Historic, Archive Document

Do not assume content reflects current scientific knowledge, policies, or practices.



ASIC SOIL MECHANICS

Reservo

 \bigcirc

R

K

B

0



UNITED STATES DEPARTMENT OF AGRICULTURE

USDA-SCS-PORTLAND, OREG. 1875

USDA, National Agricultural Library NAL Bldg 10301 Baltimore Blvd Beltsville, MD 20705-2351



SOIL MECHANICS

BY J. C. Stevenson and J. R. Talbot Engineering and Watershed Planning Unit Technical Service Center SOIL CONSERVATION SERVICE Portland, Oregon

U. S. DEFT OF MANOULTURT

A Work Book Prepared for Soil Mechanics Training

> February 1974 Revised October 1975

ENTALOGING - PHEI

MAY 271977

This manual contains basic soil mechanics information dealing mainly with engineering classification systems and volume-weight relationships. It was assembled specifically for training purposes and for future reference by trainees. It is intended for use as a basic text for the following courses conducted by the West Employee Development Unit.

- Classification, Identification and Engineering Behavior Characteristics of Soil
- Construction Inspection (sampling and testing portion)
- New Professional Engineers and Geologists Training (NPEG) (Soil Mechanics Portion).
- Qualitative Soil Mechanics and Site Evaluation.

An attempt has been made to take the best procedures from several sources and provide clear, concise information for classifying soils in the engineering systems and making basic soil mechanics calculations. The information has been updated to present accepted criteria and procedures adopted by the Service.

605244

Table of Contents

Page

SECTION 1 - INTRODUCTION

General	1-1
Types of Problems Encountered in Design	1-2
Information Required as Basis for Design	1-3
Use of Soil Mechanics in Construction	1-4
Nomenclature, Definitions and Conventions	1-5

SECTION 2 - SOIL CLASSIFICATION AND IDENTIFICATION

•

Engineering Soil Classification 2	-1
Unified Soil Classification System 2	-2
Soil Properties Used in Classification 2	-2
Definition of Soil Components 2	-3
The Plasticity Chart 2	-3
Summary of the Unified Classification System 2	-7
Method of Classification 2	-10
Field Identification of Soils 2	-22
Description of Undisturbed Soil 2	-27
AASHTO Soil Classification System 2	-30
Soil Properties Used in Classification 2	-30
Definition of Soil Components 2	-30
Summary of the AASHTO Classification System 2	-30
Method of Classification (Laboratory Test Data) 2	-33

SECTION 3 - THE SOIL-AIR-WATER SYSTEM

Soil Mass as a Multiphase System	3-1
Solid Phase	3-1
Liquid Phase	3-1
Gas Phase	3-1
Weight Relationships	3-1
Volume Relationships	3-2
Volume and Weight Relationships	3-3
Consistency and Mechanical Properties	3-6
Gradation	3-6
Grain Size and Shape	3-8
Relative Density	3–9
Soil Consistency	3-11
Consistency of Undisturbed Soils	3-13

	· · · · · · · · · · · · · · · · · · ·
Physio-Chemical Relationships	3-15
Structure and Composition of Clay Minerals	3-16
Shapes of Clay Minerals	3–17
Kaolinite	3-18
Halloysite	3-20
Montmorillonite	3-20
Illite	3-22
Soil Structure	3-22

Page '

FIGURES

2-1	A Comparison of Grain-Size Limits in the 3	2-5
	Classification Systems	
2-2	Plasticity Chart	2-6
2-3	Unified Soil Classification Group Positions	
	on Plasticity Chart	2-8
	Classification Examples	· 2-12 - 2-21
2-4	Grain Shape	2-22
2-5	Shaking Test	2-24
2-6	Breaking Test	2-25
2-7	Plasticity Test	2-26
2-8	Group Index Chart	2-34
2-9	Liquid Limit and Plasticity Index Ranges for	
	the A-4, A-5, A-6, and A-7 Subgrade Groups	2-37
2-10	AASHTO Soil Classification Group Positions	
	on Plasticity Chart	2-38
3-1	Relationship Among Soil Phases	3-4
3-2	Weight-Volume Relationships	3–5
3-3	Sample Curves	3-7
3-4	Shrinkage Diagram for a Cohesive Soil	3-12
3-5	Silica Tetrahedral Unit	3-16
3-6	Octahedral Unit	3-17
3-7	Basic Kaolinite Structure	3-18
3-8	Basic Montmorillonite Structure	3–20
3-9	Cardhouse Structure of Clay	3-24
3-10	Five Stable Units of Soil Structure	3-26

TABLES

2-1	Laboratory Criteria for Unified Soil	
	Classification System	2-4
2-2	Unified Soil Classification	2-23
2-3	Classification of Soils and Soil-Aggregate	
	Mixtures (AASHTO)	2-36
3-1	Dimensions of Typical Clay Platelets	3–19

· ii

A. GENERAL

To successfully practice the art of soil engineering, an engineer must have a knowledge of precedents, familiarity with soil mechanics, and a working knowledge of geology.

Soil has been used as a foundation and construction material throughout the history of mankind. By the time the scientific method became recognized as a fruitful approach to the solution of engineering problems, large buildings, bridges, dams, canals, tunnels and roads had been built and many had served their purpose for centuries. From this background of precedents, earthwork and foundation engineering developed primarily as an art steeped in tradition and empirical practices based on earlier successful accomplishments. No man can hope to become an artist in the practice of soil engineering without a rich background of personal experience coupled with a thorough knowledge of the experiences of his contemporaries and predecessors.

The rapidly developing science of soil mechanics provides qualitative and quantitative data on the stress-strain-time characteristics of soils. This knowledge gives the designer a feeling for the behavior of soils, under idealized conditions, that form guides to soil behavior under the more complex conditions in the field. The theories of soil mechanics provide insight into the behavior under simple ideal conditions. These aspects of soil mechanics form a framework that enables the engineer to organize, interpret, and evaluate experience. Each experience encountered by an engineer can be examined and categorized as to the type of material, the loading conditions, the hydraulic conditions, and in relation to theoretical concepts.

The procedures used to calculate bearing capacity, settlement or stability factor of safety of a slope use the framework of soil mechanics to organize experience. In fact, the procedures are valid and justified only to the extent that they have been verified by experience.

The field of soil mechanics enables an engineer to go beyond the limits of his experience. It points the way to new solution of old problems, or to the solution of previously unsolved problems. In this way, it permits the engineer to extrapolate his experience. Of course, such extrapolation involves a measure of uncertainty until the pertinent experience becomes available. Here, soil mechanics forms the guide as to what should be observed to check the design procedures as the work is executed. Geology is as basic to earthwork design as soil mechanics. Possibly the most significant role of geology is to make the engineer aware of the departures from reality inherent in the simplifying assumptions used in soil mechanics procedures. The geology of a site must be understood to permit making and evaluating the design calculations or predictions. In some instances the geologic structure or the results of geologic processes may completely override all considerations of soil mechanics (jointed clays, etc.)

Every interpretation of the results of a test boring and every interpolation between two borings is an exercise in geology.

- B. TYPES OF PROBLEMS ENCOUNTERED IN DESIGN
 - Embankments. Most of the problems encountered in the design of earth or rock embankments may be grouped into four categories:
 - a. Determination of the bearing capacity of the foundation.
 - b. Determination of the amount of settlement of the various parts of the structure caused by compression of foundation or fill materials.
 - (1) Compression caused by loads imposed by the structure.
 - (2) Compression caused by alteration of the water table.
 - c. Determination of the amount of seepage or water movement through the fill or foundation and the effect of such seepage on the stability of the structure and economic feasibility of the project.
 - d. Determination of the stability of earth slopes.
 - Reservoir basins. The majority of the problems encountered in evaluating reservoir basins may be grouped into the following categories:
 - a. Determination of the tendency for the basin to leak.
 - b. Determination of flow boundaries.
 - c. Determination of head conditions, groundwater effects, etc.
 - d. Determination of sliding stability of basin walls.

· 1-2

- 3. Channel systems. The soil engineering problems encountered in the evaluation and design of channel systems normally fit in the following groups:
 - a. Determination of the erosion stability of materials occurring in the channel bed and banks.
 - b. Determination of the sliding stability of the materials forming the channel bank.
 - c. Determination of leakage from channel sections.
 - d. Determination of uplift pressures, magnitude of structural loading and bearing capacity for channels to be lined.
- 4. Other structures. Soil engineering problems associated with the design of drop structures, diversion structures, pumping plants, etc., will generally fall in the following groups:
 - a. Determination of the bearing capacity of the foundation.
 - b. Determination of the sliding stability of the overall structure.
 - c. Determination of the earth-water loads on structural members.
 - d. Determination of settlement, both total and differential, of various parts of the structure.
 - e. Evaluation of piping problems along the structure-soil interface.
 - f. Evaluation of scour potential adjacent to hydraulic structures.
- C. INFORMATION REQUIRED AS BASIS FOR DESIGN

To approach the problems encountered in design, certain information about the character of the foundation and the construction materials is essential. The information required for analysis of foundation conditions and the design of the proposed work are:

- 1. Character and distribution of materials in the foundation to significant depth.
- 2. The location and seasonal fluctuation of the groundwater table.

- 3. The surface topography of the site.
- 4. The geologic structure of the underlying formation.
- 5. The character, location and available quantities of fill materials.

Other items that will affect the construction operations such as site seismicity, climatic conditions, diversion and control of the stream, magnitude of wave action, time restrictions on construction and the function of the facility must be studied in preparing the design.

D. USE OF SOIL MECHANICS IN CONSTRUCTION

A knowledge of engineering soil behavior is necessary to understand the effect of construction operations on soil masses. The need for this knowledge and experience, along with the exercising of sound engineering judgment by personnel in charge of construction supervision, must be continuously emphasized.

In his book "Earth Dam Projects," J. P. Justin wrote, "An entirely safe and substantial design may be entirely ruined by careless and shoddy execution, and the failure of the structure may be the result. Careful attention to the details of construction is, therefore, fully as important as the preliminary investigation and design."

In earthwork engineering, to achieve maximum economy it is generally necessary to use locally available materials. In doing this the designer assesses the behavioral characteristics of the locally available materials and blends them into an efficient protective design meeting the requirements of site conditions and project objectives.

The earthwork designed must provide a greater tolerance range of earth materials use than that provided by the designer using other construction materials. This is necessary because of the inherent variations in the engineering behavior of earth materials, from deposit to deposit as well as within a given deposit, and because generally earth structures are massive, containing large volumes of materials. Consequently, a much closer relationship between operation of investigation, design, inspection and construction is required for earthwork than is needed for other types of engineered construction.

In general, the following categories of information are needed to insure proper execution of earthwork construction:

1. Knowledge of site conditions, including regional geology, character and distribution of foundation materials, water

table location and fluctuations, character and location of. proposed fill materials, and geologic and structural conditions of rock to be encountered.

- 2. Knowledge of the intent of the designer as well as complete understanding of construction drawings and specifications.
- 3. Understanding of the engineering behavior of the soils at the site. An understanding of how these soils will respond to construction operations and various types of construction equipment.
- 4. Understanding of all the earthwork construction control tests and what the results mean with respect to engineering behavior and construction response.
- E. NOMENCLATURE, DEFINITIONS AND CONVENTIONS

Insofar as is practicable, the definitions, nomenclature and conventions contained in ASTM D-653, Standard Definitions of Terms and Symbols Relating to Soil Mechanics, prepared by a joint ASCE-ASTM committee, will be followed.







A. ENGINEERING SOIL CLASSIFICATION

The fundamental purpose of a soil classification system is to provide a language by means of which one person's knowledge of the general characteristics of a particular soil can be conveyed to another person in a brief and concise manner, without entering into lengthy descriptions and detailed analysis.

A good soil classification system should satisfy the following requirements:

- 1. It should describe the soil in well-understood terms, which convey an idea of its type and behavior.
- It should furnish an indication of soil properties and performance.
- 3. It should be applicable from visual examination both in a simplied form and, with experience, in a more refined form.
- It should employ a simple system of notations for graphic abstracts of boring logs on drawings recording boring information.

Basically there are four classifications for describing soils. These are:

- 1. Geological
- 2. Textural
- 3. Pedological
- 4. Engineering

In engineering work, it is frequently appropriate to use at least two or three, or all of these classifications for describing soils at a given project site, such as (a) the geological classification, which provides a language for stating the geological history and background of the soil; (b) the textural classification which provides names for describing the relative portions of sand, silt and clay size particles in a soil; (c) the pedological classification which reveals the nature of the soil profile as affected by the climatic and other environmental conditions under which it developed and (d) an engineering classification that groups soils on the basis of their characteristics which influence engineering performance. Great care should be exercised not to read into any classification more information than they are intended to convey. For instance, there is no engineering classification in existence that can be used directly for design values. Testing of individual materials is always necessary to determine the engineering properties of permeability, strength, and compression characteristics.

The development of a soil classification system is accomplished by the empirical correlation of the physical properties of soils with their observed behavior characteristics under such varying conditions as climate, treatment, loading, and construction methods. It is generally found that certain ranges of physical properties may be established within which the soils possessing these properties will react in approximately the same manner to a certain set of imposed conditions.

It is important that, within a given organization, the methods for selecting soil materials for a specific purpose be uniform throughout the organization. The adoption of a standard soil classification system provides a comprehensive means of communication and facilitates the review of reports and designs at all levels within the organization.

Many soil classification systems have been developed; each for the purpose of classifying soils with respect to their suitability for a specific purpose. The "Unified Soil Classification System" developed by Arthur Casagrande and modified by the Bureau of Reclamation and the Corps of Engineers is the classification system used by the Soil Conservation Service for engineering purposes.

The soil classification system developed by the U.S. Bureau of Public Roads and adopted by the American Association of State Highway and Transportation Officials is included because of its use in published soil surveys.

B. UNIFIED SOIL CLASSIFICATION SYSTEM

1. Soil Properties Used in Classification

This system identified soils according to their textural and plasticity qualities and on their groupings with respect to behavior. The system is based on those characteristics of a soil that indicate how it will behave as an engineering construction material. The soil properties that have been found most useful for this purpose and form the basis of soil identification are:

- a. Percentage of gravel, sand and fines (fraction passing No. 200 sieve).
- b. Shape of grain-size distribution curve.
- c. Plasticity and compressibility characteristics.

These properties are determined by mechanical analysis, liquid limit and plastic limit tests and with experience can be estimated with good accuracy.

2. Definition of Soil Components

ComponentSize RangeBouldersLarger than 12" diameterCobblesBetween 3" and 12" diameterGravelBetween No. 4 mesh (4.76 mm) and
3" diameterSandBetween No. 200 mesh (.074 mm) and
No. 4 meshFines*Smaller than No. 200 mesh

* Fines include silt and clay. A material is called a silt if it is non-plastic or very slightly plastic and exhibits little or no strength when air-dried. Silt fines plot below the A-line, Table 2-1. A material is called a clay if the fines exhibit plasticity within a range of water contents and has considerable strength when air-dried. Clay fines plot above the A-line, Table 2-1.

A comparison of the size of the soil components as defined in the UNIFIED, the AASHTO and the USDA textural classification systems is shown on Figure 2-1.

3. The Plasticity Chart (Figure 2-2)

The plasticity chart is a plotting of the plasticity index versus the liquid limit. The A-line on the plasticity chart is an important empirical boundary.

In general, soils with Atterberg limits that plot above the A-line behave as typical non-organic clays. Soils plotting below the

ASSIFICATION SYSTEM	LABORATORY CLASSIFICATION CRITERIA	$C_u = \frac{D}{D_{10}}$ Greater than 4 ; $C_e = \frac{(D_{30})^2}{D_{10}}$ Between 1 & 3	Not meeting all graduation requirements for GW	find the second	Of 10 Of 10 Of 10 Of 10 Of 00 Of 00 Of 00	We centre \mathbb{N}_{0}^{*} $\mathbb{N}_{0}^$	Not meeting all graduation requirements for SW	유민 성 성 성 성 성 kterberg limits below "A" Above "A" line with PI between 4 and 7 are	Atterberg limit above "A" borderline cases requiring use of dual symbols.	% G Jow SSOT DUUS OZIS	NDEX NDEX		PLAST PLAST PLAST PLAST	0 10 20 30 40 50 60 70 80 90 100	PLASTICITY CHART	elow "A" line and PI and $\frac{1}{L.L.}$ (air dry soil) < 0.75		TI. CLASSTFICATION SYSTEM
CRITERIA FOR UNIFIED SOIL C	TYPICAL NAMES	Well graded gravels, gravel-sand mixtures, little or no fines.	Poorly graded gravels, gravel- sand mixtures, little or no fines.	Silty gravels, poorly graded, gravel-sand-	Clayey gravels, poorly graded, gravel-sand- clay mixtures.	Well graded sands, gravelly sands, little or no fines.	Poorly graded sands, gravelly sands, little or no fines.	Silly sands, poorly graded sand-silt mixtures.	Clayey sands,poorly graded sand-clay mixtures.	949(Inorganic silts and very fine sands, rock flour, silty or clayey fine sands with slight plasticity.	Inorganic clays of low to medium plasticity, gravelly clays,sandy clays,silty clays,lean clays.	Inorganic silts,micaceous or diatomaceous fine sandy or silty soils, elastic silts.	Inorganic clays of high plasticity, fat clays.	Organic silts and organic silt-clays of low plasticity. LL<50	Organic clays of medium to high plasticity.	Peat and other highly organic soils	- LARORATORY CRITERIA FOR HNIFLED SC
ATORY	GROUP	GW	GР	GM	CC	SW	SP	SM	sc		ML	CL	HW	СН	OL	но	Pt	ARI.F. 2-1 -
LABOR		e. səss əų.	TO PLASTICITY CHART TO PLASTICITY CHART The No.4 sieve size. The No.4 sieve size.							01 O9		E						
			Sands and Gravels								skr	PID P	ud si	IIIS				
		Fine Grained Soils Coarse Grained Soils d ₅₀ No. 200 Mesh(0.074mm)																
		L								2_/	.							

FIGURE 2-1. A COMPARISON OF GRAIN-SIZE LIMITS IN THE 3 CLASSIFICATION SYSTEMS

American Association of State Highway and Transportation Officials Soil Classification

U.S. Department of Agriculture Soil Textural Classification

C Unified Soil Classification

Boulders	Cobbles	Cobbles	ç-	- 0.08	
Coarse Gravel	Coarse Gravel	Coarse Gravel		30.0 40.0	
Medium Grave		Fine . ravel	- N m 4	0.01	
Fine Gravel	Fine Gravel	arse band G	4-	6.0 4.0	
. es	Very Coarse Sand		01-	- 0.5	
Coar Sar	ium Coarse nd Sand	nd Sand Medi Sar		9' 9'	.e – mm
Fine Sand	Fine Med Sand Sa	Fine Sand	09 -	<u>ς</u> . ε.	rticle Siz
_	Very Fine Sand		-140 -500	-1' -80' 90'	Pa
y Silt	Silt	es(Silt or Clay)**	eve Sizes	500. 400. 800. 10. 20. 20. 20. 20.	
Colloids Cla	Clay	Fine	Sié	100. S00.	

*Colloids included in clay fraction in test reports.

** The LL and PI of "Silt" plot below the "A" line on the plasticity chart, Figure 2-2 and the LL and PI for "Clay" plot above the "A" line

Modified from PCA Soil Primer



A-line behave as plastic soils containing organic colloids or as typical inorganic silts. An exception to the above rule is the shaded area above the A-line with PI greater than 4 and less than 7, soils with Atterberg limits plotting in this zone may behave as a clay or as a silt.

The U-line on the plasticity chart represents the upper limit of the LL and PI plots for soils. At this date, (1973) no soils have been tested that plot above this line.

The plasticity chart is very useful in classifying soils for engineering purposes. The soil groupings of the unified soil classification system are shown on a plasticity chart in Table 2-1 and on Figure 2-3.

Much useful information about the behavior of a soil can be inferred from the plasticity chart.

If a mineral is ground up and the Atterberg limits of the various grain size fractions determined, the points representing such test results plot along a straight line roughly parallel to the A-line. Depending on the mineralogical compositions of the grains, the plot may be above or below the A-line.

The points representing the limits of soils from a geologically well defined sedimentary deposit will plot along a line roughly parallel to the A-line.

4. Summary of the Unified Classification System

- a. Designates soils as fine or coarse according to median size. Coarse soil if median size is larger than 0.074 mm (No. 200 mesh). Fine grained if median size is smaller than 0.074 mm (No. 200 mesh).
- b. Coarse grained soils are classified on the basis of:
 - (1) Grain size and distribution
 - (2) Quantity of fines
 - (3) Character of fines
- c. Fine grained soils are classified on basis of:

(1) Liquid limit - High if LL is greater than 50 Low if LL is less than 50

(2) Plasticity index - Clay if above A-line, silt if below A-line.





2--8

- (3) Grading is of minor importance.
- d. Soil groups and group symbols of the Unified Classification System.

Basic Symbols	Modifying Symbols
G - gravel	W - well graded
S - sand	P - poorly graded
	C - with clay fines
	M - with silt fines
C - clay	L - low liquid limit
M - silt	H - high liquid limit
0 – organic	
Pt - peat	

Each soil is classified and identified with a verbal description and a group symbol consisting of two of the above letters. The letters may be considered as initials of the name of the most typical soil in the group.

Coarse grained soils are sub-divided into:

(1) Gravel and gravelly soils; symbol G

(2) Sands and sandy soils; symbol S

The gravels and the sands are each divided into four groups:

- (a) Well graded, fairly clean material; symbol W, in combinations GW and SW.
- (b) Poorly graded, fairly clean material; symbol P, in combinations GP and SP.
- (c) Coarse material with clay fines, symbol C, in combinations GC and SC.
- (d) Coarse materials with silt fines, symbol M, in combinations GM and SM.

Fine grained soils are subdivided into:

- The inorganic silty and very fine sandy soils; symbol M, used for fine-grained non-plastic or slightly plastic soils.
- (2) The inorganic clays; symbol C.

(3) The organic silts and clays; symbol 0.

Each of these types of fine-grained soils is grouped according to its liquid limit into:

- (a) Fine-grained soils having liquid limits less than 50; symbol L, in combinations ML, CL and OL.
- (b) Fine-grained soils having liquid limits greater than 50: symbol H, in combinations MH, CH and OH.

Highly organic soils, usually fibrous, such as peat and swamp soils having high compressibility, are not subdivided and are placed in one group; symbol Pt.

The sequence of the group symbols need not be memorized but the meaning of the symbols and the sequence of the major divisions, that is, G-S-L-H should be learned.

When a material does not clearly fall into one group, boundary classification such as GW-SW or CL-ML should be used.

- 5. Method of Classification (Laboratory Test Data)
 - a. For all soils.

Step No. 1. Classify as coarse grained or fine grained soil. Coarse grained if the median size is larger than 0.074 mm (#200 sieve).
Fine grained if the median size is less than 0.074 mm (#200 sieve).

b. For fine grained soils.

Step No. 2. Enter plasticity chart with LL-PI data.

Classify as silt if LL-PI data plot below A-line (Table 2-1) or PI is less than 4. Classify as clay if LL-PI data plot above A-line with PI greater than 7. Dual classification symbols should be used if LL-PI data plot above A-line with the PI between 4 and 7.

- c. For coarse grained soils.
 - (1) Step No. 2. Classify as sand or gravel.

Sand if 50% or more of coarse fraction is sand size
(#4 to #200 sieve).
Gravel if 50% or more of coarse fraction is gravel
size (#4 to 3" sieve).

(2) Step No. 3. Classify as clean or with fines.

Clean if material contains less than 5% non-plastic fines that do not interfere with the free draining properties. In areas subject to frost action, the material should not contain more than 3% of particles smaller than 0.02 mm in size to classify as clean. A material containing less than 5% plastic fines should be classed as a borderline soil with dual symbols.

With fines if material contains more than 12% fines.

Borderline case with dual symbols if materials contain between 5-12% fines.

(3) For clean sand and gravels

<u>Step No. 4</u>. Compute the coefficient of uniformity C_u (D_{60} size divided by D_{10} size) and the coefficient of curvature C_c (D_{30} size squared divided by the product of the D_{10} size and the D_{60} size). Classify soil as well graded or poorly graded.

Sand is well graded if C_u is greater than 6 and C_c is greater than 1 and less than 3; poorly graded otherwise. Gravel is well graded if C_u is greater than 4 and C_c is greater than 1 and less than 3; poorly graded otherwise.

(4) For sands or gravels with fines.

Step No. 4. Determine character of fines.

Coarse grained with silt fines if LL-PI data plot below A-line (Table 2-1) or PI is less than 4. Coarse grained with clay fines if LL-PI data plot above A-line with PI greater than 7.

Examples illustrating the use of the Unified Soil Classification System are on pages 2-12 through 2-21. The examples cover a wide range of soil materials. The procedure used in classifying each soil is listed in the remarks space on the soil classification report sheets for each example.



2













^{2-18:}



LABORATORY NO. _





6. Field Identification of Soils

a. General

The unified soil classification system is so devised that it is possible, with experience, to classify most soils correctly on the basis of field identification methods alone. The easiest way to learn field identification is under the guidance of an experienced man. While learning the procedures, a man should systematically compare the numerical test results for typical soils in each group with the "feel" of the materials while performing the field identification methods.

The following tests were developed largely by Professor A. Casagrande, Graduate School of Engineering, Harvard University, and have been widely adopted for use in identification of soils. These tests can be performed without equipment and are simple in nature. Do not make a decision on the basis of a single test. Use all applicable tests, then identify the soil. Identification criteria are presented in Table 2-2.

- b. Test Methods
 - (1) Visual Inspection

Grain Shape. Observe and classify the sand and gravel particles as to degree of angularity or roundness. (See Figure 2-4.)



FIG. 2-4. GRAIN SHAPE
TABLE 2-2. UNIFIED SOIL CLASSIFICATION

FIELD IDENTIFICATION

FIELD IDENTIFICATION PROCEDURES

-				_					-				_				
			solls	2		CL	EAN	N GRAVEL	s	Wide ran amounts	ge of	in grain si all interm	izes edi	and subs ate partice	tan al s	tial sizes.	GW
	-		RAVELLY S	weight) is	4 inch size v	di	rt st et p	ain on a aim.		Predomin with som	ate e i	ly one siz ntermediat	e o e si	r a range zes missin	of g.	sizes	GP
ILS	veight) is o iked eye .		EL AND GI	action (by	·ger than - th size may 4 sieve size	DI	RT) ill le	r GRAVELS ave a dirt	6	Nonplast (for iden of ML be	ic tifi elo	fines or fin cation of f w.)	nes ine:	with low s see char	pla act	sticity teristics	GM
AINED SO	terial (by v le to the no		GRAVE	L.L.	$\frac{1}{100}$	st po	ain alm.	on a wet		Plastic fi character	ne: ris [.]	s (for ident tics of CL	ific or	ation of f CH below)	ines)	s see	GC
ARSE GRI	alf of mai rains visibl	ed eye.	L S rse	2	ze issification equivalent	c	LEA ill n	AN SANDS		Wide rang amounts	ge of	in grain siz all interme	e a dia	nd substan te partical	ntia siz	l zes.	SW
CO	fore than l idividual g	to the nake	ANDY SO	weight) is	4 inch si r visual clo used as	di w	et p	stain on a balm.		Predomin with som	an 1e	tly one siz intermedia	te o	r a range sizes missi	of ng	sizes	SP
	- <u>-</u>	le visible	ND AND S	action (by	naller than Fo			'Y SANDS eave a dirt	+	Nonplast (for iden of ML b	ic tif elc	fines or fi ication of (ow).	nes fine	with low s see cha	plo rac	asticity teristics	SM
		lest partic	SA	L.	E S	st p	tain alm	on a wet		Plastic fi characte	ris	s(for ident tics of CL	lific or	ation of f CH below	ine)	s see	SC
		it the small	- AYS mit)					Slight		Rapid		Low to None		None		Duli	ML
6	elght) is of e naked ey	ize is abou	S AND CI Liquid Li	ures				Medium to High	ACTION	Medium to Slow		Medium		Weak		Slight to Shiny	CL
NED SOILS	erial (by w isible to the	00 sieve s	MOT) SILT	ion Proced	Pronou Orgar	nced nic	RENGTH	Medium	AKE) REA	Slow to None	INESS	Low (Spongy)	AR THE P	None	VR THE PL	Dull to Slight	OL
INE GRAI	half of mat rains not vi	No. 2(LAYS mit)	Identificat	do O		DRY ST	Medium	ANCY (SH.	Very Slow to None	TOUGH	Medium to High	BON (NE	Weak to Strong	SHINE (NEA	Slight	MH
	More than ndividual gi		S AND CI Liquid Lir	See				Very High	DIL AT	None		High	RIE	Strong	0	Shiny	СН
			SILT (High		Pronou Orgai	nced nic	-	High		None		Low to Medium (Spongy)		Weak		Dull to Slight	ОН
		нібн	ILY ORGA	NIC	SOILS				Rec fee	adily ident I and freq	ifie jue	d by color ntly by fib	r,od rou	lor,spong s texture	у		Pt

Grain Size and Gradation. Sand and gravel sizes are readily identified by visual inspection. Individual grains below the smallest sand size cannot be seen by the naked eye and must be identified by other tests.

To estimate the gradation of <u>coarse-grained soils</u>, spread a representative sample out on a flat surface and observe the distribution or uniformity of grain sizes. Observe the proportion of fines and subject the <u>fine-grained fraction</u> to all tests described for fine-grained soils.

For gradation of <u>fine-grained soils</u>, shake the sample in a jar of water and allow it to settle out. Approximate gradation is indicated by the separation of the particles in the jar from top to bottom. Silt remains in suspension for at least one minute; clay, one hour or more.

(2) Shaking Test (Dilatancy)

The shaking test aids in the identification of finegrained soils. After removing particles larger than No. 40 sieve size, prepare a pat of moist soil with a volume of about one-half cubic inch. Add enough water if necessary to make the soil soft but not sticky.

Place the pat in the open palm of one hand and shake horizontally, striking vigorously against the other hand several times (Figure 2.5). A positive reaction



Method of shaking

FIG. 2-5. SHAKING TEST

consists of the appearance of water on the surface of the pat which changes to a livery consistency and becomes glossy. When the sample is squeezed between the fingers, the water and gloss disappear from the surface, the pat stiffens, and finally it cracks or crumbles. Shake the broken pieces until they become livery and flow together again. Distinguish between slow, medium, and rapid reaction to the shaking test.

Rapid Reaction indicates a lack of plasticity, such as is the case with a typical inorganic silt, a rock flour, or a very fine sand.

Slow Reaction indicates a slightly plastic silt or silty clay.

No Reaction indicates a clay or a peaty (organic) material.

(3) Breaking Test (Dry Strength)

The breaking test may be used to determine the dry strength of a soil, which is a measure of its cohesiveness.

After removing particles larger than No. 40 sieve size, mold a pat of soil to the consistency of putty, adding water if necessary. Allow the pat to dry completely by oven, sun, or air drying, and then test its strength by breaking and crumbling between the fingers (Figure 2-6). This strength is a measure of the character and quantity of the colloidal fraction contained in the soil. The dry strength increases with increasing plasticity.



Method of crumbling soil between fingers

FIG. 2-6. BREAKING TEST

Slight dry strength indicates an inorganic silt, a rock flour, or a silty sand. However, the sand feels gritty when the sample is powdered. The dried soil pat can be powdered with slight finger pressure.

Medium dry strength indicates a low to medium plastic inorganic clay. Considerable finger pressure is required to powder the sample.

High dry strength indicates a highly plastic, inorganic clay. The dried sample may be broken but cannot be powdered by finger pressure.

Note: Cohesion or high dry strength may be furnished by some cementing material such as calcium carbonate or iron oxide. For example, non-plastic lime rock or coral may develop high dry strength.

(4) Plasticity Tests

Plasticity is the physical property of a finegrained soil which allows it to be kneaded into a putty-like consistency at the proper moisture content.

To test plasticity of a soil, prepare a moist pat and remove the coarse particles. Start rolling it out with the palm of the hand on a flat surface into approximately 1/8 inch diameter threads (See Figure 2-7. Fold the threads into a lump, roll it out again, and repeat the process until moisture loss causes the thread to crumble just when the 1/8-inch diameter is reached. The moisture content at this stage is called the plastic limit. Note the toughness of the threads as the plastic limit is reached and test to see if the crumbled pieces can be lumped together again.

- A. Method of rolling thread.
- **B**. Thread of soil above plastic limit.
- C. Crumbling thread as plastic limit is reached.



FIG. 2-7. PLASTICITY TEST

Any soil that can be readily rolled into a thread without crumbling is plastic.

High plasticity clay forms a tough thread which can be remolded into a lump below the plastic limit and deformed under high finger pressure without crumbling.

Medium plasticity soil forms a medium tough thread, but the lump crumbles soon after the plastic limit is reached.

Low plasticity soil forms a weak thread that cannot be lumped together below the plastic limit. Plastic soils containing organic material or much mica form threads which are very soft and spongy.

(5) Odor Test

Freshly sampled organic soils usually have a distinctive odor which aids in their identification. The odor can be made more apparent by heating a wet sample.

(6) Acid Test

Drop a little hydrochloric acid on a piece of soil. A fizzing reaction indicates calcium carbonate.

(7) Shine Test

Rub a dry or slightly moist sample with the finger nail or a knife blade. A shiny surface indicates a highly plastic clay; a dull surface indicates a silt or clay of low plasticity.

7. Description of Undisturbed Soil

The identity of a soil as determined by the laboratory and field tests should be supplemented by a description of the soil in its undisturbed condition. The description should include remarks concerning the following characteristics:

a. Color

The color of a soil in both the moist and dry condition should be described as accurately as possible by use of standard color designations. Acceptable soil color descriptions are: white, gray, black, brownish-black, reddish-gray, brownish-gray, orange, red brown, yellowishbrown, olive-brown, yellow, olive, blue, green, etc.,; these terms should be further modified by the adjectives light, medium, dark, or vivid. Use of the Munsell soil color charts and plates is suggested when more precise soil color descriptions are desired.

Mottles or streaks should be indicated and their description should follow the main color designation of the soil; for example, "stiff clay, plastic when moist, friable when dry; medium blue, mottled with brown, when moist; light green, mottled with brown when dry."

b. Texture

The term "texture" refers to the distinctive appearance and "feel" of the soil, which are direct indications of the fineness and uniformity of the soil. Texture should be described by a standard adjective as listed below:

c. Consistency

The term "consistency" refers to the degree of adhesion between soil particles or the resistance to deformation or rupture under applied pressure. Consistency of the moist soil in both the undisturbed and remolded states should be described by standard adjectives as listed below:

- (1) Cohesionless soils (sands and silts)
 - Loose Poorly graded, lacks binder or cementing agent, not well compacted.
 - Dense Well graded, well compacted, may contain binder or cementing agent.

(2) Cohesive Soils (clays)

Very Soft - Easily penetrated several inches by fist..

Soft - Easily penetrated several inches by thumb.

Medium - Can be penetrated by thumb with moderate effort.

Stiff - Readily indented by thumb but penetrated only with a great effort.

Very Stiff - Readily indented by thumbnail.

Hard - Indented with difficulty by thumbnail.

Additional adjectives used in connection with the terms listed above in describing consistency of cohesive soils are: sticky, plastic, friable, brittle, jointed, stratified, varved.

d. Moisture Content.

The apparent moisture content of the soil should be described by such terms as dry, moist, wet, saturated, etc.

e. Cobbles and Boulders

The amount of each of the following sizes over 3 inches in diameter is of great importance in the selection of sources of and the use of embankment material.

- 3" to 6" this size is generally left in the material used in an embankment. The amount of this size is necessary to evaluate the suitability of the proposed fill material.
- Larger than 6" size this size of material is generally excluded from the impervious zones of an embankment. The amount of this size must be known to determine waste volumes and its possible use in other zones of the proposed work.

C. AASHTO SOIL CLASSIFICATION SYSTEM

1. Soil Properties Used in Classification

This system identifies soils according to their textural and plasticity qualities and on their groupings with respect to behavior. This system is based on those characteristics of a soil that indicate how it will behave as an engineering construction material.

The soil properties that have been found most useful for this purpose and form the basis of the AASHTO soil classification system are:

- a. Percentage of gravel, coarse sand, fine sand, and fines (fraction passing No. 200 sieve).
- b. Plasticity and compressibility characteristics.

c. Group index.

These properties are determined by mechanical analysis, liquid limit and plastic limit tests and with experience can be estimated with good accuracy.

2. Definition of Soil Components

Component

Size Range

Boulders	Larger th	an 3''	diamet	ter			
Gravel	Between N	o. 10	mesh a	and (3''		
Coarse Sand	Between N	o. 40	mesh a	and 1	No.	10 n	lesh
Fine Sand	Between N	o. 200	mesh	and	No.	40	mesh
Fines*	Smaller th	han No	. 200	mest	n		

* Fines are classed as silty if the plastic index is equal to or less than 10 and clayey if it is equal to or greater than 11.

3. Summary of the AASHTO Classification System

- a. Designates soils as fine or coarse according to percent passing the No. 200 mesh. Coarse soil, if less than 35% passes the No. 200 mesh. Fine soil, if more than 35%, passes the No. 200 mesh.
- b. Coarse grain soils are further classified on the basis of:

(1) Grain size and distribution.

- (2) Quantity of fines.
- (3) Character of fines.
- (4) Group index.
- c. Fine grained soils are classified on the basis of:
 - (1) Liquid limit high if LL is greater than 40
 low if LL is less than 40.
 - (2) Plasticity index clay if greater than 10.- silt if 10 or less.
 - (3) Group index.
- d. Soil groups and group symbols of the AASHTO classification system:
 - Granular materials containing 35 percent or less passing the No. 200 sieve.
 - (a) Group A-1. The typical material of this group is a well graded mixture of stone fragments or gravel, coarse sand, fine sand, and a nonplastic or feebly plastic soil binder. This group includes stone fragments, gravel, coarse sand, volcanic cinders, etc. without soil binder.
 - <u>1</u> Subgroup A-1-a includes those materials consisting predominantly of stone fragments or gravel, either with or without a well graded binder of fine material.
 - <u>2</u> Subgroup A-1-b includes those materials consisting predominantly of coarse sand either with or without a well graded soil binder.
 - (b) Group A-3. The typical material of this group is fine beach sand or fine desert blow sand without silty or clay fines or with a very small amount of nonplastic silt. The group may include streamdeposited mixtures of poorly graded fine sand and limited amounts of coarse sand and gravel.

- (c) Group A-2. This group includes a wide variety of "granular" materials which are mixtures of the materials falling in Groups A-1 and A-3 and the silt-clay materials of Groups A-4, A-5, A-6, and A-7. It includes all materials containing 35% or less passing the No. 200 sieve which cannot be classified as A-1 or A-3 because of fines content or plasticity or both.
 - Subgroups A-2-4 and A-2-5 include various granular materials containing 35% or less passing the No. 200 sieve and with a minus No. 40 portion having the characteristics of the A-4 and A-5 groups. These groups include such materials as gravel and coarse sand with silt contents or plasticity indexes in excess of the limitations of Group A-1, and fine sand with nonplastic silt content in excess of the limitations of Group A-3.
 - 2 Subgroups A-2-6 and A-2-7 include materials similar to those described under Subgroups A-2-4 and A-2-5, except that the fine portion contains plastic clay having the characteristics of the A-6 and A-7 group.
- (2) Silt-clay materials containing more than 35 percent passing the No. 200 sieve.
 - (a) Group A-4. The typical material of this group is a nonplastic or moderately plastic silty soil usually having 75% or more passing the No. 200 sieve. The group also includes mixtures of fine silty soil and up to 65% of sand and gravel retained on No. 200 sieve.
 - (b) Group A-5. The typical material of this group is similar to that described under Group A-4, except that it is usually of diatomaceous or micaceous character and may be highly elastic, as indicated by the high liquid limit.
 - (c) Group A-6. The typical material of this group is a plastic clay soil usually having 75% or more passing the No. 200 sieve. The group also includes mixtures of fine clayey soil and up to 64% of sand and gravel retained on the No. 200 sieve. Materials of this group usually have high volume change between wet and dry states.

- (d) Group A-7. The typical material of this group is similar to that described under Group A-6, except that it has the high liquid limits characteristic of the A-5 group, and may be elastic as well as subject to high volume change.
 - <u>1</u> Subgroup A-7-5 includes those materials with moderate plasticity indexes in relation to liquid limit and which may be highly elastic as well as subject to considerable volume change.
 - 2 Subgroup A-7-6 includes those materials with high plasticity indexes in relation to liquid limit and which are subject to extremely high volume change.
- (e) Group A-8. These soils are highly organic peats or mucks. Classification of these materials is based on visual inspection, and is not dependent on the percentage passing the No. 200 sieve, liquid limit, or plasticity index. The material is primarily composed of partially decayed organic matter, generally has a fibrous texture, dark brown or black color, and a very noticeable odor of decay.
- 4. Method of Classification (Laboratory Test Data)
 - a. For all soils.
 - (1) <u>Step No. 1</u>. Determine the group index from the formula:

GI = (F-35)[0.2 + 0.005(LL-40)] + 0.01(F-15)(PI-10)in which

F = percentage passing No. 200 sieve, expressed as a whole number. This percentage is based only on the material passing the 3-in. sieve. LL = liquid limit PI = plasticity index

The group index may also be determined by the use of Figure 2-8.

Group Index (GI) = (F-35) [0.2+0.005(LL-40)] +0.01(F-15)(PI-10) where F=% Passing No. 200 sieve, LL = Liquid Limit, and PI = Plasticity Index.

When working with A-2-6 and A-2-7 subgroups the Partial Group Index (PGI) is determined from the PI only.



FIG. 2-8. GROUP INDEX CHART

- (a) If calculated group index is negative, report group index as zero.
- (b) Report group index to the nearest whole number.
- (c) In calculating group index of A-2-6 and A-2-7 soils, use only PI portion of formula or of Figure 2-8.
- (2) <u>Step No. 2</u>. Classify as coarse grained or fine grained soil.

Coarse grained if less than 35% passing the No. 200 mesh. Fine grained if more than 35% passing the No. 200 mesh.

b. For fine grained soils.

Step No. 3. Determine soil group from Table 2-3 and Figure 2-9.

- c. For coarse grained soils.
 - Step No. 3. Determine percentages passing the No. 10, 40, and 200 mesh and determine soil groups in the coarse grained portion of Table 2-3.
 - (2) <u>Step No. 4.</u> Determine liquid limit and plasticity index and determine soil group using Table 2-3 and Figure 2-9.

Example problems on page 2-12 to 2-21 have AASHTO class shown.

The position of the soils groups in the AASHTO classification system on a plasticity chart are shown in Figure 2-10.

AASHTO CLASSIFICATION OF SOILS AND SOIL-AGGREGATE MIXTURES

•

								•
0. 200)	A- 7	A- 7-5 A-7-6		36 min.	41 min. 11 min.	Soils		us 30.
Materials passing No		A-6		36 min.	40 max. 11 min.	Clayey		n LL'min
Silt -Clay than 35%		A-5		36 m in.	41 m in. 10 max.	Soils	to Poor	greater tho
(More		A-4		36 min.	40 max. 10 max.	Silty	Fair	ubgroup is
		A-2'- 7		35 max.	41 min. 11 min.	Sand		of A-7-6 s
	2	A-2-6		35 max.	40 max. 11 min.	Gravel and		city index
200)	Ą	A-2-5	-	35 max.	41 min 10 max	or Clayey		30. Plasti
Materials bassing No.		A-2-4		35 max.	40 max. 10 max.	Silty	poog	LL minus
Granular % or less p		A-3		51 min. 10 max.	a: z:	Fine Sand	cellent to (r less than
(35	_	A - I - b		50 max. 25 max.	max.	agments, ind Sand	Ж	eaual to o
	Α-	A - † - a	50 max.	30 max. 15 max.	ک	Stone Fi Gravel o		subaroup is
General Classification	Group Classification		Sieve Analysis, Percent passing No. 10	No. 40 No. 200	Characteristics of Fraction passing No. 40 : Liquid limit Plasticity index	Usual types of Significant Constituent Materials	General Rating as Subgrade	^a Plasticity index of A-7-5

TABLE 2-3







A. SOIL MASS AS A MULTIPHASE SYSTEM

The soil mass consists of a framework of solid particles of varying size and shape enclosing voids of varying size between the particles. The voids may be filled with water, air, or partly with water and partly with air (see Figure 3-1).

1. Solid Phase

The solid phase includes all soil particles with included organic or other solid matter.

The soil particles range from boulders to fines as defined in Section 2.

2. Liquid Phase

For purposes of soil mechanics, the liquid phase is considered to be water, however under field conditions it may include dissolved salts, oil, gas or other liquids. In soil mechanics, water is defined as the material evaporated when a soil sample is dried to constant weight at a temperature of 105 to 110 degrees centigrade.

3. Gas Phase

In soil mechanics, the gas phase is considered to be air.

- 4. Weight Relationships (see Fig. 3-1 and Fig. 3-2)
 - a. Nomenclature

 W_S = Weight of the solid phase of the soil (oven dry). W_W = Weight of water. W_a = Weight of gas (assumed to be zero). W = Total weight of the mass = W_S + W_W + W_a = W_S + W_W

b. Moisture Content = w

The ratio of the weight of water to the weight of solids in the soil mass expressed as a percentage.

$$w = \frac{W_W}{W_S} \quad (100)$$

c. Specific Gravity

(1) Specific gravity of solids = G_s

Ratio of the weight in air of a given volume of soil solids at a stated temperature to the weight in air of an equal volume of distilled water at a stated temperature.

$$G_s = \frac{\gamma_s}{\gamma_s}$$

where γ_s = unit weight of solids γ_0 = unit weight of water at 4° C.

(2) Apparent specific gravity = Ga or G

Ratio of the weight in air of a given volume of the impermeable material (that is the solid matter including its impermeable voids) at a stated temperature to the weight in air of an equal volume of distilled water at a stated temperature.

$$G_a = G = \frac{\gamma_{s+v}}{\gamma_o}$$

where γ_{s+v} = unit weight of solids plus impermeable voids

(3) Bulk specific gravity (specific mass gravity) = G_m

Ratio of the weight in air of a given volume of permeable material (including both permeable and impermeable voids normal to the material) at a stated temperature to the weight in air of an equal volume of distilled water at a stated temperature.

$$G_{\rm m} = \frac{\gamma_{\rm t}}{\gamma_{\rm o}}$$

where γ_{t} = unit weight of soil mass

5. Volume Relations (see Fig. 3-1 and Fig. 3-2)

a. Nomenclature

 V_s = Volume of solid phase of soil $V_w = Volume of water$ $V_a = Volume of gas$ $V_v = Volume of voids = V_w + V_a$

V = Total volume of soil mass $V = V_s + V_a + V_w = V_s + V_v$

b. Porosity = .n

The ratio of the volume of voids to the total volume of the mass, expressed as a percentage.

n =
$$\frac{V_v}{V}$$
 (100) = $\frac{e}{1+e}$ (100)

c. Void ratio = e

The ratio of the volume of voids to the volume of solids in the soil mass.

$$e = \frac{V_V}{V_S} = \frac{n}{100-n}$$

d. Degree of saturation = S

The ratio of the volume of water to the volume of voids, expressed as a percentage.

$$S = \frac{V_w}{V_v} (100)$$

6. Volume and Weight Relationships

a. Total unit weight =
$$\gamma_t$$

The ratio of the total weight to the total volume

$$\gamma_t = \frac{W}{V} = \frac{G_s + S_e}{1 + e} \gamma_w = \frac{1 + w}{1 + e} G_s \gamma_w$$

b. Unit weight of water = $\gamma_w = 62.4$ pcf.

c. Dry unit weight =
$$\gamma_d$$

The ratio of the weight of solids to the total volume.

$$\gamma_{d} = \frac{W_{s}}{V} = \frac{G_{s}}{1+e} \gamma_{w} = \frac{G_{s}\gamma_{w}}{1+\frac{wG_{s}}{s}} \frac{\gamma_{t}}{1+w}$$



Gw = Se

RELATIONSHIP AMONG SOIL PHASES

FIGURE 3-1



FIG. 3-2 - WEIGHT-VOLUME RELATIONSHIPS

Submerged or buoyant unit weight = γ_b d.

The buoyant unit weight is the effective weight when a soil mass is submerged in a body of water.

$$\gamma_b = \frac{G_s - 1 - e(1 - S)}{1 + e} \gamma_w$$

or when S = 100%

$$\gamma_{b} = \frac{G_{s}-1}{1+e} \gamma_{w}$$

and $\gamma_b = \gamma_{sat} - \gamma_w$

Β. CONSISTENCY AND MECHANICAL PROPERTIES

1. Gradation

> The distribution of the various particle sizes throughout a mass of soil is known as the grading of the soil.

The gradation of a soil is determined by a process termed mechanical analysis. This process includes the sieve analysis for larger particles and the sedimentation or hydrometer analysis for smaller particles.

The knowledge of the grain size distribution alone does not allow an accurate estimate of such properties as permeability, strength, and cohesion. Supplemental information including density and consistency is needed to permit better estimation of these soil properties.

The effects of gradation on soil behavior can best be illustrated by discussing some typical grain size distribution curves.

The three curves shown on Figure 3-3 represent specimens each having the same range of particle sizes but different gradations.

Curve A represents a specimen with an even distribution of particle sizes. Such a material is said to be well graded. A material with this gradation may have good resistance to erosion or scour, may be readily compacted to a dense condition, may develop high shearing resistance and bearing capacity, and have relatively low permeability and be slightly compressible.



Curve B represents a material that is primarily composed of particles of one size. This material is considered to be poorly graded. Such a material contains insufficient fines to fill the voids between the larger particles and consequently will have an open porous structure even with heavy compaction. As a result it will probably be more easily displaced under load, be highly permeable, and have less supporting power and lower shearing strength.

Curve C is said to be skip-graded. The characteristics of such a material are usually dominated by the finer faction, the coarse particles being inclusions in the fine matrix. The engineering behavior of such a soil defies prediction on the basis of gradation alone.

Many attempts have been made to describe soil gradation in mathematical terms. The most common used terms are those developed by Hazen:

Effective size = D_{10} and

Uniformity coefficient, $C_u = D_{60}/D_{10}$

In common usage in recent years is the coefficient of curvature

$$C_{c} = \frac{D_{30}^{2}}{D_{10}^{D}_{60}}$$

These relationships define the position and shape of the grain size distribution curve, not the soil particles.

2. Grain Size and Shape

The shape and size of individual soil particles have been found to be of considerable importance in explaining many phenomena of soil behavior.

With reference to shape, there are two major groups of soil particles:

- a. Those that are more or less equidimensional. These are described as being bulky in shape. The bulky grains are classified as rounded, sub-rounded, sub-angular or angular. See Fig. 2-4.
- b. Those that are platey or scale-like, many of which have a thickness less than 1/100th of their diameter.

This difference in shape reflects a difference in the crystal structure and characteristics of the mineral of

which the particle is composed. Primary minerals have no plane of marked weakness in any one direction. Consequently, particles of the primary minerals are generally bulky in shape. Secondary minerals have planes of weakness in at least one direction and are subject to what is known as basal cleavage. As a result, particles of the secondary minerals are of platey shape.

The mechanical processes of rock weathering normally does not break up or grind the hard equidimensional bulky mineral grains much smaller than 0.01 mm. As a consequence most sand, gravel and coarse silt size particles are of this shape.

On the other hand, the friable flake-shaped particles of secondary minerals, although initially very small, are readily ground, broken and chemically disintegrated into still smaller particles. Hence, the very fine fractions of natural soils consist principally of flake-like particles.

The size and shape of bulky grains have an important bearing on that part of the shearing strength which is influenced by internal friction. The more angular particles offering the higher frictional strengths.

The bulky grained soil normally has greater permeability, is less compressible, and is easier to compact to a high density.

The state of compaction of natural deposits of granular soils depends upon the size of their constituent particles. A deposit of fine sand may exist in a relatively loose condition while a gravel deposit is nearly always found to be in a comparatively dense condition.

It was originally proved by Atterberg that only particles of colloid size exhibit plasticity and only then if they were plate-like instead of bulky.

The flat, platey or scaley particles have lower frictional strength, greater cohesion, lower permeability and are more compressible.

Again it is emphasized that gradation, grain size, and shape when used alone to estimate soil behavior can give misleading results. However, an experienced soil engineer may obtain valuable indications of the properties of a soil from study of these characteristics along with the Atterberg limits, the in-place density, and the geologic origin.

3. Relative Density

Deposits of granular soils are found in nature in varying states of compaction from loose to dense. Generally, the engineering properties of these materials are largely controlled by their degree of compaction.

A dense soil consisting of well rounded sand particles essentially of one size may have a higher shear strength than a loose soil composed of a well graded mixture of angular sand and gravel particles.

When subjected to a static load, even relatively loose granular soils are virtually non-compressible. If vibrations or shear stresses are applied to a loose granular soil, substantial volume changes will occur. The reduction in volume of such soils can result in excessive differential movement and much damage to a structure founded on the material. This type of volume change generally occurs almost instantaneously and is much affected by the location and intensity of the applied dynamic load. These soils when under the water table and subjected to vibratory loads or shear stresses may have a considerable build up of excess pore pressure. In such cases the effective stresses may approach zero and liquifaction can occur.

The relative density of granular soils is determined by the following formulae:

In terms of the void ratios, e

$$D_{d} = \frac{e_{\max} - e}{e_{\max} - e_{\min}} (100)$$

where Dd

= relative density, expressed as a
 percentage

- e_{max} = void ratio in loosest state
- e_{min} = void ratio in densest state
- e = void ratio at state of compaction at which the relative density is to be expressed.

In terms of dry density

$$D_{d} = \frac{\frac{1}{\gamma_{\min}} - \frac{1}{\gamma}}{\frac{1}{\gamma_{\min}} - \frac{1}{\gamma_{\max}}} (100)$$

where Y_{min} = minimum dry unit weight of soil .
Y_{max} = maximum dry unit weight of soil
.
. Y = dry unit weight of soil at which the
the relative density is to be determined

Methods of determining the maximum and minimum void ratios and dry densities are discussed in Section 4.

4. Soil Consistency

a. General

The physical properties of fine grained soils differ greatly at different water contents. A given clay may act almost like a liquid, it may behave as a plastic solid, as a semi-solid or as a solid with limited elastic properties.

Consistency is defined as the degree of resistance of a fine-grained soil to flow or to deformation in general.

If the water content of a clay slurry is reduced by slow desiccation, the clay passes from a liquid state through a plastic state and finally into a solid state. The water contents at which different clays pass from one of these states into another are very different. Therefore, the water contents at these transitions can be used for identification and comparison of different clays. The transition from one state to another does not occur abruptly as soon as some critical water content is reached. It occurs gradually over a fairly large range of water contents. Consequently, all attempts to establish criteria for these boundaries have resulted in setting somewhat arbitrary limits.

b. Atterberg Limits

The consistency limits set up by Atterberg are the most widely applied in soil mechanics. The Atterberg limit tests are discussed in Section 4.

Consider a clay soil that is mixed with sufficient water to exhibit the properties of a true liquid. In this state the degree of saturation is much greater than 100%.

If this soil is allowed to dry out without disturbance, a point will be reached where the degree of saturation with respect to the volume of the soil-air-water mass is 100%. When this point is reached, the mass begins to behave as a viscous liquid. The moisture content at this change of state is defined as the flocculation limit. This limit has little practical significance. In the viscous liquid state, the soil mass will flow slowly under its own weight. At this point with further drying out, capillary forces are set up as the water starts to retreat into the pore spaces of the soil-air-water mass. These forces exert compressive forces on the soil grains and forces them closer together causing a decrease in volume of the mass. This is illustrated by Figure 3-4. The rate of drying out and the rate of volume change are such that the degree of saturation of the mass remains at 100%.



FIG. 3-4 - SHRINKAGE DIAGRAM FOR A COHESIVE SOIL

REFERENCE : ENGINEERING PROPERTIES OF SOIL, HOGENTOGLER, MC GRAW - HILL, 1937 As this drying out and shrinkage continues, a point is reached where the properties of the soil-air-water mass change from those of a viscous liquid to those of a plastic solid. The moisture content at this point is defined as the liquid limit. With additional drying out, the capillary forces become larger and force the soil particles closer together. With the change in volume, the degree of saturation is maintained at 100% and the percent moisture versus percent volume change plot traces the line shown on Fig. 3-4.

As the drying out and shrinkage continues, a point is reached where the properties of the soil-air-water mass change to those of a semi-solid. The moisture content at this point is defined as the plastic limit. The range of moisture contents between the liquid limit and the plastic limit is called the plasticity index. This index represents the range of moisture through which the soil behaves as a plastic solid. That is, the soil can be permanently deformed under load without rupture.

As additional drying occurs, a point is reached where the capillary forces cannot force the soil particles closer together. At this point no further decrease in volume occurs with additional drying out. The moisture content at this point is defined as the shrinkage limit. This is the lowest point where the degree of saturation is 100% in the drying out process.

Below the shrinkage limit, a cohesive soil has the properties of a solid and develops its maximum bearing capacity and strength. A soil containing relatively small quantities of inactive colloidal material may be weak and friable in this state. A soil containing chemically active colloids may develop very high strength and hardness.

The ratio of the volume change of soil to its loss in moisture is called the shrinkage ratio. This ratio is equal to the bulk specific gravity of the soil.

Nonplastic soils undergo the same general volume change pattern with drying out but they do not go through the plastic state or become a solid. The total volume change of nonplastic soils is very low compared to highly plastic soils.

c. Consistency of Undisturbed Soils

Fine grained soils exist in a wide range of states of consistency under natural conditions. The consistency of undisturbed soils is dependent on the mode of deposition and on other factors that have occurred in the geologic history of the deposit such as dessication, glacial loading, weathering and unloading of thick deposits of soil by erosion, etc.

The following nomenclature describing the consistency of undisturbed soils is widely used. This nomenclature is based on the undrained, unconfined compressive strength (q_u) .

Consistency	q _u (tons/sq.ft.)
Very soft	less than 0.25
Soft	0.25 - 0.5
Medium	0.5 - 1.0
Stiff	1.0 - 2.0
Very stiff	2.0 - 4.0
Hard	more than 4.0

The consistency of undisturbed soils based on simple qualitative estimates is described in item 7c of Section 2.

d. Sensitivity of cohesive soils

When remolded at the natural moisture content, most cohesive soils exhibit a considerable loss in strength and stiffness.

Present theory is that this loss in strength is probably due to the reorientation of soil particles into less favorable positions in the soil mass and to a reduction in bonding forces between particles.

Some clays when allowed to set after remolding without a change in moisture condition will regain some of the strength lost, this process is termed thixotropy.

The sensitivity of cohesive soils is represented by the ratio of the undrained, unconfined compressive strength before remolding to the undrained, unconfined compressive strength after remolding.

 $S_t = \frac{q_u \text{ (undisturbed soil)}}{q_u \text{ (remolded soil)}}$

where S_t is the sensitivity.

The following table is used to evaluate the sensitivity of soils.

Cla	SS	ifi	Lca	tio	n
-----	----	-----	-----	-----	---

St (sensitivity)

Insensitive	less than 2
Moderately sensitive	2-4
Sensitive	4-8
Very sensitive	8-16
Slightly quick	16-32
Medium quick	32-64
Quick	greater than 64

The variation in natural moisture content of cohesive soils is an effective indication of the homogeneity of the deposit. Cohesive soils with a natural moisture content near the plastic limit will generally be relatively stiff while those near the liquid limit may be soft or if stiff, very sensitive to remolding.

The liquidity index as proposed by Terzaghi is used to relate the natural moisture content to the Atterberg Limits.

$$LI = \frac{W - PL}{LL - PL}$$

where $L_T =$ liquidity index

w = natural moisture content

PL = plastic limit

LL = liquid limit

It is generally possible to relate the liquidity index of soils to soil properties such as shear strength. This permits qualitative evaluation of the more complex properties after the natural moisture content and Atterberg Limits have been determined.

C. PHYSIO-CHEMICAL RELATIONSHIPS

To understand the behavior characteristics of fine grained soils, it is necessary to consider the chemical effects and properties as well as the physical properties of the soil particles.

The first step in understanding clay behavior is to become acquainted with the clay minerals.

Since the first x-ray diffraction studies were made on clays in 1925 much research into the factors controlling the behavior of clays has been done.

This research has established that argillaceous (clay) materials are composed essentially of extremely small particles (of the order of one or two microns or less), of one or more members of a small group of minerals that have come to be known as the clay minerals.

Chemically, the clay minerals are hydrous silicates of aluminum or iron and magnesium. Some of them also contain alkalies or alkaline earths as essential components.

The clay minerals are dominantly crystalline, i.e., the atoms composing them are arranged in definite geometric patterns.

Most of the clay minerals have sheet or layered structures, some of the clay minerals have elongate tubular or fibrous structures. Thus agrillaceous materials can be considered as being essentially made up of extremely small particles, each one of which is either a book of sheet like units or a bundle of tubes or fibers.

Individual soils or clays may contain more than one kind of book like units or mixtures of books and bundles of tubes or fibers.

1. Structure and Composition of Clay Minerals

There are two fundamental building blocks for clay mineral structures.

a. The silica tetrahedral unit (Fig. 3-5a)

In this unit four oxygens or hydroxyls having the configuration of a tetrahedron, enclose a silicon atom.

The tetrahedra are combined in a sheet structure (Fig. 3-5b) so that the oxygens of the bases of all the tetrahedra are in a common plane and each oxygen belongs to two tetrahedra.



α.



○ & ○ = OXYGENS
■ = SILICONES

FIG. 3-5 - SILICA TETRAHEDRAL UNIT

The silica tetrahedral sheet alone may be viewed as a layer of silicon atoms between a layer of oxygens and a layer of hydroxyls (tips of the tetrahedra).

b. The octahedral unit (Fig. 3-6)

In this unit an aluminum, magnesium, or iron unit is enclosed in six hydroxyls having the configuration of an octahedron.

The octahedral units are joined together into a sheet structure (Fig. 3-6b) which may be viewed as two layers of packed hydroxyls with the aluminum, iron or magnesium cation between the sheets in octahedral coordination.





b.

۵.

O & C = HYDROXYLS

= ALUMINUMS, MAGNESIUMS, ETC.

FIG. 3-6 - OCTAHEDRAL UNIT

2. Shapes of Clay Minerals

The shapes of some of the common clay minerals are:

Kaolinite	•••	sheets (often regular hexagonal sheets
Halloysite	• •	tubes of curled sheets
Montmorillonite	• •	flaky sheets
Nontronite	• •	laths
Illite	• •	irregular flakey sheets
Attapulgite	• •	narrow fibers
Chlorite	• •	sheets
Vermiculite		sheets

3. <u>Kaolinite</u>

The kaolinite structural unit is an alumina octahedral layer with a parallel superimposed silica tetrahedral layer intergrown in such a way that the tips of the silica sheet and one of the layers of the octahedral unit form a common sheet.



FIG. 3-7 - BASIC KAOLINITE STRUCTURE

The kaolin unit may be viewed as a succession of layers of oxygens, silions, oxygens and hydroxyls, aluminiums, and hydroxyls. This unit is about 7A thick, and extends indefinitely in the other two directions, i.e. the flat dimension of the sheet. The kaolinite mineral is a stacking of such 7A thick sheets, the structure is like that of a book with each leaf 7A thick. Successive 7A thick layers are held together with hydrogen bonds. The mineral cleaves fairly easily along the plane surface of the 7A units.

Table 3-1 shows the average dimensions of the kaolinite mineral.

Normally there is no substitution of one cation for another within the kaolinite structure so that the mineral has the fixed formula

(OH)₈ Al₄ Si₄ O₁₀

In some clays, the kaolinite particles are composed of 7A thick units regularly stacked one above the others. In other clays, the stacking is somewhat random.

	TABLE 3-1 D	IMENSIONS OF TYPICAL CLAY PLA	TELETS	
	Characteristic	Approximate Ran Actual Dimensi	lge of one	Specific
	Ratios of	in Angstroms		Sullace (Sq. Meters
	Dimensions	Length and Breadth	Thickness	/gr.)
Montmorillonite	100 x 100 x 1	1,000 to 5,000	10 to 50	800
Illite	20 x 20 x 1	1,000 to 5,000	50 to 500	80
Kaolinite	10 x 10 x 1	1,000 to 20,000 (2u)	100 to 1000	10 .
	TABLE 3-	-2 CLASSIFICATION OF CLAY MIN	IERALS	
Clay Mineral Group		Chemical Formula	Mineral	l Sub-Group
Kaolinite	(0H) ₈ A1	4 Si4 0 ₁₀	Kaolini Dickite	i te
Montmorillonite	(OH)4 Al	4 Sig 020.nH20	Nacrité Swellir moril	ag Mont- llonite
			Non-Swe Montr Talc	arring morillonite
Illite (Hydrous Mica Halloysite*	a) (0H)4 Ky (0H)8 A1 (0H)8 A1	r(Al4.Fe4.Mg4.Mg6) (Si8-y.Aly) 4 Si4 OlO 4 Si4 OlO	Nontror 020 Illite Halloys	aite
* Halloysite occurs and the second slo	in two forms havi owly changes to th	ng the compositions shown. The first by dehydration. Part	the first of these interesting the second se	ls more stable, e clay minerals

.

4. Halloysite

Structurally, the halloysite mineral is similar to kaolinite as it is composed of sheets about 7A thick, each sheet being made up of one silica tetrahedral layer and one alumina octahedral layer. Halloysite differs from kaolinite in that successive 7A units are more randomly stacked one above the other and a single molecular layer of water may enter between the 7A units. With a molecular layer of water between each sheet, the mineral has the composition,

$$(OH)_8 Al_4 Si_4 O_{10} 4H_2O$$

Without the water layer, the composition is the same as kaolinite.

A further structural factor of halloysite revealed by electron micrographs is that the mineral occurs in elongated units that appear to be tubes, as if the sheet layers had rolled up.

5. Montmorillonite

This mineral is made of sheet-like units, each unit being composed of two silica tetrahedral sheets and one octahedral sheet. The octahedral sheet is between the silica sheets with the tips of the tetrahedra pointed toward the octahedral sheet.

The tips of each tetrahedral sheet and a hydroxyl layer of the octahedral sheet are intergrown to form a single layer.



FIG. 3-8 - BASIC MONTMORILLONITE STRUCTURE
The montmorillonite structure can be viewed as a sheet structure with each sheet composed of successive layers of oxygens, silicons, oxygens and hydroxyls, aluminum (or irons or magnesium), oxygens and hydroxyls, silicons and oxygens.

The thickness of each sheet is about 9.5A and the dimensions in the other two directions are indefinite.

The octahedral units may be aluminum, iron or magnesium, or a combination of these elements.

If aluminum is the sole occupant of the octahedral positions, only two-thirds of the possible positions are filled.

If magnesium is the sole occupant about all possible positions are filled. Also, a small amount (normally less than 15%) of aluminum may replace silicon in the tetrahedral layers.

Therefore, it is apparent that there is considerable possible element replacement within the structure of the montmorillonites. Of great importance is the fact that there are always replacements within the structures, and they always produce a net positive charge deficiency.

This deficiency is balanced by absorbed cations which are held on the outside of the sheets. These cations in general are readily exchangeable. While it is difficult to write a general formula, common compositions of montmorillonite are:

> (OH)₄ Si₈ (Al_{3.34} Mg₆₆) O₂₀ (OH)₄ (Si_{7.34} Al₆₆) Fe₄ O₂₀ (OH)₄ (Si_{7.34} Al₆₆) Mg₆ O₂₀

Special mineral names have been applied to varieties of montmorillonite with distinctive chemical compositions. Thus

<u>Saponite</u> is used when magnesium is about the sole octahedral cation.

Nontronite is used when iron fills these positions in the octahedron.

In the montmorillonite mineral the 9.5A thick sheets are stacked one above the other like leaves of a book. There is very little bonding between successive sheets and water may enter between the sheets causing the mineral to swell. Also about 80% of the absorbed balancing cations are between these sheets. The water layer that can exist between sheets may be about any thickness depending somewhat on the nature of the absorbed cations and the water available.

If sufficient water is available, the mineral can split into individual unit layers 9.5A thick.

The average range of sizes of the montmorillonites are shown in Table 3-1.

6. Illite

The structure of illite is similar to that of montmorillonite except that there is always a substantial (20%7) replacement of silicons by aluminums in the tetrahedral layers and potassiums are between the layers serving to balance the charges resulting from the replacement of the silicon with aluminum and to tie the sheet units together. The structure does not swell with the introduction of water between successive sheets.

The population of the octahedral positions can vary from one illite to another as can the amount of aluminum replacing silicon and the balancing potassiums.

Illite is similar to the micas in structure and composition but differ from them mainly in containing less potassium, slightly higher silicon to alumina molecular ratios and in being less well ordered.

A general formula is

 $(OH)_4 \text{ Ky}(Si_{8-y} Al_y) (Al_4 \cdot Mg_6 \cdot Mg_4 \cdot Fe_4) O_{20}$

D. SOIL STRUCTURE

The mechanical properties of the so-called "cohesive soils" have been dealt with as a subject of scientific research for more than 30 years.

In the early 1920's the general opinion concerning cohesion was that it was due to some undefined action of a hypothetical amorphous clay substance.

The first major contribution toward our present knowledge of clay behavior came in 1923 when Assar Hadding proved by x-ray analyses that clays had mainly a crystalline nature.

In 1925, Karl Terzaohi published a remarkably modern discussion of the structure and bonds in cohesive soils. He drew attention to the adhesion between particles and held that the cohesion in clays was due to this interparticle adhesion. He suggested that clay particles stick together at the points of contact with forces sufficiently strong for the building of a honeycomb structure. This permits comparatively large amounts of water to be enclosed within voids built up of aggregates of minerals glued to each other by the adhesion forces. Thus each cell in the honeycomb structure was supposed to be made up of numerous single mineral grains.

In 1926, V. M. Goldschmidt published his views and the results of his research on clays. He performed a series of experiments with mixtures of clay minerals and various liquids such as water, benzene, carbon tetrachloride, liquid sulphur dioxide and ammonia. In these experiments, he found that the clay minerals exhibited plasticity only when mixed with a liquid of polar nature (i.e. opposite electrical charges on each end of the molecule). He concluded that the clay properties were dependent on both the crystal chemistry of the mineral phase and the atomic structure of the liquid phase. His theory was that the clay properties were due to crystalline minerals surrounded by a film of absorbed water molecules and that the water molecules stuck to each other and to the minerals because of their dipole moments. His work led him to the opinion that the flaky minerals in highly sensitive clays are arranged in unstable cardhouse structures (Figure 3-9). Surplus water was encased in the space between a few mineral flakes leaning on each other and the difference in clays of high and low sensitivity was due to a denser arrangement of the minerals in the clays of low sensitivity.

In 1932, Arthur Casagrande presented a theory of honeycomb structure in sensitive soils very similar to Terzaghi's theory.

In 1953, T. Wm. Lambe published a paper indicating that for undisturbed marine clays an open structure similar to Goldschmidt's cardhouse exists, whereas for fresh water clays the structure was somewhat denser and for remolded clays, a high degree of parallelism with a short range order between flakes would exist (Figure 3-9). The work of Rosenquist and Bjerrum in 1955 and 1956 support the views of Lambe.

In 1957, T. K. Tan presented the view that a clay mineral network dominated by contacts between a corner of one mineral and the plane of another, would best describe the structure of clays. This is a refinement of the Goldschmidt-Lambe hypothesis.

The Work of Rosenquist et al of the Norwegian Geotechnical Institute in 1958 and 1959 with the electron microscope has conclusively proved the validity of the cardhouse concept of clay structure.

At present there are five stable units of soil structure. These are shown on Figure 3-10.



UNDISTURBED SALT WATER DEPOSITS



UNDISTURBED FRESH WATER DEPOSITS



CARDHOUSE STRUCTURE OF CLAY

3-24 FIG. 3-9

Figure 3-10a illustrates the only stable structural unit for clean coarse grained soils. In this unit the strength that results in the stability of these types of soils is mainly because of friction. (The so-called "macro-dilatency", which is caused by the lifting of one mineral grain on the one side of the shear plane when the shear movement takes place.) This strength is proportional to the effective normal stress.

Figure 3-10b illustrates the disperse colloidal structure. This dispersed or oriented system is the structure associated with the condition of the minimum free energy in a mass of colloidal particles. A soil in this structural condition is acted on by forces of attraction or by forces of repulsion and by external forces.

This type of soil structure is about the least stable of all the colloidal structures. Distinctive features of dispersed soil structure are the approximately parallel position of adjacent particles and the almost complete absence of contact between particles.

In these clays it is the liquid rather than the solid phase that is continuous. The strength and behavior of such clays depend largely on characteristics of the colloid system.

Figure 3-10c and 3-10d show the flocculated types of "cardhouse" structure. These types of soil structure are characterized by edge-to-surface contact of the plate like colloid particles. The two factors contributing to this arrangement are:

- The overall tendency of a mass of plate-like particles deposited at random would be to take up an irregular pattern rather than to fall into an oriented arrangement.
- The natural tendency for edge-to-surface contact is enhanced by electrostatic attraction between negatively charged surfaces and positively charged edges under some conditions.

The diamond structure tends to expand under the influence of internal forces (expansive system). This system is much more dependent on external forces for stability than is the triangular system (contractive system).

Figure 3-10e shows the loose granular structure stabilized by a colloidal matrix.

In this system the clay matrix acts as a void filler and thus resists the tendency of the larger grains to achieve an interlocking particle arrangement.



INTERLOCKED GRANULAR a.



b.

DIAMOND COLLOIDAL

C.



TRIANGULAR

d.



LOOSE GRANULAR STABILIZED BY COLLOIDAL MATRIX e.

Reference: Trollope and Chan EXTERNAL FORCES

FIVE STABLE UNITS OF SOIL STRUCTURE

FIGURE 3-10





Pr.