Water in Subsurface Environment

Groundwater Engineering | CE60205

Lecture:02

Learning Objective(s)

To estimate the physical properties of water, air, and porous media

Properties of Water

- Density
 - The mass density of fresh water ρ_w varies within a narrow range
 - Weight density $\rho_w g$ equals the mass density times the gravitational acceleration at the earth's surface, g
- Water density does vary slightly with temperature, pressure, and chemistry if the concentration of solute molecules is high enough.
- The density of pure water at atmospheric pressure varies between 0.998 and 1.000 g/cm³ in the range of temperatures typical for groundwater (o°C to 20°C)
- As the temperature of a liquid rises, it usually becomes less dense as molecules move with greater velocity and molecular attraction forces are overcome to a greater extent.
- Water is an unusual liquid because the maximum density does not occur at the freezing temperature, but instead slightly above freezing at 4°C.

Property	Symbol	Dimensions †	Value
Mass density	$ ho_w$	[M/L ³]	I.00 g/cm ³ I000 kg/m ³ I.94 slugs/ft ³
Weight density	$ ho_w g$	[F/L ³]	9810 N/m³ 62.4 lb/ft³
Compressibility	β	[L ² /F]	$4.5 \times 10^{-10} \text{ m}^2/\text{N}$
Dynamic viscosity	μ	[FT/L ²]	$1.4 \times 10^{-3} \text{ N}\cdot\text{sec/m}^2$

 $^{^{\}dagger}L$ = length, M = mass, T = time, F = ML/T² = force.

- Compressibility
 - Water is often considered incompressible, but it does have a finite, low compressibility.
 - As water pressure P rises an amount dP at a constant temperature, the density of water increases $d\rho_w$ from its original density ρ_w , and a given volume of water V_w will decrease in volume by dV_w in accordance with $BdP = \frac{d\rho_w}{d\rho_w}$

$$\begin{aligned}
\rho_w &= \overline{\rho_w} \\
&= -\frac{dV_w}{V_w}
\end{aligned}$$

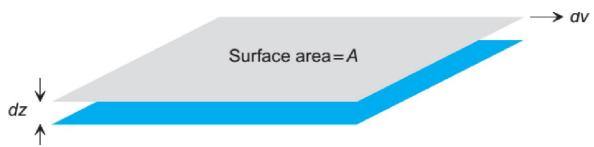
- where β is the isothermal compressibility of water.

Property	Symbol	Dimensions [†]	Value
Mass density	$ ho_w$	[M/L ³]	1.00 g/cm ³ 1000 kg/m ³ 1.94 slugs/ft ³
Weight density	$ ho_w g$	[F/L ³]	9810 N/m³ 62.4 lb/ft³
Compressibility	β	[L ² /F]	$4.5 \times 10^{-10} \text{ m}^2/\text{N}$
Dynamic viscosity	μ	[FT/L ²]	$1.4 \times 10^{-3} \text{ N}\cdot\text{sec/m}^2$

 $^{^{\}dagger}$ L = length, M = mass, T = time, F = ML/T² = force.

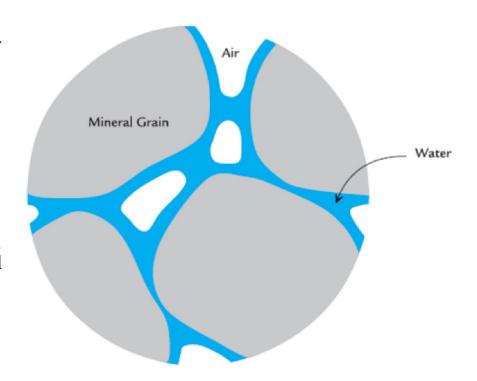
- Viscosity
 - Viscosity is friction within a fluid that results from the strength of molecule-to-molecule attractions.
 - The resisting force F

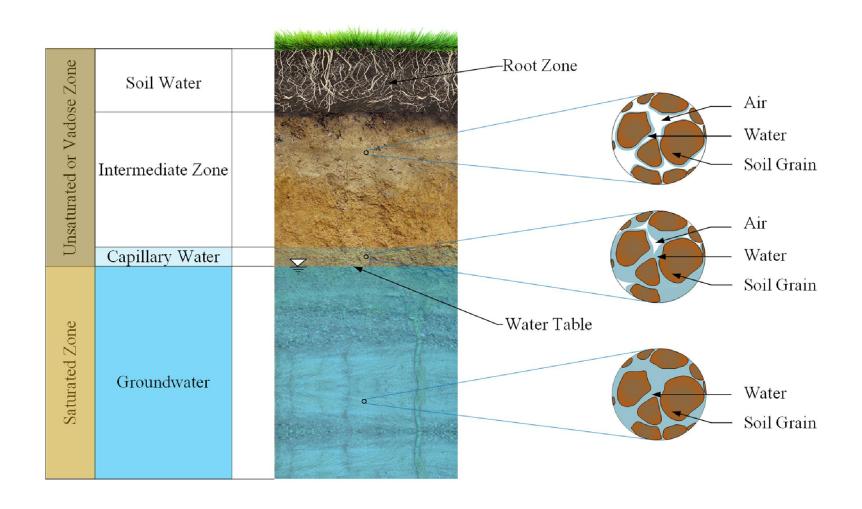
$$F = A\mu \frac{dv}{dz}$$



- This resistance to internal shear causes water to resist flow through geologic materials.
- In order to flow through pores or fractures, a packet of water must change shape and shear as it flows.
- pore size is analogous to dz
- water encounters greater viscous resistance flowing through materials with smaller pores
- The viscosity of a liquid generally decreases with increasing temperature
- The dynamic viscosity of water ranges from μ = 1.79×10⁻³ N sec/m² at 0°C to μ =1.01×10⁻³ N sec/m² at 20°C.
- Kinematic viscosity $\nu = \frac{\mu}{\mu}$

- Surface Tension and Capillarity
 - polar water molecules are attracted to each other -> a mass of water has internal cohesion that tends to hold it together
 - water drops tend to form spheres as they fall through the air
 - In pore spaces containing both air and water, the water will generally wet the mineral surfaces, leaving the central parts of the pores for the air
 - A layer of water molecules on the order of 0.1 to 0.5 μ m (10⁻⁶ m) thick is so strongly attracted to mineral surfaces that it is essentially immobile
 - the forces of attraction are not strong enough to prevent movement of water molecules and water outside that distance is free to move
 - Surface attraction forces are stronger for clay minerals -> due to the charged nature of clay mineral surfaces.





- The attraction of water to mineral surfaces causes water to pull and spread itself across the surfaces -> the pressure within the water is less than the air pressure within the pores
- As amount of water present decreases -> the pull of the mineral surface attraction forces increases -> the pressure within the water decreases -> the air—water interface develops a more contorted shape conforming to the mineral grains
- This attraction of water to mineral surfaces in partly saturated materials is called capillarity
- Capillarity allows water to wet pore spaces above the water table
- Capillary forces tend to be greater in finer-grained granular materials, due to a greater amount of mineral surface area.

Table 2.4.1 Capillary Rise in Samples of Unconsolidated Materials (after Lohman³⁴)

Material	Grain size (mm)	Capillary rise (cm)	
Fine gravel	5–2	2.5	
Very coarse sand	21	6.5	
Coarse sand	1-0.5	13.5	
Medium sand	0.5-0.2	24.6	
Fine sand	0.20.1	42.8	
Silt	0.1-0.05	105.5	
Silt	lt 0.05-0.02		

Note: Capillary rise measured after 72 days; all samples have virtually the same porosity of 41 percent.

^aStill rising after 72 days.

Properties of Air

- Air occupies some amount of the subsurface pore spaces
- The pressure in the atmosphere at the earth's surface varies with the weather and with elevation
- At sea level, atmospheric pressure averages about 1.013 ×10⁵ N/m²
- Groundwater pressures are often measured as gage pressure -> the amount of pressure in excess of atmospheric pressure
- The density of the atmosphere also varies with weather and altitude.
- \bullet At the earth's surface, the atmospheric density averages about 1.2 kg/m 3

Properties of Porous Media

• Soil Texture



Grade-I



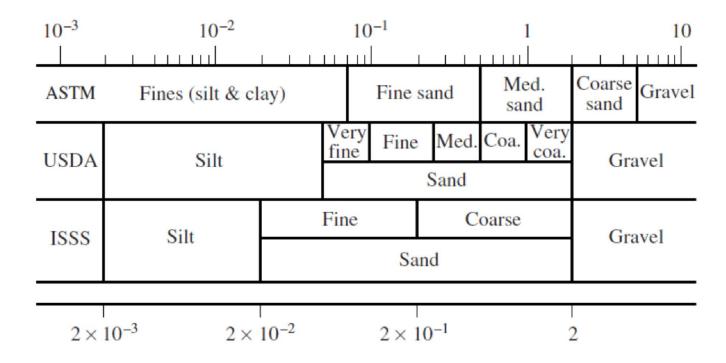
Grade-II



Grade-III

Indian Standard Sand

Soil classification

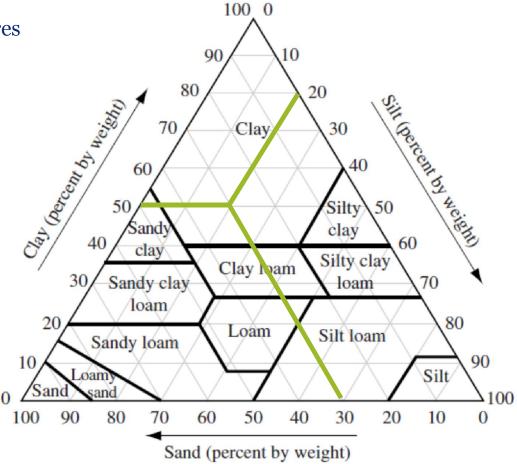


• Triangle of Soil Textures

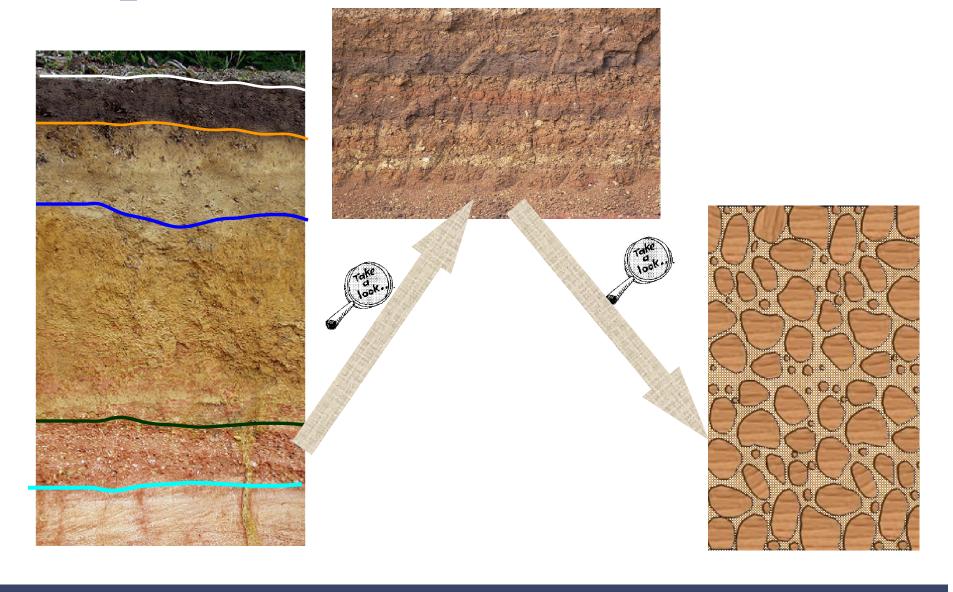
- Clay: 50%

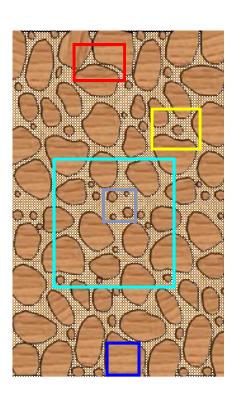
- **Sand:** 30%

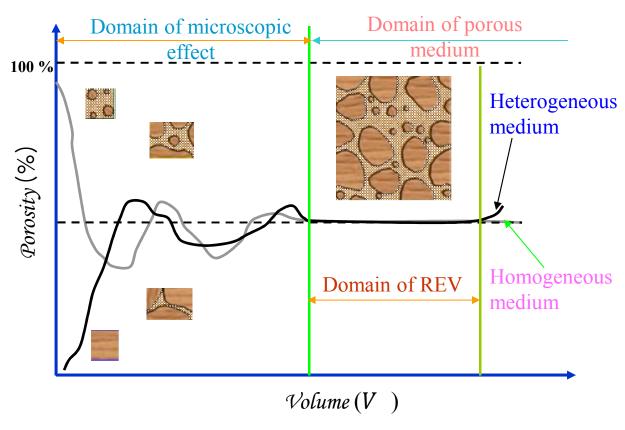
- **Silt:** 20%



USDA soil textural triangle.

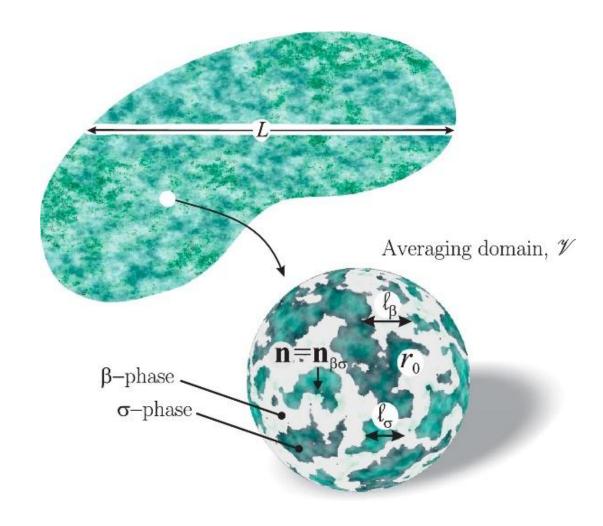




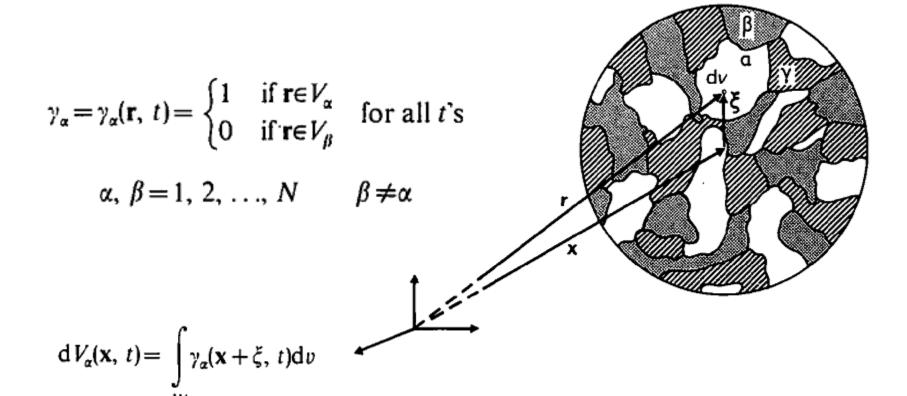


A sub-volume of a porous medium that has the "same" geometric configuration as the medium at a macroscopic scale.

 $\bullet l \ll D \ll L$



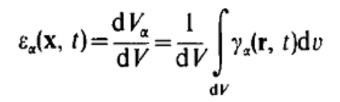
$$\mathbf{r} = \mathbf{x} + \boldsymbol{\xi}$$



$$\int_{\mathrm{d}V_{s}} f \, \mathrm{d}v = \int_{\mathrm{d}V} f \gamma_{\alpha} \, \mathrm{d}v$$

$$dA_{\alpha}(\mathbf{x}, t) = \int_{\mathbf{d}A} \gamma_{\alpha}(\mathbf{x}, \xi, t) da$$

Volume fraction



$$\sum_{\alpha} \varepsilon_{\alpha} = 1 \qquad \text{and} \qquad$$

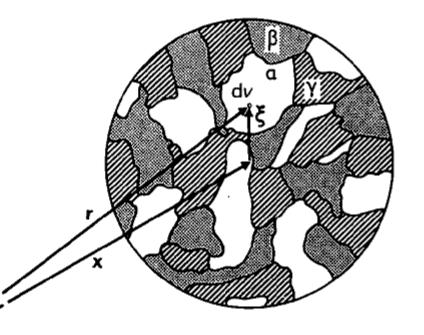
and
$$0 \le \varepsilon_x \le 1$$

Areal fraction

$$\bar{\varepsilon}_{\alpha}(\mathbf{x}, t) = \frac{\mathrm{d}A_{\alpha}}{\mathrm{d}A} = \frac{1}{\mathrm{d}A} \int_{\mathrm{d}A} \gamma_{\alpha}(\mathbf{r}, t) \mathrm{d}a$$

$$\Sigma \bar{\varepsilon}_{\alpha} = 1$$
 and $0 \leq \bar{\varepsilon}_{\alpha} \leq 1$

• In general ε_{α} and $\bar{\varepsilon}_{\alpha}$ need not be equal



• Volume average operator $\langle \ \rangle_{\alpha}$

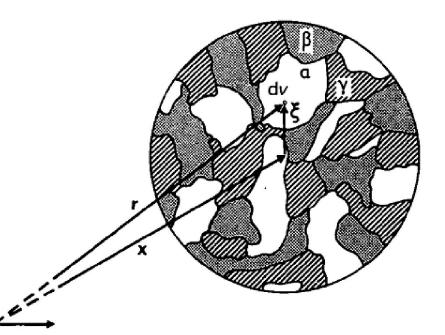
$$\langle f \rangle_{\alpha}(\mathbf{x}, t) = \frac{1}{\mathrm{d}V} \int_{\mathrm{d}V} f(\mathbf{r}, t) \gamma_{\alpha}(\mathbf{r}, t) \mathrm{d}v$$

$$\langle \rho \rangle_{\alpha} = \frac{1}{\mathrm{d}V} \int_{\mathrm{d}V} \rho \gamma_{\alpha} \mathrm{d}v$$

• Intrinsic volume average operator (

$$\langle f \rangle_{\alpha}^{\alpha}(\mathbf{x}, t) = \frac{1}{\mathrm{d}V_{\alpha}(\mathbf{x}, t)} \int_{\mathrm{d}V} f(\mathbf{r}, t) \gamma_{\alpha}(\mathbf{r}, t) \mathrm{d}v$$

$$\langle \rho \rangle_{\alpha}^{\alpha} = \frac{1}{\mathrm{d} V_{\alpha}} \int_{\mathrm{d} V} \rho \gamma_{a} \mathrm{d}v = \frac{1}{\varepsilon_{\alpha}} \langle \rho \rangle_{\alpha}$$



$$\langle f \rangle_{\alpha} = \varepsilon_{\alpha} \langle f \rangle_{\alpha}^{\alpha}$$

• Only when the mass density of the α -phase, is microscopically constant will the intrinsic volume average mass density function be equal to the microscopic mass density, or

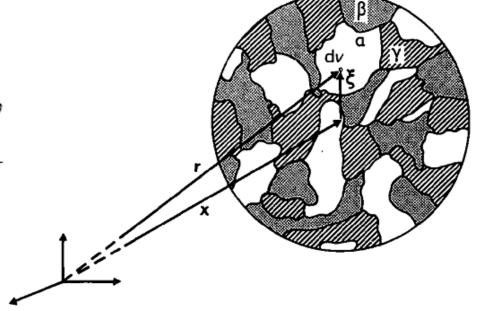
$$\langle \rho \rangle_{\alpha}^{\alpha} = \rho_{\alpha}$$

• Mass average operator $^{-\alpha}$

$$\int \rho(\mathbf{r}, t) f(\mathbf{r}, t) \gamma_{\alpha}(\mathbf{r}, t) dv$$

$$\int \rho(\mathbf{r}, t) \gamma_{\alpha}(\mathbf{r}, t) dv$$

$$\int \rho(\mathbf{r}, t) \gamma_{\alpha}(\mathbf{r}, t) dv$$



• Simplified form

$$\bar{f}^{\alpha}(\mathbf{x}, t) = \frac{1}{\langle \rho \rangle_{\alpha}(\mathbf{x}, t) dV} \int_{dV} \rho(\mathbf{r}, t) f(\mathbf{r}, t) \gamma_{\alpha}(\mathbf{r}, t) dv$$

- The porosity of a rock or soil is simply the fraction of the material volume that is pore space.
- In quantitative terms the porosity η is defined as

$$n = \frac{V_{\nu}}{V_{t}}$$

where V_{ν} is the volume of voids in a total volume of material V

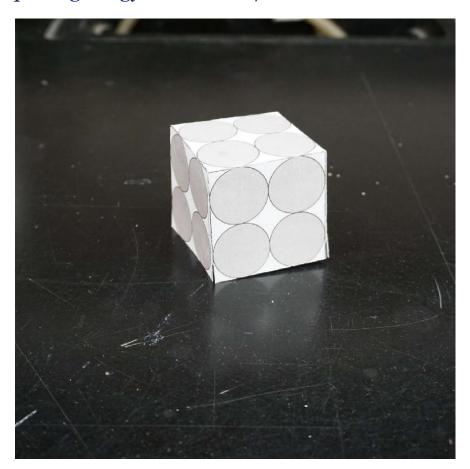
- The porosity is a dimensionless parameter in the range $0 < \eta < 1$.
- Geotechnical engineers often use a related dimensionless parameter called the void ratio *e*, which is defined as

$$e = \frac{V_{\nu}}{V_{s}}$$

where V_s is the volume of mineral solids in a given volume of material.

$$n = \frac{e}{1+e}, \quad e = \frac{n}{1-n}$$

- Home Lab
- Foldable Aquifer Project -http://aquifer.geology.buffalo.edu/
- Paper aquifer model
 - Porosity and Grain Packing



Learning Strategy

Chapter 2: Physical Properties

Section 2.1, 2.2, 2.3, 2.4



Thank you