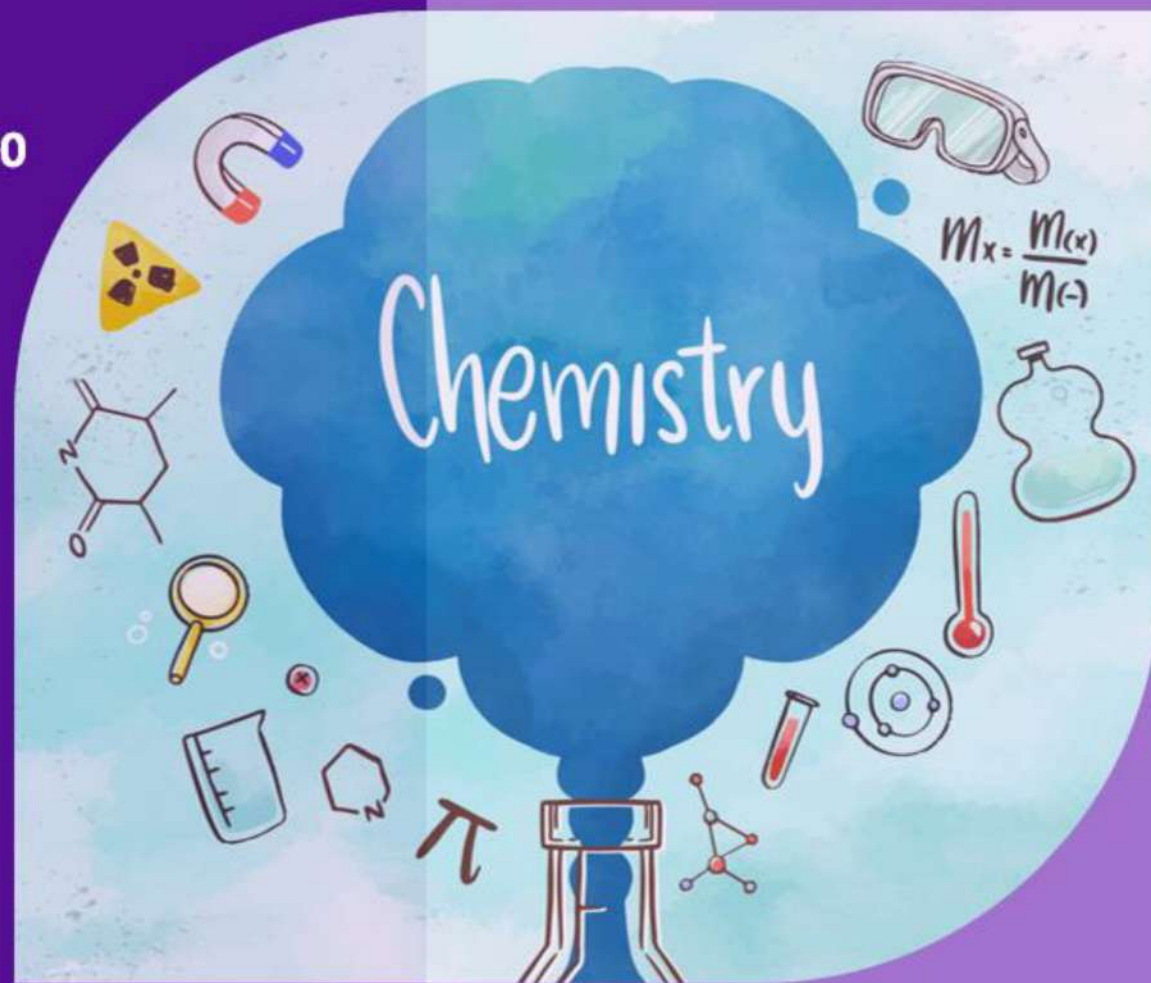


MEDICAL AND DENTAL ADMISSION PROGRAM 2020

CHEMISTRY

LECTURE : C-02

CHAPTER 03 : CHEMICAL BOND



উন্মেষ

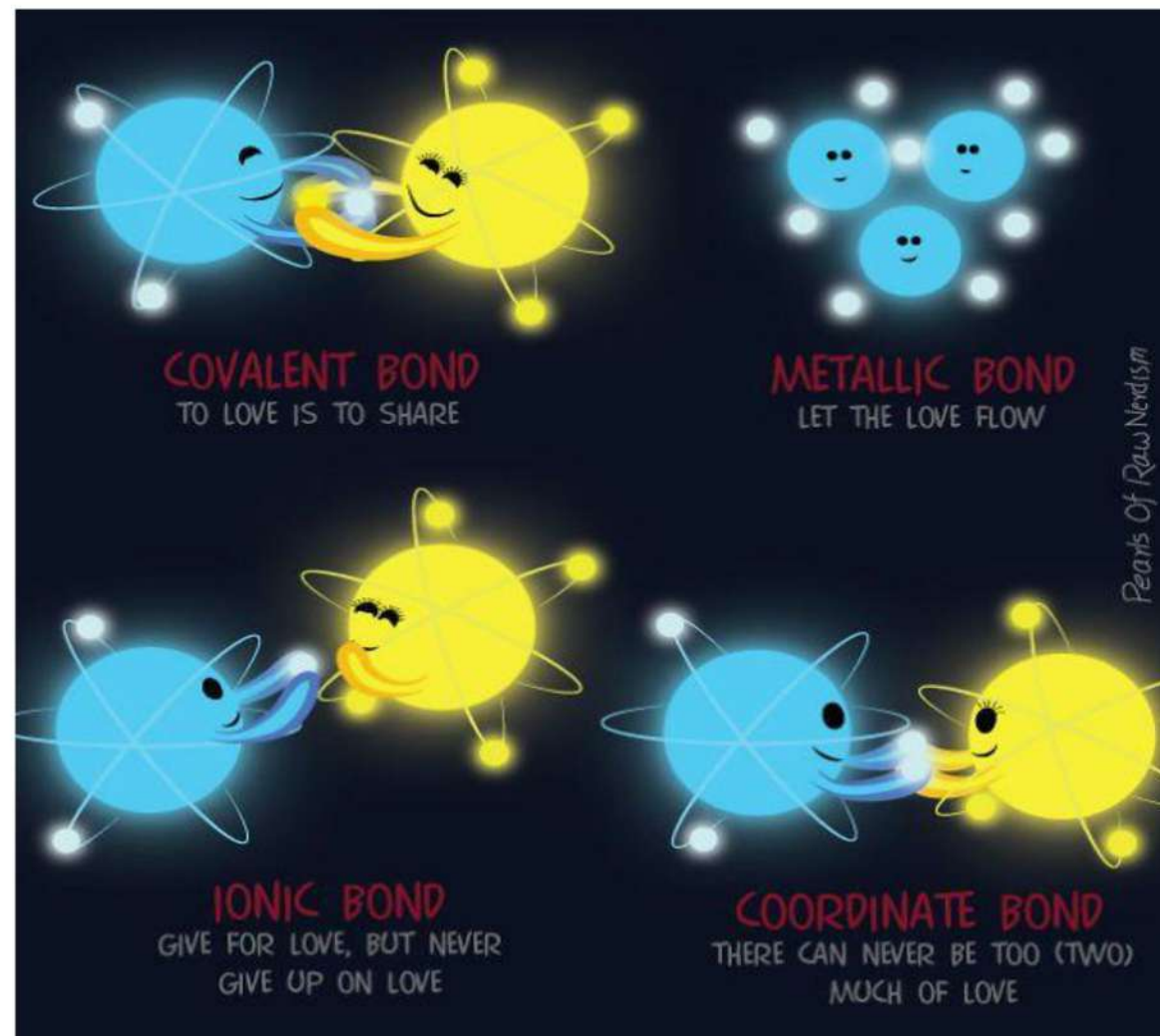
মেডিকেল এন্ড ডেন্টাল এডমিশন প্রোগ্রাম

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Important topic from this chapter for Medical and Dental admission test

Importance	Topic	Admission test years
★★★	Chemical bonds	MAT: 19-20, 14-15, 13-14, 11-12, 09-10, 07-08, 06-07, 05-06, 04-05, 00-01; DAT: 19-20, 17-18, 10-11, 08-09, 03-04, 02-03, 00-01
★★	Hybridization of orbitals	MAT: 14-15, 02-03; DAT: 04-05, 02-03
★	Polarity of covalent bonds and covalent characteristics of ionic compounds	MAT: 10-11; DAT: 01-02
★★	Nomenclature of inorganic compounds	MAT: 07-08, 05-06; DAT: 16-17, 07-08

Chemical Bond:



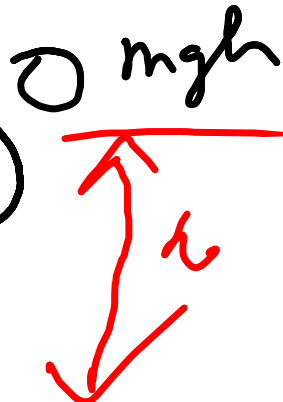
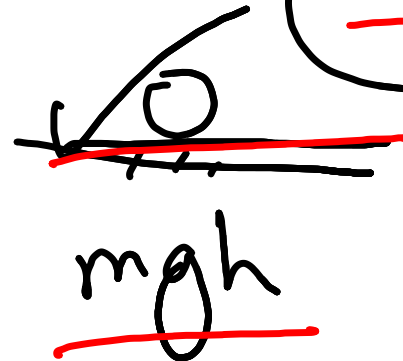
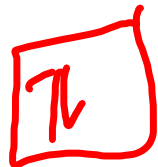
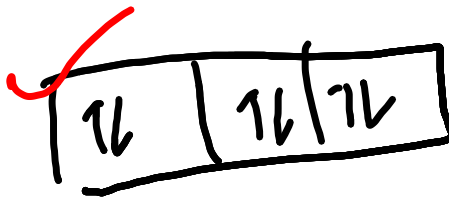
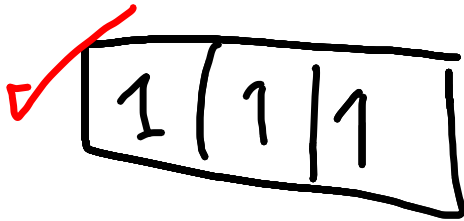
Bond formation — emission a / absorption b

**Why Bond is formed
???**

↓
H → duplet
→ octet

The tendency of achieving (stable) electronic configuration of inert gases.

The tendency of elements having minimum static energy to achieve maximum stability.



Chemical Bond:

Types: Different elements transfer or share or partially share electron of their outermost shell to form 3 types of bonds, they are-

- ✓ Ionic bond (Metal + Non-metal)
- ✓ Covalent bond (Non-metal + Non-metal)
- ✓ Co-ordinate bond

↓
Kabir

Bond

3

Intramolecular

Intermolecular



Ionic

Covalent

Co-Covalent

Metallic

V D W

D D

Hydrogen

Dispersion

Previous question

Which is not a type of chemical bond?

[MAT: 13-14, 11-12, 06-07]

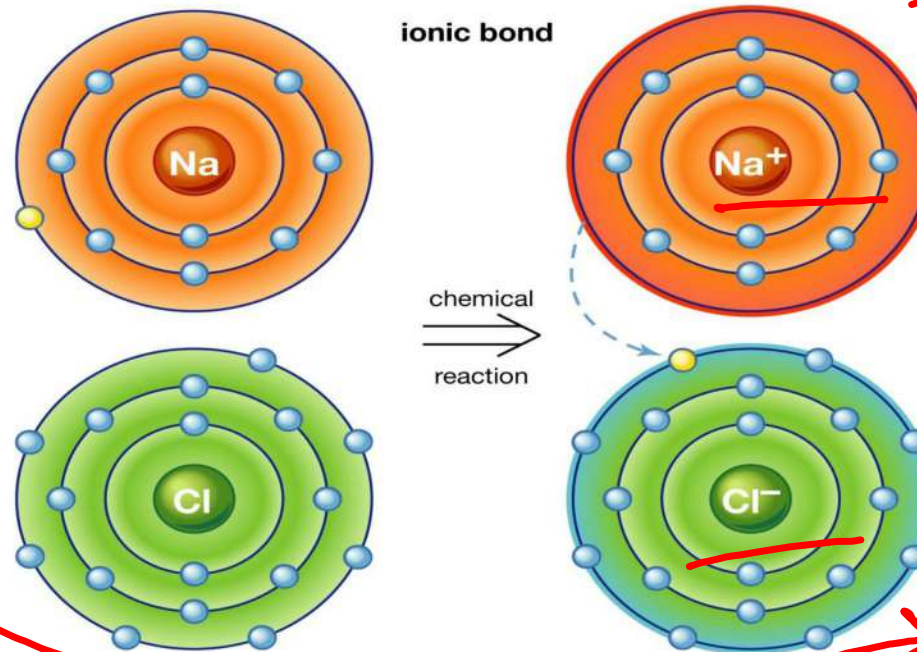
- (a) Co-ordinate ionic bond
- (b) Covalent bond
- (c) Ionic bond
- (d) Co-ordinate covalent bond

Ionic bond:

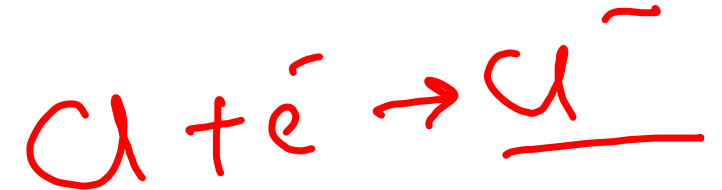
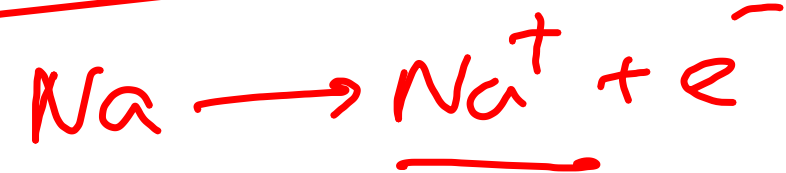
Formation of chemical bond among elements of atoms is the process of attaining more stable electronic configuration by those atoms similar to their nearest inert gas elements either by donation-acceptance of electrons

Metal

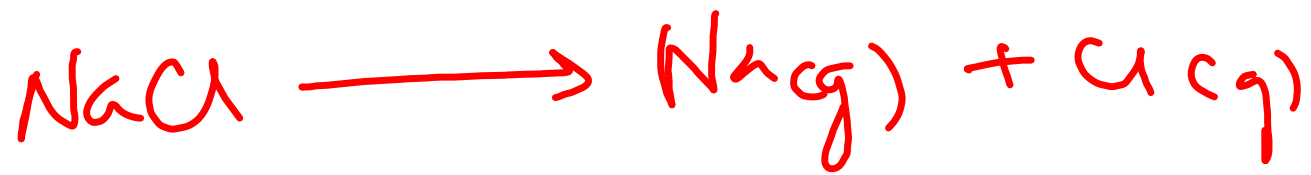
Nonmetal



↓ I. Energy



↑ e Affinity

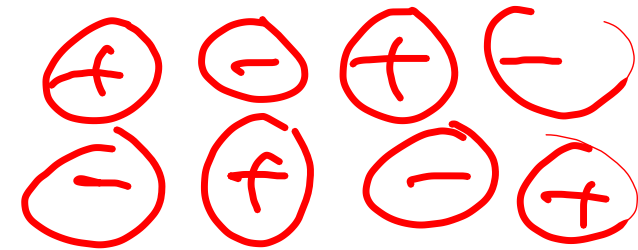
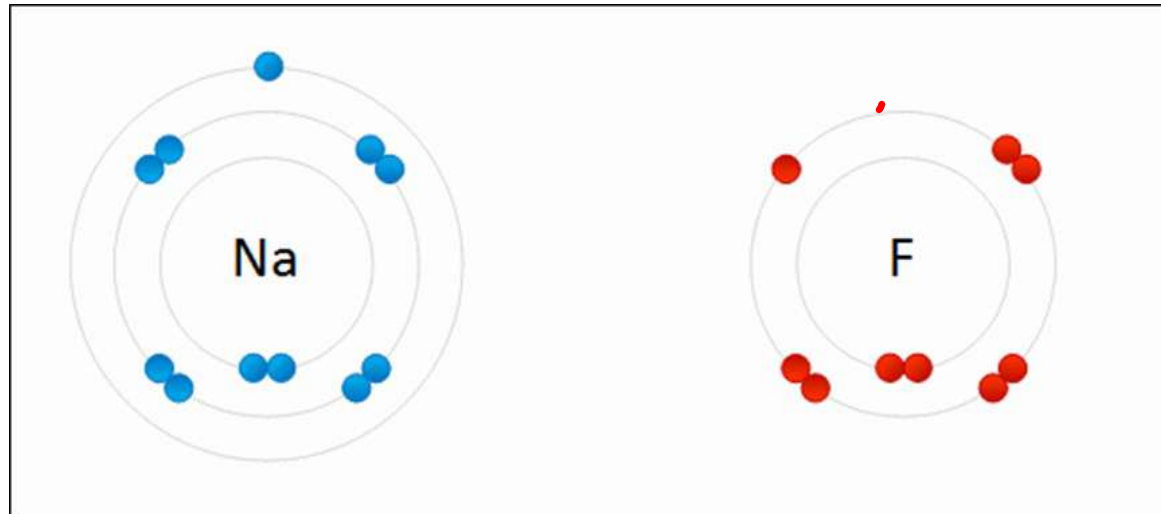


Conditions for formation of ionic bond:

- (i) **metal** atoms should have low ionization energy.
- (ii) **non-metal** atoms should have high electron affinity.
- (iii) The lattice energy of the crystals of ionic compounds must be high.



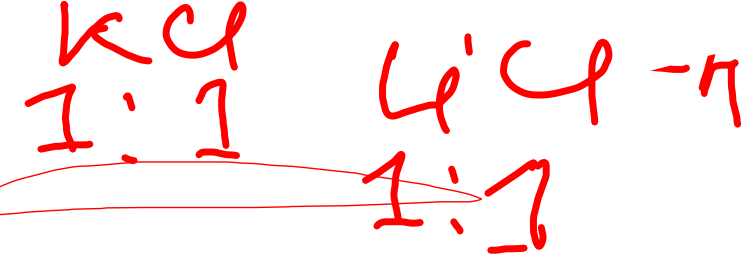
$$LE \propto \frac{q_1 q_2}{Er}$$



crystal
lattice

Properties of ionic compounds:

✓ CO ✓ SO ✓
• Polar in nature.



• Crystal. Example: NaCl takes the form of a side-centered cube.

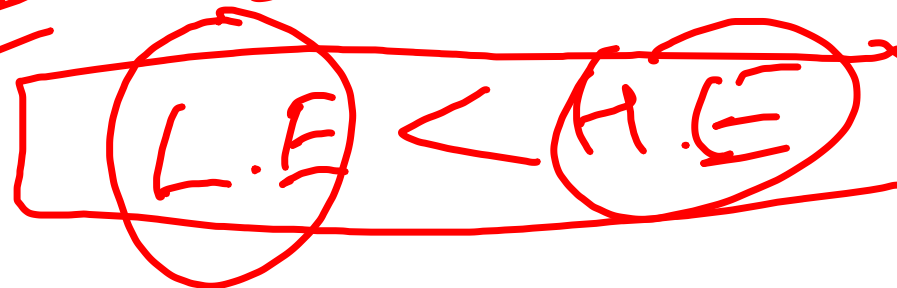
• High melting & boiling points. Example: boiling point of NaCl is 1470°C
whereas boiling point of covalent chloride CCl₄ is



• Soluble in polar solvent. Example- NaCl dissolves in water, but not 77°C in.

• Conducts electricity. (molten state)
Like dissolves like
soluble

isomorphism



Previous question

Which is not true regarding electric conductivity of ionic compounds? [MAT: 13-14]

(a) Can conduct electricity in solid state ✓

(b) Conduct electricity in solution

(c) Conduct electricity at melted state

(d) Very high melting point

Covalent bond:

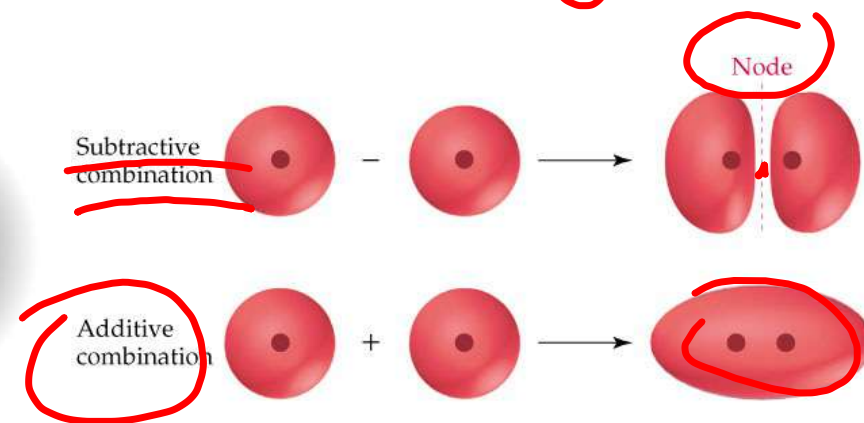
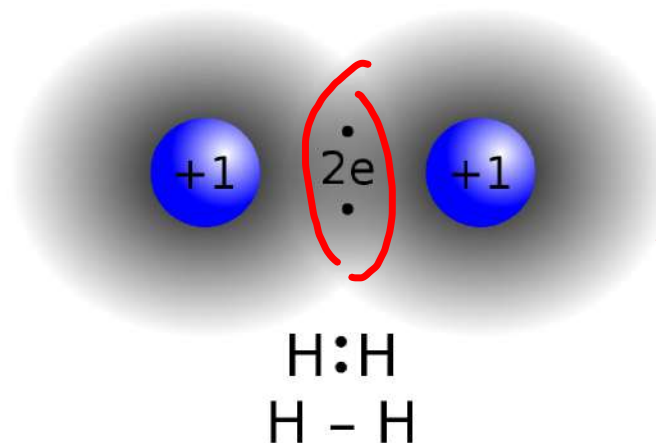
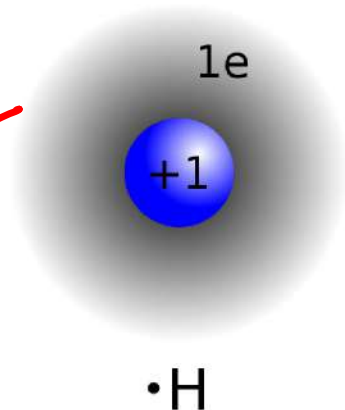
Proposer: Scientist GN Lewis.

Theories about formation:

There are two theories given based on modern wave mechanics on covalent bond structure. Such as-

- Valence bond theory,
- Molecular orbital theory.

Magnetism



BO
Bo Anti

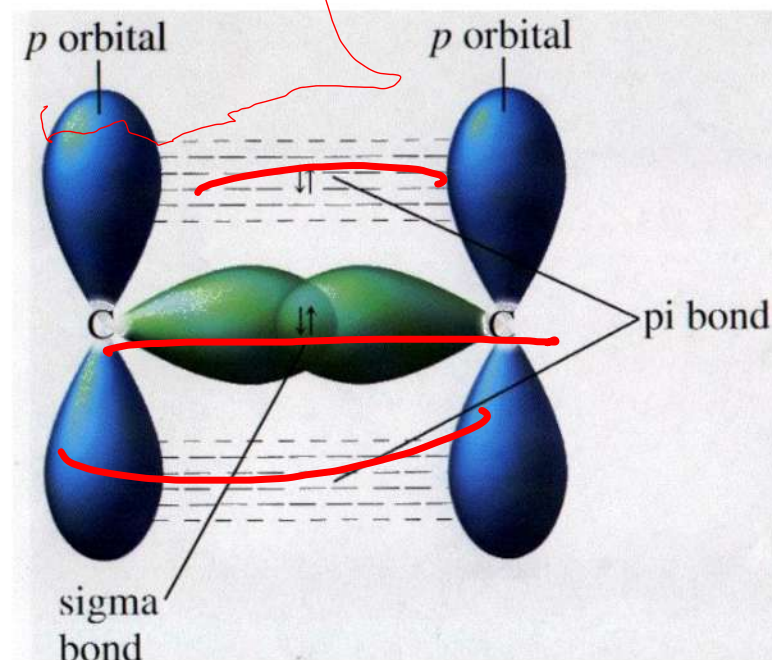
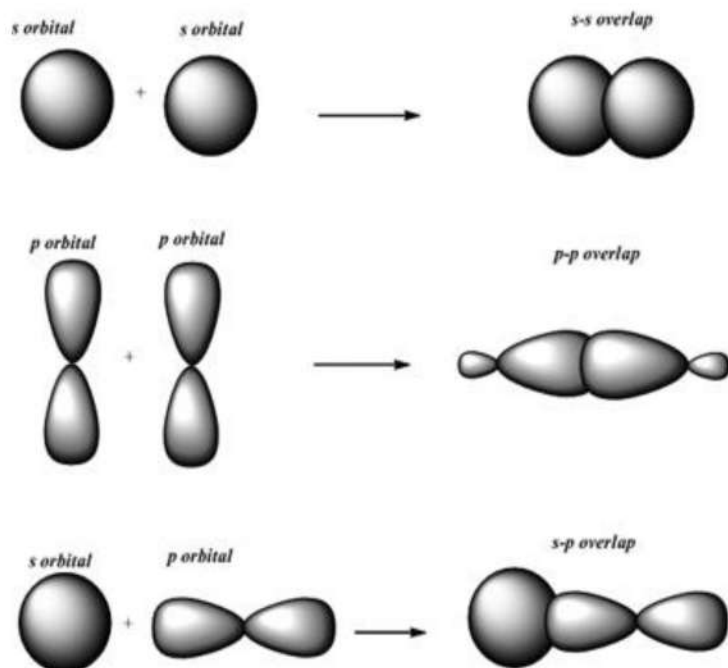
Hybridization
H₂

Classification of covalent bonds

According to overlapping of orbitals, covalent bonds are of two types. They are-

I. Sigma bond

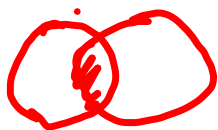
II. Pi bond



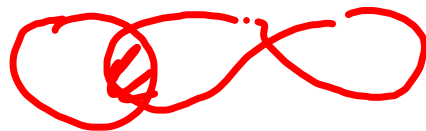
Poll Question-01

Overlapping of which element is most strong ?

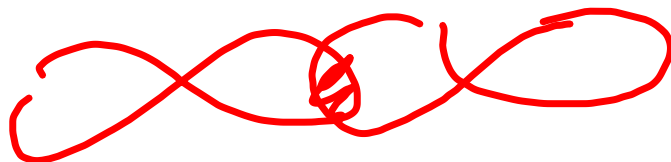
(a) H_2



(b) HCl



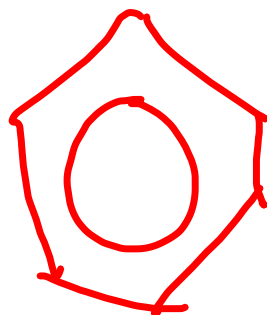
(c) F_2



Differences between sigma bond & pi bond

sigma bond	Pi-bond
This type of bond formed by direct overlapping of orbitals	This type of bond formed by partial side to side overlapping of orbitals
sigma bond is stronger than Pi-bond	Pi-bond is weaker than sigma bond.
Can occur in any orbital ,	Pi bond does not occur with hybrid orbitals but occurs with other pure orbitals except s-orbital.

Inductive effect



Mesomeris effect

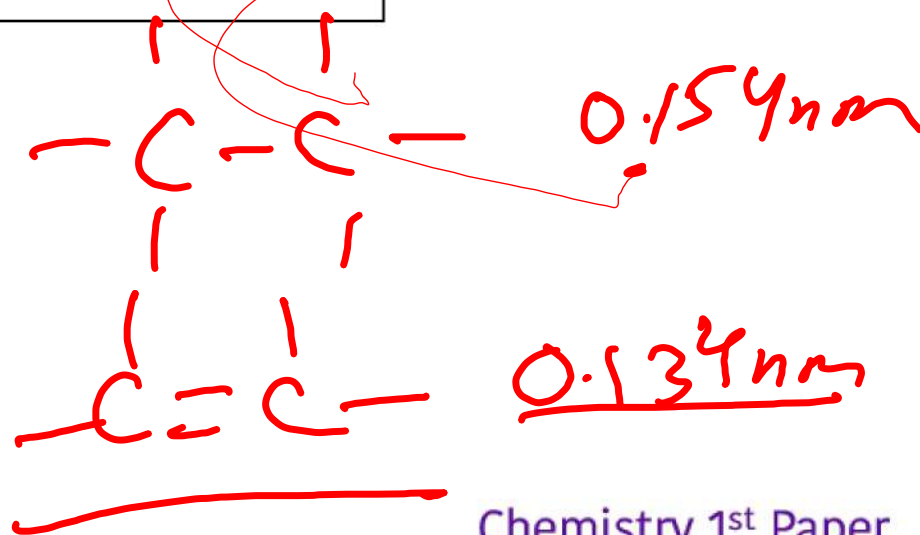
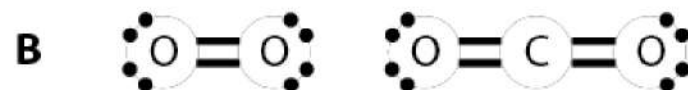


Classification of covalent bonds

How many pairs of electrons needed to share between two atoms; depending on it covalent bond is of 3 types. These are-

Bond	Example
✓ Single bond	Hydrogen and Ethane molecule
✓ Double bond	Oxygen and Ethylene molecule
✓ Triple bond	Nitrogen and Acetylene molecule

Alkane



Previous question

By which bond two hydrogen atoms are connected?

[MAT: 19-20]

(a) Ionic

(b) Hydrophilic

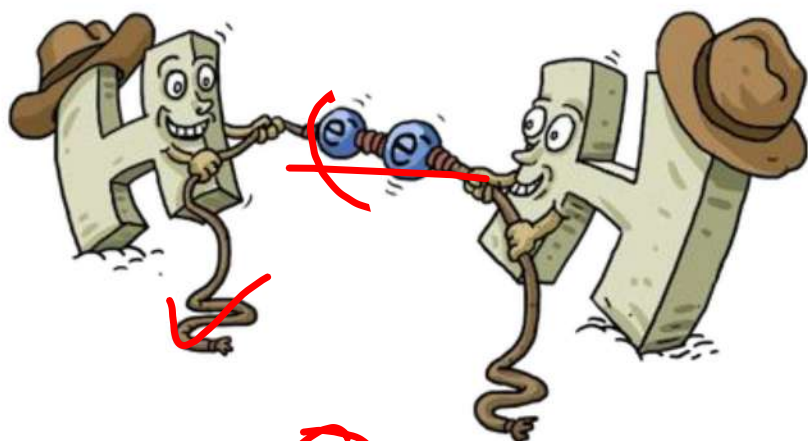
(c) Hydrophobic

(d) Covalent

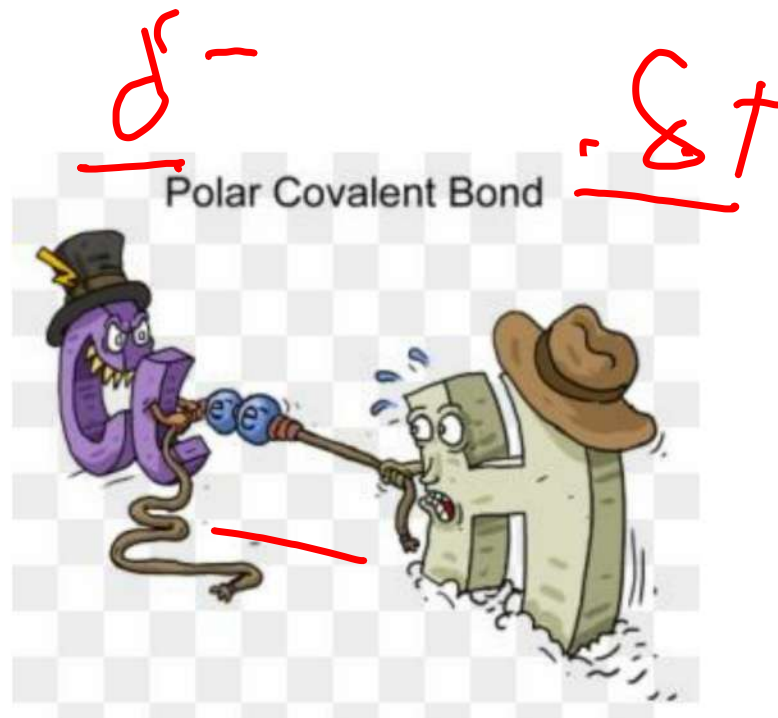
• Polarity in covalent bond •

- The formation of a dipole in a covalent compound is called polarity of the compound.
- HCl , HF , H_2O , HNO_3 , etc. are polar compounds.

Non-Polar Covalent Bond



Polar Covalent Bond

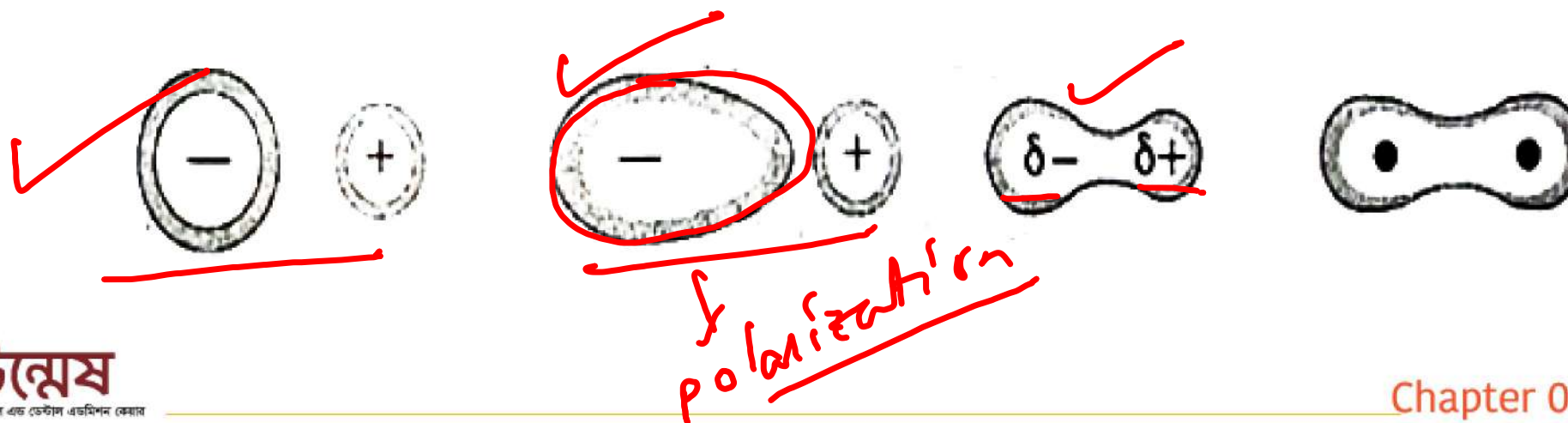


Polarity in covalent bond

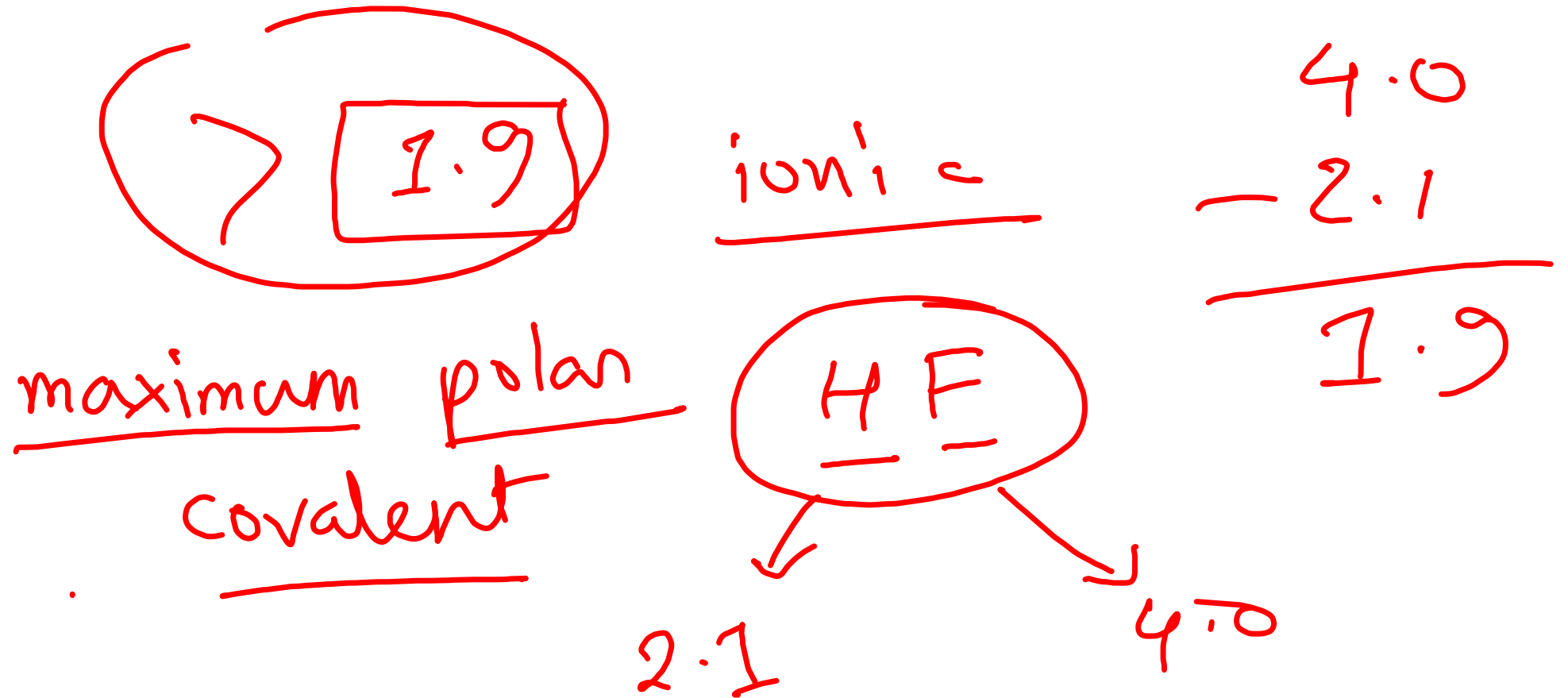
↑↑ polarization → less polar compound.

Differences in electro-negativity	Types of compound
0	Pure covalent
<u>< 0.5</u> 0.4	✓ Non-polar covalent
HCl 2.1 1.4 <u>0.5-1.9</u>	— Polar covalent
> 1.9	Ionic compound

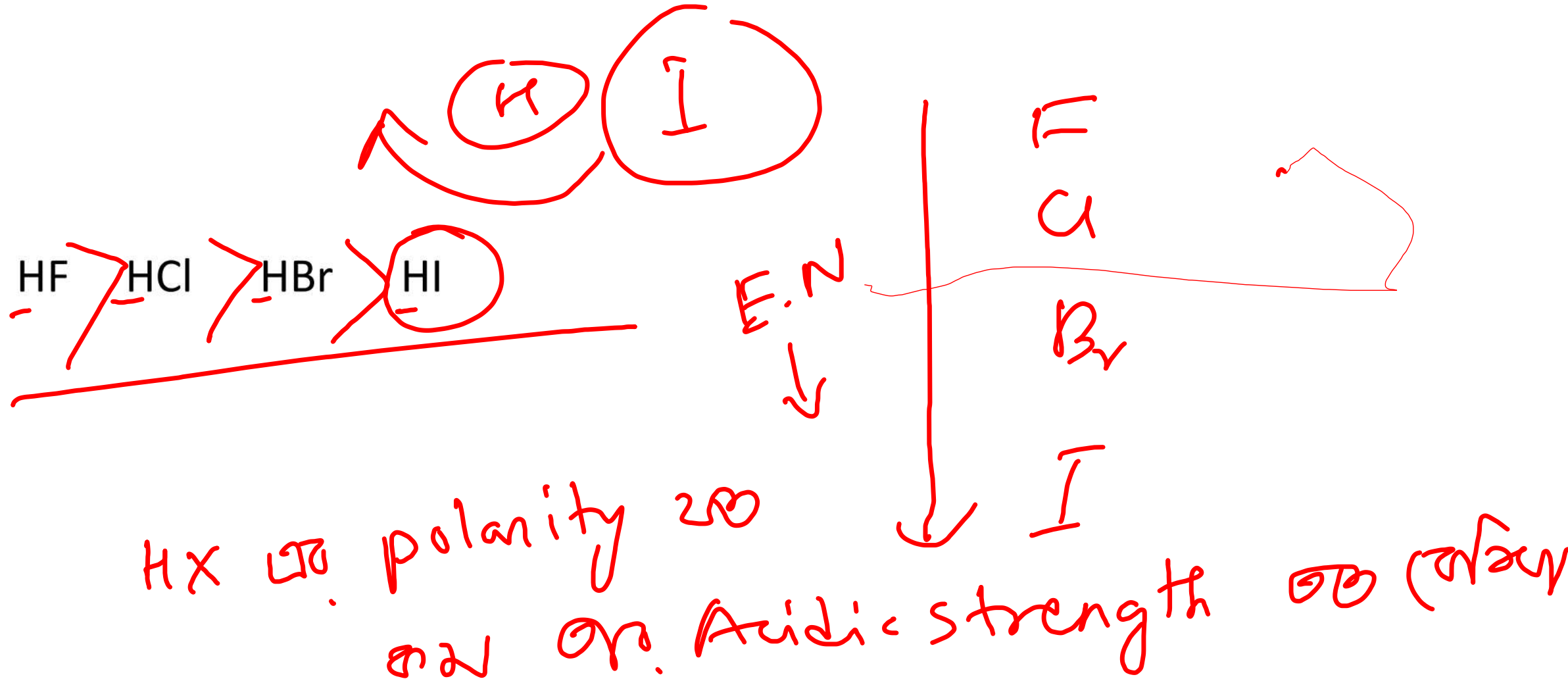
2.5 2.1
CH₄
NaCl
2.1



Why molecules become ionic if difference of electronegativity is more than 1.9?



What is the order of polarity among hydrogen halide?



Poll question-02

Which one is polar?

[DAT: 17-18]

(a) CH_4

(b) CCl_4

(c) HCl

(d) N_2

0.5 - 1.9

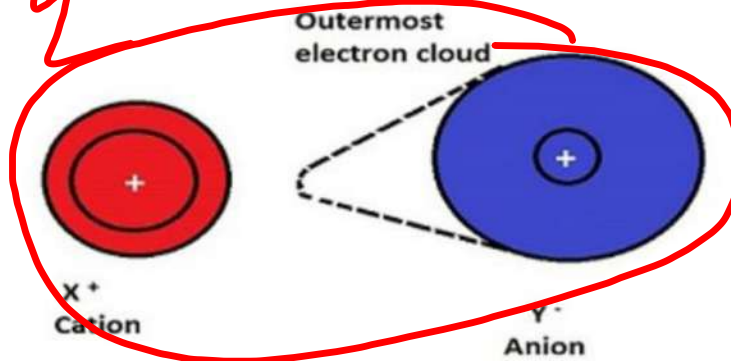
Covalent characteristics in ionic compounds

- In reality no compound is 100% ionic. Like that there is no 100% covalent compounds.

- NaCl has 80% ionic property.

Electronegativity difference, ΔEN :	0.1	0.2	0.6	0.9	1.1	1.7	2.0	2.1	3.0
Average partial ionic character :	0.5	1.0	10	19	25	50	75	80	90

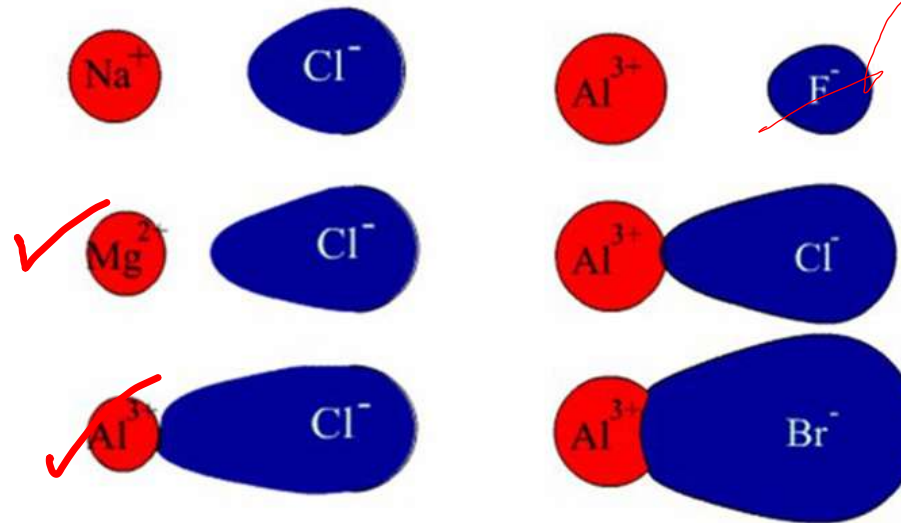
Fajan's Rules



20% covalent

Fajan's polarization rules

- With the increase of charges on cations and anions,
- With decrease of sizes of cations and increase of sizes of anions,
- Cations with electronic configuration $ns^2 np^6 nd^{1-10}$



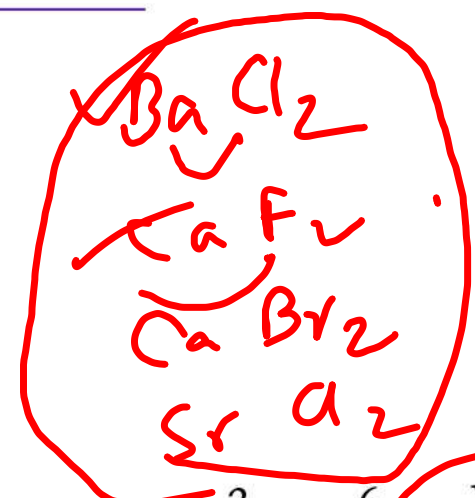
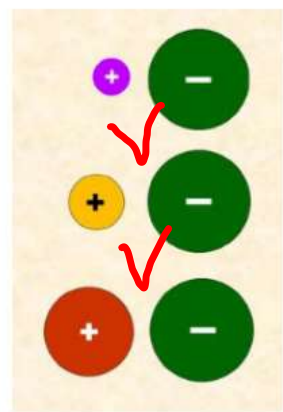
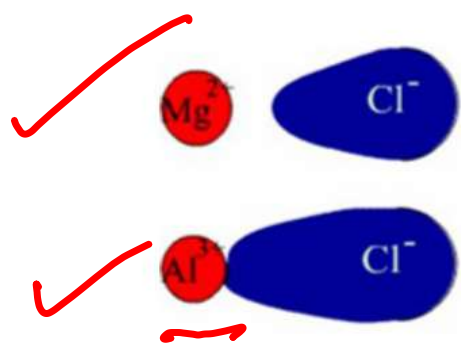
$\phi = \frac{\text{charge}}{\text{radius}}$

ionic potential \uparrow
 \Downarrow
polarization \uparrow

Application of fajans rule

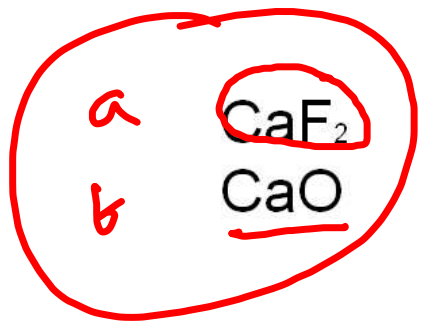
1st rule

2nd rule



$ns^2 np^6 nd^{1-10}$

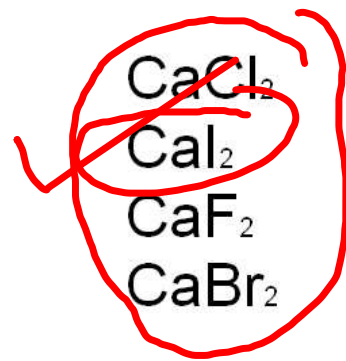
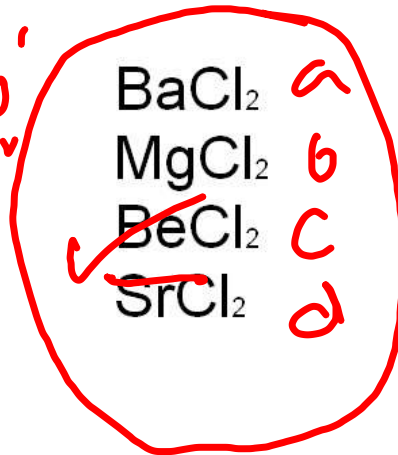
crystal solubility ↓



F⁻ 1-2

anion polarizability

polar ionic



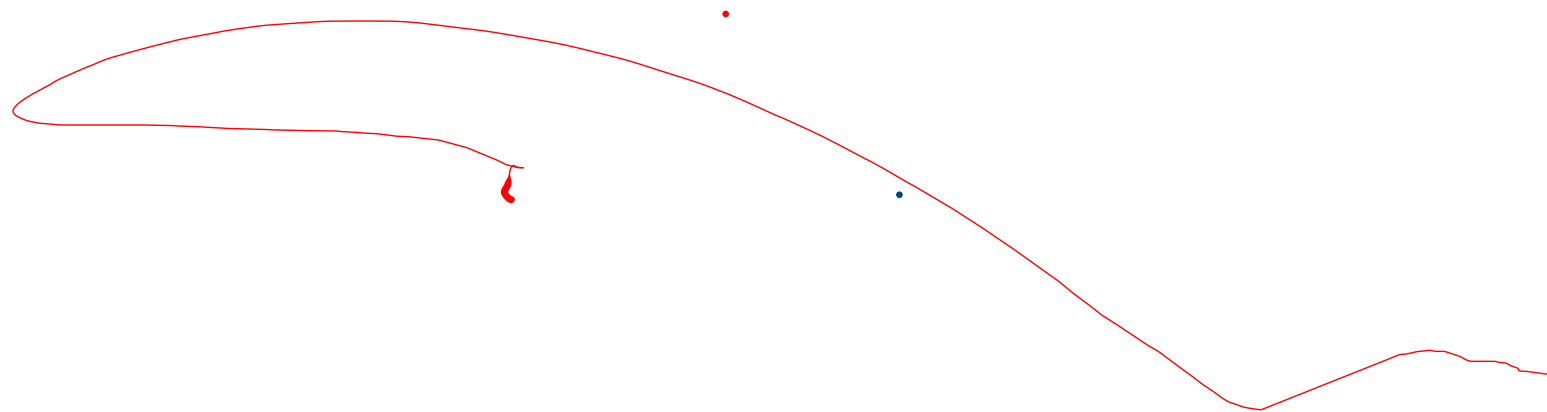
0.095 nm

0.096 nm

Poll Question-03

Which one is most covalent?

- (a) CaF_2
- (b) CaCl_2
- (c) CaBr_2
- (d) CaI_2



Ionic potential

Ionic potential of cation: $\phi = \frac{\text{charge}}{\text{radius of cation}}$ / radius of cation

With the increase in ϕ of cation-

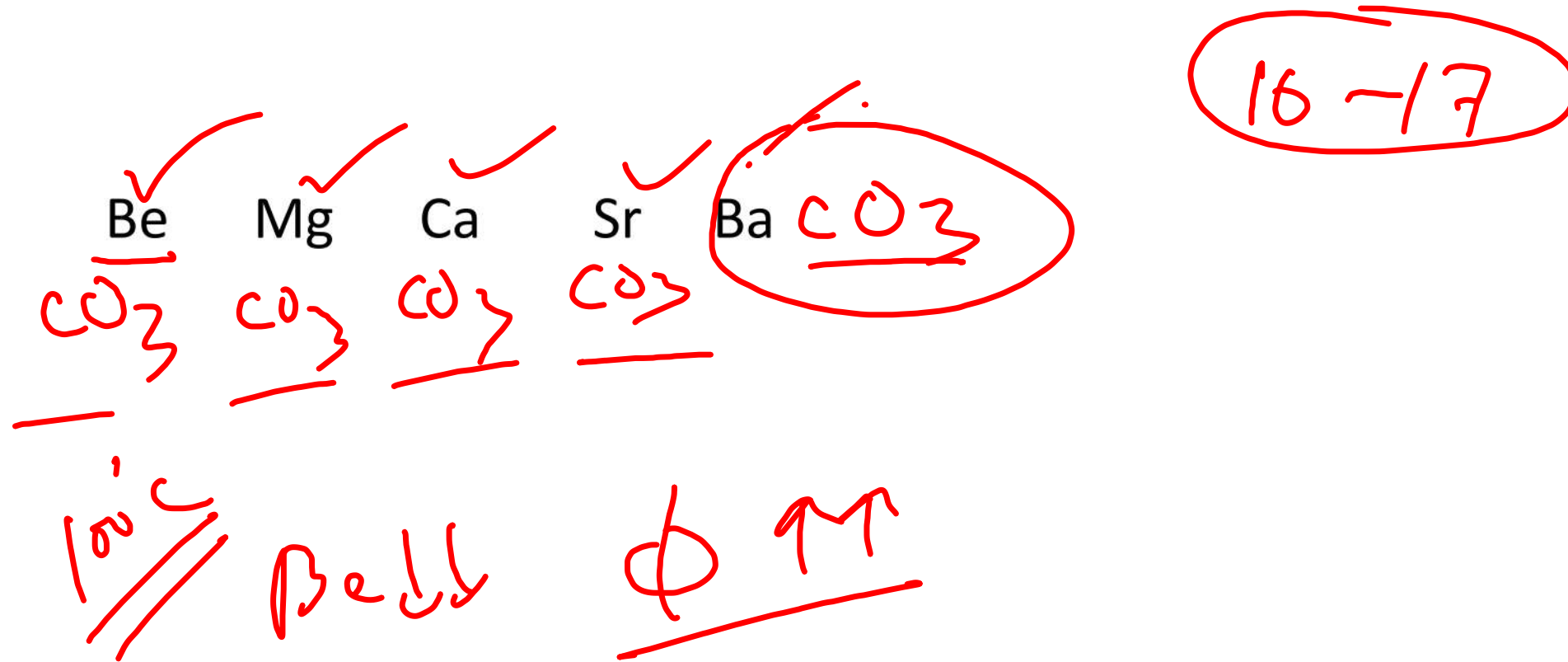
- Melting and boiling points of ionic salts gradually decrease,
- Decrease of solubility in water,
- Coloration of compounds, [↑]
- Dissociation of metallic carbonates in low heat etc. properties are expressed.

Why AgF is water soluble?

Decrease of solubility of salts in water : Among AgF, AgCl, AgBr and AgI, polarization of anion occurs the least in AgF. So, AgF is soluble in water.

$$\phi = \frac{\text{charge}}{\text{radius}}$$

which metallic carbonate has more Thermal stability ?



What will be the color of these compounds

???

d, f
element
an:

→ oxides, sulphides

Black
oxides

CuS, PbS, HgS

black

~~CdS, AgBr, AgI, PbI₂~~

ay

PbCl₂, HgCl₂, AgCl, AgF

F AgF
Cl AgCl
Br AgBr
I

I

AgI colored

white
polarization
color

Poll Question-04

Which salt is least stable? on heat



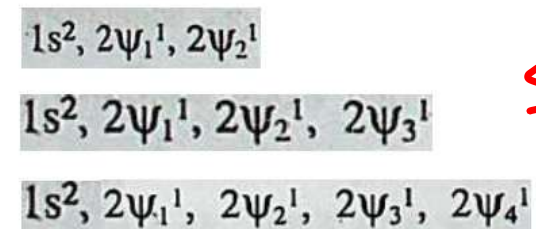
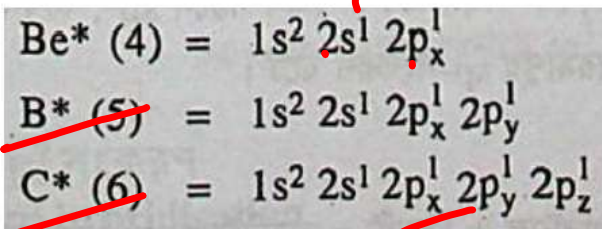
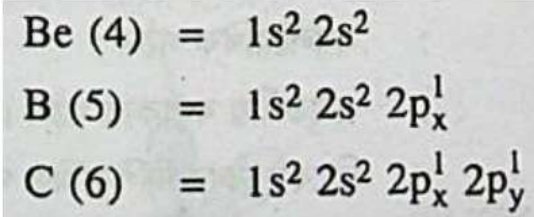
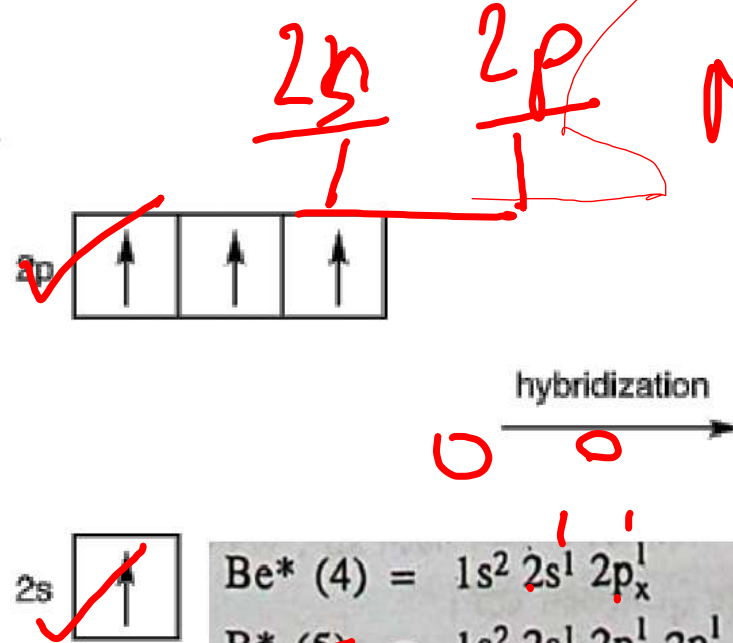
100°C $\phi = \pi$

1000°C

Hybridization of orbitals

Proposed by: Scientist Linus Pauling.

Covalent



Handwritten notes in red ink:
 - BCl_2 with a circled 2 and an arrow pointing to the hybridization process.
 - BCl_3
 - CCl_4
 - sp^3 written multiple times with arrows pointing to the hybrid orbitals.

Properties of hybridization

2nd

s p

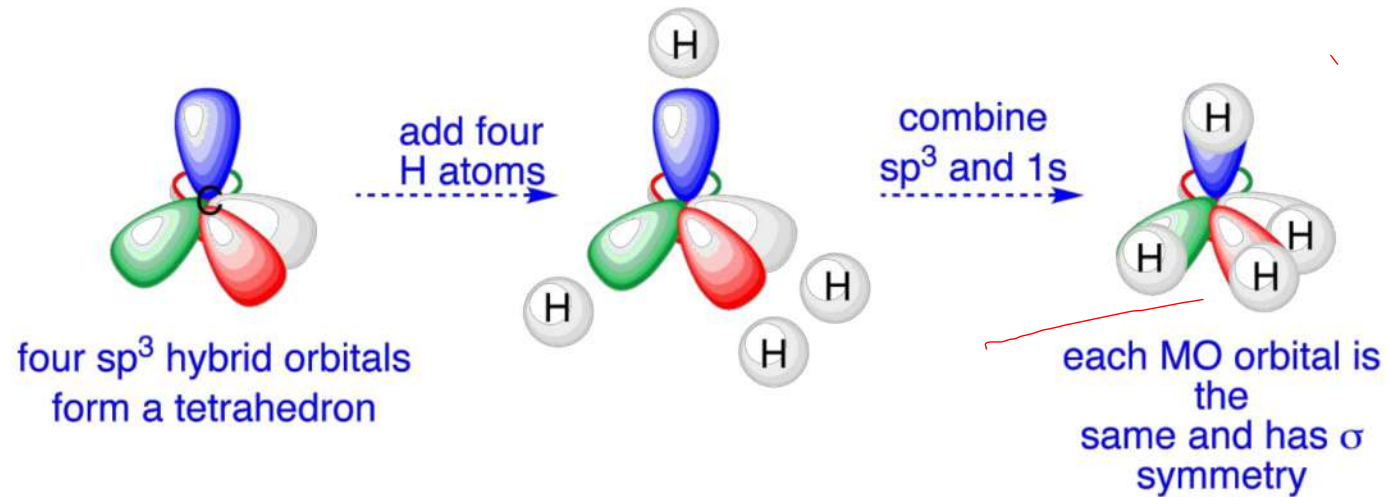
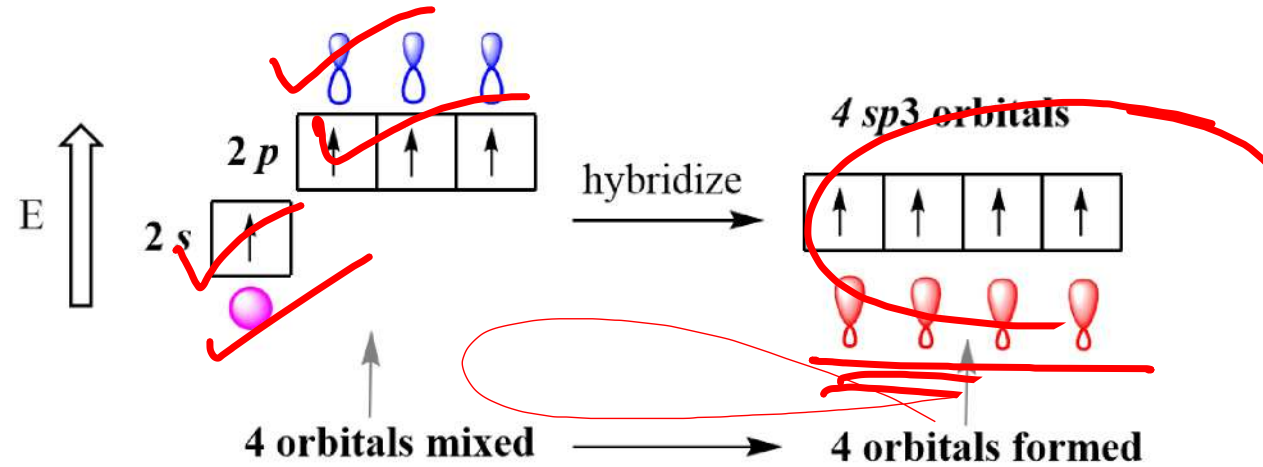
s p d

PC B O S B₂ N

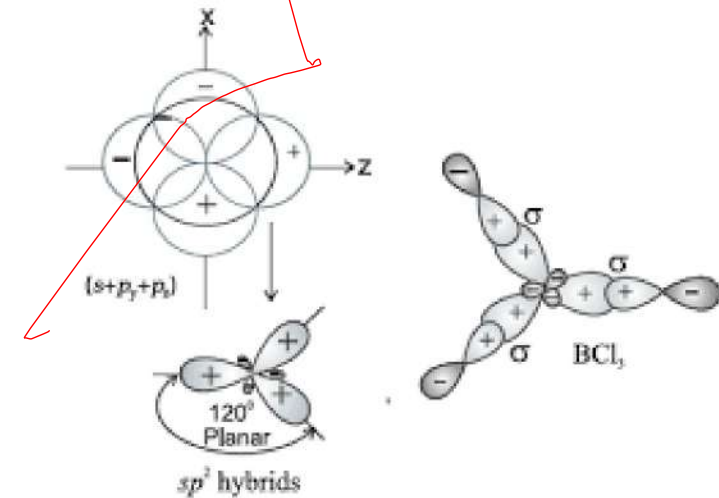
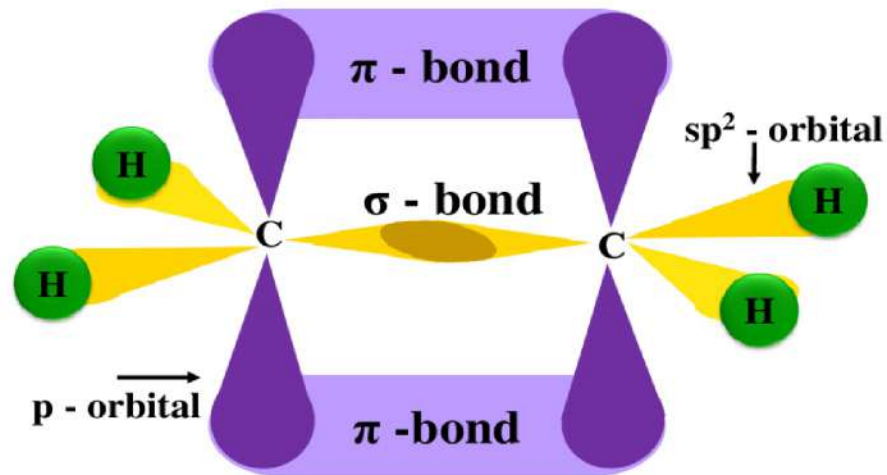
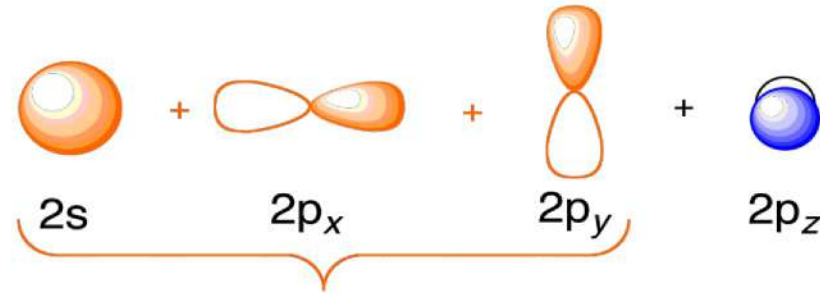
C O S N P Be

- Orbital hybridization occurs in a separated atom only.
- Generally orbitals of different subshells in a same energy level undergo hybridization.
- The number of orbitals that take part in hybridization indicates the number of hybrid orbitals of equal energy that are formed.
- As orbitals formed from hybridization are of equal energy, they create an equal angular distance by repelling themselves.
- The shapes and bond angles of molecules can be assumed from type of hybridization.
- Each hybrid orbital accommodate maximum two electrons like pure atomic orbital.
- The bonds formed by hybrid orbitals become stronger.

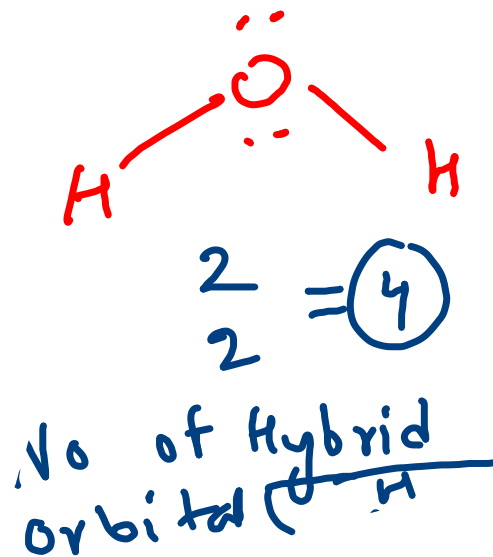
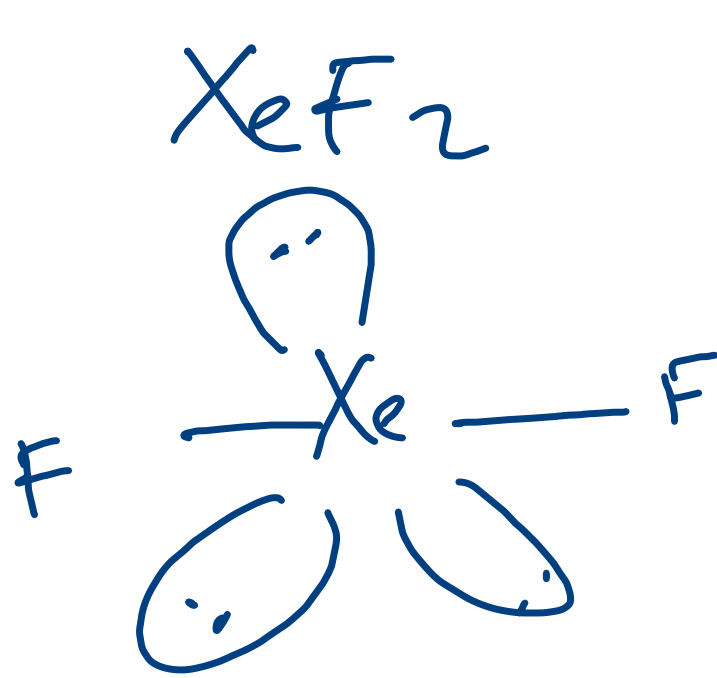
sp^3 HYBRIDIZATION



SP² HYBRIDIZATION



Determination of hybridization



Things to Consider

- no of sigma bond
- no of lone pair electron
- no of Co-ordinate bond

(Structural formula)



Determination of hybridization of central atom of compound and number of lone pair electron

Hybridization of central atom:

$$H = \frac{1}{2}(V + X - C + A)$$

Here,

V = number of electron in valence shell

X = number of univalent atoms

C = number of charge on cations

A = number of charge on anions

$$\begin{aligned} C &= 4 \\ V &= 5 \\ O &= 6 \end{aligned}$$



Number of lone pair electron in hybrid orbital :

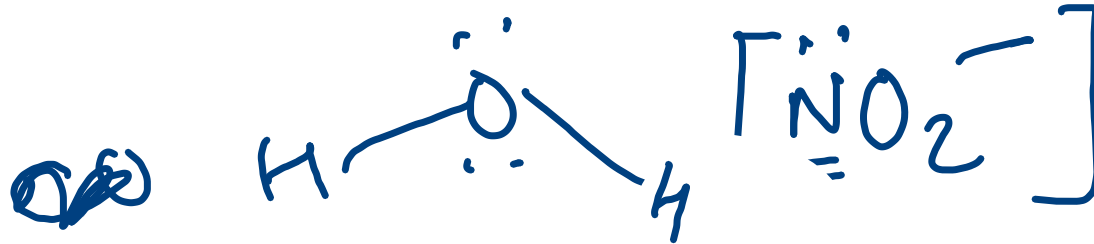
$$L = H - X - D$$

(Here, D is the number of divalent atoms)

Practice Problems

$$H = \frac{1}{2}(V + X - \text{C} + A)$$

$$L = H - X - D$$



NO

$$\frac{1}{2}(5)$$

$$2.5$$

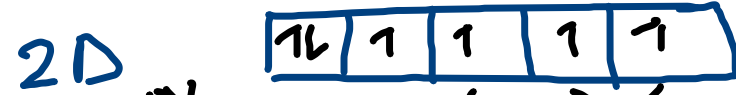
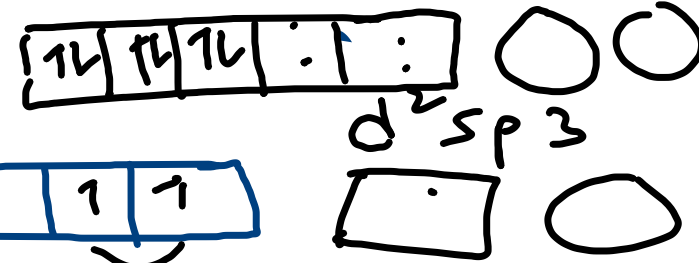


~~H₂~~
 $\frac{1}{2}(4 - 2)$
 $= 1$

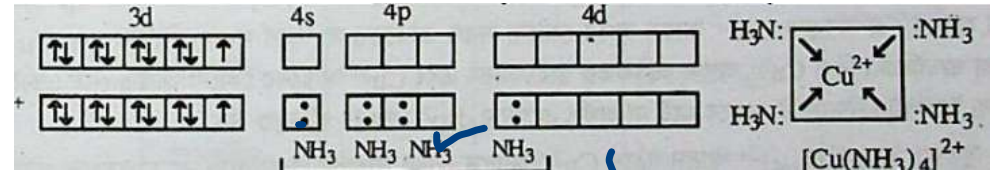
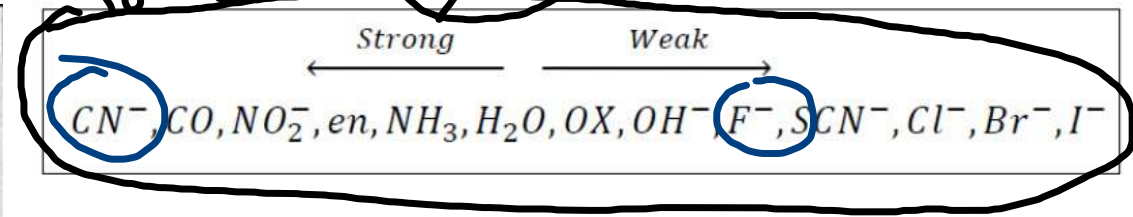


Hybridization of (Complex ions)

$6 = sp^3d^2$



2	<u>sp</u>	<u>linear</u>	Ag ⁺	NH ₃	[Ag(NH ₃) ₂] ⁺
2	<u>sp</u>	<u>linear</u>	Cu ⁺	Cl ⁻	[CuCl ₂] ⁻
4	<u>sp³</u> px, py, pz	<u>tetrahedral</u>	Co ²⁺	Cl ⁻	[CoCl ₄] ²⁻
4	<u>dsp² or, sp²d</u> ($d_{x^2-y^2}$)	<u>square coplaner</u>	Ni ²⁺ dsp ²	CN ⁻	[Ni(CN) ₄] ²⁻
4	<u>sp²d</u>	<u>square coplaner</u>	Cu ²⁺ sp ² d	NH ₃	[Cu(NH ₃) ₄] ²⁺
6	<u>d²sp³</u>	<u>octahedral</u>	Fe ²⁺	CN ⁻	[Fe(CN) ₆] ⁴⁻
6	<u>d²sp³</u>	<u>octahedral</u>	Fe ³⁺	<u>CN⁻</u>	[Fe(CN) ₆] ³⁻
6	<u>sp³d²</u>	<u>octahedral</u>	Fe ³⁺	<u>F⁻</u>	[FeF ₆] ³⁻





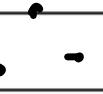




- coordination no
 - hybridization
 - color
 - shape
- John Teller Effect

"X ray diffraction analysis"

Shape of hybride orbitals

VSEPR

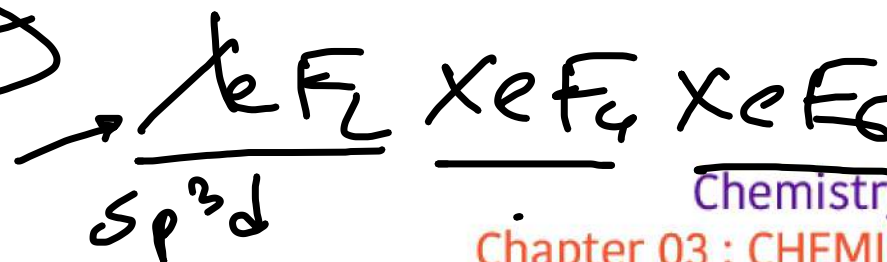
$$\frac{1}{2}(8+2) = 5$$

Hybridized orbital number	Hybridization	Shape of molecule	Bond angle	Example
Two	 sp	Coplanar straight linear	180°	BeCl ₂ , C ₂ H ₂
Three	 sp ²	Coplanar <u>triangular</u>	120°	BCl ₃
Four ^{3D}	 sp ³	<u>Tetrahedral</u>	109.5°	CH ₄ , NH ₃
Four ^{2D}	 sp ² d	<u>Coplanar square</u>	90°	[Cu(NH ₃) ₃] ²⁺
✓ Five	 sp ³ d	<u>Trigonal bipyramid</u>	90° & 120°	PCl ₅
<u>Six</u>	 sp ³ d ²	<u>Octahedral</u>	90°	SF ₆
✓ Seven	 sp ³ d ³	<u>Pentagonal bipyramid</u>	90° & 72°	IF ₇

7

7

5 + 2
linear



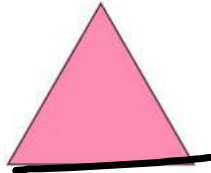
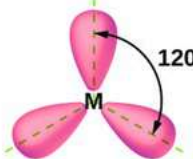
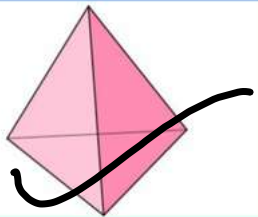
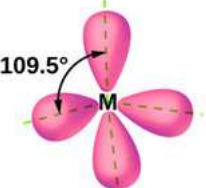
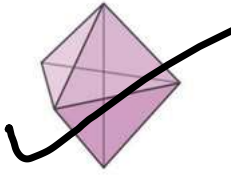
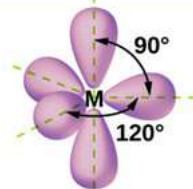
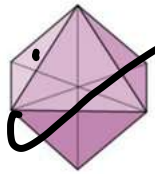
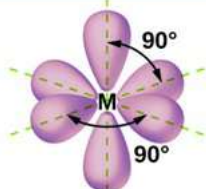


ডিনেশ

মেডিকেল এন্ড ডেন্টাল এডমিশন কোয়ার

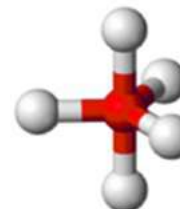
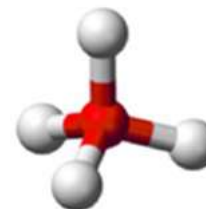
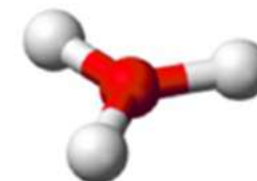
Chemistry 1st Paper

Chapter 03 : CHEMICAL BOND

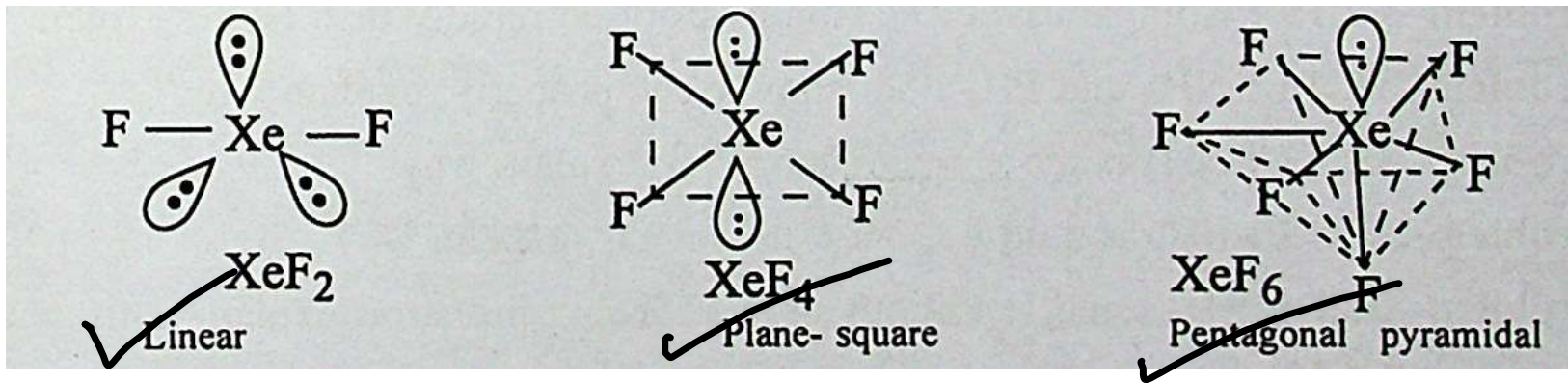
Regions of Electron Density	Arrangement		Hybridization	
2		Linear	sp	
3		Trigonal planar	sp^2	
4		Tetrahedral	sp^3	
5		Trigonal bipyramidal	sp^3d	
6		Octahedral	sp^3d^2	



VSEPR



Exception of Shape



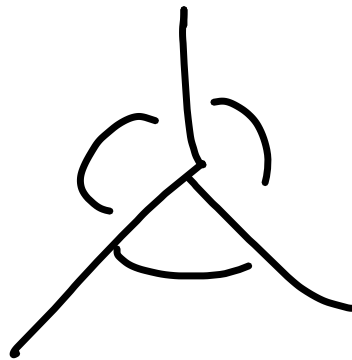
Previous question

sp^2

Bond angle in hybridized orbital is-

[MAT: 02-03, DAT: 04-05, 02-03]

- a) 180°
- b) 109°
- ☒ c) 120°
- d) 170°



Poll Question-05

What is the hybridization of central atom of HSO_4^- ?

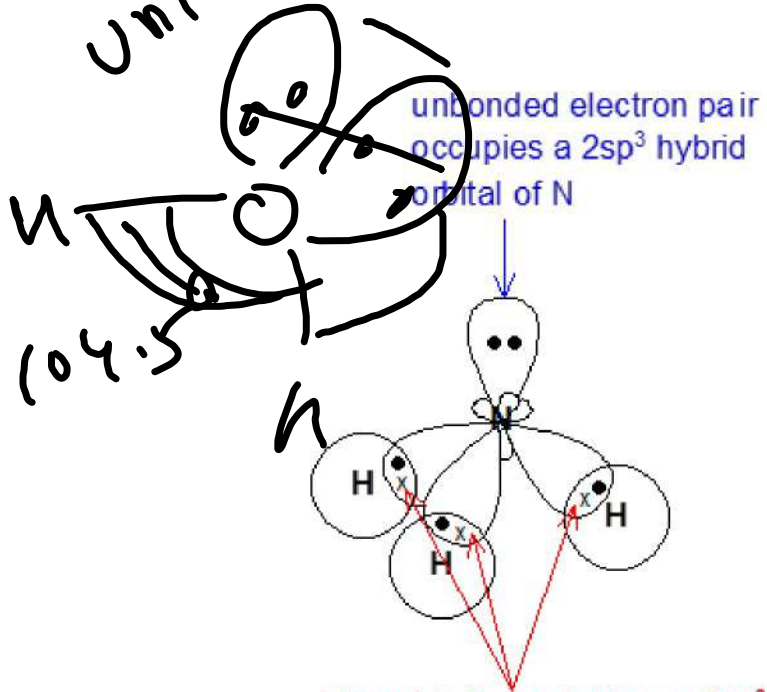
- (a) sp^2
- (b) sp^3
- (c) dsp^3
- (d) d^2sp^2

$$\frac{1}{2}(6 + 1 - 0 + 1)$$

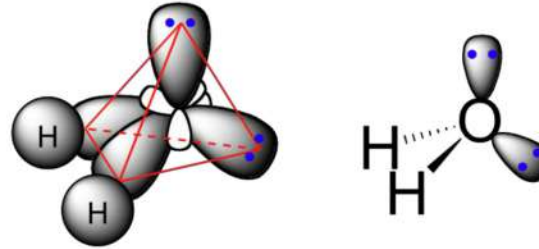
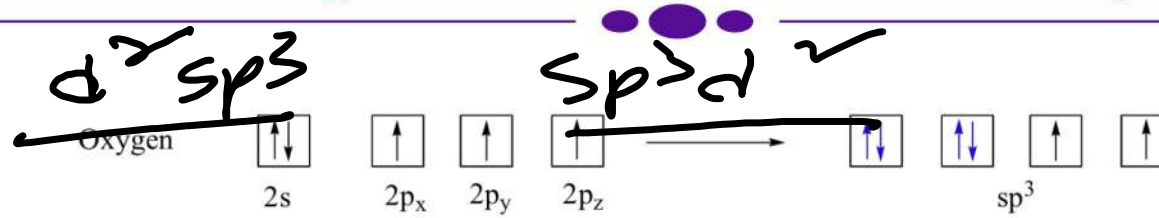
4

Effect of lone pair electron in shape of orbital

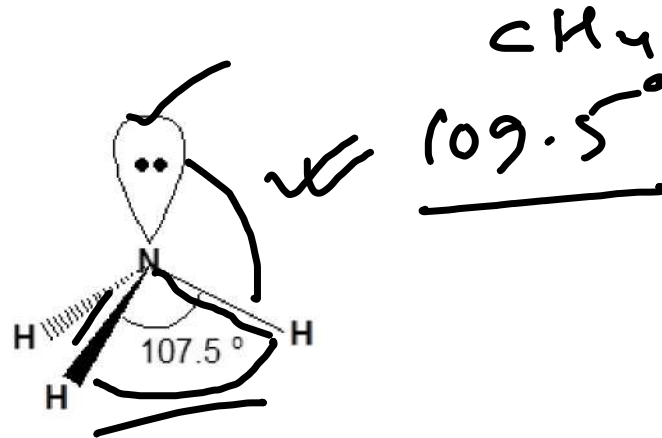
Gr. 17
Univalent



σ bond is formed when a $2sp^3$ hybrid orbital of N overlaps with the 1s orbital of H



Valence shell electron pair repulsion theory.



$\text{lp-lp} > \text{lp-bp} > \text{bp-bp}$

Special information

Molecule	Hybridization	Lone pair electron	Bond angle ✓✓	Shape of molecule
NH ₃	sp ³	1	107°	Trigonal pyramid
NF ₃	sp ³	1	102.5°	Trigonal pyramid
PH ₃	sp ³	1	94°	Trigonal pyramid
H ₂ O	sp ³	2	104.5°	V shaped
H ₂ S	sp ³	2	92°	V shaped



Distorted tetrahedral

Poll Question-06

Bond angle in ammonia molecule is?

[MAT: 14-15]

a) 180°

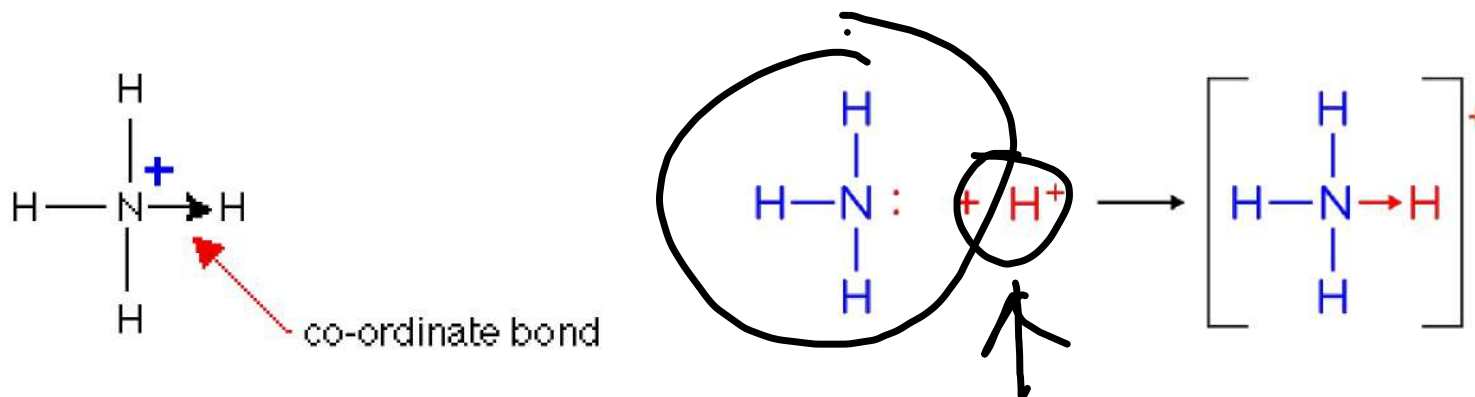
b) ☒ 107°

c) 120°

d) 170°

✓ Co-ordinate covalent bond

Definition: In the formation of a covalent bond between two atoms, each atom supplies one electron to form an electron pair which is shared by the related two atoms equally



Exmple :



complex metallic ion

Nucleophile electrophile

a —
b —

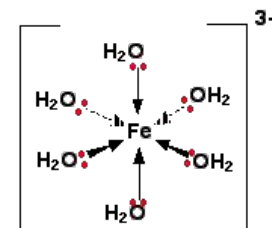
Determination of co-ordinate number:

6

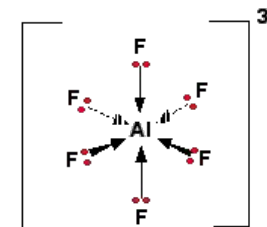
6

6

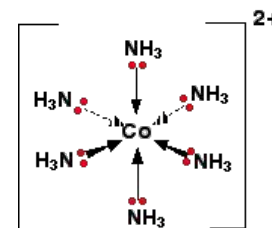
sp^3d^2



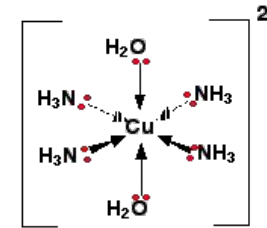
$[Fe(H_2O)_6]^{3+}$



$[AlF_6]^{3-}$



$[Co(NH_3)_6]^{2+}$



$[Cu(NH_3)_4(H_2O)_2]^{2+}$

Determination of oxidation number in complex molecule:

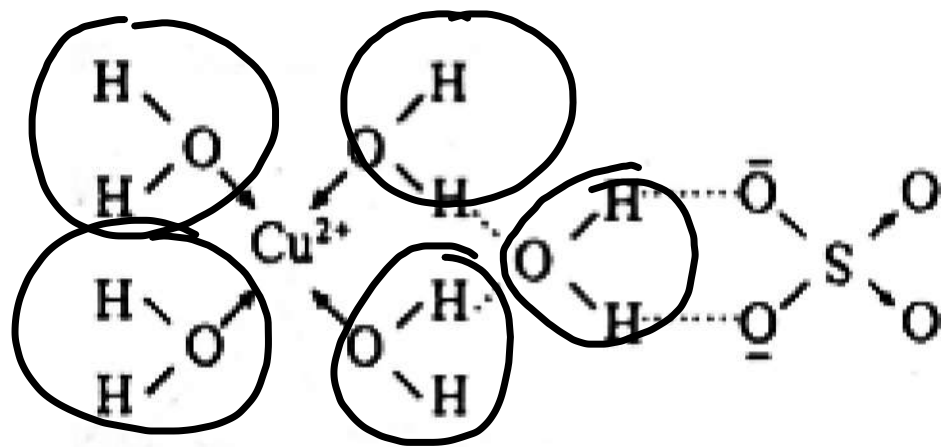
F^{-1}

$$x + (-1 \times 6) = -3$$

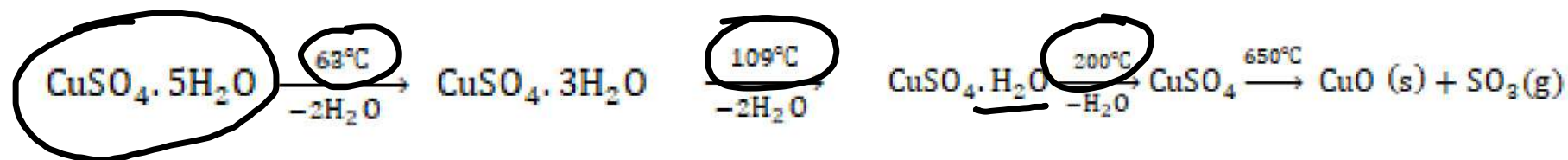
$$\therefore x = 6 - 3$$

$$= +3$$

Effect of temperature in blue vitriol



২৫৮



Dark blue crystal

Blue crystal

Bluish white crystal

White powder

Black

Different types of bonds in same compound

Type

Compound	Number of bonds	Name of bonds
\checkmark NH_4Cl	Three	• Covalent, Co-ordinate, Ionic bond
$K_4[\text{Fe}(\text{CN})_6]$	Three	• Covalent, Co-ordinate, Ionic bond
$[\text{Co}(\text{NH}_3)_6]\text{Cl}_3$	Three	• Covalent, Co-ordinate, Ionic bond
$(\text{CuSO}_4 \cdot 5\text{H}_2\text{O})$	Four	• Covalent, Co-ordinate, Ionic bond and Hydrogen bond
KOH	Two	• Ionic bond and covalent bond
$\text{H}_3\text{N} \rightarrow \text{BF}_3$	Two	• Covalent, Co-ordinate bond
H_3PO_4	Two	• Covalent, Co-ordinate bond
H_2SO_4	Two	• Covalent, Co-ordinate bond
$(\text{H}_2\text{O})_n$	Two	• Covalent and Hydrogen bond

1 > < > < <

Poll Question-07

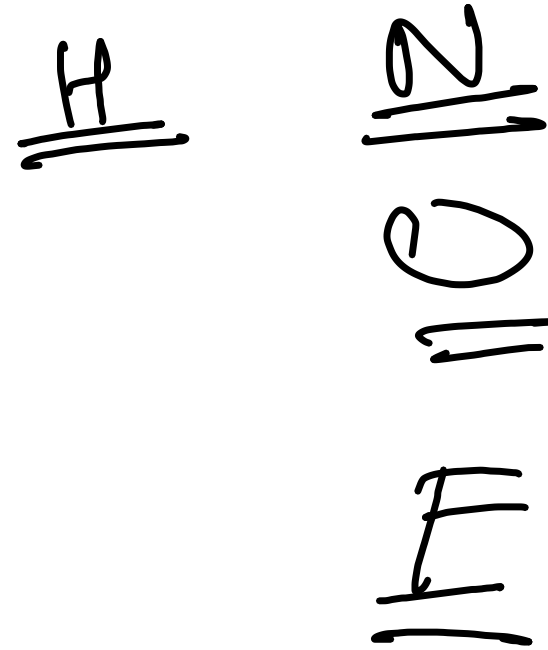
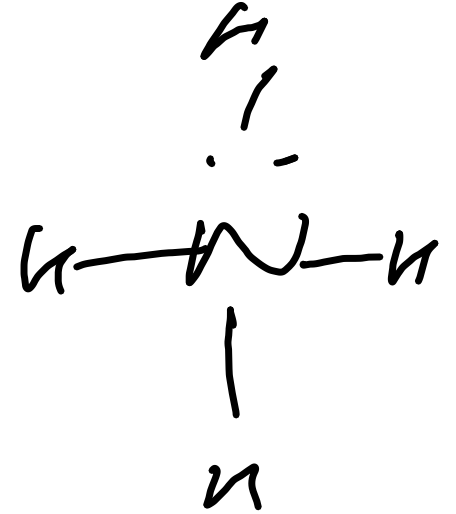
Which bond is not present in NH_4Cl molecule?

(a) Covalent bond

~~(b) Hydrogen bond~~

(c) Ionic bond

(d) Co-ordinate bond



Metallic Bond:

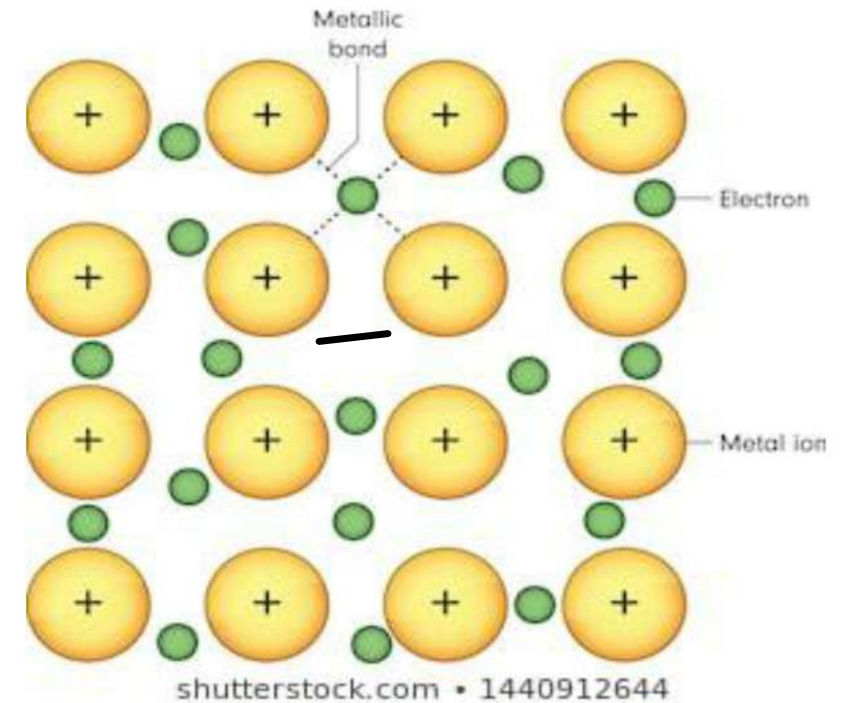
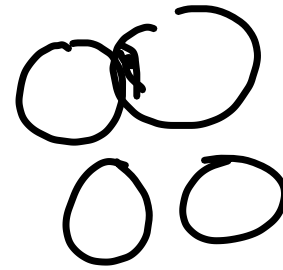
Defination: In metallic bond, shared electrons are mobile

Metallic bond is weaker than covalent bond.

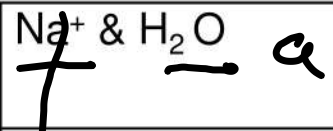




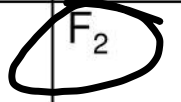
Special properties:

- Metals are crystalline in structure.
- Metals are very good conductors of electricity.
- Metals have special luster.
- Metals are malleable and soft.

a covalent radius
b metallic radius



Comparison of degree of energy between bonding and non-bonding forces

Effective attraction force	Base of bonding	Degree of energy (kJ/mol)	Example
(a) In case of bonding:			
✓ Ionic bond	Cation and anion	400 – 4000	Crystal of NaCl ✓
✓ Covalent bond	Nucleus and shared electrons	150 - 1100	H ₂ molecule ✓
✓ Metallic bond	Cation and polarized valence electron	✓ 75 - 1000	metals
(b) In case of non-bonding:			
✓ Ion-dipole attraction: ✓	Ion charge and dipole charge	10 – 50	Na ⁺ & H ₂ O 
✓ H-bond (Polar molecule) [N, O, F compounds]	Polar bond and H-dipole charged	10 – 40	H ₂ O 
✓ Dipole-dipole ✓	Dipole charges	3 - 4	HCl - HCl 
✓ Ion-induced dipole	Ion's charges and polarized e ⁻ cloud	3 – 15	Fe ²⁺ & O ₂ 
✓ Dipole-induced dipole	Dipole charge and polarized e ⁻ cloud	2 – 10	HCl & Cl ₂ 
✓ London force or dispersion force	Polarizable e ⁻ cloud	1 – 10	F ₂ 

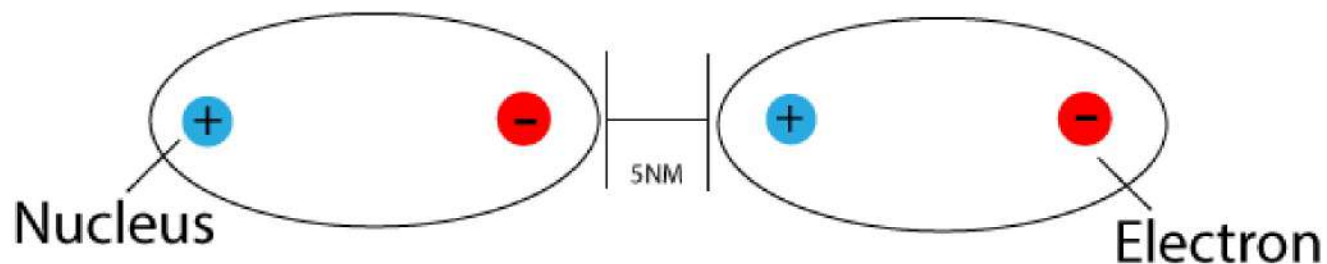
Vander Waals force

Intermolecular force among non-polar covalent compound is called Vander Walls force.

On the basis of sources, Vander Waals forces are of two types; such as:

- Permanent dipole and induced dipole attraction,
- Dispersion force or London force.

van der Waals Forces



Dipole and dipole moment

Dipole: The different charges or poles developed on two ends of a covalent compound containing highly electronegative elements are called dipole.

Types of dipole: Two types. They are- permanent dipole and temporary dipole.

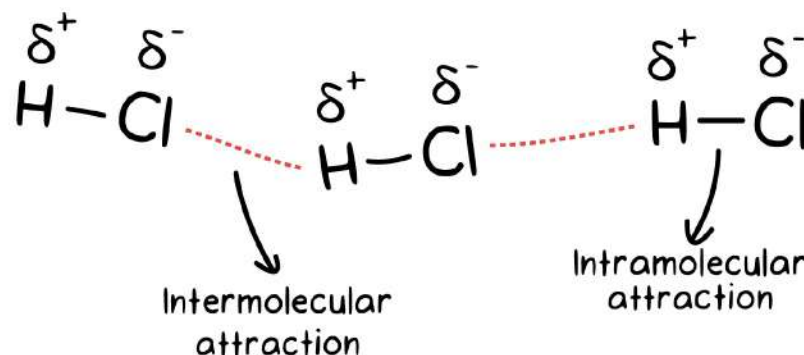
Permanent dipole: H_2O , NH_3 , HCl

Dipole moment: $\mu = Q \times r$

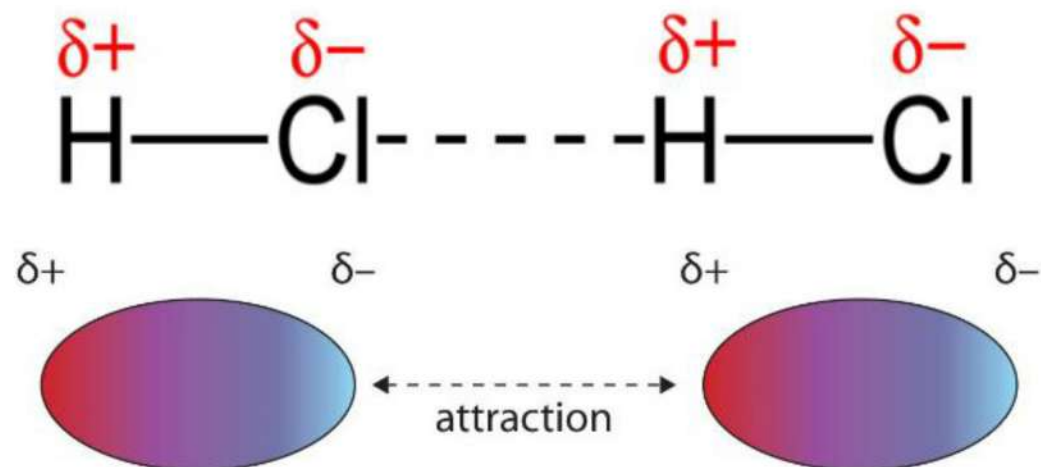
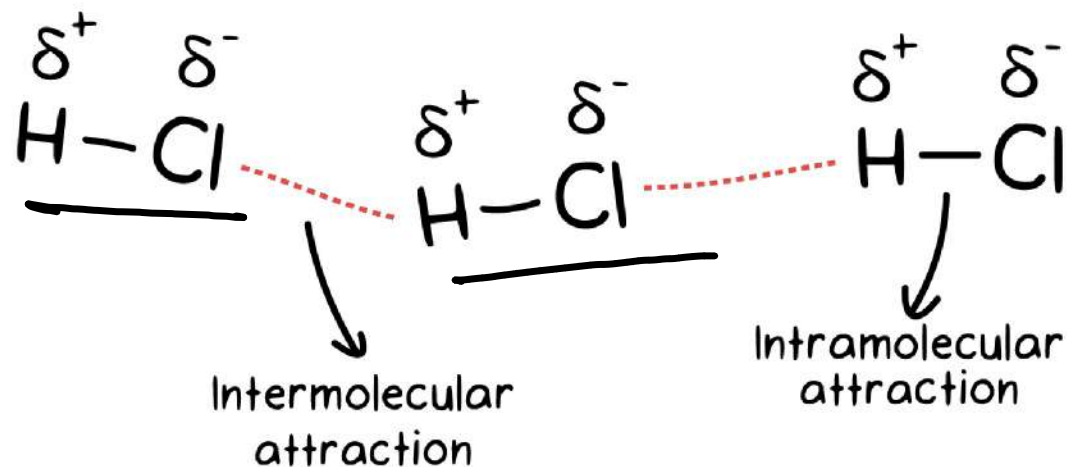
Unit of dipole moment is debye (D)

$1\text{D} = 3.336 \times 10^{-30}$ coulomb meters (Cm).

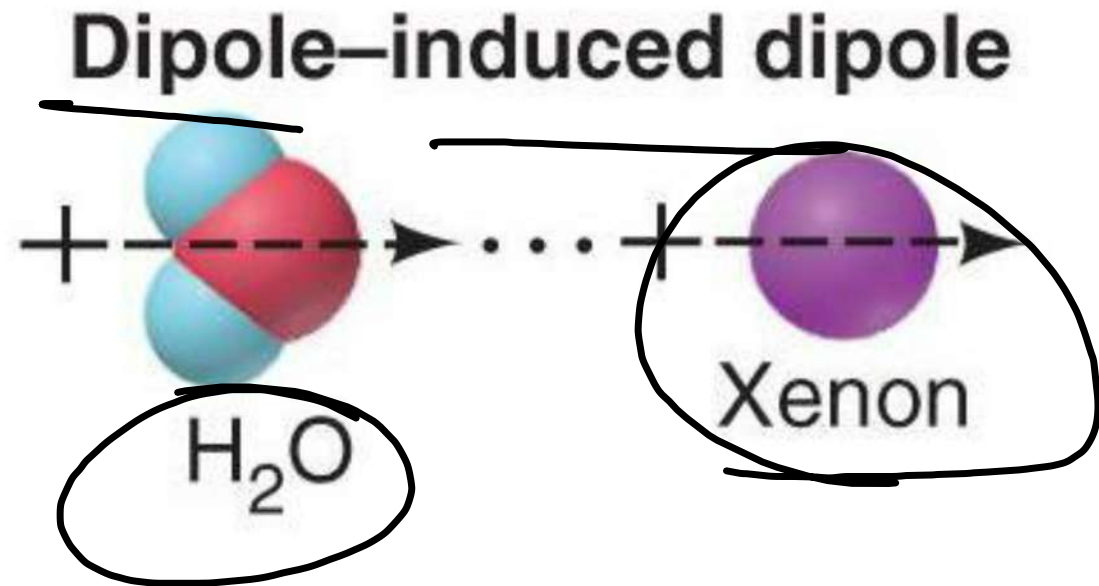
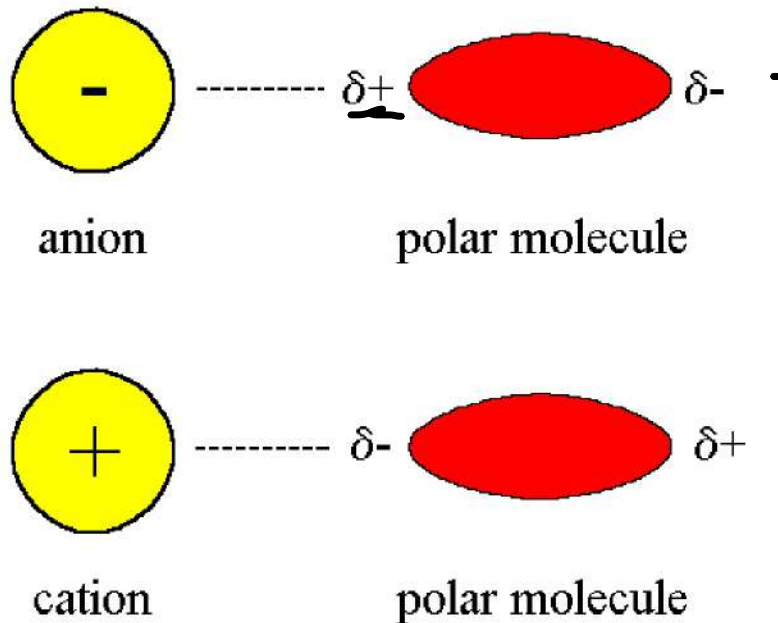
D - debye



Dipole-dipole interaction



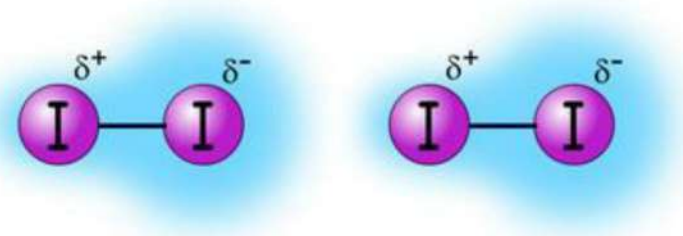
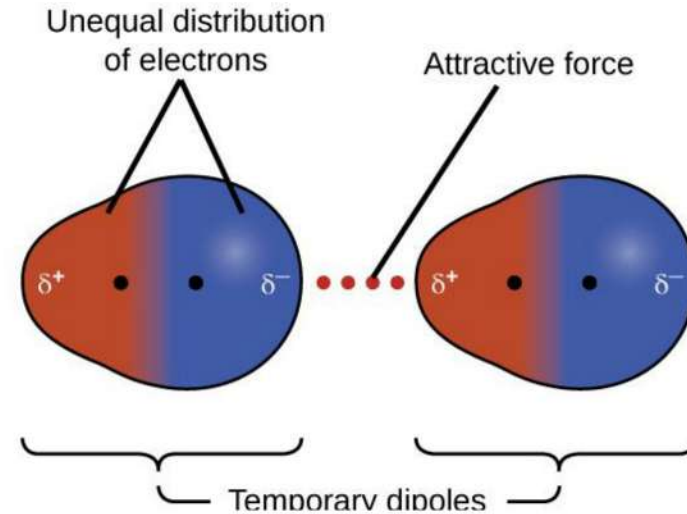
Various types of non-bonding force



Dispersion force or London force

Other name: Dispersion force is also called dipole-induced dipole attraction

F
Cl
Br
I



London force

Effectiveness: It is effective for all kinds of non-polar atoms and molecules.

Nomenclature : Named after Fritz London.

Dependence: The more the electron in an atom or molecule and the lower the nuclear attraction on its outer electrons, the greater this force becomes.

✓ Effect: 10⁸ ২

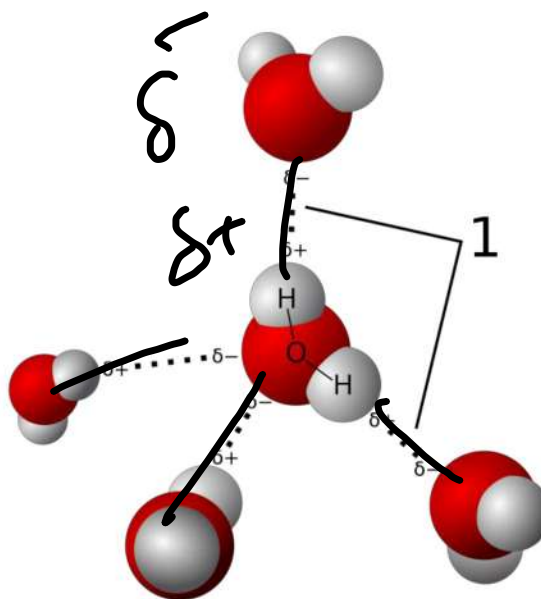
- Physical states of halogen as gas, liquid and solid are affected by dispersion force.
- Nano scale particle can make suspension due to dispersion force, example- gold suspension.

Hydrogen bond

Hydrogen bonding forms for two reasons:

- More polarization occurs between H atom with small but more electronegative N, O, F atoms etc.
- Lone pair electrons of N, O, F etc. attract H to form weak H-bond.

Example: H_2O , HF , NH_3



Water

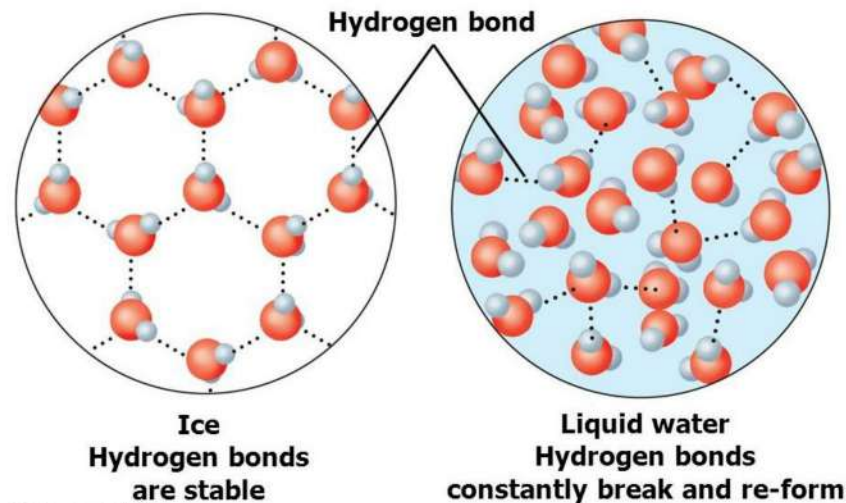
(4)

Effects of Hydrogen bond on compounds

- As effective H-bond is present in H_2O , HF , NH_3 molecules, they have higher boiling and melting points than any other hydrides.
- H_2S is non polar covalent molecule; but H_2O is polar.

H_2S — gas

H_2O — liquid



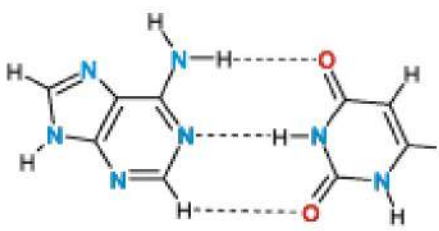
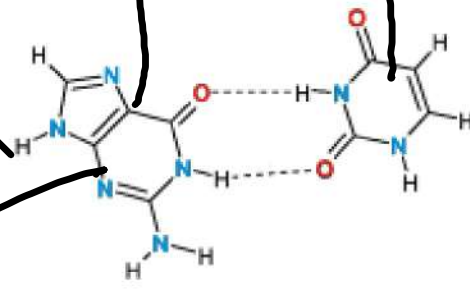
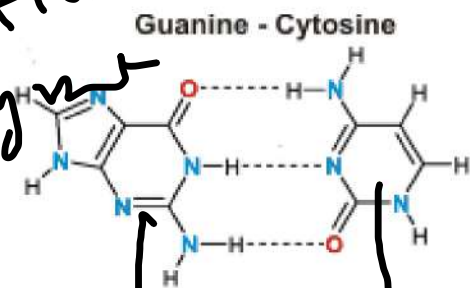
Importance of hydrogen bond

- water
- secondary and tertiary proteins chains.
- Ice floats on water
- Double-helix or two chains of nucleic acids

Hydrogen
Enzyme

Nucleic acid

Restriction
enzyme
phosphodiester
bond



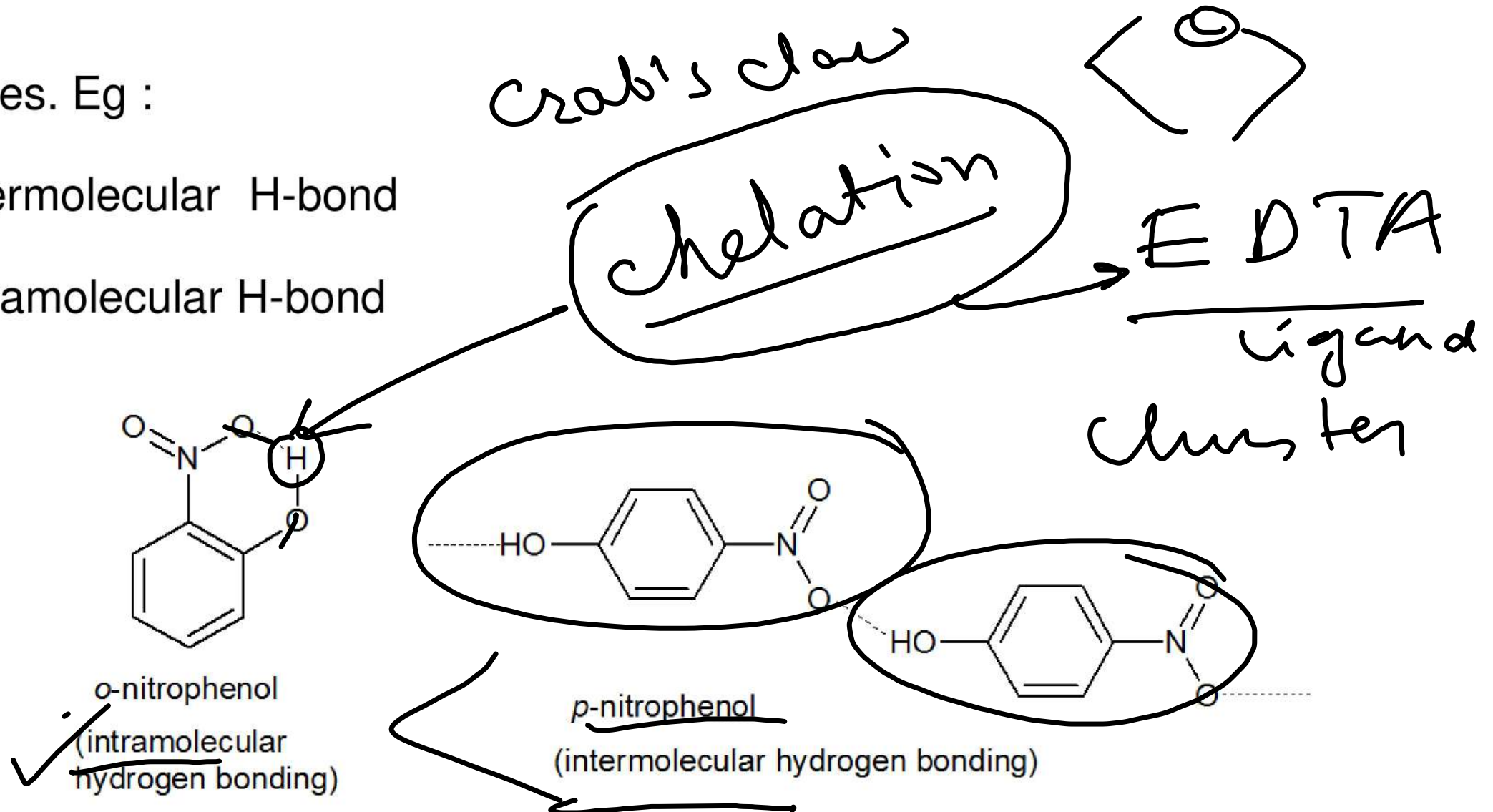
$A = T \quad A \therefore \therefore T$

$G \therefore \therefore C$

Types of hydrogen bond

2 types. Eg :

- intermolecular H-bond
- Intramolecular H-bond



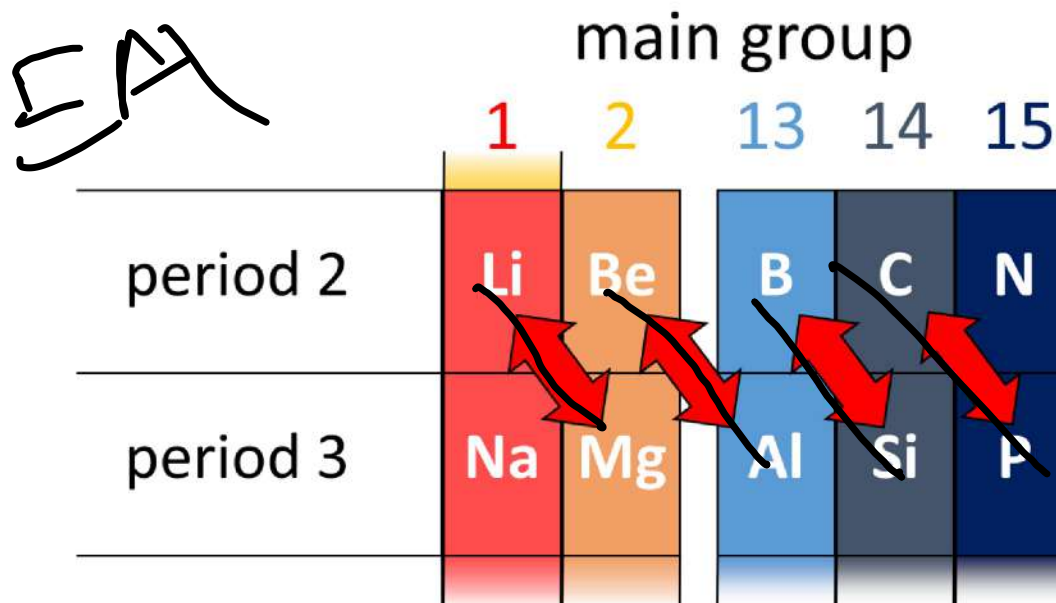
Diagonal relations of elements

Similarity in :

- Ionic radius ✓
- Ionization energy ✓
- Electronegativity ✓
- Polarization power ✓
- Charge density ✓

EA

	main group				
	1	2	13	14	15
period 2	Li	Be	B	C	N
period 3	Na	Mg	Al	Si	P



Nomenclature of inorganic compounds

Some common polyatomic ions:

Name of ion	Formula	Charge	Name of ion	Formula	Charge
Ammonium	NH_4^+	+1	Permanganate	MnO_4^-	-1
Nitrite	NO_2^-	-1	Acetate	$CH_3CO_2^-$	
Nitrate	NO_3^-		Carbonate	CO_3^{2-}	-2
Hydroxide	OH^-		Sulfite	SO_3^{2-}	
Cyanide	CN^-		Sulfate	SO_4^{2-}	
Hydrogen carbonate	HCO_3^-		Thiosulfate	$S_2O_3^{2-}$	
Hydrogen sulfate	HSO_4^-		Hydrogen phosphate	HPO_4^{2-}	
Di-hydrogen phosphate	$H_2PO_4^-$		Chromate	CrO_4^{2-}	
Hypo chlorite	ClO^-		Di-chromate	$Cr_2O_7^{2-}$	
Chlorite	ClO_2^-		Peroxide	O_2^{2-}	
Chlorate	ClO_3^-		Phosphate	PO_4^{3-}	-3
Perchlorate	ClO_4^-				

Nomenclature of oxo-acids

In case of nomenclature of multiple oxo-acids of same elements, the oxidation number of the central atom is considered as the basis.

(1) The oxo-acid in which the oxidation number of the third non-metal element is the lowest, in case of its nomenclature ('hypo' + third element + 'us' acid) this rule is followed.

Name	Formula	Oxidation Number of central atom
Hypo chlorous acid	HClO	Oxidation number of Cl <u>+1</u>
Hypo bromous acid	HBrO	Oxidation number of Br +1
Hypo iodous acid	HIO	Oxidation number of I +1
Hypo nitrous acid	H ₂ N ₂ O ₂	Oxidation number of N +1

- (2) The oxo-acid in which the oxidation number of the third element is higher than the hypo-acid, in this case (third element + 'us' acid) this rule is followed.

Name	Formula	Oxidation Number of central atom
Chlorous acid	HClO_2	+3
Bromous acid	HBrO_2	+3
Iodous acid	HIO_2	+3
Sulphurous acid	H_2SO_3	+4

- (2) The oxo-acid in which the oxidation number of the third element is higher than the us-acid, in this case (third element + 'ic' acid) this rule is followed.

Name	Formula	Oxidation Number of central atom
Chloric acid	HClO_3	+5
Bromic acid	HBrO_3	+5
Iodic acid	HIO_3	+5
Sulphuric acid	H_2SO_4	+6

- (4) The oxo-acid in which the oxidation number of the third element is higher than the ic-acid, in this case (per + third element + 'ic' acid) this rule is followed.

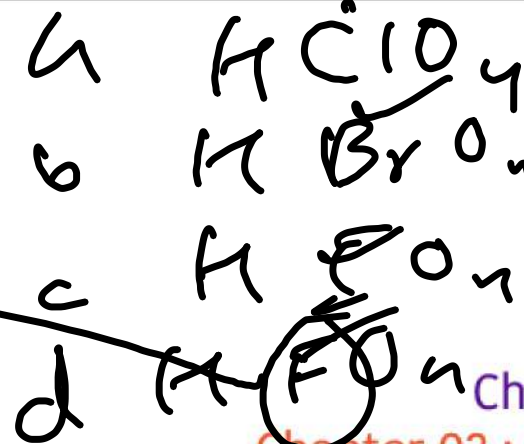
Name	Formula	Oxidation Number of central atom
Perchloric acid	HClO_4	+7
Persulphuric acid	$\text{H}_2\text{S}_2\text{O}_8$	+7

- (5) P_2O_5 is an acidic oxide. The nomenclature of acids produced by addition of different numbers of molecules with is given below-

Water	Name of the acid	Formula
✓ 3 H_2O	Ortho-phosphoric acid	$2\text{H}_3\text{PO}_4$
✓ 2 H_2O	Pyro-phosphoric acid	$\text{H}_4\text{P}_2\text{O}_7$
✓ H_2O	Meta-phosphoric acid	<u>2HPO_3</u>

Oxo-acid of halogens

Hypohalous acid	Hypo chlorous acid (HOCl), Hypo bromous acid (HOBr), Hypo iodous acid (HOI)
Halous acid	Chlorous acid (HClO_2), Bromous acid (HBrO_2), Iodous acid (HIO_2)
Halic acid	Chloric acid (HClO_3), Bromic acid (HBrO_3), Iodic acid (HIO_3)
Per halic acid	Perchloric acid (HClO_4), Per iodic acid (HIO_4)
<ul style="list-style-type: none"> Perchloric acid is the strongest acid amongst the oxo acids. There is no oxo-acid of F. 	



Nomenclature of hydrate compounds

Name	Formula
Copper sulfate penta-hydrate (Blue vitriole)	$\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$
Iron sulfate penta-hydrate (Green vitriole)	$\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$
Magnesium sulfate hepta-hydrate <i>Epsom salt</i>	$\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$
Zinc fluoride tetra-hydrate	$\text{ZnF}_2 \cdot 4\text{H}_2\text{O}$
Sodium sulfate deca-hydrate <i>Glauber salt</i>	$\text{Na}_2\text{SO}_4 \cdot 10 \text{H}_2\text{O}$
Zinc sulfate hepta-hydrate <i>white vitriol</i>	$\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$
Calcium sulphate dihydrate (Gypsum)	$\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$

Plaster of Paris \rightarrow $(\text{CaSO}_4)_2 \cdot \text{H}_2\text{O}$

Previous Question

Which one is 'pyro' acid?

[DAT: 16-17]



Previous Question

Chemical formula of Gypsum-

[DAT: 16-17]



Poll Question-08

Which is not correct formula of oxo-acid?

[MAT: 07-08]

☒ (a) Meta phosphoric acid-HPO₂



(b) Phosphinic acid- H₃PO₂

(c) Phosphonic acid-H₃PO₃

(d) Ortho-phosphoric acid-H₃PO₄

লেগে থাকো সৎ ভাবে,
স্বপ্ন জয় তোমারই হবে।



উন্মেষ

মেডিকেল এন্ড ডেন্টাল এডমিশন কেয়ার