

1. Laboratory Test

1.1 Atterberg Limits (Liquid Limit and Plastic Limit)

- ✱ Classification of soils (cohesive soils) based on behavioral difference by their water contents

- Water content
increase
- ↓
- Cohesion limit : Generally not in use
 - Sticky limit : Generally not in use
 - Shrinkage limit: Moisture content below, which no further soil volume reduction occurs.
 - Plastic limit (PL): Lower limit for plastic behavior of soils.
 - Liquid limit (LL): Upper limit for plastic behavior of soils.
- Plasticity Index (PI): Range of water content for plastic behavior of soils.

$$PI = LL - PL$$

- ✱ Purpose and usage

- Soil identification and classification (Simple method without difficulty in finding out mineral compositions)
- Quick referencing and most commonly used method for soil characteristics
 - 1) Settlement or volume change :
 - ex) $C_c = 0.009(LL-10)$ for undisturbed clays of low to medium sensitivity.
(Terzaghi and Peck)
 - 2) Undrained strength : $s_u/\sigma_v' = 0.1 + 0.0037PI$ for NC clay deposits
(Skempton)
 - 3) Friction angle : empirical correlations between PI and ϕ
(increase of PI \rightarrow decrease of ϕ')
 - 4) $K_o = 0.44 + 0.42(PI/100)$ (Massarsch, 1979)

✿ Testing methods for liquid limit

1) Conventional Method

- Determine water content at which a pat of soil in a brass cup, cut with a standard groove, and then dropped from a height of 10mm will undergo a groove closure of 12.7mm when the cup of soil is dropped 25times(=N) at the rate of 120 drops/min. (Need at least 4-5 reasonably well spaced points from about N=15 to 35.)
- Several variables
 - 1) Size of soil pat
 - 2) Rate of blows
 - 3) Length of time that soil is in cup
 - 4) Laboratory humidity and speed of performing test
 - 5) Degree of care for soil mixing
 - 6) Type of material for LL device base.
 - 7) Accuracy of height of fall
 - 8) Type of grooving tool.
 - 9) Condition of liquid limit device
- Use the soil fraction passing the No. 40 sieve (0.425mm opening)
 - For air dry soil (pulverizing and sieving , need curing time of 24-48 hours after remixing with water.
 - Minimal-to-no drying soil (wet sieving), can induce substantial problem of segregation and require a very careful mixing and long mixing time.

- One-Point Liquid Limit Method

Based on regression analysis of 767 soils,

$$\begin{aligned}LL &= w_n(N/25)^{\tan\beta} \\ &= w_n(N/25)^{0.121}\end{aligned}$$

Error is negligible if N is between 20 and 30.

2) Fall cone

- Alternative method for determining the liquid limit and maybe more reproducible.
- Liquid limit is defined as that water content at which 20mm of cone penetration with a 30° apex angle and a total mass of 80g into soil in the 55mm diameter and 40mm depth occurs in the 5s test time.
- LL is generally defined in a semilog plot made of penetration depth and water content.

⊗ Testing Method for Plastic Limit

- PL has been defined as that water content at which a soil thread just crumbles when it is rolled down to a diameter of 3mm.
- Somewhat more operator-dependent but can be reproduced within 1-3%.

1.2 Specific Gravity of Soil Solids

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* Specific gravity can be computed using any ratio of mass of substance to mass of water for equal volume.

$$G_s = \frac{M_s / V}{M_w / V}$$

- With any given mass of soil (M_s), measure the mass of flask and water (M_{bw}) and the mass of flask, water and soil (M_{bws}) with measurement of temperature.
- Mass of water with same volume of soil (M_w) can be given as

$$M_w = M_{bw} + M_s - M_{bws}$$

- Specific Gravity can be obtained as

$$G_s = M_s / (M_{bw} + M_s - M_{bws})$$

- Correction for temperature

$$G_s = \alpha M_s / (M_{bw} + M_s - M_{bws})$$

(α is given in Table in page 73, based on 20°C)

* Major point of testing approach

- Deaerating the soil water mixture and water itself in flask.
 - ⇒ Accomplished by applying a vacuum and/or heating.
- Checking the efficiency of air removal
- Any temperature error (deviation of volume of flask and density variation) and error due to the use of tap water is negligible, if 18°C ~22°C and no visible mineral in a flask.

1.3 Particle-Size Analysis

- Mechanical Sieving Method
- Hydrometer Test

1.3.1 Mechanical Method

- Particle size distribution is the most elementary and indispensable measurement of soils.
- It is one of suitability criteria of soils for any kind artificial application such as road, backfill of retaining structure, airfield, dam....
- Stacking a series of sieves (the size bracketing and not determination of individual particle sizes) does the test. (No reproducible analysis => a deviation of $\pm 2.0\%$ for (-)No.4)
- Standard sieve sizes (Table 5-1).

⇒ The No.200 sieve (0.075mm) is the smallest practical sieve size.

(Hydrometer method is used to estimate particle size < the No.200 sieve size.)

1) Key Point of Testing Method

boundary between gravels and sands

boundary between sands and silts

- Typical set of sieves: 4 – 7 sieves including No.4, 10, 40 and 200.
- Most important factors: **How to obtain a representative sample and how to break down soil lumps into individual soil particles.**
- Soaking (2 to 24hrs) and washing. (Pulverizing process is not enough to

break down soils to individual soil particles especially in fine particles.)

⇒ Can be omitted for soils with less than 5% of (-) 200 material.

- The result is presented as a semilog plot of percent finer versus particle size.

- Oven-dry soil or air-dry soil?

⇒ For air dry, must determine water content to compute dry sample mass.

⇒ Seem to be no significant difference.

- How to determine adequate quantity of soils.

⇒ The more, the better result. But it depends on the maximum grain

size.	⇒	D_{\max}	$M_{\min}(\text{g})$
		No. 10	200
		No. 4	500
		3/4 in.	1500

⇒

2) Problems and Limitations

- Obtaining a statistically representative sample.
- The presence of soil lumps.
- The practical limitations of using square sieve mesh openings for irregularly soil particles.
- The limit on the number of sieves used in a stack for the analysis.

1.3.2 Hydrometer Method

1) Principle and Key Points of Hydrometer Method

- The larger the diameter of particle, the higher the velocity of fall of particles.

➔ Stokes' law

Velocity of fall of the spheres,
$$v = \frac{2}{9} \frac{G_s - G_f}{\eta} \left(\frac{D}{2}\right)^2$$

The range of soil particles in which the equation is valid,

⇒ (Brown movements) $0.0002\text{mm} \leq D \leq 0.2\text{mm}$ (Turbulence of fluid)

- Specific gravity of soil-water suspension is changed (decreased) with time, because soil particles continue to fall down and be sedimented at bottom.
- A dispersing agent is used to neutralize the particles to prevent soils from adhering together. ⇒ Sodium metaphosphate (NaPO_3) and Sodium Silicate (A 125mL quantity of 4% solution of Sodium metaphosphate in the 1000mL of soil water suspension.)

2) Hydrometer

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- The hydrometer displays the specific gravity of the soil-water suspension

at the center of the bulb. (➔percent finer)

- The velocity of fall of the particles (from time reading +L)

$$v = L/t \quad (\text{➔particle size})$$

- Generally used type hydrometer: H152 (Calibrated to read 0 to 60grams of soils in the 1000mL soil water mixtures. ($G_s=2.65$))

3) Corrections and Calibrations

- The immersion correction :

Correction of L (=the distance particles fall) for rising the surface of suspension due to inserting the hydrometer.

$$L = L_1 + \frac{1}{2}L_2 - \frac{1}{2} \frac{V_b}{A} \quad \leftarrow \text{Figure 6-2}$$

It depends on shape of hydrometer. For 152H, $L=16.3 - .1641R$ (R =reading of hydrometer) or Table 6-7

- Meniscus correction ← Figure 6-2
- Temperature correction and corrections for water impurities and for the use of a dispersal agent :

$$R_c = R_{actual} - zero \text{ correction} + C_T$$

where C_T is presented in Table 6.3.

- Correction for specific gravity of soils

$$a = \frac{1.65G_s}{2.65(G_s - 1)}$$

$$\text{Percent finer} = \frac{aR_c}{M_s} 100\%$$

4) How to obtain the relation between % finer and particle size.

1. Taking hydrometer reading (R_{actual}) and temperature at various elapsed times (t).
2. Making corrections and calibrations as necessary.
3. Computing particle size with elapsed time and percent finer with hydrometer reading.

$$D = \sqrt{\frac{30\eta}{980(G_s - G_w)} \frac{L}{t}} = K \sqrt{\frac{L}{t}}$$

(L in cm, t in min and D in mm)

where K (Table 6-4) can be computed with temperature and G_s .

$$\text{Percent finer} = \frac{aR_c}{M_s} 100\%$$