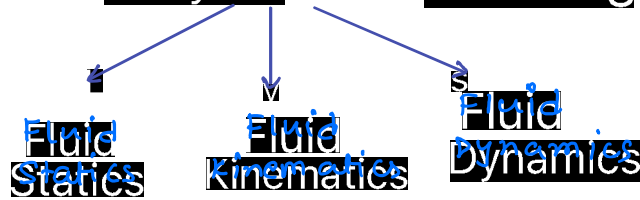


**FLUID MECHANICS**  
**JASPAL SINGH SIR**

# FLUID MECHANICS

- Fluid Mechanics is a branch of engineering, that deals with the properties of the fluid at rest as well as in motion.

- It is further analysed in following forms.



- Fluid Statics :- It deals with fluid in rest condition.

- Fluid Kinematics :- It deals with fluid in motion without considering the forces responsible for motion.

- Fluid Dynamics :- It deals with fluid in motion by considering the forces responsible for motion.

- In general matter exist in following states

(A) SOLID

(B) FLUID  $\rightarrow$  LIQUID  
 $\rightarrow$  GASES

- A fluid is a substance which exists in liquid and gaseous phase.

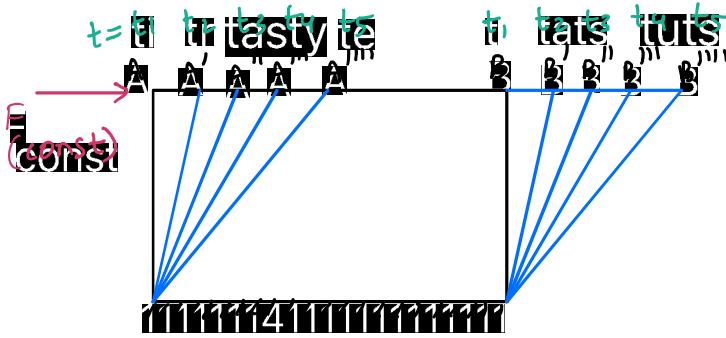
- Fluid is capable of deforming continuously under the action of shear force. Shear stress or tangential stress however small the magnitude of stress may be.

- Hence in solids where stress is proportional to strain, in fluid stress is proportional to rate of deformation or rate of strain.

NOTE :- Mere continuous deformation of the fluid means that when a constant shear stress is applied over the fluid, it will continue to deform as long as stress is present.

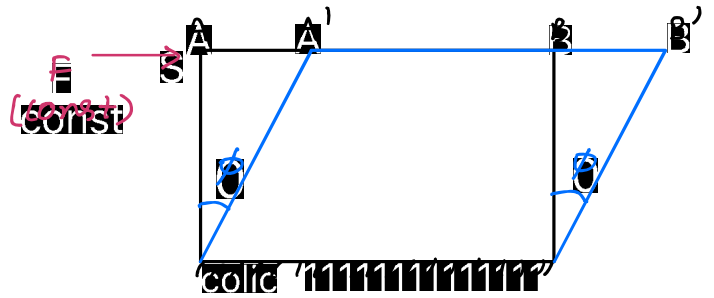
But in case of solid, deformation eventually stops at fixed strain levels.

## FLUID



$$\tau \propto \frac{d\phi}{dt}$$

## SOLID



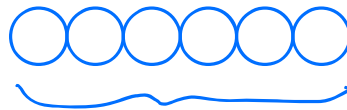
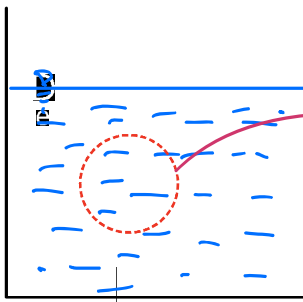
$$\phi = \text{shear strain}$$

$$F \propto \phi$$

Hence in fluid, we are more concerned with rate of deformation and in solids we are more concerned with its magnitude.

Fluid can also be stated to be in CONTINUUM for its analysis.

In macro system, when the intermolecular distances are very small as compared to dimensions of the medium (system), we can assume that the adjacent to one molecule, there is another molecule without any space in between. Hence the entire fluid mass can be considered as continuous distribution of mass, which is termed as continuum.



continuous distribution of mass: CONTINUUM.

GLASS/ BOTTLE etc.

Fluid may also be defined as the substance that is capable of flowing and it has no definite shape of its own and conforms to the shape of container.

- Fluid can be classified as :-

(a) Liquid :- It is a fluid which has a definite volume, and it changes slightly with temp and pressure. (For practical purposes it is considered to be incompressible i.e. does not show change in volume)

(b) Gas :- It is a fluid, which is compressible & possesses no definite volume & always expands up to its volume is equal to the volume of container, it is placed in.

**NOTE:-** A vapour is a gas, temp and pressure of which is very near to liquid.  
eg → steam.  
Souvik 2021

(c) IDEAL FLUID :- This are the fluids which have no viscosity, surface tension, and are incompressible.

- They do not offer resistance against flow.

- It is an imaginary or hypothetical fluid, considered only for simplicity in mathematical calculation.

eg:- Those fluids having very low viscosity may be considered as ideal fluid. eg water, air.

(d) REAL / PRACTICAL :- All fluid that exist in nature are real fluids.

- They offer resistance against flow

- possess viscosity, surface tension and compressibility.

# # PROPERTIES OF FLUID :-

## (A) MASS DENSITY / SPECIFIC MASS (ρ) :-

- It is defined as ratio of mass of fluid and its volume
- OR
- It is the mass of the fluid which possesses per unit volume.

$$\rho = \frac{M}{V}$$

UNIT

$\frac{\text{kg}}{\text{m}^3}$      $\frac{\text{g}}{\text{cm}^3}$   
(MKS)    (CGS)

Time 2

- Mass density of the fluid depends upon its Temperature and pressure.

- With increase in Temperature, molecular activity or Randomness increases thus spacing of molecules increases hence lesser number of molecules would pack in same volume that results in reduced mass density.

$$\rho \propto \frac{1}{T}$$

- With increase in pressure, large number of molecules can be forced into given volume that results in higher mass density.

$$\rho \propto P$$

- At STP,  $\rho_{\text{water}} = 999.9 \text{ kg/m}^3 \approx 1000 \text{ kg/m}^3$   
 To 0°C     $\rho_{\text{air}} = 1.292 \text{ kg/m}^3$   
 P = 1 atm

## (B) SPECIFIC WEIGHT / WEIGHT DENSITY (γ) :-

- It is the weight of the fluid per unit volume.

$$\gamma = \frac{W}{V}$$

units :-  $\text{N/m}^3$  ,  $\text{kgf/m}^3$   
 $\text{dyne/cm}^3$

$\text{dyne/cm}^3 = 10 \text{ N/m}^3$   
 like  $1 \text{ m}^3 = 1000 \text{ L}$      $1 \text{ kgf/m}^3 = 9.81 \text{ N/m}^3$

- It signifies the force exerted by gravity over the unit volume of fluid.





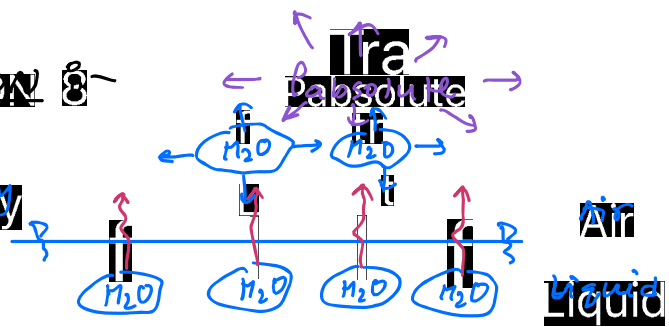
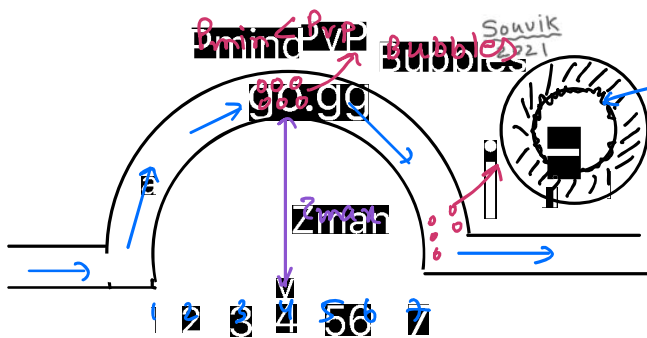
- hence,  $\rho$  can also be defined as ratio of specific weight or specific density of fluid to the specific weight or specific density of the standard fluid.

- For LIQUID standard fluid is taken as PURE WATER at 4°C.

- For GASES, sta. fluid is considered as HYDROGEN or AIR at sta. temp & pres.

- Since weight density and mass density varies with temp. must also be reported when  $\rho$  is reported.

(e) VAPOUR PRESSURE & CAVITATION :-



June 25

- At liquid air interface, continuous exchange of molecules takes place as liquid molecule escape from the surface in to gaseous form termed as vapour, due to the energy losses by them.

- This vapour molecules exert partial pressure over the liquid surface (in general space) termed as vapour pressure.

- On increase in temperature as molecular activity increases, vapour pressure also increases.

- If absolute pressure impose over the liquid becomes less than vapour pressure of liquid boiling of the liquid will start.

- Hence it can be stated that the boiling of liquid can start at any given temperature if the pressure above the liquid surface becomes less than vapour pressure.

- Liquid with a high vapour pressure evaporates readily and are termed as volatile liquid. Eg benzene

- Hg has very low VP, hence it doesn't vapourise even at very low pressure, thereby it is used in equipments for pressure measurements.

- V.P of water at different temp. is as follows

T (°C)                      VP (kPa)

10  
20  
100

1.2  
2.3  
101.3 = 1 atm.

→ Boiling start.

**cavitation** :- If in any flow system pressure at any point in the liquid becomes less than V.P, vapourisation starts resulting in pockets of dissolved gases and vapours.

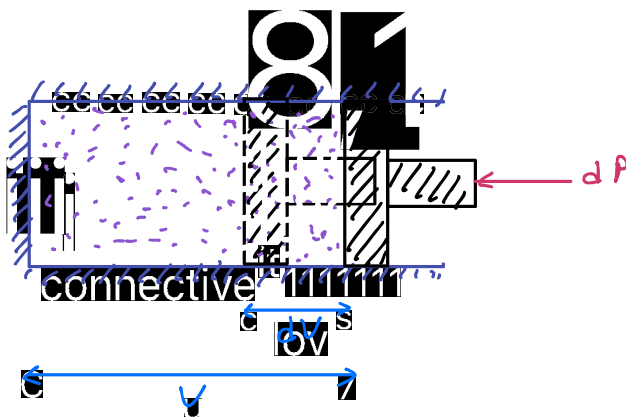
- This bubbles <sup>Sovik</sup> move with flow of liquid into the high pressure region where they burst or collapse giving rise to high impact pressure.

- Due to which VIBRATION, NOISE, EROSION SURFACE PITTING and FATIGUE FAILURE of the material takes place.

- This phenomenon is termed as cavitation.

- With increase in temp, VP increases, tendency of cavitation <sup>of vap.</sup> increases, hence tendency of cavitation also increases.

(f) **COMPRESSIBILITY AND ELASTICITY** :-



- All fluids can be compressed by application of the pressure and when this pressure is removed fluid expand to their original volume (similar to that of elastic solid)



- This property of fluid to undergo volume change on application of pressure is termed as **COMPRESSIBILITY**.

- which is quantitatively related with Bulk modulus of elasticity ( $K$ )

- **Bulk Modulus ( $K$ )** is defined as change in pressure per unit change in volume to the original volume.

$$K = \frac{\text{Volumetric stress}}{\text{Volumetric strain}} = -\frac{\Delta P}{\left(\frac{\Delta V}{V}\right)}$$

unit :-  $\text{N/mm}^2$ ,  $\text{kgf/cm}^2$ ,  $\text{gmf/cm}^2$

At NTP,  $K_{\text{water}} = 2.05 \times 10^9 \text{ N/m}^2$

$K_{\text{air}} = 1.03 \times 10^5 \text{ N/m}^2$

\* Air is 20000 or  $2 \times 10^4$  more compressible than water.

-  $K$  is not const, for fluids it increases with increase in pressure, as when a fluid mass is compressed its molecules become close together and the resistance to further compression increases.

- For eg.  $K_{\text{water}}$  almost doubles when pressure is

raised from 1 atm to 3500 atm

\* " $K$ " also varies with Temperature

• For "LIQUID", with increase in temperature, inter molecular bond b/w the particles decreases, hence resistance against vol change decreases, thus more volume change occurs. Hence  $K$  decreases.

• For "GASES", with increase in temperature, random motion of particles increases, that further increases resistance to vol change, thereby "less" volume change is observed. Hence  $K$  increases.

- Now, compressibility is quantitatively related to bulk modulus as follows.

compressibility  $\leftarrow B = \frac{1}{K}$  units :-  $\frac{m^2}{N}$

- on general,

$m = \text{const}$

$m = \rho v$

$dm = \rho dv + v d\rho$

$0 = \rho dv + v d\rho$

$-\frac{dv}{v} = \frac{d\rho}{\rho}$

$K = -\frac{dP}{\left(\frac{dv}{v}\right)} = -\frac{dP}{\left(\frac{d\rho}{\rho}\right)}$

$K = -\rho \frac{dP}{d\rho}$

$\therefore B = \frac{\rho}{d\rho} \frac{dP}{d\rho}$

→ If  $\frac{d\rho}{dP} = 0 \Rightarrow B = 0 \rightarrow$  fluid is incompressible.

→ If  $\frac{d\rho}{dP} \neq 0 \Rightarrow B \neq 0 \rightarrow$  fluid is compressible.

For liquids (water) at 20°C

$P = 1 \text{ atm}, \rho_{\text{water}} = 998 \text{ kg/m}^3$

$P = 100 \text{ atm}, \rho_{\text{water}} = 1003 \text{ kg/m}^3$

% change in  $\rho = \frac{1003 - 998}{998} \times 100 = 0.49\%$  (negligible)

Hence, liquids are considered to be incompressible.