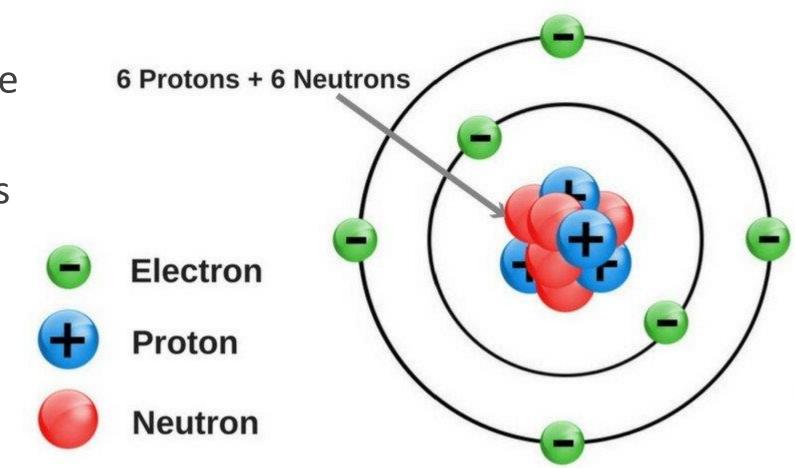
Structure of Matter

DR. AHMED MAGDY SAYED LECTURER OF DENTAL BIOMATERIALS

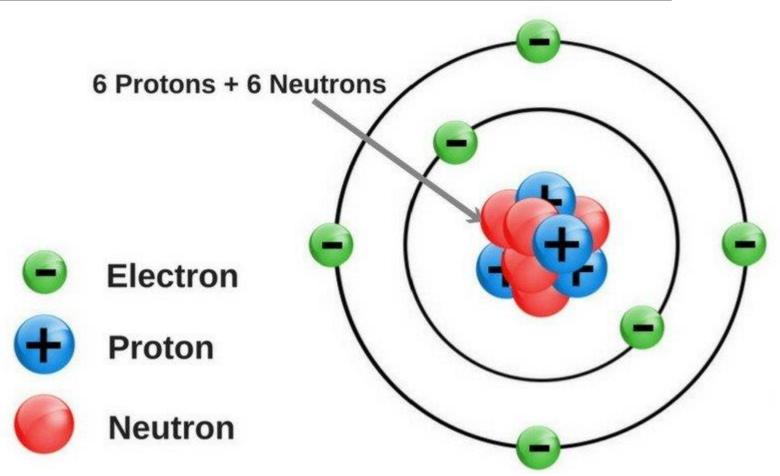
Chapter Content:

- 1. Atomic Structure.
- 2. Atomic bonds (Primary and Secondary bonds).
- 3. Inter-atomic distance.
- 4. Classification of solids (acc to bond and arrangement)
- 5. Atomic packing factor
- 6. Imperfection in crystalline solids
- 7. Polymorphism
- 8. Correlation between atomic structure and material properties

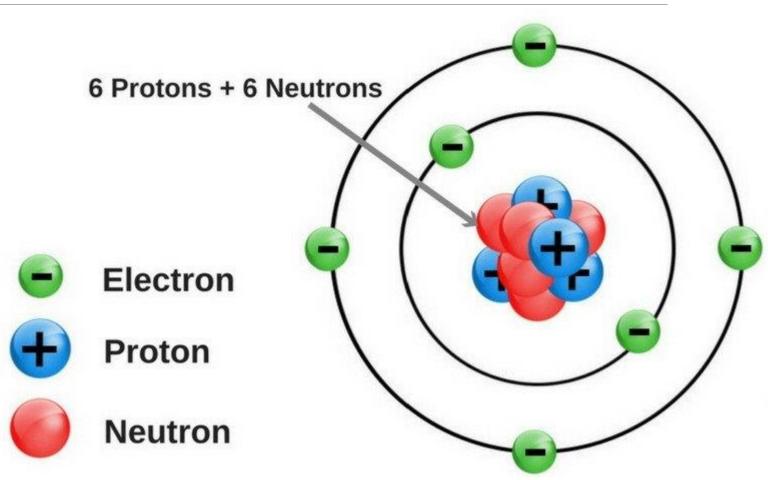
- The basic unit of any material is the atom
- The atom consists of a nucleus and surrounding electrons



- Nucleus is the core of the atom and composed of:
- 1. Positively charged protons.
- 2. Uncharged neutrons.



- Electrons surround the nucleus in shells
- 1. Negatively charged.
- 2. The most outer electrons are called VALANCE ELECTRONS
- 3. Valence electrons affect the <u>physical &</u> <u>chemical properties</u>



Definitions:

1. Atomic number = Number of electrons = number of protons.

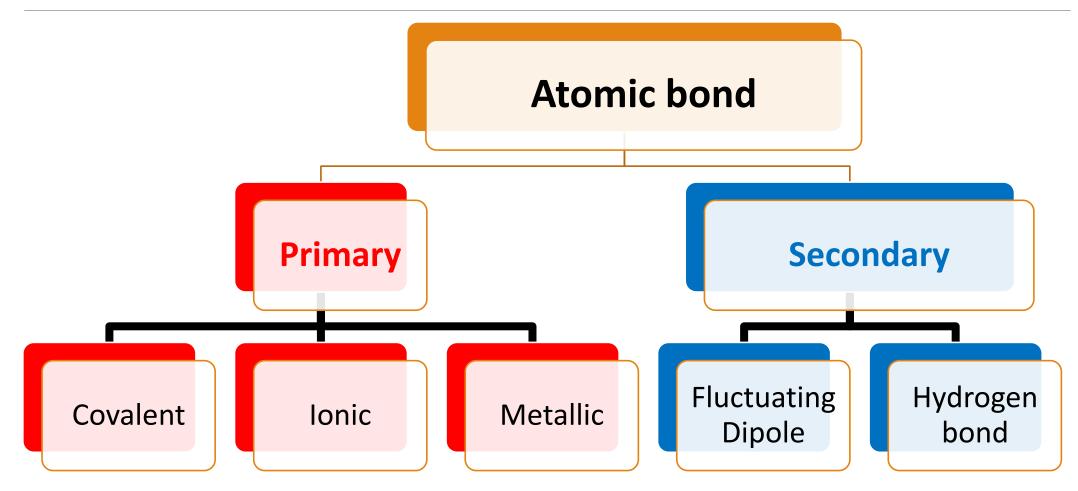
2. Atomic weight = weight of protons + neutrons.

Every element tries to reach the stable configuration by having 8 electrons in its outer shell by:

- 1. Releasing Electrons Decome +ve charged
- 2. Receiving extra electrons \bigcirc become –ve charged
- 3. Sharing electrons with other atom

Then forming the **atomic bonds**

Atomic Bonds



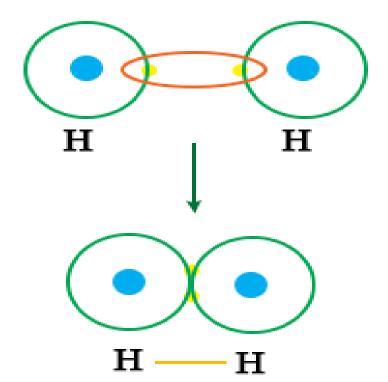
Covalent bond

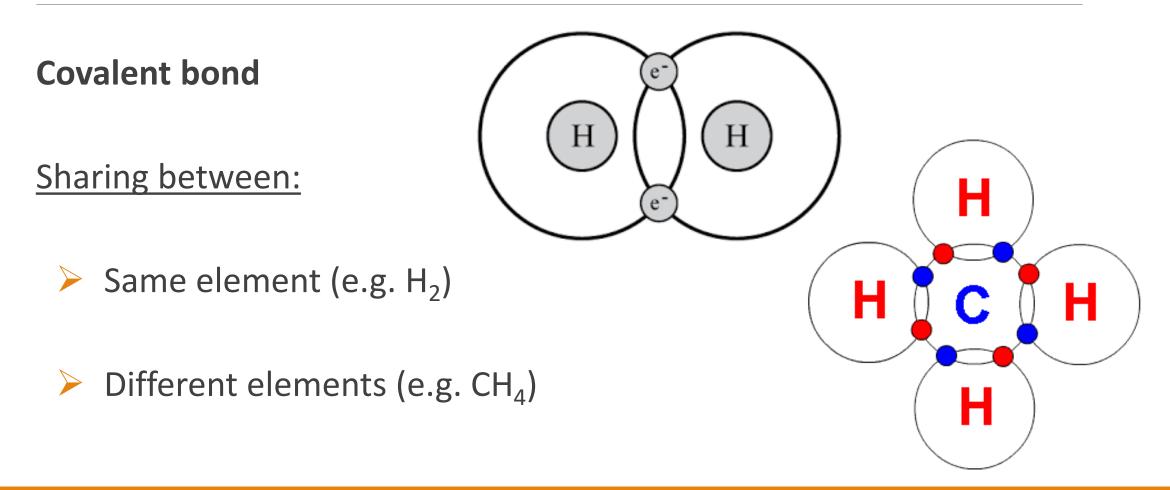
Arises by sharing electrons

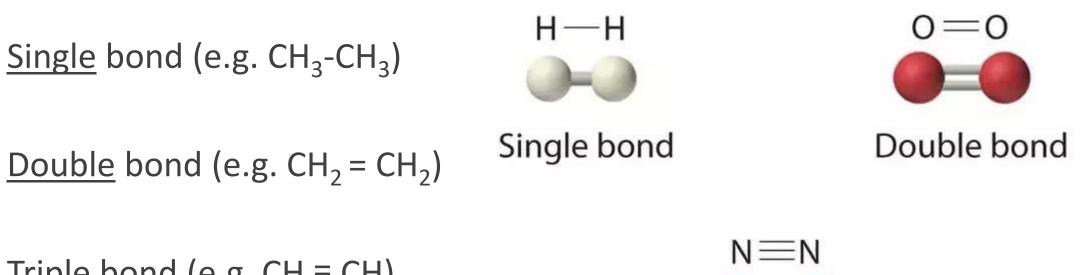
between atoms, the atoms

approach one another and

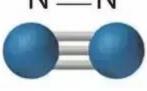
orbital overlap happened.



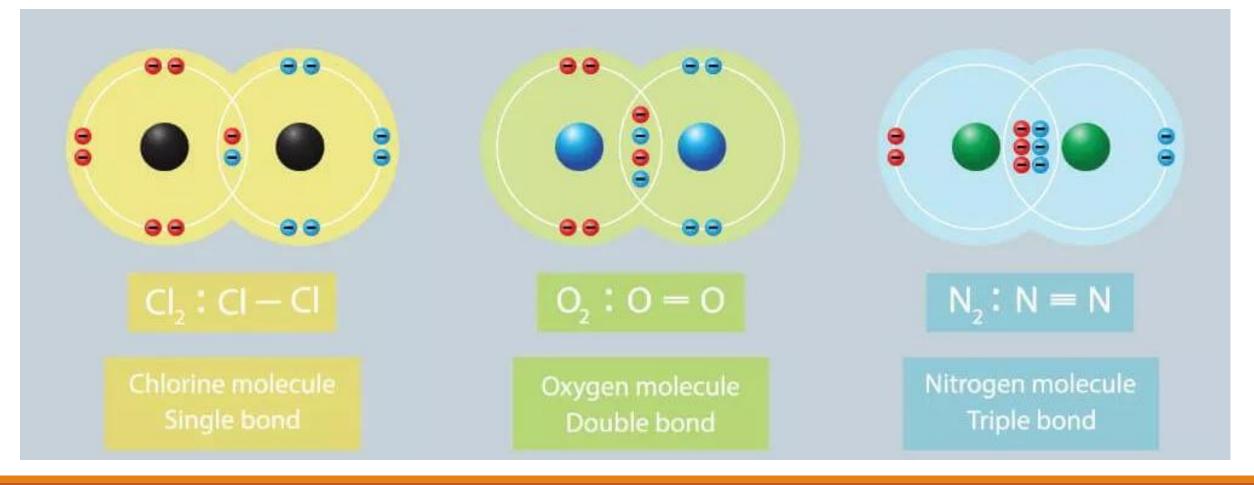


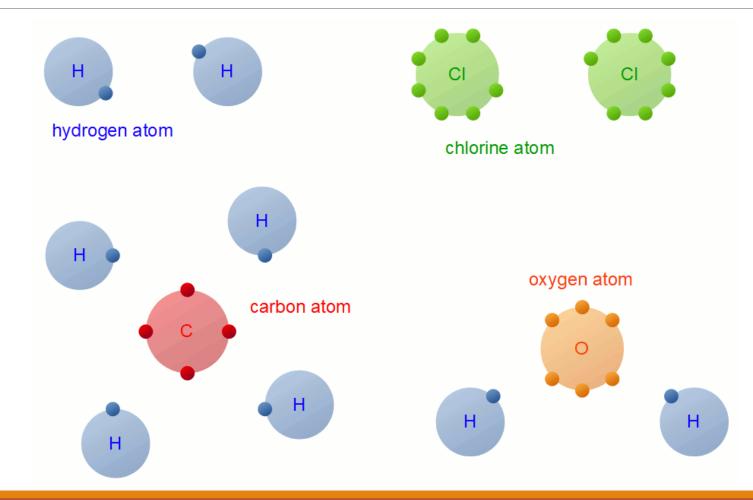


<u>Triple</u> bond (e.g. $CH \equiv CH$).

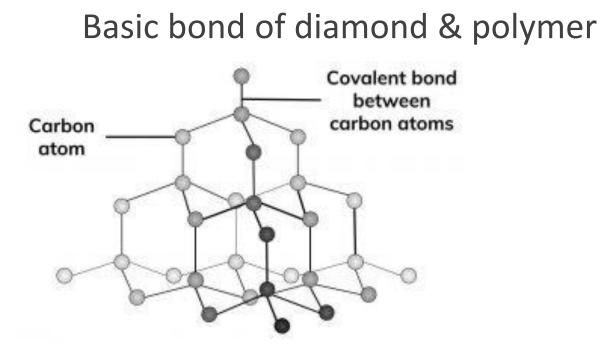


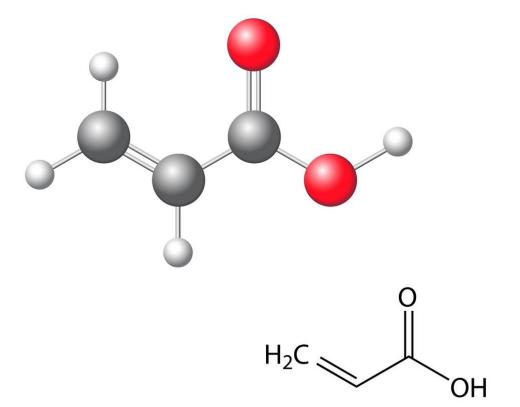
Triple bond





Examples:



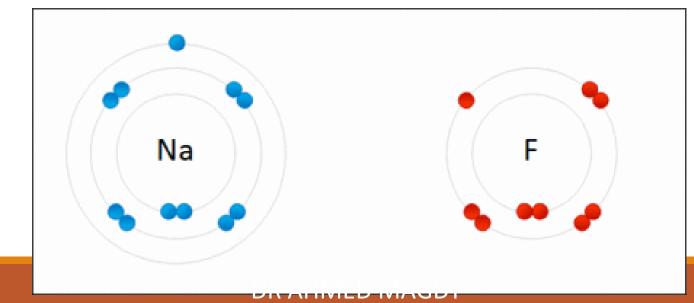


Characteristics of covalent bond:

- **1**. Highly directional bond.
- 2. High strength and hardness.
- 3. High heat resistance.
- 4. Thermal and electrical insulators.
- 5. Dissolve in organic solvents.

Occurs by:

- 1. Electron transfer from one atom (become +ve ion)
- 2. To another (Become –ve ion)
- 3. Electrostatic attraction occurs between them.



Sodium gives his outer electron to Chlorine \bigcirc Na⁺ Cl⁻.

Examples: Basic bond for Ceramics, Glasses.



Characteristics of Ionic bond:

- 1. Spherical in nature.
- 2. High strength and hardness.
- 3. High heat resistance.
- 4. Insulators as solids.
- 5. Electric conductors in solutions.
- 6. Dissolve in ionizing solvents (not in organic solvents).

Metallic bond

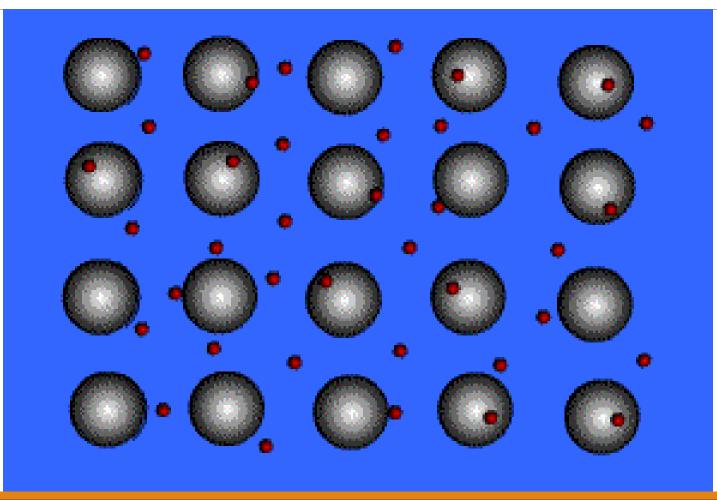
The metals have loosely held valence electrons, so they move freely

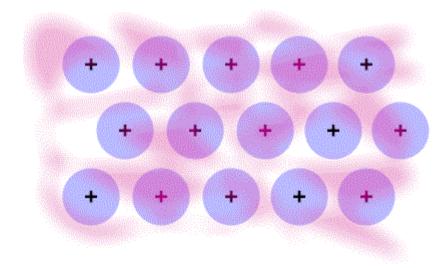
between atoms forming *cloud of electrons*.

Metallic bond

So the metallic bond results from the attraction between consisting

+ve ion cores and the surrounding cloud of free electrons.





Characteristics of Metallic bond:

- 1. High strength and hardness.
- 2. High thermal resistance.

Characteristics of Metallic bond:

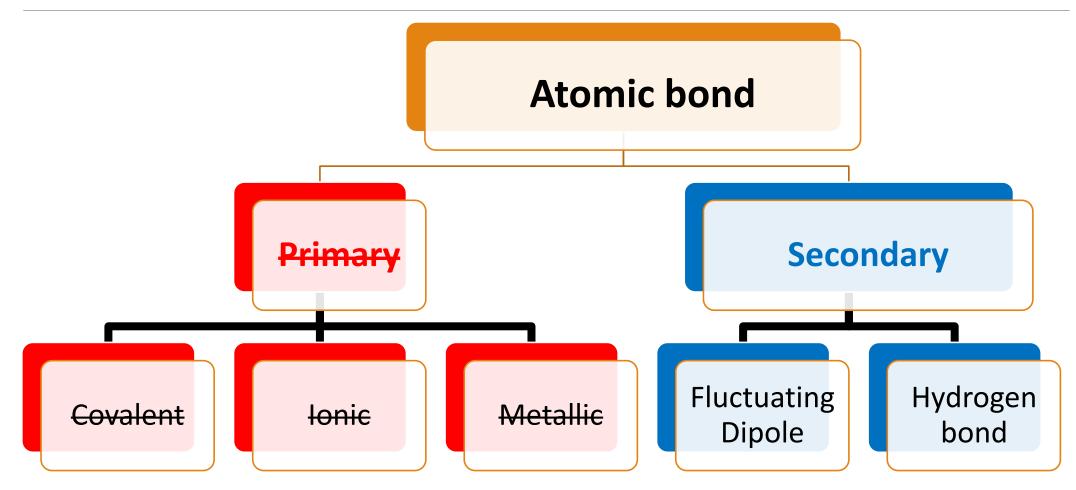
- 3. High thermal conductivity (as <u>free electrons</u> conduct heat).
- 4. High electric conductivity (as <u>free electrons</u> carry the electricity).
- 5. Opaque (as <u>free electrons</u> absorb light).
- 6. Lustrous (as <u>free electrons</u> reflect the light).

Characteristics of Metallic bond:

7. Leads to crystalline arrangement in metals

8. Leads to easy of deformability of metals.

Atomic Bonds



Fluctuating Dipole:

It developed between atoms due to asymmetry of electron distribution.

> This asymmetry gives the atom dipole character.

> It is a temporary bond.

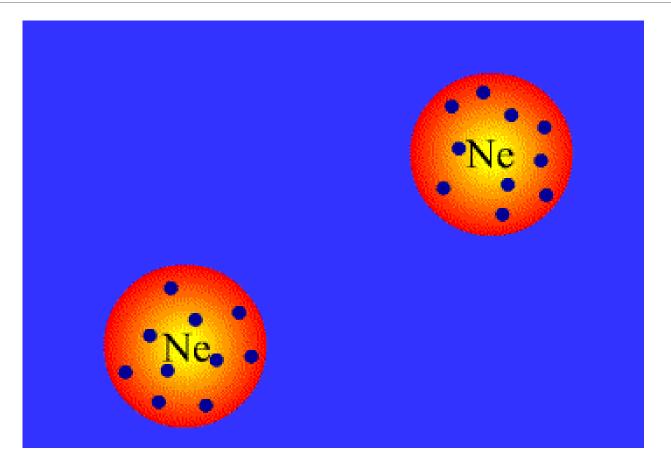
Fluctuating Dipole:

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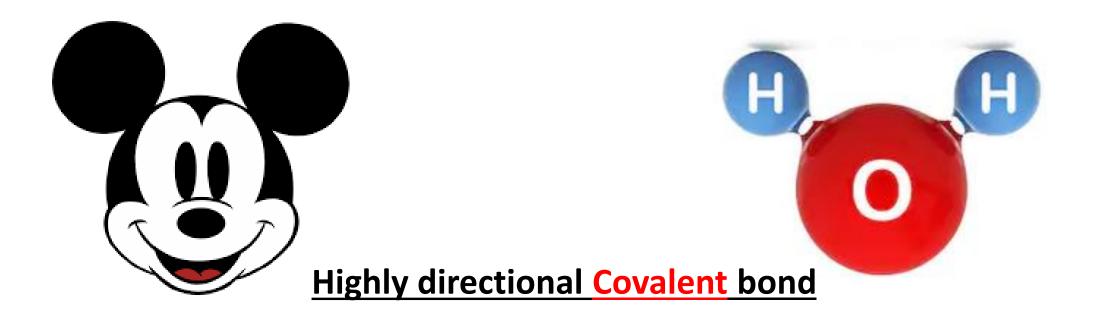
Fluctuating Dipole:



Hydrogen bond:

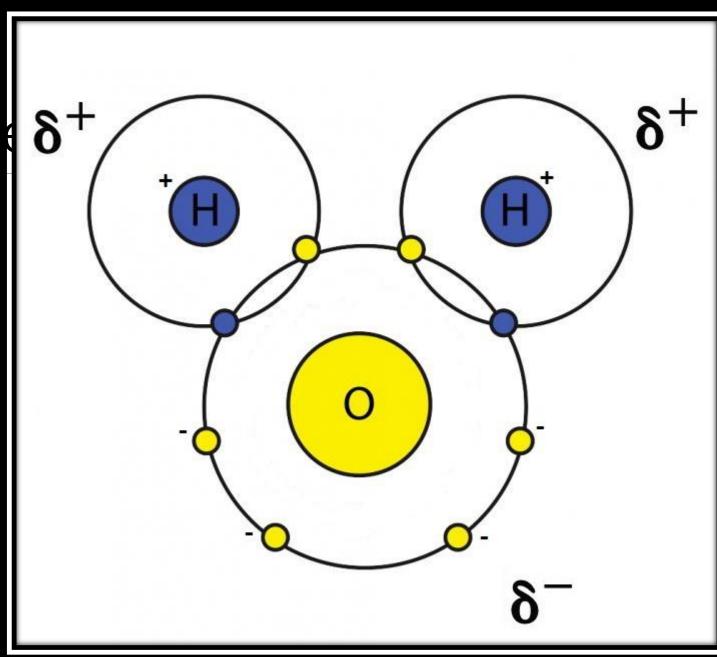
It developed between molecules when each molecule has a dipole.

Hydrogen bond:



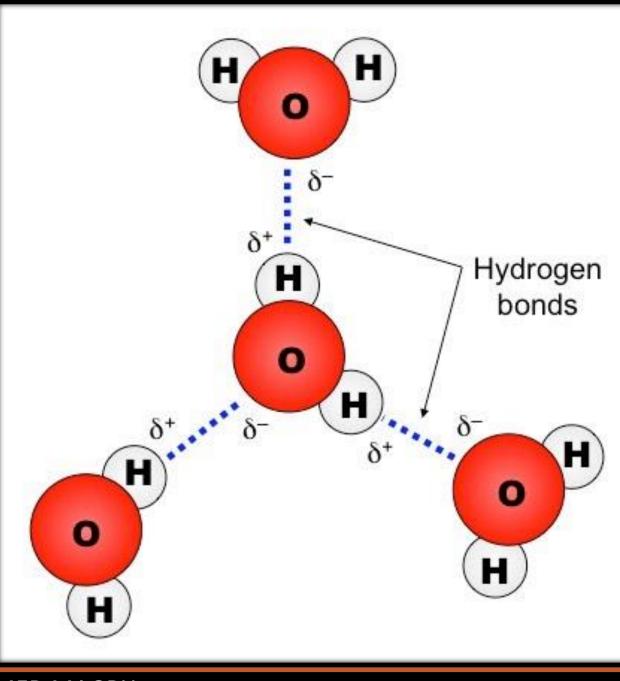
Atomic Bonds (S δ^+ ,

Hydrogen bond:



Atomic Bonds (Sec

Hydrogen bond:



Characteristics of secondary bond:

- 1. Low strength and hardness.
- 2. Low heat resistance.
- 3. High thermal expansion.

Inter-atomic Distance

It is the distance between atoms.

Resulted from 2 equal but opposite forces:

- 1. Repulsive forces \rightarrow due to electrostatic field of each atom.
- 2. Attractive forces \rightarrow different types of atomic bonds.

Inter-atomic Distance

Factors affecting interatomic Distance:

- 1. Temperature
- 2. Number of adjacent atoms:
- **3**. Type of the bond:
- 4. External forces

Factors affecting interatomic Distance:

1. <u>Temperature</u>

Heat increases I.A.D (as it increases energy of atoms)



2. <u>Number of adjacent atoms:</u>

More adjacent atoms will increase I.A.D (as less specific

attraction to any neighboring atom).



3. Type of the bond:

Increase number of shared electrons in covalent bond will decrease I.A.D (as it is means more strong bond).

 $CH \equiv CH$ is stronger than $CH_2 = CH_2$, stronger than CH_3 - CH_3 .







4. Any external forces

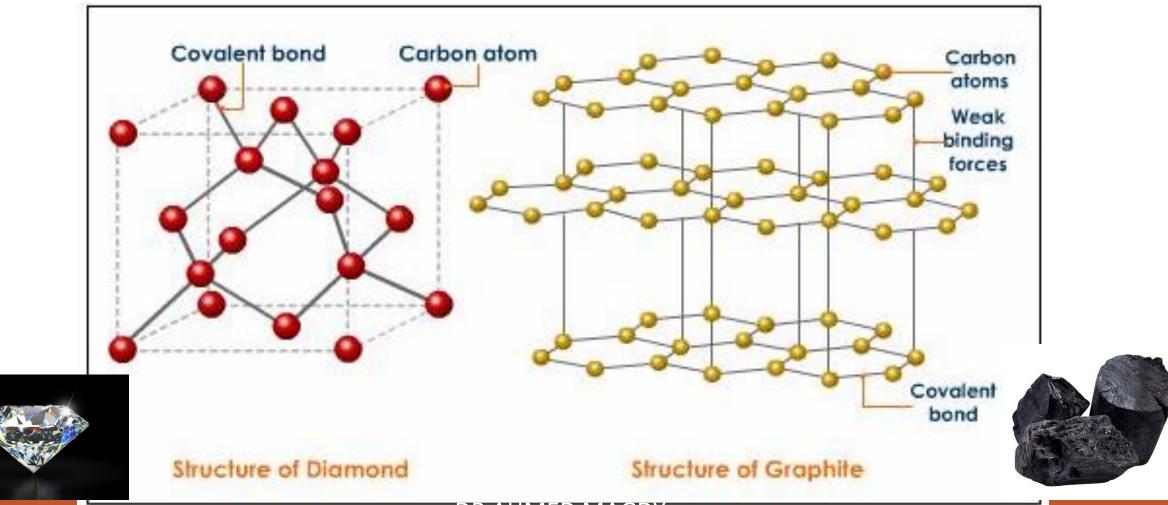
May displace atoms and change I.A.D.



According to intermolecular bond:

Bond between atoms = inter-atomic bond = intra-molecular bond Must be primary bond Bond between molecules= inter-molecular bond May be primary or secondary Intra جوة Inter بين

	Atomic Solids	Molecular Solids
Bonds between atoms	Primary	Primary
Bonds between molecules	Primary	Secondary
Properties	High strength and	Low strength and
	hardness	hardness
Example	Diamond	Polymers



According to Arrangement of Atoms:

Amorphous



Crystalline 100

Amorphous Solids	Crystalline Solids
Atoms are randomly distributed or with	Atoms are regularly arranged with
very short arrangement	repetition in 3D (called space lattice or
	<u>crystal lattice</u>)

Amorphous Solids	Crystalline Solids
Have high internal energy	Have low internal energy
<image/>	

Amorphous Solids	Crystalline Solids	
Have no definite melting temperature (gradually	Have definite melting	
soften by heating and gradually harden by cooling)	temperature	
Glass Transition temperature:		
The temperature at which the amorphous solids		
start to soften or harden		

Amorphous Solids	Crystalline Solids
Examples: Wax, Glass	Metals

The atoms arrange themselves in a repeated manner.

The smallest repeated unit in a crystal lattice is called unit cell.

Unit cell may be one of 7 main patterns and subdivided to make 14 possible patterns.

Cubic System:

Axes: a = b = c

Angles between axes: 90°

It is subdivided into:

- 1. Simple cubic system
- 2. Body centered cubic
- **3**. Face centered cubic

Cubic system

Simple cubic system	Body centered cubic	Face centered cubic
8 atoms at the corner	8 atoms at the corner	8 atoms at the corner
	+ One atom at the center	• One atom at each face
1 atom (8 x 1/8)	2 atoms (8 x 1/8) + 1	4 atoms (8 x 1/8) + (6 x 1/2)

Hexagonal System:

Axes: $a = b \neq c$

Angles between axes: $\alpha = \beta = 90^{\circ}$, $g = 120^{\circ}$

It is subdivided into:

- 1. Simple hexagonal system
- 2. Hexagonal closed packed



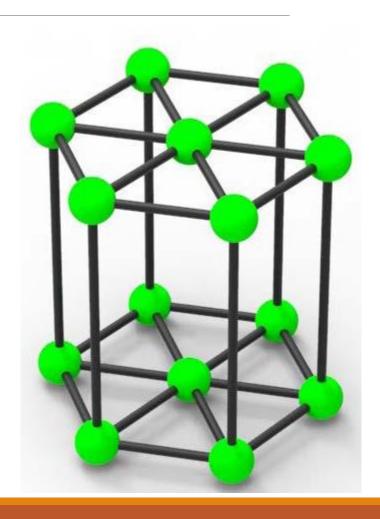
Hexagonal System:

Axes: $a = b \neq c$

Angles between axes: $\alpha = \beta = 90^{\circ}$, g = 120°

It is subdivided into:

- 1. Simple hexagonal system
- 2. Hexagonal closed packed



Simple Hexagonal System	Hexagonal closed Packed
6 atoms at the top (6 x 1/6)	6 atoms at the top (6 x 1/6)
6 atoms at the bottom (6 x 1/6)	6 atoms at the bottom (6 x 1/6)
One at the upper face (1 x ¹ / ₂)	One at the upper face (1 x ¹ / ₂)
One at the lower face(1 x ¹ / ₂)	One at the lower face (1 x ¹ / ₂)
	3 atoms at the center
1 + 1 + ½ + ½ 3 atoms	$1 + 1 + \frac{1}{2} + \frac{1}{2} + 3$ 6 atoms

Atomic Packing Factor

It is the fraction of space occupied by the atoms

APF = <u>Volume of atoms inside unit cell</u> Volume of unit cell

Simple cubic system = 0.54.

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Body Centered Cubic = 0.68.
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Face Centered Cubic = 0.74.

Hexagonal Closed Packed = 0.74.

Atomic Packing Factor

Clinical importance:

Materials with \uparrow APF \bigcirc have \uparrow densities and strength properties.

Theoretical calculation of strength is much higher than actual strength.

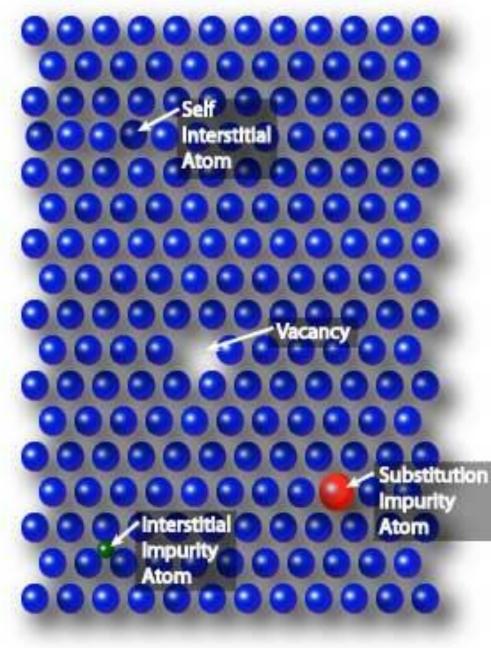
This is due to the presence of defects in the crystalline system

Types of Crystalline Defects:

- 1. Point defect.
- 2. Line Defect.
- 3. Plane Defect.

- 1. Point Defects:
 - a) Vacancy \rightarrow missing atom.
 - b) Self interstitial atom \rightarrow extra atom from same metal.
 - c) Interstitial impurity \rightarrow extra atom from another metal.
 - d) Substitutional impurity \rightarrow replaced atom from another metal.

Imperfections of Crystallir



2. Line Defects:

It is the most common type.

Dislocation:

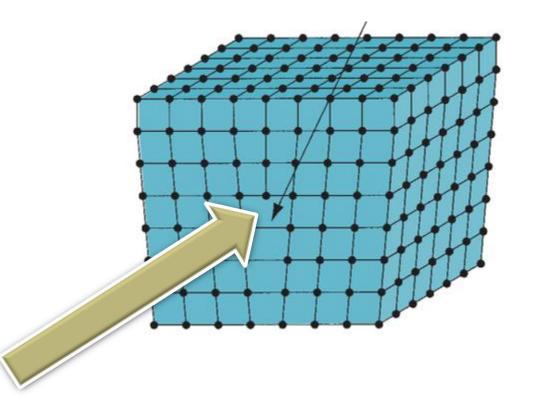
It is the displacement of a raw of atoms from their normal positions in the lattice

<u>Clinical importance</u>: plastic deformation in metals occurs by motion of dislocations

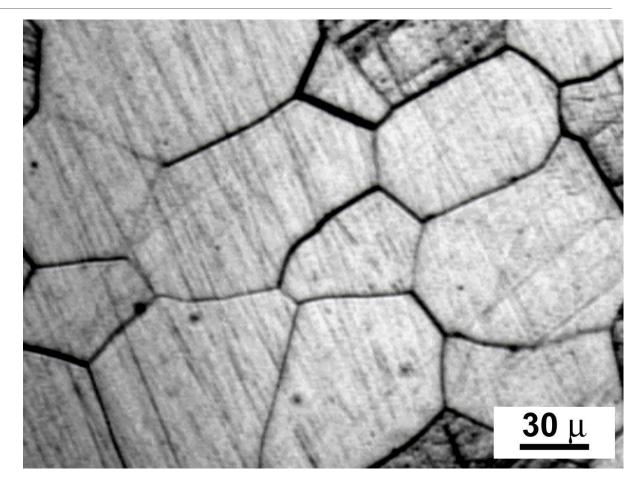
2. Line Defects:

Clinical importance:

Helps in plastic deformation in metals



Plane (planer) Defects:
Like Grain boundaries.



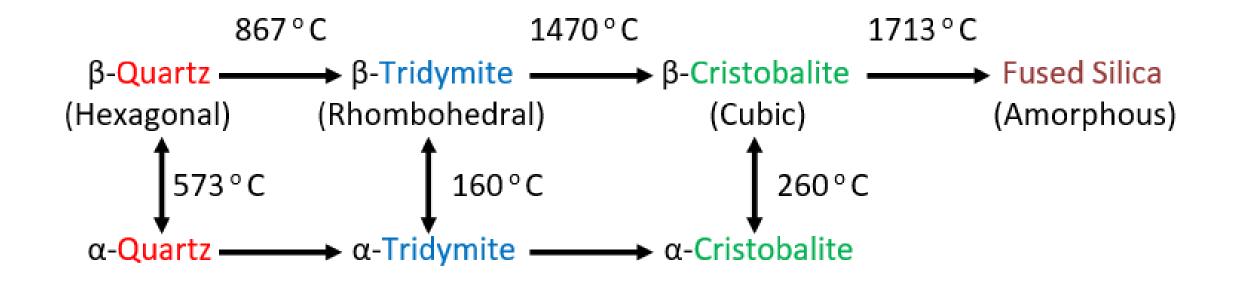
Materials that have the <u>same chemical</u> composition but found naturally in different physical forms.

The polymorphic forms have different physical properties as density,

freezing point, optical properties, conductivities ... etc

If the material is **inorganic**, the polymorphism is called **allotropy**.

If the material is **organic**, the polymorphism is called **isomerism**.



Displacive transformation	Reconstructive transformation
No breakdown of atomic bonds	Breakdown of atomic bonds
	followed by reconstruction of
	new space lattice
Accompanied by expansion	No Expansion
Rapid transformation	Slow transformation
Occurs at lower temperatures	Occurs at higher temperature

1. Density:

Controlled by:

- a) Atomic weight.
- b) Atomic radius.
- c) Atomic packing factor.

- **2.** Bond strength leads to:
 - a) High strength and hardness.
 - b) High melting and boiling temperature.
 - c) Low coefficient of thermal expansion.

3. Electrical and Thermal conductivity:

Depends on nature of atomic bond

- a) Metallic solids conduct heat and electricity.
- b) Ionic solution conduct electricity, while ionic solids are electrical insulators.
- c) Covalent solids are insulators.

4. Crystalline solids have low internal energy

Because they send their internal energy in arranging their atoms.





5. FCC is more ductile than BCC due to higher atomic packing factor.





 FCC is more ductile than HCP due to symmetry of the cubic system.





