### **UNIT: 4 CHEMICAL BONDING AND MOLECULAR STRUCTURE**

### **Important Points**

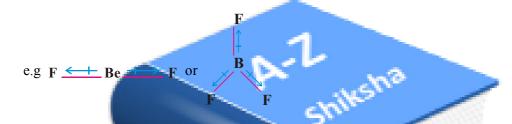
In this unit, the study of chemical bonding and formation of molecule from the atoms are included. The smallest particle of element is atom and the smallest particle in a compound is molecule. The force or the binding that keeps the atoms in the molecule combined during the formation of molecule is called chemical bonding. The concepts like that of Kossel-Lewis, VSEPR principle, valence bond theory, molecular orbital theory have been presented. In chemical bonding, it has more relation with orbitals around the nucleus and especially the valence orbitals. We do not think about the nucleus but we take into consideration the effect due to its positive charge. Scientists Lewis and Kossel have mentioned the approach of chemical bonding. In this, the atom obtains the octet either by losing or by gaining the electron, which is chemically inert. This is called law of octet. Such bonds are called ionic bonds e.g. NaCl. Also, some atoms share electrons with each other and obtain octet structure resulting into stable covalent molecule. e.g. Cl<sub>2</sub>. To explain the structures of such molecules he mentioned dot and cross symbols and explained the stability of the molecules. Such a bond is called covalent bond. The approach of Kossel Law is explained in detail in the unit.

When any bond is formed, the distance between their atoms is called bond length and the angle is)child, boomalbitan (te) Asymptotic bond lengths of single (

metrical shapes to molecules viz.  $180^{0}$ - linear,  $109^{0}28^{1}$  alent bond which can be of three types. (1) Polar aged towards the more electronegative atom and  $+\delta$  in electronegative atoms are shown. As a result the ites of the two atoms are same or the difference med by both the atoms sharing the electrons. In cosharing a pair of electrons and the second atom electron pair. viz.  $F_3B \leftarrow NH_3$ . Bond  $\leftarrow$  indicates length, bond angle, bond enthalpy (bond energy) is h, more will be the stability and so more energy will alpy may be different according to bond formation. In we have studied in detail and also the formula to py evolved in formation of compound is the mathons of atoms. It is explained in the unit by discussing

a pair instead of being localised on any molecule it is in the molecule can be shown at different positions a. Such structures are called resonance structures and onance forms is called resonance energy. This can on dioxide, benzene etc. As we have seen earlier, structures like linear, tetrahedral etc, can be obtained on the basis of bond angle. This study can be used to show the shapes of the molecule by hybridisation of atoms in them, geometrical structures etc. viz. linear  $BeCl_2 - 180^0$ , trigonal  $BCl_3 - 120^0$ , tetrahedral  $CH_4 - 109^028'$ .

Lewis approach being insufficient to explain the shapes of molecules, Sidgwick and Powell proposed one principle which is known as VSEPR principle which was developed by Nyholm and Gillespie and they proposed certain assumptions. In this it is important to note that when non-bonding electron pairs are there, then they show deviation in geometrical structure and bond angle due to repulsion between electron pairs. e.g. Molecule of water has sp³ hybridisation and so its bond angle must be  $109^028'$  but it becomes  $104^0$  30' due to repulsion by two non-bonding electron pairs. Hence, it is called distored tetrachedral. The polarity of bond is a vector quantity. Hence, if a polar bond is formed due to difference in electronegativities but another bond of the same type is formed in its opposite direction, then polar bond will be formed but the resultant polarity of the molecules becomes zero and molecule will be non-polar.



The dipole momentes of polar substances can be calculated for which both the charges  $+\delta$  and the distance

the polar bond and only in polar solven in water. Napthalene consideration the lin them: (1) Valence quantum mechanics developed by Paulin

In the assumpt charged nuclei of the Coulomb's Law if a molecule will be for overlaps can be form be shown and then the trons of valence or be with one electron in the geometrical structure. In such valence both

attains axial overlap of internuclear axis. The stability of this bond is more than that of  $\pi$  bond. In the  $\pi$ -bond the axis of the atomic orbitals undergoing overlapping remains parallel to each other and is perpendicular to internuclear axis.  $\pi$ - bonds are less stable in comparison to  $\sigma$ -bonds or they are weaker. Valence bond theory is based on overlapping of valence orbitals. It explains properties like the geometrical shapes, the bond angle etc. very simply but cannot explain magnetic properties.

Scientists Mulliken and Hund suggested molecular orbitals like atomic orbitals and proposed molecular orbital theory. Amongst its important points, the idea that atomic orbitals can also form molecular orbitals was taken into consideration. As many atomic orbitals take part in the formation, same number of molecular orbitals, their energy, symmetry etc. were taken into consideration. The formation of these types of atomic orbitals can be shown in the formation of homonuclear molecules like H<sub>2</sub>, Be<sub>2</sub>, F<sub>2</sub> etc. and heteronuclear molecules like CO, NO etc. Molecular orbitals are formed by linear combination of atomic orbitals—LCAO principle. On the basis of these types of combination two types of molecular orbitals are formed which are known as Bonding Molecular Orbitals (BMO) and Anti-Bonding Molecular Orbitals (ABMO). In the formation of rules these types of BMO and ABMO the principles like Hund's rule of maximum spin, Pauli's exclusion principle, Aufbau principle etc. which are applicable in formation of atomic orbital are also obeyed and maintained. In the unit the molecular orbital diagrams of construction of molecular orbitals from the atomic orbitals for formation of homonuclear molecules from H<sub>2</sub> to Ne<sub>2</sub> elements as well as for formation of heteronuclear molecules like CO, and NO are shown. From these diagrams, important property like bond order can be calculated. Bond order

=  $\frac{1}{2}$  {electron in bonding molecular orbitals – electrons in anti – bonding orbital} viz. for N<sub>2</sub> mol-

ecule bond order  $=\frac{1}{2}(10-4)=3$  i.e. there will be triple bond  $N \equiv N$ . In the same way, in NO

molecule bond order will be  $=\frac{1}{2}[10 \quad 5] = 2.5$ . Here, we will note that if the value of bond order

becomes zero, the bond will not be formed e.g. He<sub>2</sub>. If the value of bond order is integer, the bond will describe (ex) and trained ording to the integer 1, 2, or 3, there will be single (

#### on, then molecule will attain unstable

magnetic properties e.g. In  $O_2$  molecule and in  $N_2$  molecule, all the electrons are y is superior to valence bond theory in

and. The first element of 15, 16, and 17, ments of the group it can form covalent a. Afterwards the molecule like

H — F combines leattle ethato we the hydrogen bond

After knowing about ionic bond, covalent bond, co-ordinate covalent bond, we shall study the special type of bond present in metals which is called metallic bond. As there are 1, 2 or 3 electrons in the outermost orbit of the metals, they are not able to form covalent bonds. Their ionisation energy is less and attraction of electron towards the nucleus is less. One, two or three electrons are arranged around the nucleus of the atom. Hence, the positively charged nucleus or kernel is there. The electrons around it have attraction towards other nuclei of the atoms in the lattice. Thus, the electron instead of being localised for any one atom, remains delocalised in the whole metal crystal. For this theory Electron Sea model was proposed. In this, the atomic kernel is imagined as floating in the sea, delocalised electrons are arranged around kernel possessing positive charge. Because of this type of metallic bonds, the specific properties of metals, like density, duetility, malleability etc. are different.

Co-ordinate covalent bond is a type of covalent bond as seen earlier. The characteristic in it is that from the two atoms undergoing sharing of electrons, only one of the atoms provides a pair of electrons, and is shared by both the atoms. Hence, it is called co-ordinate covalent bond. e.g. In BF<sub>3</sub>, three F atoms were bonded with B-atom through three covalent bonds but the octet of B is not complete. Similary in NH<sub>3</sub> molecule, three H atoms are bonded with N through three covalent bonds. But N has one non-bonding pair of electrons, which it gives to BF<sub>3</sub> molecule and is shared by both the molecules. Hence  $F_3B \leftarrow NH_3$  Co-ordinate covalent bond is formed. In this, the molecule which gives pair of electrons is shown by arrow  $(\rightarrow)$  from the molecule which donates it towards the molecule or atom which accepts and shares gained electron pair. You will study more about co-ordinate covalent bond in the unit of complex salts in Standard-12.

# M.C.Q.

(1)	Which of the following	g is ionic?		
	(a) HCl	(b) CHCl <sub>3</sub>	(c) IF <sub>5</sub>	(d) KI
(2)	When molecule is form	n by chemical bonding b	etween atoms then	
	(a) nucleous of combin	ning atoms are participat	e	
	(b) valence electrons a	and inner cell electrons a	are participate	
	(c) only valence electron	ons of combining atoms	are participate	
	(d) only inner cell elec	trons of combining aton	ns are participate	
(3)	Which factor is not res	ponsible for the formation	on of ionicbond?	
	(a) crystal lattice energ	y	(b) density	
	(c) ionisation enthalpy	_	(d) electron gain enth	alpy
(4)	According to valence-	bond theory which mag	netic property oxygen	possess?
	(a) Paramagnetic	(b) Ferrimagnetic	(c) Diamagnetic	(d) Anti Ferromagnetic
(5)	Who was proposed va	lence-bond theory?	ch3	
	(a) Mulliken	(b) Lenus Pauling	(c) Hittler and Londa	n (d) Hund
(6)		= $CH_2$ molecule $C^3$ -	C <sup>2</sup> single bond carb	ons has which type of
	hybridization?			
	(a) $sp^2 - sp^3$	(b) sp - sp <sup>2</sup>	(c) $sp^3 - sp$	(d) sp3 -sp3
(7)	Which of the following	g pair of species is isoele		
	(a) $NO_3^-$ , $SO_3$	3	(c) $CO_3^2$ , $ClO_3$	(d) $NO_3^-$ , $CO_3^{2-}$
(8)	Which of the following	g sentence is incorrect for	or covalent bond?	
	(a) Strenght of covaler	nt bond depenas upon o	verlapping at atomic or	rbitals.
	(b) Covalent bond is n	ot directional.		
	(c) There is sharing of	electrons between atom	s bonded by covalent	bond
	(d) Covalent bond is for	ormed between atoms h	aving less difference in	their electronegativity.
(9)	Which of the following	g compound possesses of	eovalent bond?	
	(a) MgCl <sub>2</sub>	(b) NaH	(c) BF <sub>3</sub>	(d) CsCl
(10)	Which of the following	g molecule possesses po	lar and nonpolar coval	ent bond?
	(a) NH <sub>4</sub> Cl	(b) CCl <sub>4</sub>	(c) $H_2O_2$	(d) HCN
(11)	Which of the following	g compound does not po	ossesses coordinate co	valent bond?
	(a) CO	(b) SO <sub>2</sub>	(c) HNO,	(d) HNO <sub>3</sub>

	(a) They do not possesses particular geometical structure					
	(b) They may be polar or nonpolar					
	(c) Their boiling and m	elting point is low				
	(d) Generally they are	insoluble in water				
(13)	Which of the following	g possesses ionic and co	valent bond?			
	(a) CO <sub>2</sub>	(b) $H_2SO_4$	(C) NH <sub>4</sub> Cl	(D) NaI		
(14)	Whhat is Geometrical	Structure of CIF <sub>3</sub> molecular	ule?			
	(a) Triogonal bipyramie	d(b) Corn shpae	(c) sea-saw	(d) T-shape		
(15)	Which of the following	g molecule possesses line	ear structure?			
	(a) SO <sub>2</sub>	(b) CO <sub>2</sub>	(c) H <sub>2</sub> O	$(d) C_2 H_4$		
(16)	Correct structure of SI	F <sub>4</sub> is				
	<i>p</i> -					
	(A) (D) F F		F (D)			
	F	The Francisco	FEF			
(17)	Numbers of possible r	esonating structure of ca	arbonate iong is			
	(a) 9	(b) 6	(c) 3	(d) 2		
(18)	Which of the following	molecule has not zero c	lipol movement?			
	(a) NF <sub>3</sub>	(b) BF <sub>3</sub>	(c) CO <sub>2</sub>	(d) BeF <sub>2</sub>		
(19)	Which of the following	molecule possesses hig	hest dipolspace moven	nent?		
	(a) CCl <sub>4</sub>	(b) CHCl <sub>3</sub>	(c) CHCl <sub>2</sub>	(d) CH <sub>3</sub> Cl		
(20)	Which of the following	molecule possesses dip	ol movement?			
	(a) trans - 1, 2 - dichlo	ro ethene	(b) trans pent - 2 - en	ne		
	(c) 2, 2- dimethyl prop	pane	(d) 2, 2, 3, 3- tetra m	ethyl butane		
(21)	Which of the following	g molecule has lowest bo	ond space angle?			
	(a) NH <sub>3</sub>	(b) SO <sub>2</sub>	(c) H <sub>2</sub> O	$(d) H_2 S$		
(22)	Which orbital has high	est energy?				
	(a) $\sigma(2Px)$	(b) <i>π</i> * (2Py)	(c) σ(2S)	(d) $\sigma^*$ (1S)		

(12) Which of the following characteristic is not for covalent compound?

(23)	) Which is the paramagnetic species ?				
	(a) $\overline{CN}$	(b) $_{\mathrm{O}_{2}}^{-}$	(c) NO <sup>+</sup>	(d) CO	
(24)	Which of the following respectively?	g statement is incorrect	when $N_2$ and $O_2$ are co	envert into N <sub>2</sub> <sup>+</sup> and O <sub>2</sub> <sup>+</sup>	
	(a) In $O_2^+$ , O - O bond	order increases.	(b) In $N_2^+$ , N - N bo	nd become weaker.	
	(c) N <sub>2</sub> <sup>+</sup> become param	agnetic	(d) Increasing dimagr	netism in $O_2^+$	
(25)	According to VSEPR t	heory geomety of which	h block elements can b	e explain ?	
	(a) s	(b) p	(c) d	(d) f	
(26)	Atoms complete octet proposed by which scient		during the bond form	ation. This postulate was	
	(a) Powel	(b) Lewis	(c) Sigdwick	(d) Mulliken	
(27)	Crystal formation is wh	nich type of reaction?			
	(a) endothermic and ex	othermic	(b) endothermic		
	(c) exothermic	, ,	(d) no heat change of	ecurs	
(28)	Lattice energy of ionic	compound depends upo	on which factor?		
	(a) Size of ion		(b) Size of ion and ch	narge	
	(c) charge on ion		(d) Arrangement of ic	on	
(29)	Which is correct order	for C - O bond length i	in $CO$ , $CO_3^{2-}$ , $CO_2$		
	(a) $CO_3^{-2} < CO_2 < CO$		(b) $CO_2 < CO_3^{2-} < CO_3^{2-}$	CO	
	(c) $CO < CO_2 < CO_3^{2-}$		(d) $CO < CO_{3}^{2-} < CO_{3}^{2-}$	$O_2$	
(30)	Maximum how many r	numbers of hydrogen bo	ond can be form by H <sub>2</sub> O	) molecule ?	
	(a) 2	(b) 4	(c) 3	(d) 1	
(31)	In buta 1, 3 - diene				
	(a) only one sp hybridi	sed carbon atom			
	(b) only sp <sup>2</sup> hybridised	carbon atoms			
	(c) Two sp <sup>3</sup> and two sp	<sup>2</sup> hybridised carbon ator	ms		
	(d) sp, sp <sup>2</sup> and sp <sup>3</sup> hybr	ridized carbon atoms			

(32)	Which of the following statement is irrelevant for sigma bond?					
	(a) strength of sigma bond is not related with overlapping of atomic orbitals.					
	(b) $\sigma$ - bond can form by overlapping of S - P orbitals.					
	(c) $\sigma$ - bond ca	n form by c	overlapping o	of end of ato	mic orbitals	of inner center axis.
	(d) This type o	f overlappin	g is also kno	wn as axial	overlaping	
(33)	In which molec	cule inter mo	olecular hydr	ogen bond c	an be form	?
	(a) methanol	(b)	ethelene gly	col (c)	p - nitrophe	enol (d) phenol
(34)	In which molec	cule intra mo	olecular hydr	ogen bond c	an be form	?
	(a) o - nitro pho	enol (b)	aniline	(c)	ethylene gly	col (d) all of these
(35)	Which of the fo	ollowing pa	ir possesses	very strong	H - bond ?	
	(a) CH <sub>3</sub> COCH	I <sub>3</sub> and CHC	$l_3$	(b)	HCOOH at	nd CH <sub>3</sub> COOH
	(c) H <sub>2</sub> O and H	2		(d)	SiH <sub>4</sub> and Si	Cl <sub>4</sub>
(36)	Which of the fo	ollowing rela	ation is corre	ct?		
	(a) Bond order	rα Bond e	energy α Bo	nd length	α stability	
	(b) Bond orde	era Bond	$\frac{1}{\text{length}} \alpha$	$\frac{1}{\text{energy}}$ st	ability	
	(c) Bond orde	rα Bond e	nergy α - F	l Bond length	$\alpha$ stability	
	(d) Bond orde	erα Bond	$\frac{1}{\text{length}} \alpha = \frac{1}{1}$	Bond energy	-α stability	
(37)	Molecule:	$H_2$	F <sub>2</sub>	Cl <sub>2</sub>	$\mathrm{Br}_2$	
	Bondlength:	74pm	144pm	199pm	228pm	
	Mention more	stable mole	cule from abo	ove		
	(a) Cl <sub>2</sub>	(b)	$H_2$	(c)	$\operatorname{Br}_2$	(d) $F_2$
(38)	In water bond	angle is 104	№ 30 because			
	(a) Oxygen ato	m is sp <sup>3</sup> hyb	oridised			
	(b) Repulsion b	oetween lon	e pair electio	n and bondi	ng pair elec	tron
	(c) Oxygen has	high electro	onegetivity.			
	(d) H <sub>2</sub> O molec	ule possess	es "V" - shaj	pe.		

(39)	(39) In which of the following strong H-bond is present?			
	(a) F - HF	(b) O - HN	(c) O - HO	(d) O - HF
(40)	Which is correct orde	er for bond dissociation	energy in $O_2$ , $O_2^+$ , $O_2^-$	$\frac{1}{2}$ and $O_2^{2-}$
	(a) $O_2 > O_2^+ > O_2^{2-} >$	$\mathrm{O}_2^-$	(b) $O_2^+ > O_2^- > O_2^-$	> O <sub>2</sub> <sup>2-</sup>
	(c) $O_2^- > O_2^{2-} > O_2^+ >$	$\mathrm{O}_2$	(d) $O_2^- > O_2^{2-} > O_2$	> O <sub>2</sub> <sup>+</sup>
(41)	O, P, Q, R elements e	lectronic configuration is	s given below	
	$O = 1s^2$	$P = 1s^2, 2s^2, 2p^2$		
	$Q = 1s^2, 2s^2, 2p^5$	$R = 1s^2, 2s^2, 2p^6$		
	Which atom has stron	ng behaviour of electrova	alent bond ?	
	(a) O	(b) P	(c) Q	(d) R
(42)	In which molecule bo	ond distorsion is more ac	cording to VSEPR the	eory?
	(a) SO <sub>2</sub>	(b) NH <sub>3</sub>	(c) O <sub>3</sub>	(d) H <sub>2</sub> O
(43)	Which of the following	ng species is more stable	? <b>K</b> ST	
	(a) O <sub>2</sub>	(b) Ne <sub>2</sub> <sup>+</sup>	(c) O <sub>2</sub>	(d) F <sub>2</sub>
(44)	Number of nonbondi	ng electron pair in XeF <sub>6</sub> ,	, XeF <sub>4</sub> and XeF <sub>2</sub> respe	ectively
	(a) 2, 3, 1	(b) 1, 3, 2	(c) 3, 2, 1	(d) 1, 2, 3
(45)	On keeping two cube	of ice on each other wh	ich become one cube	which factor is responsible
	for it?			
	(a) Van-der waals attr	raction (b) Hydrogen l	bond (c) Dipole attra	action (d) Covalent bond
(46)	Determine lattice ener	rgy of LiF <sub>(S)</sub> according to	given data.	
	$(i) \operatorname{Li}_{(S)} \to \operatorname{Li}_{(g)}$		155.2 KJ mol $^{-1}$ ( $\Delta_{ m S}$	H)
	(ii) $F_{2(g)} \rightarrow 2F_{(g)}$		75.2 KJ $\Delta \left( \frac{\text{H}_{\text{D}}}{2} \right)$	
	(iii) $\operatorname{Li}_{(g)} \to \operatorname{Li}_{(g)}^+ + \overset{-}{e}$		520.0 KJ molΔ <sup>1</sup> H(	i )
	(iv) $F_{(g)}^{+\bar{e}} \rightarrow \overline{F}_{(g)}$		$-33.0~{ m KJ}\Delta(~{ m Hg}~)$	
	(v) $\operatorname{Li}_{(s)} + \frac{1}{2} \operatorname{F}_{2(g)} \to \operatorname{Li}$	$iF_{(S)}$	-504.1 KJ mol ⋅ (∠	<b>,</b> H)
	(a) $-86.7  \text{KJ}  \text{mol}^{-1}$	(b) 86.7 <i>KJ mol</i> <sup>-1</sup>	(c) $-867$ KJ mol $^{-1}$	(d) 867 <i>KJ mol</i> <sup>-1</sup>

(47)	Which of the following statment is incorrect for metallic bond?				
	(a) There is attraction between delocalised electrons and atomic karnel				
	(b) Directionl property	is shown by metal			
	(c) Delocalised electron	n can change their positi	on easily in crystal		
	(d) Explanation of met	callic bond can be given	by 'electron sea model	S - Orbital	
(48)	Why lattice energy of	NaCl > KBr ?			
	(a) When size of negat	ive ion decrease in ionic	crystal then lattice end	ergy increases.	
	(b) When volume of p more and hence latice	_	is small than then inte	erionic attraction become	
	(c) In ionic crystal whe	en size of positive ion de	ecrease, then lattice ene	ergy increases.	
	(d) All of given				
(49)	Number of H - bond for	orm by unpaired electro	ns of liquid NH <sub>3</sub> , H <sub>2</sub> O	and HF respectively are	
	(a) 3, 4, 2	(b) 4, 4, 2	(c) 3, 2, 1	(d) 1, 2, 1	
(50)	Which of the following	g pair is not in order for	boiling point for 14, 1:	5, 16 and 17 group?	
	(a) $H_2O > H_2S$	(b) HF > HCl	(c) $CH_4 > SiH_4$	(d) $NH_3 > PH_3$	
(51)	Which of the following	g compound possesses ic	onic bond?		
	(a) CH <sub>4</sub>	(b) SiCl <sub>4</sub>	(c) BF <sub>3</sub>	(d) MgCl <sub>2</sub>	
(52)	Which of the following diatomic malecule or io		and ABMO electrons	is correct for stability of	
	(a) $N_a > N_b$	(b) $N_b > N_a$	(c) $N_a + N_b = 0$	$(d) N_a = N_b$	
(53)	At what distance van-	derwaals attraction exist	?		
	(a) $4.5 \times 10^{-10}$ m	(b) 0.45nm	(c) 4.5 A°	(d) Given all	
(54)	What is bond energy o	of H-bond ?			
	(a) 40 J mol <sup>-1</sup>	(b) 40 KgJ mol <sup>-1</sup>	(c) 40 cal. mol <sup>-1</sup>	(d) 40 Kg cal mol <sup>-1</sup>	
(55)	In which molecule inte	r molecular H-bond is p	ossible?		
	(a) CH <sub>3</sub> COCH <sub>3</sub>	(b) CH <sub>4</sub>	(c) SiH <sub>4</sub>	(d) NH <sub>3</sub>	
(56)	Which of the following	g characterstic does not p	possesses by metal?		
	(a) luminus			(b) ductility	
	(c) increase in conduct	ance by increase in temp	perature	(d) malleability	

(57)	7) On which factor conductance of metals responsible?				
	(a) ions	(b) delocalised	(c) atomic kernel	(d) number of atoms	
(58)	Which of the following	g figure shows electron-s	sea model?		
	A) 000 000 000	(B) (B-(B)-(B-(B)-(B)-(B)-(B)-(B)-(B)-(B)-(B			
(59)	According to which go living organism?	roup, hydrogen bond is	form in protein molec	ule present in musecls of	
	(a) -CO-	(b) -COOR	(c) -CONH-	(d) -COOH	
(60)	On which factor van-c	der waalls attraction force	ce does not depend?		
	(a) numbers of molecu	les	(b) contact surface an	rea of molecules	
	(c) shape of molecules	(d) numbers of electron	n in molecules		
(61)	Practicol dipal movembe the percentage of ic		bond length of HCl is	s 1.275 A° than what will	
	(a) 7	(b) 17	(c) 43	(d) 21	
(62)	Which sentence is cor	reet with respect to bone	d enthalpy ?		
	(a) As bond order is m	nore, then bond dissociat	tion enthalpy is less		
	(b) As atomic volume	is more, then bond energ	gy is more.		
	(c) As bond enthalpy i	s more, then stability of	molecule or ion is less.		
	(d) As number of nonl	oonding election pair on	bonded atom then bon	nd enthalpy is less.	
(63)	which of the following	orbitals form bonding o	orbital?		
	(a) + + S - Orbital P - Orbital	(b) P - Orbital a/f - Orbital	(c) + + + P - Orbital P - Orbital	(d) + - Orbital P - Orbital	
(64)	Mention number of be	onding electron pairs and	d nonbonding electron	pairs in NO <sub>3</sub> ion	
	(a) 3, 1	(b) 2, 2	(c) 4, 0	(d) 1, 3	
(65)	How many numbers o	f bonding and nonbond	ing electron pairs in CC	O <sub>2</sub> ?	
	(a) 4, 4	(b) 2, 4	(c) 4, 2	(d) 2, 2	

(66)	Mention proper order	of bond length given be	elow.	
	(a) $N_2 < N_2^{2-} < N_2^{-}$	(b) $N_2^{2-} < N_2^- < N_2$	(c) $N_2^- < N_2 < N_2^{2-}$	(d) $N_2 < N_2^{2-} < N_2$
(67)	Show paramagnetic co	ompound given below.		
	(a) O <sub>3</sub>	(b) KO <sub>2</sub>	(c) N <sub>2</sub> O	(d) $Na_2O_2$
(68)	Which species posses	ses pyramidal shape?		
	(a) OsF <sub>2</sub>	(b) SO <sub>3</sub>	(c) BrF <sub>3</sub>	(d) $SiO_3^{2-}$
(69)	Which of the following	ng does not possesses bo	ond order as CO?	
	(a) NO	(b) NO	(c) N <sub>2</sub>	(d) $\bar{CN}$
(70)	Which rule is violated	l in the given electronic o	configuration?	
	ı			
		/()		
	2P	$\mathbb{Q}^{-}\mathbb{Q}^{\prime}$	2P	
			hill	
	(a) Authou	(b) Pauli	(a) Hand	(d) Cirron all
(71)	(a) Aufbau	(b) Pauli	(c) Hund	(d) Given all
(71)		ving molecule double bo	000000000000000000000000000000000000000	
	(a) S <sub>2</sub>	(b) O <sub>2</sub>	(c) C <sub>2</sub>	(d) H2C = CH2
(72)	Mention AB <sub>4</sub> F <sub>2</sub> type m	nolecule.		
	(a) BrF <sub>5</sub>	(b) XeF <sub>4</sub>	(c) $SF_6$	(d) XeOF <sub>4</sub>
(73)	Which of the followin	g is the correct order for	r lone pair and bonding	g pair electrons?
	Lp = Lone pair and B	p = Bonding pair		
	(a) Lp - Lp > Lp - Bp	p > Bp - Bp	(b) $Lp - Bp > Lp - I$	$\mathbb{L}p > \mathrm{Bp} - \mathrm{Bp}$
	(c) $Bp - Bp > Lp - Lp$	p > Lp - Bp	(d) $Lp - Lp > Bp - B$	Bp > Lp - Bp
(74)	Which theory is usefu	l to determine geometric	al structure of molecul	es?
	(a) molecular orbital th	heory	(b) VSEPR theory	
	(c) Resonance theory	(d) Quantam mechani	cs	
(75)	The one outermost ek	ectron present in Na eler	ment at	

(a) one comer of simple cube

(c) center of simple cube

(b) eight corner of simple cube

(d) each corner of simple cube

(76)	6) In which molecules / ion have not all the equal bonds?				
	(a) SF <sub>4</sub>	(b) BF <sub>4</sub>	(c) XeF <sub>4</sub>	(d) SiF <sub>4</sub>	
(77)	Which of the following	has maximum bond ang	gle ?		
	(a) NH <sub>3</sub>	(b) CH <sub>4</sub>	(c) CO <sub>2</sub>	(d) $H_2O$	
(78)	Which of the following	g have equal bond order	?		
	(a) $O_2^-$	(b) CN-	(c) NO	(d) B and C	
(79)	The type of bond pres	ent in CuSO <sub>4</sub> .5H <sub>2</sub> O			
	(a) covalent and co-ord	dinate covalent	(b) electrovalent and	covalent	
	(c) electrovalent and co-ordinate covalent				
	(d) electrovalent, covalent and co-ordinate covalent				
(80)	0) Which of the following statement is wrong				
	(a) sp² hybrid orbitals are formed from two p - atomic and one s - orbitals				
	(b) hybridization is the mixing of atomic orbitals prior to their combing into molecular orbitals				
	(c) d <sup>2</sup> p <sup>2</sup> hybrid orbitals are all at go to one an other				
	(d) d <sup>2</sup> sp <sup>3</sup> hybrid orbital	s are directed towords t	the corners of a regula	tetrahedron	
(81)	CO <sub>2</sub> is isostructual with	, ,			
	(a) SnCl <sub>2</sub>	(b) HgCl <sub>2</sub>	(c) C <sub>2</sub> H <sub>2</sub>	(d) SO <sub>2</sub>	
(82)	NH <sub>3</sub> has a higher boiling	g point than expected b	ecause		
	(a) its density decrease	s on freezing			
	(b) with water it forms	NH <sub>4</sub> OH			
	(c) it has strong inter m	nolecular covalent bond	s ?		
	(d) it has intermolecula	r hydrogen bonds.			
(83)	The molecule with zero	dipole moment is			
	(a) chloroform		(b) methyl chloride		
	(c) carbon tetrachlorid	e	(d) methylene chlorid	e	
(84)	Molecular shaper of S	F <sub>4</sub> , CF <sub>4</sub> , XeF <sub>4</sub> are			
	(a) the same with 1, 1 a	and 1 lone pairs of elect	crons respectively		
	(b) different with 1, 0 a	and 2 lone pairs of electronic	rons respectively		
	(c) different with 0, 1	and 2 lone pairs of elect	trons respectively		
	(d) different with 2, 0 a	and 1 lone pairs of elect	rons respectively		

(85)	) Which of the following has the regular tetrahedral structure?					
	(a) SF <sub>4</sub>	(b) $[Ni(CN)_4]^{2-}$	(c) BF <sub>4</sub>	(d) XeF <sub>4</sub>		
(86)	In OF <sub>2</sub> , number of bon	d pairs and lone pairs of	f electrons are respecti	vely		
	(a) 2, 8	(b) 2, 6	(c) 2, 9	(d) 2, 10		
(87)	In $O_2^-$ , $O_2$ , $O_2^{2-}$ molecular	ular species the total nur	nber of antibonding el	ectrons respectively are		
	(a) 7, 6, 8	(b) 1, 0, 2	(c) 6, 6, 6	(d) 8, 6, 8		
(88)	Match the following an	nd choose the correct An	iswer			
	Column - I	Column -II				
	(i) $sp^3 d^2$	(a) $Ni[(CN)_4]^{2-}$				
	(ii) sp <sup>3</sup> d	(b) SnCl <sub>2</sub>				
	(iii) dsp <sup>2</sup>	(c) IC $\bar{l_4}$				
	(iv) sp <sup>2</sup>	(d) TeCl <sub>4</sub>				
	(a) $i \rightarrow d$ , $ii \rightarrow a$ , $iii -$	$\rightarrow$ c, iv $\rightarrow$ b	(b) $i \rightarrow c$ , $ii \rightarrow d$ , $iii$	$i \rightarrow a$ , $iv \rightarrow d$		
	(c) $i \rightarrow b$ , $ii \rightarrow c$ , $iii \rightarrow c$	$\rightarrow$ d, iv $\rightarrow$ a	(b) $i \rightarrow c$ , $ii \rightarrow d$ , $iii$ (d) $i \rightarrow a$ , $ii \rightarrow b$ , $iii$	$\rightarrow$ c, iv $\rightarrow$ d		
(89)	Among the following hyridisation is	compounds, the one the	hat is polar and has tl	ne central atom with sp <sup>2</sup>		
	(a) HClO <sub>2</sub>	(b) BF <sub>3</sub>	(c) H <sub>2</sub> CO <sub>3</sub>	(d) SiF <sub>4</sub>		
(90)	Match the following					
	Set A		Set B			
	(1) stability of bond		(p) Bond energy			
	(2) Molecular orbital th	neory	(q) Bond order			
	(3) octet rule		(r) Variable Valency			
	(4) Valence bond theor	У	(s) Electronic concep	t of valency		
	(a) $1 \rightarrow q$ , $2 \rightarrow p$ , $r$ ,	$3 \rightarrow p, 4 \rightarrow s$	(b) $1 \rightarrow p, q, 2 \rightarrow p$	$p, 3 \to r, 4 \to s$		
	(c) $1 \rightarrow p, q, 2 \rightarrow r,$	$3 \rightarrow s, 4 \rightarrow r$	(d) $1 \rightarrow p, q, 2 \rightarrow q$	$q, 3 \rightarrow s, 4 \rightarrow r$		
(91)	Bond strength increase	s with				
	(a) Bond length increas	sing				
	(b) Antibonding eletron	s being higher in numbe	r			
	(c) Bond order increas	ing	(d) Bond angle increa	asing		

(92)	$O_2^{2+}$ will have					
	(a) Bond order lower than O	2				
	(b) Bond order higher than $O_2$					
	(c) Bond order lower t	han H <sub>2</sub>				
	(d) Bond order higher	than N <sub>2</sub>				
(93)	In a molecule number	of electrons in BMO are	e more as compared to	ABMO, hence		
	(a) a bond will be form	ed				
	(b) no bond will be for	med				
	(c) information is not su	ufficient				
	(d) none of the above					
(94)	The bond angle in the a	mmonium ion is equal t	o			
	(a) 90°	(b) 104°	(c) 120°	(d) 109°.281		
(95)	The correct order of di	pole moment is				
	(a) $CH_4 < NF_3 < NH_3$	< H <sub>2</sub> O	(b) $NF_3 < CH_4 < NH$	$_{3}$ $<$ $\mathrm{H}_{2}\mathrm{O}$		
	(c) $NH_3 < NF_3 < CH_4 <$	H <sub>2</sub> O	(d) $H_2O < NH_3 < NF$	CH <sub>4</sub>		
(96)	The correct order of the	e O – O bond length in	$O_2$ , $H_2O_2$ and $O_3$ is			
	(a) $O_2 > O_3 > H_2O_2$			>		
	(c) $H_2O_2 > O_3 > O_2$	(d) $O_2 > H_2 O_2 > O_3$				
(97)	The bond order of $O_2^-$	is				
	(a) 1.0	(b) 1.5	(c) 2.5	(d) 0.5		
(98)	Choose the incorrect st	tatement.				
	(a) $\sigma$ bond is weaker the	nan π bond				
	(b) $\pi$ bond is weaker than $\sigma$ bond					
	(c) $\pi$ bond is present a	along with a $\sigma$ bond				
	(d) $\sigma$ bond can be pres	ent alone				
(99)	Which of the following	is not paramagnetic?				
	(a) NO	(b) S <sup>2-</sup>	(c) O <sub>2</sub>	(d) $N_2^-$		
(100)	Which one of the follow	wing compound has sp <sup>2</sup>	hybridization?			
	(a) CO <sub>2</sub>	(b) SO <sub>2</sub>	(c) CO	(d) $N_2O$		

## **Answer key**

1 d	2 c	3 c	4 b	5 c	6 b
7 d	8 b	9 c	10 c	11 c	12 a
13 c	14 d	15 b	16 a	17 c	18 a
19 d	20 b	21 d	22 b	23 b	24 c
25 b	26 b	27 c	28 b	29 b	30 b
31 b	32 a	33 c	34 c	35 b	36 c
37 b	38 b	39 a	40 b	41 c	42 d
43 c	44 d	45 c	46 a	47 b	48 d
49 d	50 c	51 d	52 b	53 d	54 b
55 d	56 c	57 b	58 d	59 c	60 a
61 b	62 d	63 d	64 c	65 a	66 b
67 b	68 a	69 b	70 d	71 c	72 b
73 a	74 b	75 a	76 a	77 c	78 d
79 c	80 a	81 c	82 d	83 c	<b>&gt;</b> 84 b
85 c	86 a	87 a	88 b	89 c	90 d
91 c	92 b	93 a	94 d	95 a	96 c
97 d	98.a	99 b	100 b	1	

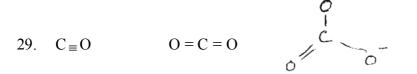
## **Hints**

- 1. Electronegative difference in KI is more
- 2. ACC to V.B.T, O<sub>2</sub> contain all e<sup>-</sup> paired. So they are diamagnetic
- 6. Here C C contain double and triple bond
- 7.  $NO_3^-$  and  $CO_3^{-2}$  contains 32 e<sup>-</sup> so they are iso electric ions Both possess sp<sup>2</sup> hybridization
- 10. In  $H_2O_2$ , O H is polar

O - O is non-polar

- 11. H-O-N = 0
- 16. Struchure A is stable
- 18. In NF<sub>3</sub>, polarity of N F bond and non-bonding e<sup>-</sup> pair are in opposite direction
- 19. In CH<sub>3</sub>Cl, dipole moment is due to C-Cl and C-H

trans - pent - 2 - ene possess magnetic moment



- 30. In H<sub>2</sub>O, O Contain two lone pair of e<sup>-</sup> H Contain positive change
- 40. Bond order  $O_2^+ = 2.5$   $O_2 = 2$   $O_2^- = 1.5$   $O_2^{-2} = 1$
- 42. In H<sub>2</sub>O, O contain two lone pair of e<sup>-</sup> so repulsion is more

46. 
$$\Delta_f H^o = \Delta_{sub} H^o + \Delta_D H^o + \Delta_i H^o + \Delta_{eg} H^o + \Delta_u H^o$$

61. Theorectical dipole momentum

$$\begin{aligned} &H = q \times d & \left[ 1 \ D = 1 \times 10^{-18} \, \text{esa cm} \right] \\ &= 4.8 \times 10^{-10} \times 1.275 \times 10^{-8} \\ &= 6.12 \times 10^{-10} \, \text{e.s.u.cm} \\ &= 6.12 \, D \end{aligned}$$

% of ionic character = 
$$\frac{\text{Practical}(\mu)}{\text{Theoretical}(\mu)}$$
$$= \frac{1.03}{6.12} \times 100$$
$$= 16.83$$
$$\approx 17\%$$