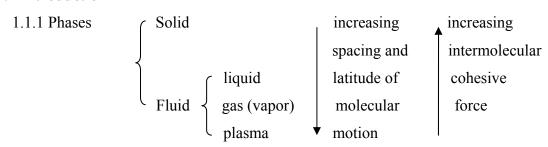
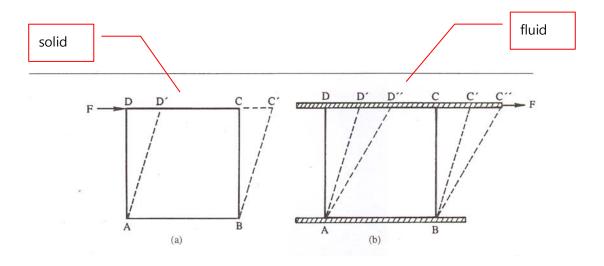
Chapter 1 Fluid Characteristics

1.1 Introduction



1.1.2 Fluidity

Fluid	 Solid deform by an amount proportional to the stress applied stress			
 deform continuously under shearing (tangential) stresses no matter how small the stress stress ∝ time rate of angular deformation (strain, displacement) 				
Newtonian fluid	Non-Newtonian fluid			
• shear stress is <u>linearly proportional</u> to rate of angular deformation starting with zero stress and zero deformation	• variable (<u>nonlinear</u>) proportionality between stress and deformation rate			
 constant of proportionality ≡ µ, dynamic viscosity → Fig. 1.1 water, air 	• proportionality = f (length of time of exposure to stress, magnitude of stress)			
[Cf] Analogy between Newtonian fluid and solids obeying Hooke's law of constant modulus of elasticity	plastics: paint, jelly, polymer solutions→ Rheology			



Elastic Solid – perfect memory



Fluid – zero memory

1.1.3 Compressibility

- 1) compressible fluid: gases, vapors → thermodynamics
- 2) incompressible fluid: liquid (small compressibility), water

1.1.4 Continuum approach

- dimensions in fluid space are large compared to the molecular spacing to ignore discrete molecular structure
- neglect void
- Consider a small volume of fluid ΔV containing a large number of molecules, and let Δm and v be the mass and velocity of any individual molecule

$$\rho = \lim_{\Delta V \to \varepsilon} \frac{\sum \Delta m}{\Delta V}$$

$$\vec{u} = \lim_{\Delta V \to \varepsilon} \frac{\sum v \Delta m}{\sum \Delta m}$$

 \mathcal{E} = volume which is sufficiently small compared with the smallest significant length scale in the flow field but is sufficiently large that it contains a large number of molecules

[Cf] Molecular approach

- molecular point of view
- well developed for light gases

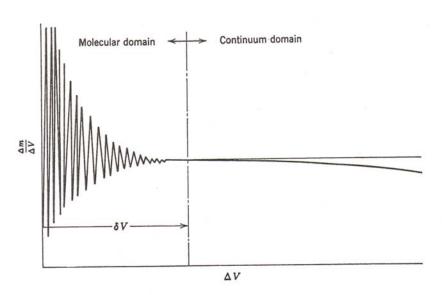


Figure 1.1 Density at a point.

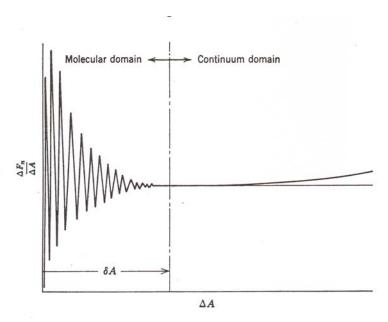
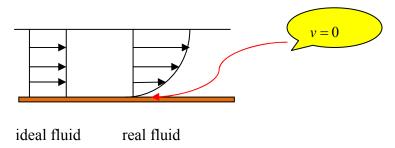


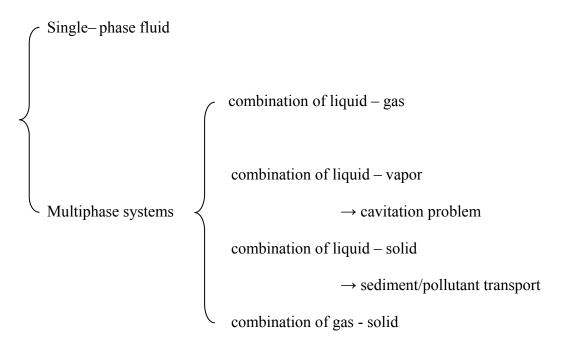
Figure 1.3 Normal stress at a point.

1.1.5 No-slip condition at rigid boundary

- 1) behavior of continuum type viscous fluids
- 2) zero relative velocity at the boundary surface (proven by experiments)



1.1.6 Multiphase system



1.2 Units of Measurement

- SI system: metric system

– English system: ft-lb system

* Newton's 2nd law of motion

F = ma

$$F = \text{force(N)}$$
; $m = \text{mass(kg)}$; $a = \text{acceleration(m/sec}^2)$

$$F \rightarrow 1 \text{kg} \cdot \text{m/sec}^2 = 1 \text{ N}$$

$$W = mg$$

W = weight; g = gravitational acceleration

1.3 Properties and States of Fluids

- 1) extensive properties ~ depend on amount of substance
 - → total volume, total energy, total weight
- 2) intensive properties ~ independent of the amount present
 - \rightarrow volume per unit mass, energy per unit mass weight per unit volume (specific weight, γ) pressure, viscosity, surface, tension
- 1.3.1 Properties of importances in fluid dynamics
 - (1) Pressure, $p \sim \text{scalar}$

$$p = F / A (N/m^2)$$

$$p_{\rm gauge} = p_{\rm absolute} - p_{\it atm}$$

Forces on a fluid element

Body force: act without physical contact

Surface force: require physical contact for transmission

1) body force

gravity force

- 2) surface forces
- normal stress

tensile stress (unusual for fluid)

 $\begin{array}{ccc} & & & & \\ & & & \\ tangential\ stress & \rightarrow & shear\ stress \end{array}$

(2) Temperature, T

two bodies in thermal equilibrium → same temperature

(3) Density, ρ

$$\rho = \text{mass / volume} = \frac{M}{V}$$



volume \propto (pressure, temperature)

(4) Specific weight, γ

 γ = weight / volume

[Re] Flow of a continuous medium

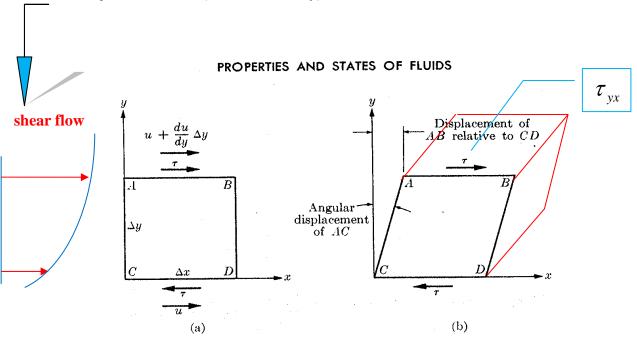
- ~ Fluids are treated as homogeneous materials.
- ~ Molecular effects are disregarded.

mass density
$$\rho(x, y, z, t) = \lim_{\Delta V \to 0} \frac{\Delta M}{\Delta V}$$

velocity vector
$$v = \lim_{\Delta t \to 0} \frac{\Delta s}{\Delta t}$$

- (5) Viscosity, μ
 - ~ due to molecular mobility
 - ~ whenever a fluid moves such that a <u>relative motion</u> exists between

adjacent volumes (different velocity)



Stress, $\tau \propto$ time rate of angular deformation

i) displacement of AB relative to CD in Δt

$$\left(u + \frac{du}{dy}\Delta y\right)\Delta t - u\Delta t = \frac{du}{dy}\Delta y \Delta t$$

ii) strain = relative displacement = angular displacement

$$\left[\frac{du}{dy}\Delta y\Delta t\right]/\Delta y = \frac{du}{dy}\Delta t$$

iii) time rate of strain (= time rate of angular displacement of AC)

$$\frac{du}{dy}\Delta t / \Delta t = \frac{du}{dy}$$

$$\tau \propto \frac{du}{dy}$$

$$\tau_{yx} = \mu \frac{du}{dy}$$

where

 τ_{yx} = shear stress acting in the x - direction on a plane

whose normal is y-direction (N/m^2)

$$\frac{du}{dy}$$
 = rate of angular deformation (1 / sec)

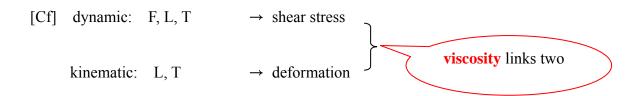
 μ = dynamic molecular viscosity

$$\mu = \frac{\tau}{\frac{du}{dy}} = \frac{N/m^2}{\frac{m/s}{m}} = N \cdot s/m^2$$

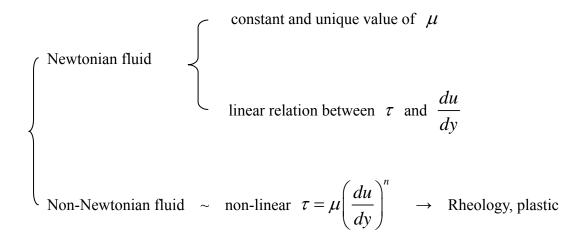
=
$$(kg \cdot m / s^2) \cdot \frac{s}{m^2} = kg / m \cdot sec = kg/m \cdot s$$

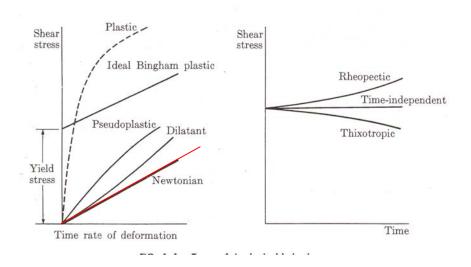
 \spadesuit Kinematic viscosity, ν

$$v = \frac{\mu}{\rho} = \frac{\text{kg/m} \cdot \text{s}}{\text{kg/m}^3} = \text{m}^2/\text{s}$$
 \rightarrow kinematic dimensions \rightarrow Fig. 1.4



Types of Fluid





[Cf] Stress-strain relationship for solid

$$\tau_{yx} = G \frac{d\xi}{dy}$$

 $d\xi$ = relative station displacement of AB

$$\frac{d\xi}{dy}$$
 = angular deformation (shear strain)

G =modulus of elasticity in torsion

 $\frac{du}{dy} \qquad \frac{d\xi}{dy}$ velocity

displacement

 \spadesuit μ = function of (temperature, pressure)

	Liquid	Gas		
major factor	intermolecular	exchange of		
for viscosity	cohesion	momentum		
	decrease			
when temperature		increase molecular activity		
	cohesive force			
is increasing		→ increase shear stress		
	→ decrease viscosity			

[Re] Exchange of momentum

fast-speed layer (FSL)



molecules from FSL speed up molecules in LSL $\,$

molecules from LSL slow down molecules in FSL

Two layers tend to stick together as if there is some viscosity between two.

low-speed layer (LSL)

Water:

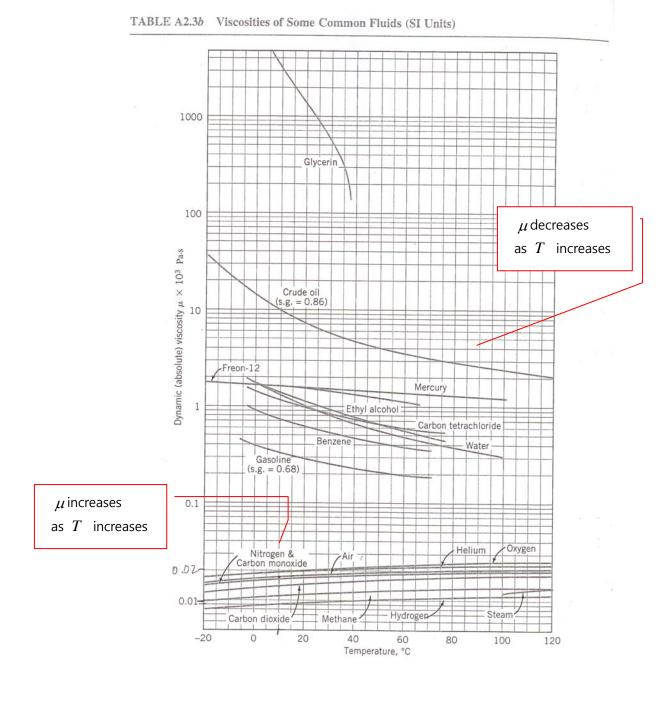
$$\mu = 1.0 \times 10^{-3} \, \frac{N}{m^2} \, s$$

TABLE A2.1 Approximate Properties of Some Common Liquids at Standard Atmospheric

sure (cont.)

-				SI Units			
	<i>T,</i> °C	$_{ m kg/m^3}^{ ho,}$	s.g.,	E, kPa	$\begin{array}{c} \mu \times 10^4 \\ \mathrm{Pa} \cdot \mathrm{s} \end{array}$	σ, N/m	p _v , kPa
Ethyl alcohol	20	788.6	0.79	1, 206 625	12.0	0.022	5.86
Freon-12	15.6	1 345.2	1.35	_	14.8	_	
	-34.4	1 499.8	-	_	18.3	-	_
Gasoline	20	680.3	0.68	-	2.9	-	55.2
Glycerin	20	1 257.6	1.26	4 343 850	14 939	0.063	0.000 014
Hydrogen	-257.2	73.7	_	_	0.21	0.0029	21.4
Jet fuel (JP-4)	15.6	773.1	0.77	_	8.7	0.029	8.96
Mercury	15.6	13 555	13.57	26 201 000	15.6	0.51	0.000 17
Oxygen (Liquid)	315.6 - 195.6	12 833 1 206.0 V	12.8	=	9.0 2.78	0.015	47.2 21.4
Sodium	315.6	876.2		_	3.30	_	- 0
	537.8	824.6	_	_	2.26	_	_166
Water ^b	20	998.2	1.00	2,170,500	10.0	0.073	2.34
Sea water ^b	20	1024.0	1.03	2,300,000	10.7	0.073	2.34

 $[^]b The$ specific heat of liquid water is approximately 25 000 ft·lb/slug·°R or 4 180 J/kg·K.



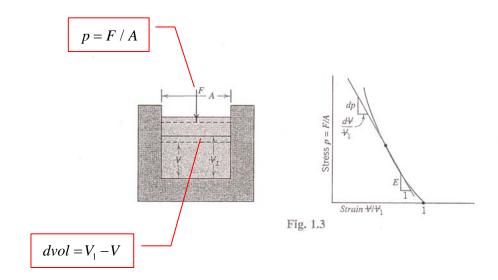
1–13

- (6) Specific heat, c
 - = ratio of the quantity of heat flowing into a substance per unit mass to the change in temperature
- (7) Internal energy, u specific internal energy = energy per unit mass, J/kg kinetic + potential energy \rightarrow internal energy
- (8) Enthalpy specific enthalpy $= u + p / \rho$
- (9) Bulk modulus of elasticity and Compressibility
 - 1) Compressibility, C
 - = measure of change of volume and density when a substance is subjected to normal pressures or tensions
 - = % change in volume (or density) for a given pressure change

$$C = -\frac{dvol}{vol} / dp = \frac{d\rho}{\rho} \frac{1}{dp}$$

2) Bulk modulus of elasticity, E_{v}

$$E_{v} = \frac{1}{C} = -\frac{dp}{dvol/vol} = \frac{dp}{d\rho/\rho}$$



(10) Vapor pressure, p_v

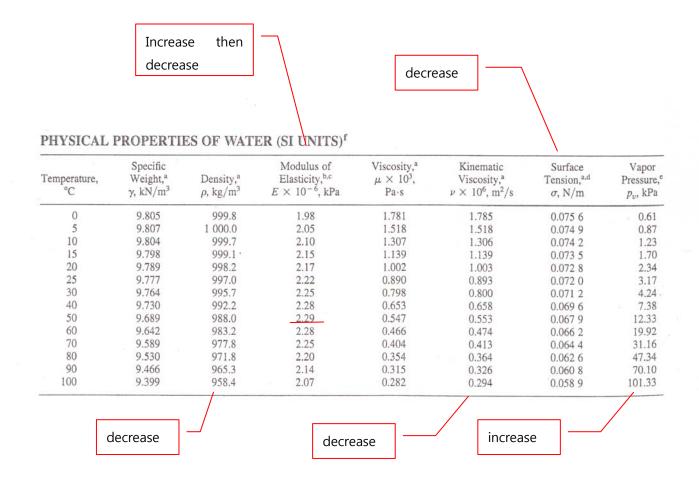
- = pressure at which liquids boil
- = equilibrium partial pressure which escaping liquid molecules will exert above any free surface
- ~ increases with temperature
- ~ The more volatile the liquid, the higher its vapor pressure.

(11) Surface energy and surface tension, σ

At boundaries between gas and liquid phase, molecular attraction introduce forces which cause the interface to behave like a membrane under tension.

$$\sigma = \frac{\text{(force)} \times \text{(distance)}}{\text{area}} = \frac{\text{work}}{\text{area}} = \frac{\text{force}}{\text{length}}$$

~ water: decrease with temperature



[Appendix 1] Coordinate Systems

- i) Cartesian (x, y, z)
- ii) Cylindrical (R, θ, z)

$$x = R\cos\theta$$

$$y = R \sin \theta$$

$$z = z$$

iii) Spherical (r, θ, ω)

$$x = r \sin \theta \cos \omega$$

$$y = r \sin \theta \sin \omega$$

$$z = r \cos \theta$$

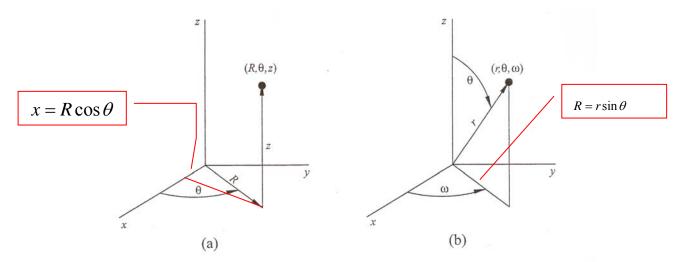


FIGURE A.1 Relationship between cartesian coordinates and (a) cylindrical coordinates and (b) spherical coordinates.

[Appendix 2] Tensor

Scalar – quantity with magnitude only

Vector – quantity with magnitude and direction

Tensor – an order array of entities which is invariant under coordinate transformation, this includes scalars and vectors

• Rank (order) of tensors

0th order – 1 component, scalar (e.g., mass, length, pressure)

1st order - 3 components, vector (e.g., velocity, force, acceleration)

2nd order - 9 components, (e.g., stress, rate of strain, turbulent diffusion coeff.)

• Example of 2nd order tensor

~ stress acting on a fluid element

Stress tensor =
$$\begin{bmatrix} \sigma_{xx} & \tau_{xy} & \tau_{xz} \\ \tau_{yx} & \sigma_{yy} & \tau_{yz} \\ \tau_{zx} & \tau_{zy} & \sigma_{zz} \end{bmatrix}$$

 σ = normal stress, τ = shear stress



 τ_{yx} = shear stress in xz - plane

and in x - direction

