CHEMICAL BONDING AND MOLECULAR STRUCTURE

TYPES OF BONDS

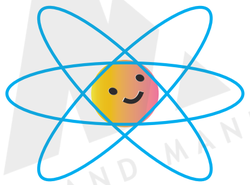
IONIC/ELECTROVALENT BOND
- Strong electro static force of attraction between positive and negative ions.
- Crystalline in Nature
- High M.P and B.P
- Soluble in Polar Solvents. Eg: NaCl, $MgCl_2$ etc.

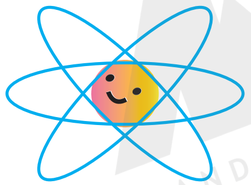
HYDROGEN BOND
- Bond formed when the -ve end of one molecule attracts the +ve end of H.
1. Intermolecular: H-Bonding occur within one single molecule.
2. Intermolecular: H Bonding between two different molecules of same or different compounds.

COVALENT BOND
- Bond formed by mutual sharing of e^-.
- Low M.P. and B.P.
- Bad conductor of electricityv
- Insoluble in Polar Solvents but Soluble in Non-Polar Solvent. Ex: CH_4, H_2, Cl_2.

CO-ORDINATE BONDING
- Bond formed by one sided sharing of electrons. ie: one atom donates a
- Pair of e^- while other accepts it.
- Bad conductors of electricity.
- Sparingly Soluble in Polar Solvents but readily Soluble in Non-Polar Solvents. eg:- NH_4^+

TYPE OF CO-VALENT BOND
- Polar covalent bond Eg: NH_3, $CHCl_3$
- Non-Polar covalent bond Eg: Cl_2, CO_2.





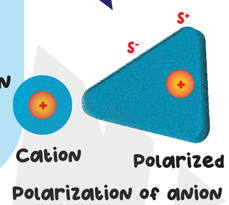
FAJAN'S RULE

No compounds is 100% ionic or 100% covalent

. Covalent Nature $\propto$ Charge on cation

. Covalent Nature $\propto \frac{1}{\text{size of cation}}$

• Polarising Power $\propto$ Covalent C



Cation — Polarized anion

Polarization of anion by cation

THEORIES OF CO-VALENT BOND

KOSSEL LEWIS APPROCH

Atoms can combine either by transfer of e- or by sharing of valence e- in order to have an Complete octet in their valence shell.

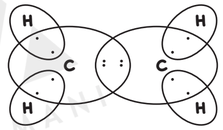
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Octet Rule

LEWIS SYMBOLS

Valence e- are represented by dots around the element.

H: Na: :Cl: ·O·. :N:



VALENCE BOND THEORY (VBT)

A covalent bond is formed by the overlapping of two half filled atomic orbitals.

Type of overlapping

Sigma(σ) Sidewise overlapping

• S-S overlapping

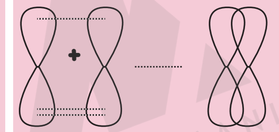


• S-P overlapping



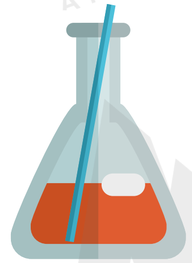


Pi(π) Sidewise overlapping



HYBRIDISATION

Concept of mixing atomic orbital to form new hybrid





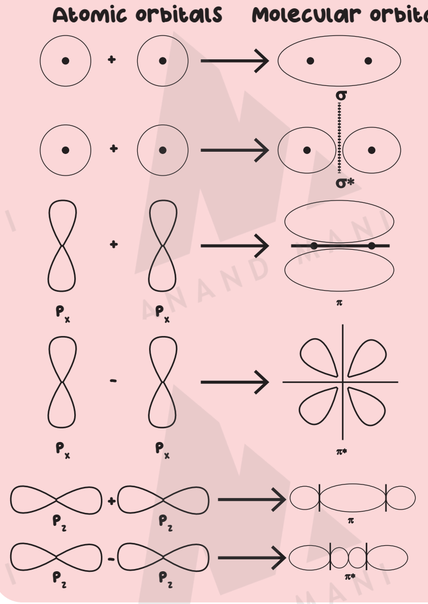
VSEPR THEORY

The Shape of a molecule depends upon the numbers of valence Shell e^- (B.P or L.P) surrounding in the central atom.

Decreasing order of repulsive interaction: lp - lp > lp - bp > bp - bp

Type of Molecule	No. of Bonding Pair	No. of Lone Pair	Arrangement of e- Pair	Shape	Example
AB_2E	2	1	Trigonal planer	Bent	SO_2O_3
AB_3E	3	1	Tetrahedral	Trigonal Pyramidal	NH_3
AB_3E_2	2	2	Tetrahedral	Bent	H_2O
AB_4E	4	1	Trigonal bi-pyramidal	See Saw	SF_4
AB_3E_2	3	2	Trigonal bi-pyramidal	T-Shape	ClF_5
AB_5E	5	1	Octahedral	Square Pyramid	XeF_5
AB_4E_2	4	2	Octahedral	Square Planner	XeF_4

Formation of Molecular Orbitals



MOLECULAR ORBITAL THEORY

- MOT States that each atom tends to combine together and form molecular orbitals
- No. of molecular orbitals = No. of atomic orbital combined.

MOT

- Bonding molecular orbitals
- Anti bonding atomic orbitals

ELECTRONIC CONFIGURATION

Electron filling order upto 14 electrons

$$\sigma_{1s} < \sigma^*_{1s} < \sigma_{2s} < \sigma^*_{2s} < \pi_{2p_x} \equiv \pi_{2p_y} < \sigma_{2p_z} < \pi^*_{2p_x} \equiv \pi^*_{2p_y} < \sigma^*_{2p_z}$$

Electron filling order for more than 14 electrons

$$\sigma_{1s} < \sigma^*_{1s} < \sigma_{2s} < \sigma^*_{2s} < \sigma_{2p_z} < \pi_{2p_x} \equiv \pi_{2p_y} < \pi^*_{2p_x} \equiv \pi^*_{2p_y}$$

BOND PARAMETERS

FORMAL CHARGE

$$FC = V - N - \frac{B}{2}$$

- Bond length: distance between the Nuclei of two bonded atom

bond length $\propto \frac{1}{\text{bond order}}$

BOND ORDER & BOND ANGLE

BOND ORDER: No. of Bond between the two atoms

BOND ANGLE: Angle between the orbitals containing bonding e^- pair around central atom.

BOND ENTHALPY

Amount of energy required to break one mole of bonds.

DIPOLE MOMENT

Product of the magnitude of the charge and distance between centres of positive and negative charge.

μ = charge × Distance Between two atoms