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_2 . 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 called bond angle. As you know the bond lengths of single (-) bond, double (=) bond and triple (=) bond are different. The bond angle gives geometrical shapes to molecules viz.180⁰- linear, $109^{0}28'$ tetrachedral. You will study in detail about covalent bond which can be of three types. (1) Polar covalent bond in which the electron remains dragged towards the more electronegative atom and $+\delta$ charge on electropositive atom and $-\delta$ charge on electronegative atoms are shown. As a result the molecule becomes polar. If the electronegativities of the two atoms are same or the difference between them is less, than non-polar bonds, formed by both the atoms sharing the electrons. In coordinate covalent bond, one of the two atoms sharing a pair of electrons and the second atom completes the octet with the help of this gained electron pair. viz. $F_3B \leftarrow NH_3$. Bond \leftarrow indicates co-ordinate covalent bond. Over and above, bond length, bond angle, bond enthalpy (bond energy) is also an important concept. Shorter the bond length, more will be the stability and so more energy will be required to break it. Thus, the values of enthalpy may be different according to bond formation. The number of bonds is called bond order which we have studied in detail and also the formula to determine it. Born-Haber showed that the enthalpy evolved in formation of compound is the mathematical results of the enthalpies of several reactions of atoms. It is explained in the unit by discussing the formation of compound like NaCl.

Sometimes, it so happens that the electron pair instead of being localised on any molecule it localises towards other molecule. Thus, the bonds in the molecule can be shown at different positions in the compounds having same molecular formula. Such structures are called resonance structures and energy associated with the changes of these resonance forms is called resonance energy. This can be studied through the molecules of ozone, carbon dioxide, benzene etc.

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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 $\text{BeCl}_2 - 180^\circ$, trigonal $\text{BCl}_3 - 120^\circ$, tetrahedral CH₄ - $109^\circ 28'$.

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 109⁰28' but it becomes 104⁰ 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 $-\delta$ and the distance between them is to be utilised. More the value of dipole moment, more will be the polar bond and more will be the ionic bond. One important aspect is that polar substances dissolve only in polar solvents and non-polar substances dissolve in nonpolar solvents. e.g. NaCl will dissolve in water. Napthalene will dissolve in benzene. New hypotheses have been presented affter taking into consideration the limitations of the principles for the approach of covalent bond. Two are main from them : (1) Valence Bond Theory and (2) Molecular Orbital Theory. These concepts are based on quantum mechanics. Heitler and London first of all gave the idea of valence bond theory and it was developed by Pauling and Slater.

In the assumptions of valence bond theory the attraction - repulsion forces between positively charged nuclei of two atoms and the electrons arranged in the orbits around them. According to Coulomb's Law if attractive forces are more than repulsive forces then the bond will be formed and molecule will be formed. In this theory, on the basis of the overlapping of valence orbitals different overlaps can be formed. In this type of overlapping the excitation of electrons in valence orbitals can be shown and then formation of molecule by covalent bond with other atoms. viz. In carbon, the electrons of valence orbital $1s^2 2s^2 2p^2$ will be excited to give $1s^2 2s^1 2p_x^{-1} 2p_y^{-1} 2p_z^{-1}$ containing four orbitals with one electron in each and four hydrogen atoms, and hence will give stable molecule like CH₄. The geometrical structure, and bond angle can be expressed from the hybridisation associated with it. In such valence bonds, two types of bonds– σ and π are also observed. σ bond is a covalent bond; it



attains axial overlap of internuclear axis. The stability of this bond is more than that of π bond. In the π -bond the axis of the atomic orbitals undergoing overlapping remains parallel to each other and is perpendicular to internuclear axis. π - bonds are less stable in comparison to σ -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_2 , Be_2 , F_2 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 (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 for formation of homonuclear molecules from H_2 to Ne_2 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 N₂ mol-

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

molecule bond order will be = $\frac{1}{2}$ [10 - 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_2 . If the value of bond order is integer, the bond will be formed and according to the integer 1, 2, or 3, there will be single (-), double (=) or triple (=) bonds respectively. If the value of **bond order is fraction, then molecule will attain unstable structure.** The molecular orbital theory can explain the magnetic properties e.g. In O₂ molecule two unpaired electrons are there and so it is paramagnetic and in N₂ molecule, all the electrons are paired and so it is diamagnetic. Thus molecular orbital theory is superior to valence bond theory in this matter.

Above this, one important phenomenon is hydrogen bond. The first element of 15, 16, and 17, groups N, O, F being higer electronegative than the other elements of the group it can form covalent molecules like NH_3 , H_2O and HF with hydrogen. Afterwards the molecule like H - F molecule through hydrogen bond

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 H_{3} , $H_{2}O$ possess hydrogen bonds and so their properties are different from those of other elements in the group. Hydrogen bond is of two types (1) Intermolecular and (2) Intramolecular hydrogen bonds. When hydrogen bond is formed between two molecules it is called intermoleculer hydrogen bond e.g. p-chlororphenol and between two groups in the same molecule, it is called intramoleculer hydrogen bond e.g. o-chlorophenol. Intermolecular hydrogen bond is stronger than intramolecular hydrogen bond. The presence of hydrogen bond is the reason for specific properties of the compounds. Viz the retaining of water in the soil, drying of terrylene clothes is faster than that of cotton clothes.

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, ductility, 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 BF₃, three F atoms were bonded with B-atom through three covalent bonds but the octet of B is not complete. Similary in NH₃ 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 BF₃ 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 coordinate covalent bond in the unit of complex salts in Standard-12.

M.C.Q.

(1)	Which of the following is ionic ?						
	(a) HCl	(b) CHCl ₃	(c) IF ₅	(d) KI			
(2)	When molecule is form	n by chemical bonding b	between atoms then				
	(a) nucleous of combined	(a) nucleous of combining atoms are participate					
	(b) valence electrons and inner cell electrons are participate						
	(c) only valence electro	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 energy	gy	(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	alence-bond theory ?					
	(a) Mulliken	(b) Lenus Pauling	(c) Hittler and Londa	ın (d) Hund			
(6)	In H - C \equiv C - CH	= CH_2 molecule C^3 -	C ² single bond carb	ons has which type of			
	hybridization?						
	(a) $sp^2 - sp^3$	(b) sp - sp ²	(c) $sp^3 - sp$	$(d) sp^3 - sp^3$			
(7)	Which of the followin	g pair of species is isoele	ectronic and same strue	cture ?			
	(a) NO_3^- , SO_3	(b) SO_3 , CO_3^{2-}	(c) CO_3^{2-} , $ClO_3^{}$	(d) NO_{3}^{-} , CO_{3}^{2-}			
(8)	Which of the followin	g sentence is incorrect fo	or covalent bond?				
	(a) Strenght of covalent bond depenas upon overlapping at atomic orbitals.						
	(b) Covalent bond is n	ot directional.					
	(c) There is sharing of	electrons between atom	is bonded by covalent	bond			
	(d) Covalent bond is formed between atoms having less difference in their electronegativity.						
(9)	Which of the following	g compound possesses c	covalent bond?				
	(a) MgCl ₂	(b) NaH	(c) BF_3	(d) CsCl			
(10)	Which of the following	g molecule possesses po	lar and nonpolar coval	lent bond?			
	(a) NH_4Cl	(b) CCl_4	(c) H_2O_2	(d) HCN			
(11)	Which of the following	g compound does not p	ossesses coordinate co	valent bond ?			
	(a) CO	(b) SO ₂	(c) HNO ₂	(d) HNO ₃			
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(12) Which of the following characteristic is not for covalent compound?

(a) They do not possesses particular geometical structure

- (b) They may be polar or nonpolar
- (c) Their boiling and melting point is low
- (d) Generally they are insoluble in water
- (13) Which of the following possesses ionic and covalent bond?

(a)
$$CO_2$$
 (b) H_2SO_4 (C) NH_4Cl (D) NaI

- (14) Whhat is Geometrical Structure of ClF_3 molecule ?
 - (a) Triogonal bipyramid (b) Corn shpae (c) sea-saw (d) T-shape
- (15) Which of the following molecule possesses linear structure?

(a) SO_2 (b) CO_2 (c) H_2O (d) C_2H_4

(16) Correct structure of SF_4 is





(23)	Which is the paramagnetic species ?				
	(a) \bar{CN}	(b) \overline{O}_2	(c) NO ⁺	(d) CO	
(24)	24) Which of the following statement is incorrect when N ₂ and O ₂ are convert into N ₂ ⁺ and O respectively ?				
	(a) In O_2^+ , O - O bond	l order increases.	(b) In N_2^+ , N - N bo	nd become weaker.	
	(c) N_2^+ 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 sci	in valence shell electron entist ?	during the bond form	ation. This postulate was	
	(a) Powel	(b) Lewis	(c) Sigdwick	(d) Mulliken	
(27)	Crystal formation is wh	hich type of reaction ?			
	(a) endothermic and ex	tothermic	(b) endothermic		
	(c) exothermic		(d) no heat change of	ccurs	
(28)) Lattice energy of ionic compound depends upon 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 in CO, CO_3^{2-}, CO_2				
	(a) $CO_3^{-2} < CO_2 < CO$		(b) $CO_2 < CO_3^{2-} < CO_3^{2-}$	20	
	(c) $CO < CO_2 < CO_3^{2-}$		(d) $CO < CO_{3}^{2-} < CO_{3}^{2-}$	O_2	
(30)) Maximum how many numbers of hydrogen bond can be form by H_2O 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 ² hybridised carbon atoms				
	(c) Two sp ³ and two sp	² hybridised carbon atom	ms		
	(d) sp, sp ² and sp ³ hyb	ridized carbon atoms			

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(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) σ - bond can form by overlapping of S - P orbitals. (c) σ - bond can form by overlapping of end of atomic orbitals of inner center axis. (d) This type of overlapping is also known as axial overlaping (33) In which molecule inter molecular hydrogen bond can be form? (a) methanol (b) ethelene glycol (c) p - nitrophenol (d) phenol (34) In which molecule intra molecular hydrogen bond can be form? (d) all of these (a) o - nitro phenol (b) aniline (c) ethylene glycol (35) Which of the following pair possesses very strong H - bond? (a) CH₂ COCH₂ and CHCl₂ (b) HCOOH and CH₃ COOH (c) H_2O and H_2 (d) SiH_4 and $SiCl_4$ (36) Which of the following relation is correct? (a) Bond order α Bond energy α Bond length α stability $\frac{1}{\text{Bond length}}\alpha = \frac{1}{\text{energy}}\alpha$ stability (b) Bond order α (c) Bond order a Bond energy $\alpha = \frac{1}{\text{Bond length}} \alpha$ stability $\frac{1}{\text{Bond length}} \alpha = \frac{1}{\text{Bond energy}} \alpha$ stability (d) Bond ordera Br₂ (37) Molecule : F_2 Cl, Η, Bondlength : 74pm 144pm 199pm 228pm Mention more stable molecule from above (a) Cl₂ (c) Br_2 (b) H₂ (d) F_{2} (38) In water bond angle is 104° 30 because (a) Oxygen atom is sp³ hybridised (b) Repulsion between lone pair election and bonding pair electron (c) Oxygen has high electronegetivity. (d) H_2O molecule possesses "V" - shape.

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(39) In which of the following strong H-bond is present? (a) F - H....F (b) O - H....N (c) O - H....O(d) O - H....F (40) Which is correct order for bond dissociation energy in O_2 , O_2^+ , O_2^- and O_2^{2-} (a) $O_2 > O_2^+ > O_2^{2-} > O_2^-$ (b) $O_2^+ > O_2 > O_2^- > O_2^{2-}$ (c) $O_2^- > O_2^{2-} > O_2^+ > O_2^+$ (d) $O_2^- > O_2^{2-} > O_2 > O_2^+$ (41) O, P, Q, R elements electronic configuration is given below $O = 1s^2$ $P = 1s^2 \cdot 2s^2 \cdot 2p^2$ $Q = 1s^2$, $2s^2$, $2p^5$ $R = 1s^2$, $2s^2$, $2p^6$ Which atom has strong behaviour of electrovalent bond ? (c) Q (a) O (b) P (d) R (42) In which molecule bond distorsion is more according to VSEPR theory? (a) SO₂ (b) NH₃ (c) O_{3} (d) H₂O (43) Which of the following species is more stable? (a) O_{2}^{-} (b) Ne_{2}^{+} (c) O_{2} (d) F_{2} (44) Number of nonbonding electron pair in XeF_6 , XeF_4 and XeF_2 respectively (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 which become one cube which factor is responsible for it? (a) Van-der waals attraction (b) Hydrogen bond (c) Dipole attraction (d) Covalent bond (46) Determine lattice energy of $LiF_{(S)}$ according to given data. 155.2 KJ mol $^{-1}$ (Δ_{c} H) (i) $Li_{(S)} \rightarrow Li_{(g)}$ 75.2 KJ $\Delta \left(\frac{H_{D}}{2} \right)$ (ii) $F_{2(\alpha)} \rightarrow 2F_{(\alpha)}$ $(iii) Li_{(g)} \rightarrow Li^+_{(g)} + e$ 520.0 KJ mol Δ^1 H(;) (iv) $F_{(g)}^{+\bar{e}} \rightarrow \overline{F}_{(g)}$ $-33.0~\text{KJA}(\ \mbox{H}_{g}\)$ (v) $\operatorname{Li}_{(s)} + \frac{1}{2} F_{2(g)} \rightarrow \operatorname{Li}F_{(S)}$ -504.1 KJ mol $(\Delta_f H)$ (a) $-86.7 \,\text{KJ}\,\text{mol}^{-1}$ (c) -867 KJ mol -1 (d) 867 KJ mol -1(b) 86.7 KJ mol $^{-1}$ 87

(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 can change their position easily in crystal						
	(d) Explanation of met	tallic 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	ositive and negative ion energy increases.	is small than then inte	erionic attraction become		
	(c) In ionic crystal whe	en size of positive ion de	ecrease, then lattice end	ergy increases.		
	(d) All of given					
(49)	Number of H - bond f	form by unpaired electro	ons of liquid NH_3 , H_2C	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) $CH_4 > SiH_4$	$(d) \operatorname{NH}_3 > \operatorname{PH}_3$		
(51)	Which of the following	g compound possesses is	onic bond ?			
	(a) CH ₄	(b) SiCl ₄	(c) BF_3	(d) $MgCl_2$		
(52)) Which of the following relation between BMO and ABMO electrons is correct for stability of diatomic malecule or ion ?					
	(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) 4.5×10^{-10} m	(b) 0.45nm	(c) 4.5 A ^o	(d) Given all		
(54)	What is bond energy of	of H-bond ?				
	(a) 40 J mol ⁻¹	(b) 40 KgJ mol ⁻¹	(c) 40 cal. mol ⁻¹	(d) 40 Kg cal mol ⁻¹		
(55)	In which molecule inte	er molecular H-bond is p	ossible ?			
	(a) CH ₃ COCH ₃	(b) CH ₄	(c) SiH ₄	(d) NH ₃		
(56)	Which of the following	g characterstic does not	possesses by metal?			
	(a) luminus			(b) ductility		
	(c) increase in conduct	ance by increase in tem	perature	(d) malleability		

(57) On which factor conductance of metals responsible?

(a) ions(b) delocalised(c) atomic kernel(d) number of atoms(58) Which of the following figure shows electron-sea model?



- (59) According to which group, hydrogen bond is form in protein molecule present in musecls of living organism ?
 - (a) -CO- (b) -COOR (c) -CONH- (d) -COOH
- (60) On which factor van-der waalls attraction force does not depend ?
 (a) numbers of molecules
 (b) contact surface area of molecules
 (c) shape of molecules (d) numbers of electron in molecules
- (61) Practicol dipal movement of HCl is 1.03D. If bond length of HCl is 1.275 A° than what will be the percentage of ionic nature in HCl?
 - (a) 7 (b) 17 (c) 43 (d) 21
- (62) Which sentence is correct with respect to bond enthalpy ?
 - (a) As bond order is more, then bond dissociation enthalpy is less
 - (b) As atomic volume is more, then bond energy is more.
 - (c) As bond enthalpy is more, then stability of molecule or ion is less.
 - (d) As number of nonbonding election pair on bonded atom then bond enthalpy is less.
- (63) which of the following orbitals form bonding orbital?



(a) 4, 4 (b) 2, 4 (c) 4, 2 (d) 2, 2

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(66) Mention proper order of bond length given below.

(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 compound given below. (c) $N_{2}O$ (d) Na₂O₂ (a) O_{3} (b) KO, (68) Which species possesses pyramidal shape? (d) SiO_3^{2-} $(b) SO_{2}$ (a) OsF, (c) BrF_3 (69) Which of the following does not possesses bond order as CO? (c) N_{2} (a) NO^{+} (b) NO^{-1} (d) \overline{CN} (70) Which rule is violated in the given electronic configuration? 2P(a) Aufbau (b) Pauli (c) Hund (d) Given all (71) In which of the following molecule double bond possesses two pispace bond? (a) S_{2} $(b) O_{2}$ (c) C_{2} (d) $H_2C = CH_2$ (72) Mention AB_4F_2 type molecule. (b) XeF_4 (c) SF_6 (d) $XeOF_4$ (a) BrF_{5} (73) Which of the following is the correct order for lone pair and bonding pair electrons? Lp = Lone pair and Bp = Bonding pair

- (a) Lp Lp > Lp Bp > Bp Bp (b) Lp Bp > Lp Lp > Bp Bp
- (c) Bp Bp > Lp Lp > Lp Bp (d) Lp Lp > Bp Bp > Lp Bp

(74) Which theory is useful to determine geometrical structure of molecules ?

- (a) molecular orbital theory (b) VSEPR theory
- (c) Resonance theory (d) Quantam mechanics
- (75) The one outermost electron present in Na element at
 - (a) one corner of simple cube (b) eight corner of simple cube
 - (c) center of simple cube (d) each corner of simple cube

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(76)	(76) In which molecules / ion have not all the equal bonds ?					
	(a) SF ₄	(b) BF ₄ ⁻	(c) XeF ₄	(d) SiF ₄		
(77)	77) Which of the following has maximum bond angle?					
	(a) NH ₃	(b) CH ₄	(c) CO ₂	(d) H ₂ O		
(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 $CuSO_4$.5 H_2O				
	(a) covalent and co-or	dinate covalent	(b) electrovalent and	covalent		
	(c) electrovalent and c	o-ordinate covalent				
	(d) electrovalent, cova	lent and co-ordinate cov	valent			
(80)	Which of the following	statement is wrong				
	(a) sp ² hybrid orbitals	(a) sp ² hybrid orbitals are formed from two p - atomic and one s - orbitals				
	(b) hybridization is the	mixing of atomic orbital	ls prior to their combin	ig into molecular orbitals		
	(c) d^2p^2 hybrid orbitals are all at go to one an other					
	(d) d ² sp ³ hybrid orbita	ls are directed towords t	he corners of a regula	r tetrahedron		
(81)	CO_2 is isostructual with	1				
	(a) SnCl ₂	(b) HgCl ₂	(c) $C_2 H_2$	(d) SO_2		
(82)	NH ₃ has a higher boilir	ng point than expected b	ecause			
	(a) its density decrease	es on freezing				
	(b) with water it forms	NH ₄ OH				
	(c) it has strong inter n	nolecular covalent bonds	s ?			
	(d) it has intermolecula	r hydrogen bonds.				
(83)	The molecule with zero	o dipole moment is				
	(a) chloroform		(b) methyl chloride			
	(c) carbon tetrachlorid	e	(d) methylene chlorid	le		
(84) Molecular shaper of SF_4 , CF_4 , XeF_4 are						
	(a) the same with 1, 1 and 1 lone pairs of electrons respectively					
	(b) different with 1, 0 a	and 2 lone pairs of electr	rons respectively			
	(c) different with 0, 1	and 2 lone pairs of elect	rons respectively			
	(d) different with 2, 0 a	and 1 lone pairs of electr	rons respectively			

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(85)	Which of the following has the regular tetrahedral structure?				
	(a) SF ₄	(b) $[Ni(CN)_4]^{2-}$	(c) BF_{4}^{-}	(d) XeF ₄	
(86)	In OF ₂ , number of bon	nd pairs and lone pairs o	of electrons are respect	ively	
	(a) 2, 8	(b) 2, 6	(c) 2, 9	(d) 2, 10	
(87)	In O_2^- , O_2^- , O_2^{2-} molec	ular species the total nu	mber of antibonding e	lectrons 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 Ar	nswer		
	Column - I	Column -II			
	(i) $sp^{3}d^{2}$	(a) Ni $[(CN)_4]^{2-}$			
	(ii) sp ³ d	(b) SnCl ₂			
	(iii) dsp ²	(c) IC $\overline{l_4}$			
	(iv) sp ²	(d) TeCl ₄			
	(a) $i \rightarrow d$, $ii \rightarrow a$, $iii \rightarrow a$	\rightarrow c, iv \rightarrow b	(b) $i \rightarrow c$, $ii \rightarrow d$, ii	ii \rightarrow a, iv \rightarrow d	
	(c) $i \rightarrow b$, $ii \rightarrow c$, $iii \rightarrow d$, $iv \rightarrow a$ (d) $i \rightarrow a$, $ii \rightarrow b$, $iii \rightarrow c$, $iv \rightarrow d$				
(89)	Among the following compounds, the one that is polar and has the central atom with sp hyridisation is				
	(a) HClO ₂	(b) BF ₃	(c) H_2CO_3	(d) SiF_4	
(90)	Match the following				
	Set A		Set B		
	(1) stability of bond		(p) Bond energy		
	(2) Molecular orbital theory		(q) Bond order		
	(3) octet rule		(r) Variable Valency		
	(4) Valence bond theory		(s) Electronic concept of valency		
	(a) $1 \rightarrow q, 2 \rightarrow p, r,$	$3 \rightarrow p, 4 \rightarrow s$	(b) $1 \rightarrow p, q, 2 \rightarrow p, 3 \rightarrow r, 4 \rightarrow s$		
	(c) $1 \rightarrow p, q, 2 \rightarrow r$,	$3 \rightarrow s, 4 \rightarrow r$	(d) $1 \rightarrow p, q, 2 \rightarrow q, 3 \rightarrow s, 4 \rightarrow r$		
(91)	Bond strength increase	es with			
	(a) Bond length increase	sing			
	(b) Antibonding eletron	ns being higher in numbe	er		
	(c) Bond order increas	sing	(d) Bond angle incre	asing	
		_	`		
		<u>92</u>	/		

(92) O_2^{2+} will have

	(a) Bond order lower than O	2			
	(b) Bond order higher	than O ₂			
	(c) Bond order lower	than H ₂			
	(d) Bond order higher	than N ₂			
(93)	In a molecule number	of electrons in BMO ar	e more as compared to	ABMO, hence	
	(a) a bond will be form	ned			
	(b) no bond will be for	med			
	(c) information is not s	ufficient			
	(d) none of the above				
(94)	The bond angle in the a	ammonium ion is equal	to		
	(a) 90°	(b) 104°	(c) 120°	(d) $109^{\circ}.28^{1}$	
(95)	The correct order of di	pole moment is			
	(a) $CH_4 < NF_3 < NH_3$	< H ₂ O	$(b) NF_3 < CH_4 < NH$	$I_3 < H_2O$	
	(c) $NH_3 < NF_3 < CH_4$	< H ₂ O	(d) $H_2O < NH_3 < NF_4 < CH_4$		
(96)	The correct order of th	ne O – O bond length ir	O_2, H_2O_2 and O_3 is		
	(a) $O_2 > O_3 > H_2O_2$	(b) $O_3 > H_2 O_2 > O_2$			
	(c) $H_2O_2 > O_3 > O_2$	(d) $O_2 > H_2O_2 > O_3$			
(97)	The bond order of O_2^-	is			
	(a) 1.0	(b) 1.5	(c) 2.5	(d) 0.5	
(98)	Choose the incorrect s	tatement.			
	(a) σ bond is weaker t	han π bond			
	(b) π bond is weaker	than σ bond			
	(c) π bond is present	along with a σ bond			
	(d) σ bond can be presented as the presented of the pr	sent alone			
(99)	Which of the following	g is not paramagnetic ?			
	(a) NO	(b) S ²⁻	(c) O_2^-	(d) N_2^-	
(100)) Which one of the follo	wing compound has sp ²	hybridization ?		
	(a) CO ₂	(b) SO ₂	(c) CO	(d) N_2O	

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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 с	18 a
19 d	20 b	21 d	22 b	23 b	24 c
25 b	26 b	27 с	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	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		

Answer key

Hints

- 1. Electronegative difference in KI is more
- 2. ACC to V.B.T, O_2 contain all e⁻ paired. So they are diamagnetic
- 6. Here C C contain double and triple bond
- 7. NO_3^- and CO_3^{-2} contains 32 e so they are iso electric ions Both possess sp² 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_3 , polarity of N F bond and non-bonding e pair are in opposite direction
- 19. In CH_3Cl , dipole moment is due to C-Cl and C-H

20. etty C=0

trans - pent - 2 - ene possess magnetic moment

- 21. Compound $SO_2 H_2O H_2S NH_3$ Bond angle 119.5° 104.5° 92.5° 106.5° 29. $C \equiv O$ O = C = O
- 30. In H_2O , O Contain two lone pair of e^- H Contain positive change
- 40. Bond order $O_2^+ = 2.5$ $O_2 = 2$ $O_2^- = 1.5$ $O_2^{-2} = 1$
- 42. In H₂O, O contain two lone pair of e⁻ 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

 $H = q \times d \qquad \begin{bmatrix} 1 & D = 1 \times 10^{-18} \text{ esa cm} \end{bmatrix}$ = 4.8×10⁻¹⁰×1.275×10⁻⁸ = 6.12×10⁻¹⁰ e.s.u.cm = 6.12 D

% of ionic character = $\frac{Practical(\mu)}{Theoretical(\mu)}$

$$= \frac{1.03}{6.12} \times 100$$

= 16.83
\$\approx 17\%\$