Hydrogen bonding would not affect the boiling (1*) HI (2) NH ₃	point of : (3) CH₃OH	(4) H ₂ O	
K-7.	Which of the following compound has maximu (1) HF (2) PH₃	m number of H-bonds pe (3*) H₂O	er mole ? (4) OF ₂	
K-8.	Water (H ₂ O) is liquid while hydrogen sulphide (H ₂ S) is a gas because : (1) water has higher molecular weight (2) hydrogen sulphide is weak acid (3*) water molecular associate through hydrogen bonding (4) sulphur has high electronegativity than oxygen			
Sect	ion (L) : Intermolecular forces			
L-1.	Which of the following has the highest boiling (1) H ₂ (2) Ne	point? (3*) Xe	(4) CH ₄	
L-2.	Among the following molecules the dipole-dipole the melting and boiling point is : (1) ICI (2*) Br ₂	le attraction does not play (3) H ₂ S	y most important role in determing (4) CO	
Page				

L-3.	At ordinary temperature and pressure, among halogens chlorine is a gas, bromine is a Liquid and iodine is a solid. This is because : (1) The specific heat is in the order $Cl_2 > Br_2 > l_2$ (2*) Intermolecular forces among molecule of chlorine are the weakest and those in iodine the strongest (3) The order of density is $l_2 > Br_2 > Cl_2$ (4) The order of stability is $l_2 > Br_2 > Cl_2$				
L-4.	Which of the following (1) metallic bonding (3*) non-metallic coval		ne bonding within a layer (2) ionic bonding (4) v	r of the graphite structure ?	
L-5.	Among the following, v (1) HBr	v are maximum in : (2) LiBr	(3) LiCl	(4*) AgBr	
L-6.	Which of the following (1) Covalent bond	bonds/forces is weakest (2) Ionic bond	? (3) Metallic bond	(4*) London force	
L-7.	In which molecule is th (1) H ₂ O	e London dispersion forc (2*) Br ₂	e likely to be most impor (3) SO ₂	tant in determining boiling point? (4) CIF ₃	
Secti	Section (M) : Metallic bond				
M-1	Iron is harder than soc (1) iron atoms are sma (3) metallic bonds are	aller.	(2) iron atoms are mor (4*) metallic bonds are		

- M-2._ The enhanced force of cohesion in metals is due to :
 - (1) The covalent linkages between atoms
 - (2) The electrovalent linkages between atoms
 - (3) The lack of exchange of valency electrons
 - (4*) The delocalization of valence electron between metallic kernels.

Exercise-2

Marked Questions may have for Revision Questions.

OBJECTIVE QUESTIONS

- 1. When two atoms combine to form a molecule :
 - (1*) energy is released
 - (3) energy is neither released nor absorbed

(2) energy is obserbed

- (4) energy may either released or absorbed
- 2. Which condition favours the bond formation ?
 - (1) Maximum attraction and maximum potential energy
 - (2) Minimum attraction and minimum potential energy
 - (3*) Minimum potential energy and maximum attraction
 - (4) None of the above
- **3.** Most favourable conditions for electrovalent bonding are :
 - (1*) low ionisation potential of one atom and high electron affinity of the other atom(2) high electron affinity and high ionisation potential of both the atoms

•				
		y and low ionisation pote ential of one atom and lo	ntial of both the atoms w electron affinity of the	outer atom.
4.	Which one of the follo (1*) S-F in SF2	wing bonds is least cova (2) S-F in SF₄	lent ? (3) S-F in SF₀	(4) S-S in S₅
5.	The correct order of ir (1) SiCl₄ > AlCl₃ < CaC (3) AlCl₃ < CaCl₂ < KC		cter of the following is : (2*) KCl < CaCl₂ < AlC (4) None of these	Cl₃ < SiCl₄
6.		and most covalent comp PbO, P4O10, SnO2 (2*) PbO, Mn2O7	ounds respectively from t (3) CrO₅, P₄O	-
7.		ing species is the octet r (2) SF ₆		(4*) CO ₂
8.	. ,	efers to cation. s to anion. re efficient to polarise an	iion. inert gas configuration ar	e more Ionic.
9.	In which of the followin (1) Rb	ng will have highest hydr (2) Cs	ation ? (3*) Na	(4) K
10.	If a molecule X₂ has a (1) 1s₂ 2s₂ 2p₅	triple bond, then X will h (2*) 1s ₂ 2s ₂ 2p ₃	ave the electronic config (3) 1s ₂ 2s ₁	uration : (4) 1s ₂ 2s ₂ 2p ₁
11.	Which of the following (1) BH₄⁻	does not contain coordi (2) NH4+	nate bond ? (3*) CO ₃ 2-	(4) H₃O⁺
12.	The species which is (1) NO	diamagnetic is : (2) NO ₂	(3) CIO ₂	(4*) N2O4
13.	N ₂ O is isoelectronic w	ith CO₂ and N₃⁻, which is	the structure of N ₂ O ?	
	(1) N N	(2) N–O–N	(3) N O	(4*) N ≡ N → Ö:
14.		H	ectively in the given struc B – H N – H (3) – 2, + 2	ture? (4) 0, 0

15. What is the formal charge on N in $(1^*) + 1$ (2) - 1 (3) 0 (4) + 2

16.	In PO₄ ^{3–} ion, the form (1*) – 0.75, 1.25	al charge on each oxyge (2) –0.75, 1.0		order respectively are [DMT 2004] 6 (4) – 3, 1.25
17.		•	onating structure will be = N ²⁻ ←→→ H – N ⁻ – N (III)	
	(1) I	(2*) I I	(3) I I F	(4) Both (I) and (III)
18.	The correct order of (1) C−C < C=C < C≡	-	(3) C=C <c≡c<c−c< td=""><td>(4) C=C<c−c<c≡c< td=""></c−c<c≡c<></td></c≡c<c−c<>	(4) C=C <c−c<c≡c< td=""></c−c<c≡c<>
19.	Which of the followin (a) 2 py + 2 py $\rightarrow \pi 2$		assuming z-axis to be the (b) 2 pz + 2 p	
	(c) $2 px + 2 px \rightarrow \pi 2$		(d) 1 s + 2 py $\rightarrow \pi$ (1	•••
	(1) 'a' & 'b'	(2) 'b' & 'd'	(3*) only 'd'	(4) None of these
20.	(1) π-bond is formed (2) Generally π-bond (3) π- bond may be f	g statements is correct a when a sigma bond is no are formed from hybrid o ormed by the overlapping om lateral overlap of p-at	ot formed. orbitals. g of s-atomic orbitals.	
21.	p-p overlapping will t (1) H₂	e observed in the molect (2) HBr	ule of : (3) HCl	(4*) Cl ₂
22.	Identify the correct m (i) XeF₂ (ii) N₃ ⁻ (iii) PCI₅ (s) anion (iv) I₂CI₅ (ℓ) cation (1) (i – a), (ii – b), (iii (3) (i – b), (ii – c), (iii	(a) Central at (b) Central at (c) Central at (d) Central at − c), (iv – d)		on and octahedral. and linear geometry.
23.	In a change from PC (1) sp² to sp³	I ₃ → PCI₅, the hybrid state (2) sp ³ to sp ²	e of P changes from : (3*) sp³ to sp³d	(4) sp ³ to dsp ²
24.	The type of hybrid or (1) sp, sp², sp³ and s	•	om in CIO⁻, CIO₂⁻, CIO₃⁻ p and sp³ (3*) c	and ClO₄⁻ is / are : only sp³ (4) only sp
25.	Correct statement re	garding this reaction :		
	BF₃ + NH₃ (1) hybridisation of o (3) hybridisation of N		(2*) hybridisation of o (4) none of these.	nly B changes
26.	The hybridisation of ((1*) sp , sp² , sp	central atoms in N₃⁻, NOC (2) sp , sp , sp³	Cl and N₂O are respective (3) sp² , sp , sp	ely : (4) sp² , sp² , sp
27.	The shape of O_2F_2 is (1) C_2F_2	similar to : (2*) H₂O₂	(3) H ₂ F ₂	(4) C ₂ H ₂
20	Which of the following	a malagulas is triassal hi	nuromido!?	
<u> 2ŏ.</u>		<u>g molecules is trigonal bi</u>	pyramidar?	

CHEMISTRY FOR JEE CHEMICAL BONDING (1) BF₃ (2) CH₄ (3*) PCI5 (4) SF₆ 29. The structure of $IC\ell_2^+$ would be : C C (1) (3) (2)(4*) None of these 30. The structure of IO₂F₂⁻ is analogous to : (1) SF₄ (2) XeO_2F_2 (3) F₂SeO₂ (4*) (1) and (2) both 31. The bond angle in H_2O is nearly 105° whereas bond angle in H_2S is nearly 92°. This is because : (1*) Electronegativity of oxygen is greater than that of sulphur (2) Oxygen is a gas whereas sulphur is solid (3) Sulphur contains d-orbitals whereas oxygen does not (4) The number of lone pairs present on oxygen and sulphur is not equal 32. In XeF₂, XeF₄ and XeF₆ (g) the number of lone pairs on Xe respectively are : (1) 2, 3, 1 (3) 4, 1, 2 (2) 1, 2, 3 (4*) 3, 2, 1 33. Incorrect order about bond angle is/are : (1) $H_2O > H_2S > H_2Se > H_2Te$ (2) $C_2H_2 > C_2H_4 > CH_4 > NH_3$ (4) $CIO_2 > H_2O > H_2S > SF_6$ (3^*) SF₆ < NH₃ < H₂O < OF₂

- 34.
 The ONO angle is maximum in :
 (1) HNO3
 (2*) NO2+
 (3) HNO2
 (4) NO2
- **35.**Which of the following is electron deficient compounds ?
(1) NaBH4(2*) B_2H_6 (3) AI_2CI_6 (4) LiAIH4
- **36.** Which of the following statement is incorrect ?
 - (1) During N_{2^+} formation, one electron is removed from the bonding molecular orbital of N_2 .
 - (2) During O_{2^+} formation, one electron is removed from the antibonding molecular orbital of O_{2^-}
 - (3*) During $O_{2^{-}}$ formation, one electron is added to the bonding molecular orbitals of O_{2} .
 - (4) During CN^{-} formation, one electron is added to the bonding molecular orbitals of CN.
- **37.** According to Molecular orbital theory which of the following is correct ?

(1) LUMO level for C ₂ molecule is σ_{2p_x} orbital	(2) In C_2 molecules both the bonds are π bonds
(3) In $C_{2^{2-}}$ ion there is one σ and two π bonds	(4*) All the above are correct

- **38.** Which of the following species will have the minimum bond energy? (1) N_2 (2) N_2^- (3) N_2^+ (4*) N_2^{2-}
- **39.** The sequence that correctly describes the relative bond strengths partaining to oxygen molecule and its cation or anion is
- 40. Which of the following statements is correct about N₂ molecule ?
 (1) It has a bond order of 3
 (2) The number of unpaired electrons present in it is zero and hence it is diamagnetic
 - (3) The order of filling of MOs is $\pi(2px) = \pi(2py)$, $\sigma(2pz)$

	(4*) All the above the	nree statements are correc	t	
41.	The bond order of (1) 3/2	HeH+ is : (2*) 1	(3) 1/2	(4) 2
42.	(1) The dipole mor	ing statements is true? nent of NF₃ is zero nent of NF₃ is more than NH	· · ·	nent of NF₃ is less than NH₃ ent of NH₃ is zero
43.		ing orders are correct abou I₂Cl₀ > B₂H₀ (iii) Me₃N-BF₃ (2) (i), (ii) and (iv) only	•	(iv) SiH₃OCH₃ > (SiH₃)₂O
44.	Which of the follow (1) SOCI2	ing would be expected to h (2) OF ₂	ave a dipole moment o (3*) SeF₀	of zero on the basis of symmetry? (4) CIF₅
45.	The dipole momen ionic character in H (1) 10.5		and the interatomic spa	acing is 1.41 Å. The percentage o (4) 13.5
46.	. ,	true ? hyl ether and C₂H₅OH is ec highly soluble in water.	qual. (2*) Diethyl (4) Diethyl ether is L	[RPMT 2002] ether have dipole moment. _ewis acid.
47.	H₂O boils at higher (1) ionic bonds	temperature than H ₂ S, bec (2) covalent bonds	ause it is capable of fo (3*) hydrogen bonds	-
48.	Weakest hydrogen (1) O–H N	bond is : (2*) N – H N	(3) F – H F	(4) All are equally strong
49.	Which one among (1) HSO₅⁻	the following does not have (2) C₅H₅OH	e the hydrogen bond ? (3) H₂SO₄	(4*) H ₂ S
50.	The correct order o	f the viscosity in the followi	CH₂OH	state is :
	CH₃OH(ℓ) (I)	CH₂OH │ CH₂OH(ℓ) (II)	СНОН СН ₂ ОН (ℓ) (III)	
	(1) > >	(2*) > >	(3) > >	(4) > >
51.	(1) NO₂ group at p-(2) intramolecular h(3*) there is intermolecular	p-nitrophenol is higher tha position behaves in a differ hydrogen bonding exists in plecular hydrogen bonding as a higher molecular weig	rent way from that at o- p-nitrophenol in p-nitrophenol	
52.	•	s than that of water becaus	se of :	

- (1) presence of Van der Wall interaction.
- (2) crystal modification of ice.
- (3*) open porous cage like structure of ice due to H-bonding.
- (4) different physical states of these.

53.	NH ₃ has a much higher boiling point than PH ₃ because : (1*) NH ₃ forms hydrogen bond. (2) NH ₃ has a larger molecular weight. (3) NH ₃ undergoes umbrella inversion. (4) NH ₃ contains ionic bonds where are PH ₃ contains covalent bond.					
54.	Which of the	following		unds would have H₃OH, N₂O₄, CH	significant intermolecul	ar hydrogen bonding ?
	(1) HF, N ₂ O ₄		(2) HI	F, CH4, CH3OH	(3*) HF, CH₃OH	(4) CH ₃ OH, CH ₄
55.	Which of the t (1) persulpha		-	ot have intramole loral hydrate	cular hydrogen bond ? (3) o-nitrophenol	(4*) fumeric acid
56.	(i) N⊦ fuEu gkbMªkbM	I₃ , (ii) P [ds]]oFk I₃ , (ii) P > (i) > (ii	H₃, (iii) A ukad dk v H₃, (iii) A ii) > (ii)	oiling points of th sH₃ (iv) SbH₃ , (v qjksgh Øe fuEu gSA sH₃ (iv) SbH₃ , (v	l	
57. 58.	 The boiling point of ICI is nearly 40°C higher than that of Br₂ although the two substance have the same relative molecular mass. This is because : (1) ICI is ionic compound . (2) I–CI bond is stronger than Br–Br bond. (3*) ICI is polar covalent molecule while Br₂ is non polar. (4) Ionization energy of iodine is less than that of Br. The nature of intermolecular forces among benzene (C₆H₆) molecules is : (1) dipole dipole attraction (2*) Iondon dispersion forces (3) ion dipole attraction (4) hydrogen bonding 					
59.	 Which of the following is false ? (1) V are responsible for the formation of molecular crystals. (2) Branching lowers the boiling points of isomeric organic compounds due to reduction in the van der Waals force of attraction. (3) In graphite, v act between the carbon layers. (4*) Boiling point of NH₃ is greater than SbH₃. 					
60.	Match the spectrum Column-I (A) $IO_2F_2^-$ (B) F_2SeO (C) SO_2 (D) XeF_5^+ A (1*) p (2) p (3) p (4) q	ecies giv B q q s p	ven in col C r s r r r	umn-I with the ty Column-II (p) sp ³ d (q) sp ³ (r) sp ² (s) sp ³ d ² D s r p s	rpe of hybridisation give	n in column-II.

61. Assertion : NF₃ has little tendency to act as a donor molecule.

Reason : The highly electronegative F atoms attract electrons and these moments partly cancel the moment from the lone pair.

(1*) Both assertion and reason are correct, and the reason is the correct explanation for the assertion

- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect

62. Assertion : Molecules having different hybridisation can have same shape.
 Reason : The shape of a molecule does not depend on the hybridisation but it depends on the energy factor

- (1*) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect
- $\begin{array}{lll} \textbf{63.} & \textbf{Assertion:} & \text{Aluminium chloride in acidified aqueous solution forms octahedral } [Al(H_2O)_6]_{3^+} \text{ ion.} \\ & \textbf{Reason:} & \ln [Al(H_2O)_6]^{3_+} \text{ complex ion, the 3d orbitals of Al are involved and the hybridisation state of Al is sp^3d^2.} \end{array}$
 - (1*) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
 - (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
 - (3) The assertion is incorrect, but the reason is correct
 - (4) Both are assertion and reason are incorrect
- **64.** Assertion : NO₃⁻ and PO₃⁻ have similar formula type but differ structurally i.e. they have different type of hybridisation.

Reason : NO₃⁻ ion exists as free ion. On the other hand PO₃⁻ exists as cyclic or linear polymeric structure.

- (1*) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect
- **65.** Assertion : The increasing order of acidic character of CO_2 , N_2O_5 , SiO_2 and SO_3 is $SO_3 > N_2O_5 > CO_2 > SiO_2$

Reason : As electronegativity difference (E–O) decreases acidic character of the oxide increases.

- (1) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3*) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect
- **66. Assertion :** In graphite, on increasing the temperature the conductivity decreases along the layers of carbon atoms.

Reason : Graphite cleaves easily, because the force of attraction between the layers is weak van der Waal's force.

- (1) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2*) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect
- **67.** Assertion : Molecular species like SF₆, PF₅ , I_{3}^{-} and XeF₂ violate the octet rule.

Reason : Compounds with an expanded octet are called hypervalent compounds.

- (1) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2*) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect
- **68.** Assertion : π is stronger while σ is a weak bond.

[AIEEE-2003, 3/225]

Reason : Atoms rotate freely about π bond.

- (1) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4*) Both are assertion and reason are incorrect
- **69.** Assertion : Carbon has unique ability to form pπ–pπ multiple bonds with itself and with other atoms of small size and high electronegativity.

Reason : Heavier elements of group 14th do not form $p\pi$ – $p\pi$ multiple bonds with itself because their atomic orbitals are too large and diffuse to have effective overlapping.

- (1) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2*) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect
- **70.** Assertion : Bond energy has order like C-C < C=C < C=C.

Reason : Bond energy increase with increase in bond order.

- (1*) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect
- 71. Assertion : Nitrogen molecule is paramagnetic.

Reason : N₂ molecule have unpaired electron.

- (1) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4*) Both are assertion and reason are incorrect
- **72.** Assertion : $[SiC\ell_6]^{2-}$ does not exist.

Reason : Interaction between lone pairs of chloride ions and Si⁴⁺ is not very strong and six large chloride ions can not be accommodated around Si⁴⁺ due to limitation of its size.

- (1*) Both assertion and reason are correct, and the reason is the correct explanation for the assertion
- (2) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion
- (3) The assertion is incorrect, but the reason is correct
- (4) Both are assertion and reason are incorrect

Exercise-3

PART - I : JEE (MAIN) / AIEEE PROBLEMS (PREVIOUS YEARS)

1.	Which of the following c	ompounds has the small	lest bond angle in its mo	lecule ? [AIEEE-2003, 3/225]
	[AIEEE-2003, 3/225]			
	(1) SO ₂	(2) H ₂ O	(3*) H ₂ S	(4) NH₃

2. An ether is more volatile than an alcohol having the same molecular formula. This is due to:

- (1) dipolar character of ethers. (2) alcohols having resonance structures.
- (3) inter-molecular hydrogen bonding in ethers. (4*) inter-molecular hydrogen bonding in alcohols.
- **3.** The pair of species having identical shapes for molecules of both species is : [AIEEE-2003, 3/225] (1) CF₄, SF₄ (2*) XeF₂, CO₂ (3) BF₃, PCI₃ (4) PF₅, IF₅.

4.	Which of the following pair of molecules will have permanent dipole moments for both members?					
	(1) SiF ₄ and NO ₂	(2) NO ₂ and CO ₂	(3^*) NO ₂ and C) 3	[AIEEE-2003, 3/2 (4) SiF₄ and	-
5.	The correct order of b (1) H ₂ S < SiH ₄ < NH ₃ < (3*) H ₂ S < NH ₃ < SiH ₄) in H₂S, NH₃, BF₃ and S (2) NH₃ < H₂S < SiH₄ < (4) H₂S < NH₃ < BF₃ <	: BF₃	[AIEEE-2004, 3/2	25]
6.	species? (1) bond length in NO ⁻	is 2.5 while that in NO ⁺ is ⁺ is greater than in NO ⁺ is equal to that in NO	3. Which of the following (2*) bond length in NO (4) bond length is unp	is greate	[AIEEE-2004, 3/2 er than in NO ⁺	
7.	The states of hybridiza	ation of boron and oxyger	n atoms in boric acid (H₃l	BO₃) are	respectively : [AIEEE-2004, 3/2	251
	(1*) sp ² and sp ²	(2) sp ² and sp ³	(3) sp^3 and sp^2	(4) sp ³	³ and sp ³	.23]
8.	(1) XeF4	wing has the regular tetra (2) SF₄ 5, S = 16, Ni = 28, Xe = 5	(3*) BF ₄ -		[AIEEE-2004, 3/2 (4) [Ni(CN) ₄] ^{2–}	25]
9.	Which one of the follow (1) acetone	wing does not have sp² h (2) acetic acid	•	е	[AIEEE-2004, 3/ (4) acetamide	/225]
10.	Lattice energy of an ic	nic compound depends u	ipon :		[AIEEE-2005, 3/	/225]
	(1) charge on the ions(3) packing of ions onl	•	(2) size of the ions only (4*) charge on the ion		of the ion	
11.		gy and other considerati highest melting point ? (2*) NaCl	ons, which one of the fo	ollowing (4) Rb	[AIEEE-2005, 3/2	
12.	(1) The same with 2, 0(2) The same with 1, 1(3) Different with 0, 1	s of SF4, CF4 and XeF4 and and 1 lone pairs of elect and 1 lone pair of electr and 2 lone pairs of electro and 2 lone pairs of electro	trons on the central atom ons on the central atoms ons on the central atom,	, respectiv	tively. vely.	25]
13.	Which one of the follow (1) He2+	wing species is diamagne (2*) H ₂	etic in nature ? (3) H₂⁺	[AIEEI (4) H ₂ -	E-2005, 1½/225] ⁻.	
14.	The number and type (1) one sigma, one pi	of bonds between two ca (2*) one sigma, two pi	rbon atoms in calcium ca (3) two sigma, one pi		re : [AIEEE-2005, 3/ o sigma, two pi	225]
15.	Which of the following (1*) O22-	molecules/ions does not (2) B ₂	contain unpaired electro (3) N ₂ +	ons?	[AIEEE-2006, 3/1 (4) O ₂	65]
16.	A metal, M forms chlor chlorides is correct ? (1) MCl ₂ is more volati (3*) MCl ₂ is more ionic	. ,	tion states. Which of the t Cl₂ is more soluble in anh Cl₂ is more easily hydroly	[AIEEI	E-2006, 3/165] ethanol than MCl₄	hese

17.	In which of the followin (1*) SF₄	g molecules/ions are all (2) SiF₄	the bonds not equal? (3) XeF₄	[AIEEE-2006, 3/165] (4) BF₄⁻
18.	Which of the following I (1) O – H O	hydrogen bonds is the st (2) O – H F	rongest ? (3) F – H H	[AIEEE-2007, 3/120] (4*) F – H F
19.	•			h one of the following sequences hic species, K ⁺ , Ca ²⁺ , Mg ²⁺ , Be ²⁺ ? [AIEEE-2007, 3/120]
	(1*) K+ < Ca ²⁺ < Mg ²⁺ < (3) Mg ²⁺ < Be ²⁺ < K+ <		(2) Ca ² < Mg ²⁺ < Be ²⁺ (4) Be ²⁺ < K ⁺ < Ca ²⁺ <	< K+
20.	In which of the following has changed ?	g ionization processes, t		ased and the magnetic behaviour EE-2007, 3/120]
	(1) $O_2 \longrightarrow O_2^+$	(2) $N_2 \longrightarrow N_2^+$	$(3) C_2 \longrightarrow C_2^+$	$(4^*) \text{ NO} \longrightarrow \text{NO}^+$
21.	Which one of the follow (1) CN⁻ and CN⁺	/ing pairs of species hav (2) O₂⁻ and CN⁻	e the same bond order? (3) NO ⁺ and CN ⁺	(4*) CN ⁻ and NO ⁺
22.		•••		hat of C – F in CF ₄ is 515 kJ mol- pared to that of C – F is : [AIEEE-2009, 4/144]
	 (2*) significant pπ - pr interaction between C a (3) lower degree of pπ 	and F in CF4.	and F in BF₃ wherea B and F in BF₃ than that	een C and F in CF4. s there is no possibility of such at between C and F in CF4.
23.	Using MO theory predic	ct which of the following	species has the shortes	t bond length ? [AIEEE-2009, 4/144] [AIEEE-2009, 4/144]
	(1) O ₂ +	(2) O ₂ -	(3) O ₂ ^{2–}	(4*) O ₂ ²⁺
24.	Among the following th	e maximum covalent cha	aracter is shown by the	compound : [AIEEE-2011, 4/120]
	(1) FeCl ₂	(2) SnCl ₂	(3*) AICI3	(4) MgCl2
25.	The hybridisation of orb (1) sp, sp2, sp3	bitals of N atom in NO _{3–} , (2*) sp2, sp, sp3	NO ₂₊ and NH ₄₊ are res (3) sp, sp ₃ , sp	pectively : [AIEEE-2011, 4/120] (4) sp2, sp3, sp
26.	The structure of IF7 is : (1) square pyramid	(2) trigonal bipyramid	(3) octahedral	[AIEEE-2011, 4/120] (4*) pentagonal bipyramid
27.	The number of types of (1) One sigma, one pi (3) Two sigma, two pi	f bonds between two car	bon atoms in calcium ca (2) Two sigma, one pi (4*) One sigma, two p	
28.	Which of the following I (1) XeF₄	has maximum number o (2) XeF₅	f lone pairs associated ((3*) X	with Xe ? [AIEEE-2011, 4/120] XeF ₂ (4) XeO ₃
29.	The molecule having s	mallest bond angle is :		[AIEEE-2012, 4/120]

	(1) NCl₃	(2) AsCl₃	(3*) SbCl₃	(4) PCl₃
0.		owing pairs the two speci	es are not isostructural ?	[AIEEE-2012, 4/120 (4) AIF ₆ ^{3–} and SF ₆
1.	Which one of the fo	ollowing molecules is exp	ected to exhibit diamagne	etic behaviour?
	(1*) C2	(2*) N2	(3) O ₂	[JEE(Main) 2013, 4/120] (4) S ₂
2.	In which of the follo	owing pairs of molecules/	ions, both the species are	not likely to exist ? [JEE(Main) 2013, 4/120]
	(1) H_2^+, He_2^{2-}	(2) H_2^- , He_2^{2-}	(3*) H_2^{2+} , He2	(4) H_2^- , He_2^{2+}
3.	Which of the follow (1) lodine	ring exists as covalent cry (2*) Silicon	vstals in the solid state ? (3) Sulphur	[JEE(Main) 2013, 4/120] (4) Phosphorus
4.	Stability of the spe	cies Li ₂ , Li_2^- and Li_2^+ incr	eases in the order of :	[JEE(Main) 2013, 4/120]
5.	Among the followir	ng oxoacids, the correct d	lecreasing order of acid st	-
	(1) HOCI > HCIO2 (3*) HCIO4 > HCIO		(2) HCIO4 > HOCI > (4) HCIO2 > HCIO4 >	
6.	The correct statem	ent for the molecule, CsI	з is : [ч	JEE(Main)-2014, 4/120]
	(1) it is a covalent molecule. (2^*) it contains (2*) it contain			and I_3^- and lattice I ₂ molecule.
7.	The intermolecular is:	interaction that is depen	dent on the inverse cube	of distance between the molecule
	(1) ion-ion interacti(3) London force	on	(2) ion-dipole interac (4*) hydrogen bond	tion [JEE(Main)-2015, 4/12
8.	Which one has the (1) He	highest boiling point ? (2) Ne	(3) Kr	[JEE(Main)-2015, 4/120] (4*) Xe
9.	The species in whi	ch the N atom is in a stat	e of sp hybridization is :	[JEE(Main)-2016, 4/120]
	(1) ^{NO} ₂ ⁻	(2) ^{NO} ₃ ⁻	(3) NO ₂	(4*) ^{NO} ⁺ ₂
0.	 (1) Water can act b (2*) There is exten (3) Ice formed by h 	ollowing statements abou both as an acid and as a l sive intramolecular hydro neavy water sinks in norm ed to oxygen during photo	base. Igen bonding in the conde Ial water.	[JEE(Main)-2016, 4/120] nsed phase.
1.	In the following rea (a) ZnO + Na₂O – - (b) ZnO + CO₂ – →		ly acting as a/an :	[JEE(Main)-2017, 4/12
	(U) ZIIU + UU2 - →			
			agnetic?	

1.	Which of the following has u_1 (1) N_2 (2*)	•	[JEE(Main) 20′ (3) N₂²+	14 Online (09-04-14), 4/120] (4) O2 ²⁻
2.	The number and type of bon (1) One σ -bond and one π -b (3) Two σ -bonds and two π -l	ond (2	are : [JEE(Main) 20 2*) One σ-bond and tw 4) Two σ-bonds and or	
3.	arrangements?		[JEE(Main) 2	hown in which of the following 2014 Online (11-04-14), 4/120] (4) O ₂ > O ₂ ⁻ > N ₂
4.	Which one of the following d	oes not have a pyrami		
	(1) (CH ₃) ₃ N (2*)	(SiH₃)₃N (3	נ ונדב(Main) 3) P(CH₃)₃	2014 Online (11-04-14), 4/120] (4) P(SiH₃)₃
5.	Which one of the following m $(1) N_2$ (2^*)		etic? [JEE(Main) 20 3) CO	14 Online (19-04-14), 4/120] (4) O₃
6.	respectively are :		•	st and the least ionic character 014 Online (19-04-14), 4/120] (4) RbCl and MgCl₂
7.	•	nolecular orbital (MO) o	of H ₂ , electron density i ws destructive interfer	s increased between the nuclei. ence of the combining electron 15 Online (10-04-15), 4/120]
	 (1*) Assertion is correct, reas (2) Assertion is incorrect, reas (3) Assertion and reason are (4) Assertion and reason are the assertion. 	ason is correct. e correct, but reason is	not the correct explan	
8.	The geometry of XeOF₄ by b (1) pentagonal planar (2) o	• •		015 Online (10-04-15), 4/120] (4) trigonal bipyramidal
9.	Which of the alkaline earth n	netal halides given belo	-	lent in nature ? 15 Online (11-04-15), 4/120]
	(1) SrCl ₂ (2) (CaCl ₂ (3	B*) BaCl ₂	(4) MgCl ₂
10.	Which of the following composite (1) $H_4P_2O_5$ (2) (? [JEE(Main) 20 8*) H ₄ P ₂ O ₆	15 Online (11-04-15), 4/120] (4) H ₄ P ₂ O ₇
11.	Choose the incorrect formula (1) X ₂ O ₃	a out of the four compo (2*) X ₂ Cl ₃	ounds for an element > (3) X ₂ (SO ₄) ₃	K below : (4) XPO₄
12.	Molecular AB has a bond len atom (absolute magnitude) is 4/120]	•		0. The fractional charge on each Iain) 2015 Online (11-04-15),
	-	0.05 (3	3) 0	(4) 1.0
13.	The group of molecules havi	ng identical shape is :	[JEE(Main) 2016	Online (09-04-16), 4/120]

	(1) PCI ₅ , IF ₅ , XeO ₂ F ₂ (2) BF ₃ , PCI ₃ , XeO ₃ (3*) CIF ₃ , XeOF ₂ , ^{XeF₃+} (4) SF ₄ , XeF ₄ , CCI ₄
14.	Which intermolecular force is most responsible in allowing xenon gas to liquefy? [JEE(Main) 2016 Online (09-04-16), 4/120]
	(1*) Instantaneous dipole-induced dipole (2) Ionic
	(3) Ion-dipole (4) Dipole-dipole
15.	Assertion: Among the carbon allotropes, diamond is an insulator, Whereas, graphite is a good conductor of electricity.
	Reason: Hybridization of carbon in diamond and graphite are sp ³ and sp ² , respectively.
	[JEE(Main) 2016 Online (10-04-16), 4/120]
	(1*) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion.
	(2) Assertion is incorrect statement, but the reason is correct.
	(3) Both assertion and reason are correct, and the reason is the correct explanation for the assertion.(4) Both assertion and reason are incorrect.
	(1*) Both assertion and reason are correct, but the reason is not the correct explanation for the assertion.
	(2) Assertion is incorrect statement, but the reason is correct.
	(3) Both assertion and reason are correct, and the reason is the correct explanation for the assertion.(4) Both assertion and reason are incorrect.
16.	The bond angle H–X–H is the greatest in the compound : [JEE(Main) 2016 Online (10-04-16), 4/120]

16.	The bond angle H–X–F	I is the greatest in the co	mpound : [JEE(Ma	ain) 2016 Online (10-04-16), 4/120]
	;kSfxd ftlesa lsrq cU/k H–j	X−H cU/k vf/kdre gS ====	[JEE(Ma	in) 2016 Online (10-04-16), 4/120]
	(1) NH₃	(2) PH₃	(3*) CH4	(4) H ₂ O

PART - II : JEE (ADVANCED) / IIT-JEE PROBLEMS (PREVIOUS YEARS)

* Marked Questions may have more than one correct option.							
1.	Which of the following a NO_{3}^{-} , CO_{3}^{2-} , C	[JEE-2003, 3/84]					
	(A*) NO ₃ - , CO ₃ 2-	(B) SO₃ , NO₃⁻	(C) CIO ₃ ⁻ , CO ₃ ²⁻	(D) CO:	3 ^{2–} , SO ₃ .		
2.	Among the following the (A*) CH₃CI	e molecule with the highe (B) CH ₃ Cl ₂	est dipole moment is : (C) CHCl₃	(D) CCI	[JEE–2003, 3/84] ₄		
3.	Which of the following represent the given mode of hybridisation sp ² – sp ² – sp – sp from left to rig [JEE-2003, 3/8 CH						
	(A*) H₂C=CH–C≡N	(B) HC≡C–C≡CH	(C) H ₂ C=C=C=CH ₂	(D) H ₂ N	CH ₂		
4.	The number of lone pair(s) of electrons in XeOF ₄ is : [JEE-2004						
	(A) 3	(B) 2	(C*) 1	(D) 4			
5.	Amongst the following t (A) H ₂ S ₂ O ₃	he acid having –O–O– b (B) H₂ S₂ O₅		(D*) H2	[JEE–2004, 3/84] S ₂ O ₈		

CHEMICAL BONDING

6.	According to molecular orbital theory, which one of the following statements about the molecular species O ₂ ⁺ is correct ? [JEE-2004, 3/84] (A) It is paramagnetic and has less bond order than O ₂				
	(B*) It is paramagneti(C) It is diamagnetic a	and has less bond ord c and more bond order and has less bond orde and has more bond ord	than O2 r than O2		
7.	In which of the followi	ng the maximum numb	er of lone pairs is present	on the central atom ? [JEE–2005, 3/84]	
	(A) [ClO₃] [_]	(B) XeF4	(C) SF4	(D*) I₃⁻	
8.	Which of the following ?	silicate is formed wher	n three oxygen atoms of [Si	iO₄] ^{4–} tetrahedral units are shared [JEE–2005, 3/84]	
	(A*) Sheet silicate (C) Three dimensiona	Il silicate	(B) Pyrosilicate (D) Linear chain silica		
9.	The species having b (A*) NO⁻	ond order different from (B) NO ⁺	n that in CO is : (C) CN⁻	[JEE–2007, 3/162] (D) N ₂	
10.	Among the following, (A) Na ₂ O ₂	the paramagnetic com (B) O₃	pound is : (C) N₂O	[JEE–2007, 3/162] (D*) KO ₂	
11.	The percentage of p- (A) 25	character in the orbitals (B) 33	forming P – P bonds in P₄ (C) 50	is : [JEE–2007, 3/162] (D*) 75	
12.	 Statement-1 : Boron always forms covalent bond, because Statement-2 : The small size of B³⁺ favours formation of covalent bond. [JEE-2007, 3/162] (A*) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1. (B) Statement-1 is True, Statement-2 is True; Statement-2 is NOT a correct explanation for Statement-1. (C) Statement-1 is True, Statement-2 is True. 				
13.		gap in germanium is sm nergy spread of each ge	nall, because ermanium atomic energy le	evel is infinitesimally small. [JEE–2007, 3/162]	
	(B) Statement-1 is Tru (C*) Statement-1 is T		; Statement-2 is NOT a co se	explanation for Statement-1. rrect explanation for Statement-1	
14.	is:		-	ature of the diatomic molecule B2	
	(A*) 1 and diamagnet(C) 1 and paramagne		(B) 0 and diamagnetic(D) 0 and paramagnet		
15.	The shape of XeO ₂ F ₂ (A) trigonal bipyramid (C) tetrahedral		(B) square plannar (D*) see-saw	[JEE–2012, 3/136]	
16.	Assuming 2s-2p mixir	ng is NOT operative, th	e paramagnetic species ar		
	(A) Be ₂	(B) B ₂	(C*) C ₂	[JEE(Advanced) 2014, 3/120] (D) N ₂	
17.	Match the orbital over answer using the cod			en in List-II and select the correct	

	List-I					List-II
P.		Þ			1.	<i>p–d</i> π antibonding
Q.	8X	1			2.	<i>d–d</i> σ bonding
R.	¥X				3.	$p-d\pi$ bonding
S.		Þ			4.	<i>d–d</i> σ antibonding
Code :		Р	Q	R	S	
	(A)	2	1	3	4	
	(B)	4	3	1	2	
	(C)	2	3	1	4	
	(D)	4	1	3	2	

Additional Problems For Self Practice (APSP)

PART - I : PRACTICE TEST PAPER

This Section is not meant for classroom discussion. It is being given to promote self-study and self testing amongst the Resonance students.

Max. Marks : 120

Important Instructions

- **1.** The test is of **1 hour** duration.
- 2. The Test Booklet consists of 30 questions. The maximum marks are 120.
- 3. Each question is allotted 4 (four) marks for correct response.
- 4. Candidates will be awarded marks as stated above in Instructions No. 3 for correct response of each question. ¼ (one fourth) marks will be deducted for indicating incorrect response of each question. No deduction from the total score will be made if no response is indicated for an item in the answer sheet.
- There is only one correct response for each question. Filling up more than one response in any question will be treated as wrong response and marks for wrong response will be deducted accordingly as per instructions 4 above.

1.	If the electronic configuration of an element is $ls^2 2s^2 2p^6 3s^2 3p^6 3d^2 4s^2$, then the electrons involved in bonding will be.						
	(1) 3p ⁶	(2) 3p ⁶ 4s ²	(3) 3p ⁶ 3d ²	(4*) 3d ² 4s ²			
Sol.	Element belongs to d-b	lock in d-block elements	(n - 1) d and ns electron	take part in the bonding.			
2.	Two ice cubes are pressed together until they form one block. Which of the following force is primarily						
	responsible for holding the cubes together?						
	(1) Dipole-dipole intera	action	(2) Van der Waals force	es			
	(3) Ionic interaction		(4*) Hydrogen bonding				
3.	A sigma bond may be f	ormed by the overlap of	2 atomic orbitals of atom	s A and B. If the bond is formed			
along as the x-axis, which of the following overlaps is acceptable ?							
	(1) s orbital of A and p_z orbital of B		(2) p_x orbital of A and p_y orbital of B				
	(3) p_z orbital of A and p	∝ orbital of B	of B (4^*) p _x orbital of A and s orbital of B				
4.	NH ₃ and BF ₃ combine r	eadily because of the for	mation of :				
	(1) a covalent bond	(2) a hydrogen bond	(3*) a coordinate bond	(4) an ionic bond			
5.	Maximum bond energy	is in :					
	(1) F ₂	(2*) N ₂	(3) O ₂	(4) equal in all			
6.	The structure of XeF2 ir	nvolves hybridization of t	he type :				
	(1) sp ³	(2) sp ³ d ³	(3*) sp ³ d	(4) sp ³ d ²			
7.	The bond angle and hybridization in ether (CH ₃ OCH ₃) is :						
	(1)106° 51', sp ³	(2) 104°31', sp ³	(3) 109°28' sp₃	(4*) >109°28' sp₃			

Max. Time : 1 Hr.

8.	Which of the following ha	ave maximum boilling p (2) CF4	ooint (3*) CCl₄	(4) CD4	
9.	In which of the following (1) XeF₄	molecule / ion all the b (2) BeF ₄ -	onds are not equal ? (3*) C ₂ H ₄	(4) SiF₄	
10.	The correct order of incre (1) H ₂ O > Cl ₂ O > F ₂ O	-			
11.	Which of the following is $(1^*) O_2^-$	paramagnetic ? (2) CN⁻	(3) CO	(4) NO+	
12.	Which of the following is (1) NF₃	non-polar : (2*) BF₃	(3) PF₃	(4) SF4	
13.	Strongest hydrogen bond (1) O – H S	•	(3*) F – H	- F (4) F – H O	
14.	(1*) Identical bond position	Resonance structure of a molecule should not have(1*) Identical bond position(2) Identical arrangements of atoms(3) Nearly the same energy content(4) The same number of paired electrons			
15.	The paramagnetic property of oxygen is well explained by :(1*) Molecular orbital theory(2) Resonance theory(3) Valence bond theory(4) VSEPR theory				
16.	 Which of the following statement is correct regarding molecular orbital theory (MOT) : (1*) Energy of bonding orbital is less than anti-bonding orbital. (2) Energy of bonding orbital is more than anti-bonding orbital. (3) Bonding orbitals are monocentric. (4) Bonding orbital follow n + l rule 				
17.	PF₂Cl₃ in non polar becar (1) P – Cl bond is non-po (3) P – Cl bond is polar		(2*) Its dipole moment (4) P & Cl have equal e		
18.	The hybrid states of cent (1) sp ² , sp ³ , sp ²	ral atom in diaborane, (2*) sp ³ , sp ³ , sp ²	diamond and graphite ar (3) sp ³ , sp ³ , sp		
19.	Which of the set of speci (1) NO ₂ +, NO ₂ , NO ₂ -	es have same hybridiz (2) ClO₄⁻, SF₄, XeF₄		hapes:- (4) SO4 ⁻² , PO4 ⁻³ , ClO4 ⁻	
20.	The bonds present in № (1) Only ionic (3) Only covalent	O₅ are :	(2*) Covalent & coordir (4) Covalent & ionic	nate	

21.	The correct statement for the reaction-				
	$NH_3 + H^+ \longrightarrow NH_4^+$				
	(1) Hybridisation state	is changed	(2*) Bond angle increa	ses	
	(3) NH₃ act as a Lewis	acid	(4) Regular geometry i	s changed	
22.	The correct order of decreasing polarisabality of ions is :				
	(1) Cl⁻ > Br⁻ > l⁻ > F⁻	(2) F ⁻ > I ⁻ > Br ⁻ > CI ⁻	(3) F ⁻ > Cl ⁻ > Br ⁻ > l ⁻	(4*) I⁻ > Br⁻ > CI⁻ > F⁻	
23.	CCl4 is more covalent	than LiCl because :			
	(1*) There is more pol	arization of CI in CCI4	(2) There is more pola	rization of CI in LiCI	
	(3) CCl4 has more wei	ght	(4) None of the above		
24.	An ionic compound A ⁺	⁻ B ⁻ is most likely to be fo	rmed when -		
	(1) Ionization energy of	of A is low	(2) Electron affinity of E	B is high	
	(3) Electron affinity of	B is low	(4*) Both (1) and (2)		
25.	Which of the following	statements regarding HC	CIO3 :		
	(1) oxidation state of c	hlorine is +5	(2) it has two pπ-dπ bo	onds	
	(3) it has two type of C	CI–O bond	(4*) all of these		
26.	The correct sequence	of increasing covalent ch	naracter is represented by	у -	
	(1) BeCl₂ < NaCl < LiC		(2*) NaCl < LiCl < BeC	l ₂	
	(3) BeCl ₂ < LiCl < NaC		(4) LiCl < NaCl < BeCl	2	
27.	Correct order bond ler	ngth is :			
	(1) N – H > P – H > SI	о – Н	(2*) N – H < P – H < S	b – H	
	(3) P – H > N – H > SI	о – Н	(4) $Sb - H > N - H > F$	P-H	
28.	Which of the follwing s	species contains three bo	nd pairs and one lone pa	air around the central atom ?	
	(1) NH ₂ -	(2*) PCI ₃	(3) H ₂ O	(4) BF ₃	
29.	During change of O_2 to O_2^- ion, the electron adds on which one of the following orbitals ?				
	(1) σ* 2p₂ orbital		(2) $\sigma 2p_z$ orbital		
	(3*) π* 2p _× / π* 2p _y orbi	tal	(4) π 2p _x / π 2p _y orbital		
	· · · · · · · · · · · · · · · · · · ·				
30.	The number of S-S bo	onds in sulphur trioxide tri	mer (S₃O൭) is :		
	(1) 3	(2*) 0	(3) 1	(4) 2	

PART - II : PRACTICE QUESTIONS

Practice Questions: 20-50 depending on chapter length.

1.	If the atomic number of element X is 7 the lewis diagram for the element is :					
	(1) X•	(2) • * •	•ו (3*)	(4)		
2.	Which of the following	Lewis diagram is incorre :Ċi: :Ċi – Ċ – Ċi: :Ċi – Ċ – Ċi: (2) :Ċi:		н н		
	(1) Na⁺[:Ö – C:]⁻	(2) : <u>C</u> !:	(3*) :Ö: C:::Ö:	(4) $H - N - N - H$		
3.	If the electronic config bonding will be.	uration of an element is	ls² 2s² 2p ⁶ 3s² 3p ⁶ 3d² 4s	s ² , then the electrons involved ir		
	(1) 3p ⁶	(2) 3p ⁶ 4s ²	(3) 3p ⁶ 3d ²	(4*) 3d ² 4s ²		
4.	A Bond formed betwe	en two same atoms canr	not be			
	(1*) Ionic	(2) Covalent	(3) Metallic Bond	(4) None of these		
5.	The correct order of d	ecreasing polarisabality of	of ions is :			
	(1) CI [_] > Br [_] > I [_] > F [_]	(2) F ⁻ > I ⁻ > Br ⁻ > CI ⁻	(3) F ⁻ > Cl ⁻ > Br ⁻ > l ⁻	(4*) I⁻ > Br⁻ > CI⁻ > F⁻		
6.	CCl4 is more covalent	than LiCl because :				
	(1*) There is more pol	arization of CI in CCI4	(2) There is more polar	(2) There is more polarization of CI in LiCI		
	(3) CCl₄ has more wei	ght	(4) None of the above			
7.	The correct sequence	of increasing covalent cl	haracter is represented by	/ -		
	(1) BeCl ₂ < NaCl < LiC	CI	(2*) NaCl < LiCl < BeC	2		
	(3) BeCl ₂ < LiCl < NaC		(4) LiCl < NaCl < BeCl ₂			
8.	NH₃ and BF₃ combine	readily because of the fo	ormation of :			
	(1) a covalent bond	(2) a hydrogen bond	(3*) a coordinate bond	(4) an ionic bond		
9.	Which of the following	leads to bonding?		[Kerla CEE 2003]		
	s-orbital	p-orbital d-orbital	s-orbital p-orbital	d-orbital d-orbital		
		(2*)				
		(2) 0	(3)	(4) • • • • •		
10.		ng molecule / ion all the b	•	() 		
	(1) XeF ₄	(2) BeF ₄ -	(3*) C ₂ H ₄	(4) SiF₄		
11.	-	•		ir around the central atom?		
	(1) NH₂ [−]	(2*) PCI₃	(3) H ₂ O	(4) BF ₃		
12. Which does not form linear structure ?				[RPMT 2003]		
	(1) CO ₂	(2*) NO ₂	(3) C ₂ H ₂	(4) BeCl ₂		
13.	Structure of ICI-4 is :					
	(1) trigonal	(<u>2) dis</u>	storted trigonal bipyramid			

•	(3) octahedral	(4*) sq	uare planar	•			
14.		The correct order towards bond angle is : (1) Bond angle does not depend on hybridisation.(2) sp < sp ² < sp ³					
	(3) $sp^2 < sp < sp^3$		(4*) $sp^3 < sp^2 < sp$				
15.	Which is not true abou	ıt CH₄ molecule ?					
	(1) Tetrahedral hybridisation		(2) 109.5° bond angle				
	(3) Four sigma bonds		(4*) One Ione pair of	electrons on carbon			
16.	The structure of XeF2	involves hybridization of t	he type :				
	(1) sp ³	(2) sp ³ d ³	(3*) sp ³ d	(4) sp ³ d ²			
17.	Which of the set of spe	ecies have same hybridiza	ation state but different	shapes:			
	(1) NO ₂ +, NO ₂ , NO ₂ -	(2) ClO₄⁻, SF₄, XeF₄	(3*) NH4+, H3O+, OF2	(4) SO ₄ ⁻² , PO ₄ ⁻³ , CIO ₄ ⁻			
18.	The correct statement	for the reaction-					
	$NH_3 + H^+ \longrightarrow NH_4^+$						
	(1) Hybridisation state	is changed	(2*) Bond angle incre	ases			
	(3) NH₃ act as a Lewis	acid	(4) Regular geometry	is changed			
19.	Among the following p	Among the following pairs in which the two species are not isostructural is :					
	(1*) SiF₄ and SF₄	(2) IO₃⁻ and XeO₃	(3) BH_4^- and NH_4^+				
20.	Which among the follo	owing molecules have sp	³ d hybridisation with c	one lone pair of electrons on the			
	central atom ?						
	(i) SF4	(ii) [PCl ₄]+	(iii) XeO ₂ F ₂	(iv) CIOF₃			
	(1) (i), (ii) and (iii) only	(2*) (i), (iii) and (iv) only	(3) (i) and (iii) only	(4) (iii) and (iv) only			
21.	The bond angle and h	ybridization in ether (CH ₃ 0	OCH₃) is :				
	(1)106° 51', sp ³	(2) 104°31', sp ³	(3) 109°28' sp₃	(4*) >109°28' sp₃			
22.	During change of O2 to	During change of O_2 to O_2^- ion, the electron adds on which one of the following orbitals ?					
	(1) $\sigma^* 2p_z$ orbital		(2) σ 2p _z orbital				
	(3*) π* 2p _x / π* 2p _y orbi	tal	(4) π 2p _x / π 2p _y orbita	al			
23.	Which of the following	statements are correct ?					
	Ö						
	(I) O structure	(I) of structure is not allowed because octet around 'O' can not be expanded.					
	(II) H ₂ O ₂ is ionic comp	ound					
	(III) In B ₂ molecule, the	e highest occupied molect	ular orbital is σ molecu	lar orbital.			
	(IV) The lp–bp repulsio	on is stronger than bp–bp	repulsion.				
	(1) (I) and (III)	(2) (II) and (III)	(3*) (I) and (IV)	(4) (III) and (IV)			

24.	Which of the following is paramagnetic ?					
	(1*) O ₂ -	(2) CN⁻	(3) CO	(4) NO+		
25.	5. The boiling point of a compound is raised by -					
	()	ar hydrogen bonding	(2) High volatility			
	(3) Intramolecula	r hydrogen bonding	(4) Non-polarity			
26.	Which of the following have maximum boiling point					
	(1) CH ₄	(2) CF ₄	(3*) CCI ₄	(4) CD ₄		