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LATERITE SOILS AND THEIR STABILIZATION

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Synopsis

Origin, occurrence, and correct identification of laterite rock and soil are briefly discussed. The peculiar properties of laterite soils are outlined and suggested methods of dealing with them are indicated.

Results are reported of experimental studies on four laterite soils and one intrazonal non-laterite soil, with special attention to their susceptibility to stabilization. The latter varied over a wide range. Of the stabilizers used, Portland cement gave best results in some cases and aniline-furfural (2:1) in others. The presence of organic matter seems to play a detrimental role in many cases.

The possible use of "oiled earth" properly fortified with anti-oxidants and bactericidal substances for low-cost roads in laterite areas is briefly discussed.

Of the important groups of the tropical and subtropical soils of the world, the laterite soils occupy a unique place, in regard to both their extensive occurrence and peculiar properties. They are widely distributed in such areas as India, Indonesia, Indo-China, Malaya, Burma, Western Australia, Madagascar, Central Africa, the Guianas, Brazil, and Cuba (Fig. 1). From a world-wide political and economical point of view¹ study of these soils is of vital interest because: (1) they normally possess good tilth and excellent drainability, with plenty of solar energy and water available; with an adequate supply of fertilizers they are capable of excellent yields and may well be destined to contribute in a major degree to the food supply of the world; and (2) a great need exists for a suitable network of low-cost roads in these areas already in their present under-developed condition and even more so if their proper agricultural development is to proceed.

From a purely scientific point of view, the peculiar engineering properties of laterite soils as extreme products of soil genesis call for an extensive investigation and possible elucidation, not only for their own sake but also for a better understanding of the properties of less extreme soil types. The present work is confined to the engineering characteristics of these soils, especially those of importance in low-cost road construction.

The development of the science of soil stabilization has given a scientific footing to an understanding, though as yet more or less qualitative, of soil behavior in highway and airport structures (31). Most of the available information, however, pertains to soils of the temperate zones. That soils of the different climatic zones vary extensively in their properties is well-known. The necessity for a possible special approach to the study of the tropical and subtropical soils has been emphasized by many authors (13, 27, 32, 33).

¹See Point IV of the Truman Program, and the British plans for assisting in the economic development of Southeastern Asia and of Africa.

PREVIOUS INVESTIGATIONS

Ever since the word laterite was introduced, first in geology and later

in pedology, there has been a sea of controversy over its correct nomenclature and identification. Laterite owes its origin to the Latin word *later* meaning "brick," and was first discussed in 1807 by Buchanan (3) purely on its field occurrence: "It is one of the most valuable materials for building. It is full of cavities and pores, and contains a large quantity of iron in the form of red and yellow ochres. In the mass, while excluded from the air, it is so soft that any iron instrument cuts it, and is dug up in square masses with a pick-ax, and immediately cut into the shape wanted with a trowel or a knife. It soon after becomes hard as brick, and resists the air and water much better than any brick I have known in India. . . The most proper name would be Laterite from *Lateritis*." ²

In 1911, Fermor (9), in a most explanatory paper, defines laterite as being formed by a process which causes the superficial decomposition of the parent rock, removal in solution of combined silica, lime, magnesia, soda and potash, and accumulation of hydrated iron, aluminum, titanium, and rarely, manganese. The latter were termed by him "lateritic constituents." A residual rock with 90 percent or more of lateritic constituents is termed a true laterite. This true laterite is to be distinguished from the lesser groups, viz., lithomargic laterites and lateritic lithomarges with 50 to 90 percent and 25 to 50 percent of lateritic constituents respectively. The controversy over the identification of bauxite as a variety of laterite was answered by Clark (7) in 1916 to the effect that there is no dividing line between bauxites and laterites - one shades into the other. Later,

²This term used by Buchanan, meaning literally "brick disease", throws a peculiar light on the problem whether "laterite" refers primarily to the use of the material in construction or to its brick-like appearance. In several Indian dialects laterite is called "brick-stone" (*itica cullu*), in the Tamil language it is called *shuri cull* or "itch-stone" because its surface appearance resembles certain cutaneous disorders. The Latin-derived English term happens to cover both appearance and function.

the term bauxite came to be used only for commercial aluminum ores. Campbell (4) in 1917 called a material laterite only if it contained uncombined alumina in the form of the hydroxide.

In 1926 Harrassowitz (14) first characterized laterites as red soils enriched on the surface with oxides of iron and aluminum. According to him the European red beds are of fossil laterite origin. Marbut's (21) "normal" laterites are those formed under the influence of good drainage free from the action of high ground water. He calls ground-water laterites the soils falling within Fermor's first group. Mohr (23) believes that the laterite crust is the result of eluviation followed by erosion. He mentions five stages in the full development of a typical laterite, viz., fresh ash soil, tarapan (juvenile soil), brownish yellow lixivium, red lixivium, and laterite. Periodic variation of ground-water level (descent and ascent) appears to be a most important factor in laterite formation.

LATERITIC SOILS

One of the important reasons for the confusion regarding the correct identification of laterites is the large area of the tropical and subtropical zones involved and the different degrees of laterization encountered in the various parts. Martin and Doyne (21) used the silica-alumina ratio as a classification criterion:

<u>Soil Type</u>	<u>SiO₂/Al₂O₃</u>
Laterite soil	1.33 or less
Lateritic soil	1.33 - 2.00
Non-lateritic soil	2.00 and over

The SiO₂/Al₂O₃ rather than the SiO₂/R₂O₃ ratio was employed for analytical reasons since iron may sometimes be in the form of ferrous oxide. This leads to the paradox that formalistically the presence of iron is non-essential for a soil to be termed laterite; this, of course, is in direct contradiction to the original definition of Buchanan and overlooks the important role which

iron oxides play in the rock named laterite by him. Moreover, the presence of iron in laterite soils is one of the most important factors that influence their engineering properties. It certainly is more appropriate to employ as classification criterion the silica sesquioxide ratio:

<u>Soil Type</u>	<u>SiO₂/R₂O₃</u>
Laterite soil	1.33 or less
Lateritic soil	1.33 - 2.00
Non-lateritic soil	2.00 and over

- (4) Non-accumulation of organic matter
- (5) Distinctive red color.

B. Characterization of the residual weathering product "laterite".

- (1) A tropical climate subject to alternations of dry or wet seasons or monsoons.
- (2) A level, or very gently sloping, elevated land surface which is not subject to appreciable mechanical erosion (abrasion by rain and wind)
- (3) The chemical and mineralogical com-



▨ Laterite and Lateritic Soils
(AFTER BOL'SHOI SOVETSKI ATLAS MIRA: JOFFE)

Figure 1. Distribution of Laterite and Lateritic Soils in the World.

It should be useful at this point to juxtapose the presently accepted characterization of lateritic and laterite soils and the conditions stipulated in the revised theory of C. S. Fox for the tropical residual weathering product "laterite" (12, 19).

A. Characterization of lateritic and laterite soils.

- (1) Disintegration and decomposition of the parent material in the direction of the end products of weathering.
- (2) Release and removal of silica.
- (3) Separation of sesquioxides and fixation in the profile.

position of the exposed rocks to be suitable for a supply of the lateritic constituents, alumina and ferric oxide.

- (4) The texture of the rock to be (or rapidly become during weathering) sufficiently porous for the entry of percolating water, so that the conditions for chemical action will be at a maximum.
- (5) The infiltrating water to remain in the interstices of the rock for long periods annually, i. e., during the wet monsoon, but eventually to drain away in the dry period thus giving maximum play to chemical erosion
- (6) The infiltrating water to contain

either an acid or alkaline substance with which to react on the rock components as well as to constitute an electrolyte and allow electro-kinetic phenomena to operate.

- (7) These annual processes to be in operation for at least a geological epoch of roughly a million years

This comparison shows that it is relatively easy for a rock to become a lateritic or laterite soil, but that the chances to become a "laterite" are severely restricted.

The general acceptance of absence of organic matter due to intense biologic and inorganic oxidation assumed for the red soils of tropical areas has been questioned by some authors who testify to high organic content of some of these soils. Joffe (19) states that a red soil with organic content either is an immature specimen of laterite or falls in a category between laterites and podsols. This explanation, however, hardly touches the nucleus of the problem or organic matter in laterite soils.

PROPERTIES OF LATERITE SOILS

As has been excellently dealt with elsewhere (13), laterite soils in their natural state are granular in structure and are possessed of low plasticity and excellent drainability. In this state they can carry heavy loadings. When remolded in the presence of water they often become clayey and plastic to the depth disturbed (34). An extraordinary influence of the change in the natural moisture content on the plasticity and density characteristics of a certain Hawaiian lateritic clay of volcanic origin has been reported; depending upon the natural moisture content, the plasticity index of this clay varied from 245 to zero (13). Thus, it is easy to see that the conventional subgrade soil test results may not at all reveal the true characteristics of members of the laterite soil group.

POSSIBLE METHODS OF APPROACH

Plasticity, density, and structural characteristics of soils are only the

external manifestations of the granulometry and the inherent affinity of the individual particles or aggregates to one another under different conditions of moisture and mechanical stress. Consequently, the nature, mechanical strength, and the permanency of cementation in these soils must be understood.

The granulating effect of organic material is well known. However, the resulting aggregations are usually soft and friable, rather than mechanically strong. It is felt that the organic matter in laterite soils may be more important for cementation because of its reducing effect giving the iron greater mobility than for its specific cementing or water-repelling power. Baver (1) makes the following statement concerning organic matter and cementation in laterite soils:

"The only group of soils in which a correlation has not been observed between organic matter and aggregation is the lateritic soils, where dehydrated oxides of alumina and iron are responsible for stable aggregate formation."

In connection with the problem of organic matter it is of interest that the previously discussed Hawaiian clay which exhibited such unique consistency properties contained about 20 percent of organic matter (15).

The existing situation indicates a great need for a fundamental study of the structure of laterite soils with special emphasis on the actual type and amount of organic matter present including the role of micro-organisms (1, 13, 28, 34). An integral part of such a study must be concerned with swell and shrinkage properties, with cracking patterns, as related to the degree of laterization and to the type and amount of organic matter, as well as with the effects of remolding (13, 27, 28) and of prevailing moisture conditions.

PROBLEMS IN SUBGRADE COMPACTION AND GRANULAR SOIL STABILIZATION

The engineering soil problems encountered in laterite areas as related

to the nature of the soil material as well as to the environmental conditions have been thoroughly studied and ably presented by Woollorton (33). He was the first to point out the importance of swelling and shrinking for tropical soils which by many "temperate-zone soil scientists" were believed to possess great volume constancy. His work combined with concepts normally employed in Portland cement concrete design led to Winterkorn's work on volume relationships in soil stabilization (27). Application of the volume principle leads, for granular stabilization of laterite soils, to clay contents greater than those given by the ASTM Standard specifications. Obviously, the latter represent a specific solution of the general problem; but being specific they are applicable only for the average illite clay soils and for the average climatic conditions of the United States.

Specifically, Woollorton (34) suggested: For no over-all swelling of a coarse granular system: Plastic Index \times Fines Content $>$ Volume of voids available between granular aggregates to accommodate swelling, or a modification thereof when some swelling is permissible. In place of the plastic index, shrinkage volumes may be employed as indicators of suitability of the binder portion, viz., (a) liquid limit-shrinkage limit, where exposure conditions permit slaking, or (b) field moisture equivalent-shrinkage limit where environmental conditions do not favor slaking.

It is necessary to caution here that the fines content as determined by the ordinary standard method of mechanical analysis for soils may be greatly misleading; well developed laterite soils with their granular structure are not easily dispersed by the common dispersing agents. Secondly, the consistency limits may show large variations in particular cases, depending upon the moisture content of the sample and the extent to which it was remolded during the tests. Special tests or modifications of the standard tests may have to be developed to serve as indices for the plasticity and swelling characteristics that form the basis for proper design.

With respect to compaction, both in the laboratory and in the field, the task is to search for and adopt suitable mixing and compaction methods that would not destroy the granular structure and yet yield sufficient density.

In the field extra care is needed if heavy machinery is employed. Experience has shown that primitive manual compaction of certain laterite soils has yielded better airfields than compaction of the same soils with standard heavy-compaction machinery and procedure.

So far consideration has been given mainly to the problems encountered in the engineering use of laterite soils. This might give the impression that all laterites and laterite soils give trouble. This is far from the truth. The physical and chemical properties of the group of materials encountered range from excellent to poor for engineering purposes. The problem is one of recognizing the good, eliminating the poor, and improving the intermediate ones. This is well stated in a recent paper by Christophe (6) treating specifically with the utilization of African laterites as road construction materials.

Many of Christophe's statements are so pertinent from a road building as well as from a scientific point of view that they are given in the following as abstracted from the original French.

The term laterite is applied to any rock colored red to dark maroon which is either in the hardening or in the decomposition stage as a result of environmental variations. Laterites range from friable soils to hard rock similarly as lime stone may range from marl to marble. The use of laterites must be preceded by a study of their properties.

The silty parts of the foundation soils must be eliminated. In granular soil stabilization higher amounts of lateritic clays than of other clays are allowed. The pH influences the clay content as determined by sedimentation analysis. The interpretation of sedimentation curves requires great care. The strength of compacted laterites may go up to 2,000 kg per sq. cm. and higher.

At the Ivory Coast a very excellent laterite rock is found of strength properties comparable to those of medium porphyry. In equatorial Africa and especially at L'Oubangui-Charu pudding stones predominate. In Togo amorphous laterites abound, in the Haute Volta strong lateritic gravels are found. In the Sudan, under the present dry climate, hills of laterite rock are eroded

to a considerable extent. In laterite soils one can always find more or less rock-like aggregations.

The affinity of hard laterite to asphaltic bitumen is good, similar to the affinity of bitumen to Basalt and lava, but the presence of silt hinders the proper contact between the hard laterite and bitumen. Laterite used in bituminous construction should not contain too much loam or clay. Design must be based on the results of preceding tests. Pougnaud, chief engineer of Abidjan, has treated laterite by impregnation and subsequent surface treatment. The oldest job is 13 years old and still in good condition. Treatment with cut-backs yields good results if it is thoroughly mixed with the soil. Improvement of granulometry is recommended to obtain greatest strength. This can be done by breaking amorphous laterite down to sand size. The expense of this is compensated by the resulting smaller bitumen requirements, for the base and surface treatment.

Laterite clays make excellent binders for stabilized soil roads. In the case of soils composed of pisoliths (lateritic gravel) and fines, compaction alone is sufficient, and results in bearing values of 60 or more. Hydrocarbon binders have good adhesion to laterite rocks. Excellent bituminous concrete has been made with such rock. Amorphous laterites should be comminuted and well-mixed in order to obtain a homogeneous material. Compact laterite rocks have sufficient strength to be used as a construction material for highways as broken stone, gravel, etc. However, it is indispensable to judiciously choose the borough pit.

The statements by Christophe obviously refer to actual "laterite" or material coming very close to it. The experiments described in the following deal with the more troublesome laterite - lateritic and non-lateritic soils occurring in tropical regions.

EXPERIMENTAL INVESTIGATION

Five soils, two laterite, two lateritic and one non-lateritic, were employed in the experimental investigation (Table 1). The latter was concerned with: (1) physical and physico-chemical tests and (2) stabilization tests with external additives.

Consistency limit and thermal analysis tests were made on all materials. The Matanzas soil was studied in greater detail, with respect to mechanical analysis, Proctor compaction, permeability, base exchange capacity, etc. Only a limited number of tests were performed on the other soils.

The stabilization tests consisted of preparation of 2-by 2-in. cylindrical

soil-stabilizer specimens of Proctor density, moist or dry curing for 7 days, exposure to four cycles of freezing and thawing³, four cycles of wetting and drying⁴, and 7 days immersion in water, respectively, with subsequent determination of the compressive strength in a Carver hydraulic press. The susceptibility to stabilization is rated on the basis of compressive strength values, combined with a qualitative judgment of the nature of failure etc.

STABILIZERS USED

The stabilizers used and the nomenclature employed are given in the relevant tables of results of the stabilization tests. Only Portland cement, MC-3 cut-back, and aniline-furfural (2:1) were used with the Havana and Catalina soils while these and a variety of other stabilizers were employed with the Matanzas soil. The tests with the Nipe soil were confined to the respective soil-cement system. The data reported for the Guinea soil were obtained during an investigation undertaken for Portuguese Guinea, which had as object the utilization of local low-cost vegetable materials in low-cost road and air-drome construction. The common stabilizers, Portland cement and asphaltic cut-backs, had been employed originally only for purposes of comparison; however, the failure of these materials to stabilize the soil in question prompted a search for the underlying reasons.

PREPARATION OF THE SPECIMENS

Soil (passing through No. 10 sieve) was mixed with predetermined quantities of water and stabilizer in a Hobart mixer and compacted in a Dietert machine. Solid stabilizers such as

³Freezing and thawing: 4 cycles, each cycle consisting of 8 hours of freezing and 15 hours of thawing.

⁴Wetting and drying: 4 cycles, each cycle consisting of 3 hours of wetting and 20 hours of drying.

TABLE 1
DESCRIPTION OF THE SOILS USED FOR EXPERIMENTAL INVESTIGATIONS

Soil Name Given	Description of the Soil	Profile Characteristics	Organic Matter Present	Parent Material	Clay Mineral from Thermal Analysis
Matanzas	Dark red, well granulated laterite soil from Cuba	A ₂ horizon 0 15 to 1 10 ft below ground level	Yes (rather high)	Lime Rock (coco')	Hydrated Halloysite with some gibbsite
Havana ^a	Ashy gray, clayey tropical soil from Cuba (Probably similar to the 'C' horizon soil of the Matanzas series)	B ₃ horizon 0 50 to 2 75 ft below ground level	Yes (rather low)	Lime Rock (coco')	Mixture of illite and montmorillonite
Catalina	Yellowish-red lateritic soil from Puerto Rico, not well granulated	B and C horizons 1-5 ft below ground level	Yes (rather low)	Andesitic tuff & tuffaceous shale	Kaolinite
Guinea	Light red, sandy lateritic soil from Guinea Africa	--	Yes (very high)	--	Kaolinite
Nape	Dark red, well granulated laterite soil from Puerto Rico	B ₂ horizon-- 2 to 5 ft below ground level	Yes (rather low)	Serpentine Rock	Kaolinite with gibbsite

^a Contains more than 50 percent CaCO₃

Portland cement and lime were first mixed dry with the soil; subsequently, water in an amount calculated to satisfy the moisture requirement of the soil and the hydration needs of the stabilizer was added. In the case of semi-solid and liquid stabilizers, the concept of optimum liquid content was used. The reported percentages of the stabilizers are based on the dry weight of the soil.

The preparation of soil-cement specimens at the standard optimum moisture content in the earlier attempts posed a problem due to the fact that the soil-water-cement system presented a rather sticky consistency, hardly conducive to obtaining well defined samples. A close observation of the mixing process showed the existence of a gradual change of the mix from the granular texture to a rather sticky consistency. This was much pronounced with an increase in the speed and time of mixing.

Best specimens were obtained by employing the lowest speed of the mixer and molding the specimens when the mix presented a very loose granular appearance with uniform moisture distribution.

Observations of the behavior of the laterite and lateritic soils in the preparation of specimens with a large number of stabilizers brought out the importance of avoiding over-mixing, not only for soil-cement, but also for the other systems. Excessive mixing destroys the desirable granular structure of the soil. However, with bituminous materials the dispersion was poor and did not improve significantly, even with continuous and intimate mixing.

The non-lateritic Havana soil alone showed a uniform mix with the bituminous stabilizer.

Aniline and furfural were added separately in the respective order to the soil-water system and the mixing continued after the addition of each.

TABLE 2

PHYSICAL CHARACTERISTICS OF THE MATANZAS SOIL

A Granulation Characteristics

1 Dry Sieve Analysis Characterizing Natural Granulations

Sieve No.	Percentage Passing (by weight of total)
4	77.1
10	59.1
40	23.95
200	2.50

2 Aggregate Analysis in an Elutriator

Size of Aggregates	Percentage by Weight
> 0.1 mm	84.45
0.1 - 0.05 mm	7.85
0.05 - 0.02 mm	3.73
< 0.02 mm	3.97

3 Mechanical Analysis on Soil Passing No. 200 Sieve, Sedimentation Method

Size and Description of Particles	Without Treatment	After Treatment	After Removal ^a
	With H ₂ O ₂	With H ₂ O ₂	of Fe and Al Oxides
Sand >50 μ	6.9	31.4	37.4
Coarse silt 5 μ - 50 μ	28.3	61.97	38.64
Fine silt 5 μ - 2 μ	12.8	3.17	15.30
Clay < 2 μ	51.9	3.46	9.59

B Consistency Limits

Description of test	Fresh Soil Moisture Content 20.5 Percent		Soil air-dried for a long time (Moisture Content 4.7 Percent)	
	Remolded	With Minimum Handling	Remolded	With Minimum Handling
Liquid limit	53.0	46.2	52.6	45.9
Plastic limit	31.2	31.2	32.1	32.1
Plasticity Index	21.8	15.0	20.5	13.8
Shrinkage Limit	20.8	31.5	20.32	26.8

C Other Tests

- Specific gravity

In Water	2.90	In tetrahydronaphthalene	2.94
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- Compaction tests

Proctor	30 %	Modified AASHO	25.5 %
Optimum moisture content			
Maximum dry density in lb per cu ft	88		95.25
- Permeability coefficient from consolidation test at 4.8 tons per sq ft
 3×10^{-3} cm per sec

^aJeffries "Rapid Method for the Removal of Free Iron Oxides in Soil Prior to Petrographic Analysis" Proc. Soil Science Society of America 211-212 (1946)

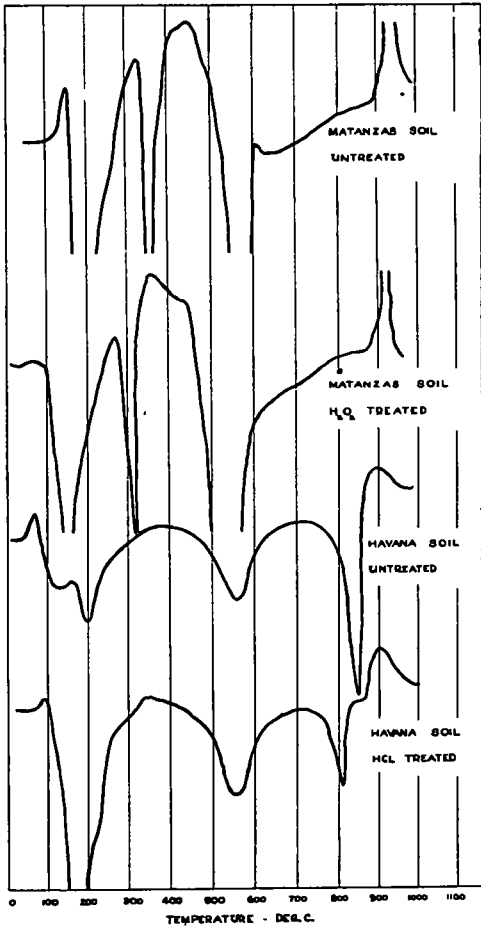


Figure 2. Differential Thermal Curves of -200 Fractions of the Soils.

CURING

All the Portland cement and lime specimens were moist cured. One-half of the number of specimens containing organic cementing or waterproofing agents were moist cured, the other half were dry cured: the same procedure was followed with the sodium-silicate specimens.

EXPOSURES

Separate sets of the lime and Portland-cement specimens were subjected to the three kinds of exposure tests mentioned previously. The only exception was the Guinea soil which failed

to set with moist curing for 7 and 14 days, respectively.

All the other specimens were immersed in water for 7 days before testing for compressive strength.

COMPRESSIVE STRENGTH TESTS

On completion of the exposure tests the compressive strength tests were performed at room temperature and the nature of failure noted.

PHYSICAL AND PHYSICO-CHEMICAL PROPERTIES

The results of the tests are given in Tables 2, 3, and 4.

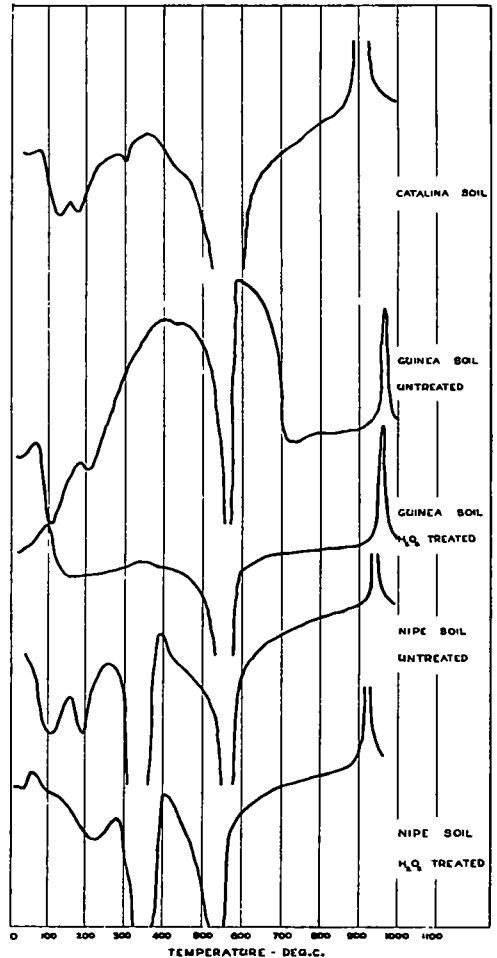


Figure 3. Differential Thermal Curves of -200 Fractions of the Soils.

MATANZAS SOIL

The results of the sieve analysis and aggregate analysis signify a very high degree of stable aggregation of the primary particles of the soil in its natural condition.

No proper dispersion could be obtained employing agents such as sodium silicate, sodium hydroxide, sodium carbonate, ammonium hydroxide, or a mixture of sodium hydroxide and sodium oxalate.

CONSISTENCY LIMITS

A study of the effect of change in the initial moisture content of the soil and

the effect of remolding on its consistency properties was made. The change in the initial moisture content did not affect the values to any significant extent. On the other hand, the role played by the structure is indicated clearly by the increase of the plasticity index by 40 to 50 percent in the case of the remolded soil, as compared with the value obtained when structural disturbance due to the test itself was held at a minimum.

The other tests conducted on this soil show a high specific gravity of 2.94, high permeability of 3×10^{-3} cm per sec., a low affinity for water, and a low base exchange capacity of 6.6 me per 100 grams. The thermal curve

TABLE 3

PHYSICO-CHEMICAL PROPERTIES OF THE MATANZAS SOIL

	3.3% H ₂ SO ₄	30% H ₂ SO ₄	50% H ₂ SO ₄
1 Hygroscopicity at 25 C	22.4	5.7	2.8
2 pH value of soil-water slurry (1 l)	6.30		
3 Base exchange capacity by conductometric titration	6.6 me 100 gms		
4 Carbon content by combustion analysis	2.0 Percent		
5 Clay mineral from thermal curve	Hydrated halloysite with some gibbsite		

TABLE 4

A. Physical and Physico-Chemical Characteristics of Havana, Catalina, Guinea and Nipe Soils

No	Name of Soil	L L	P L	P I	S L	Clay Mineral From Thermal Curve
1	Havana	62.2	36.7	25.5	26.5	Combination of illite and montmorillonite (non-lateritic)
2	Catalina	72.0	53.7	18.3	25.9	Kaolinite (lateritic)
3	Guinea	--	--	--	--	Kaolinite (lateritic)
4	Nipe	48.3	38.0	10.3	25.7	Kaolinite with some gibbsite (laterite)

B. Results of Additional Tests on the Guinea Soil

1	Wet Sieve Analysis	Passing Sieve No	Percent by Weight
		4	100
		10	95
		40	38
		200	12
2	Maximum dry density of Proctor Compaction	113.5 lb. per cu ft	
3	Optimum moisture	10.3 Percent	

shows presence of gibbsite in an essentially hydrated halloysite, the presence of gibbsite probably indicating a fairly high degree of laterization (Fig. 2, Table 3).

HAVANA, CATALINA, NIPE AND GUINEA SOILS

The limited number of tests performed on these soils shows reduced values of plasticity indices with higher degrees of laterization as seen from the thermal curves. The Guinea soil had only 12 percent passing the No. 200 sieve, and an optimum moisture content of 10.3 percent.

The thermal analysis shows the Havana soil to contain a combination of illite and montmorillonite clay minerals, which are typical for non-laterite soils. The Catalina and the Guinea soils show typical kaolinite clay minerals indicative of their lateritic character. The Nipe soil shows kaolinite with gibbsite, the latter an indicator of a higher degree of laterization (Figs. 2 and 3).

SUSCEPTIBILITY TO STABILIZATION WITH ADDITIVES

A number of investigators have reported good stability characteristics of systems of red and yellow soils stabilized with Portland cement in amounts as low as 6 to 10 percent (5, 30). Studies have revealed that soils with high iron and aluminum contents are easily stabilized by means of bituminous materials and by silicate of soda, the latter to be supplemented with a waterproofing agent (20, 29).

The results of the present series of stabilization tests are given in Tables 5, 6, 7, 8, and 9. In the following discussion, only the most significant features are pointed out.

MATANZAS SOIL (TABLE 5)

Fairly good compressive strength values were obtained at comparatively high percentages of cement. Steady and optimum strength conditions seemed to be reached around a cement content of

14 percent. Further increase to 16 and 18 percent did not improve the strength values appreciably. It is even possible that a lesser quantity of cement (12 to 14 percent) may show satisfactory stability characteristics.

Stabilization with lime was a total failure; in fact, decreasing values of compressive strength accompany increase in lime content. It is possible that for lime stabilization a longer period must be allowed for the carbonation of the calcium hydroxide. Furthermore, if a soil contains acid organic matter, lime may stimulate the growth of bacteria with detrimental action on mechanical resistance of the system.

The dry-cured specimens failed completely while the moist cured ones showed very poor compressive strengths. It is, of course, realized that field performance of a soil-bitumen system cannot be judged correctly from compressive strength values alone; however, such data are important for comparison of relative effectiveness. The complete failure in one case and the very poor strength values obtained in the other, show the ineffectiveness of the bituminous stabilizer used or the method adopted.

Of the dry cured specimens, those containing 10 percent of tar alone withstood the immersion exposure, but showed rather poor strength. The moist cured specimens showed fairly satisfactory values. Though in general, stabilization with tar alone may not be adequate, treatment with the latter resulted in distinctly better stability characteristics than treatment with corresponding quantities of MC-3 cut-back.

Both the dry cured and moist cured specimens showed satisfactory values, though a feeble tendency for slaking was noticed during immersion of the dry-cured specimens. This, however, did not significantly affect the strength characteristics. The poor results, amounting practically to total failures with the use of the other stabilizers listed in Table 5 do not warrant separate or detailed discussion. Noteworthy tendencies, if any, are brought

TABLE 5

RESULTS OF THE STABILIZATION TESTS WITH MATANZAS SOIL

Type of exposures		F-T	4 cycles of freezing and thawing		
		W-D	4 cycles of wetting and drying		
		IMM	7 days immersion in water		
Description of Soil-Stabilizer Specimens	Percent of Stabilizer	Compressive Strength in P S I	Remarks		
I Inorganic Stabilizers					
(a) Portland cement - moist cured					
	8	258	} Shear failures		
F-T	14	481			
	16	506			
	18	550			
	8	132	} Shear failures		
W-D	14	296			
	16	299			
	18	314			
	8	150	} Shear failures		
IMM	14	313			
	16	351			
	18	354			
(b) Lime - moist cured					
	8	30.0	} Crumbly shear failures		
F-T	14	15.7			
	16	12.0			
	18	8.0			
	8	--	} Slaked and failed on wetting for the second cycle		
W-D	14	--			
	16	--			
	18	--			
	8	40.0	} Crumbly shear failures		
IMM	14	52.0			
	16	31.0			
	18	41.0			
Note All compressive strength values listed in the following parts of Table 5 were obtained after 7 days immersion of the specimens in water					
(c) Silicate of soda (N-Brand 40 % Concentrated Solution) - dry cured-					
	4	--	} Stabilizer added directly to soil		
	4	--	} Stabilizer added in an aqueous medium		
Silicate of soda - moist cured					
	4	12	} Stabilizer added directly to soil		
	4	--	} Stabilizer added in an aqueous medium		

TABLE 5 (Cont'd)

Description of Soil-Stabilizer Specimens	Percent of Stabilizer	Compressive Strength in P S I.	Remarks
II Organic Stabilizers			
(a) MC-3 cut-back - dry cured	6	--	} Swelled, cracked and failed
	8	--	
	10	--	
MC-3 cut-back - moist cured	6	10	} After exposure all specimens were wet but did not show failure
	8	12 5	
	10	13	
(b) Tar RT-6 - dry cured	6	--	} Both 6 and 8 percent samples swelled, cracked and failed
	8	--	
	10	8	} Cracked slightly and showed some compressive strength
Tar RT-6 - moist cured	6	30	} Plastic failures
	8	33	
	10	33	
(c) Aniline-furfural (2 l) - dry cured	2	46.0	} Very feeble slaking was noticed, failed in crumbly shear
	3	54 0	
	4	43.0	
Aniline-furfural (2 l) - moist cured	2	46 0	} Crumbly shear
	3	52.0	
	4	70 0	
(d) Resins (abiectic acid) - dry cured	0.5	17.0	} Considerable slaking was noticed in all cases. The 1 percent specimens were of poor mold
	0.75	18 0	
	1.00	7 0	
Resins (abiectic acid) - moist cured	0.5	8	} Crumbled
	0.75	8	
	1.0	14	
(e) Abietic salt Resin 321 - dry cured	0.5	0	} All the specimens slaked considerably
	0.75	3	
	1.0	5	
Abietic salt Resin 321 - moist cured	0.5	11	} Crumbled
	0.75	11	
	1.0	10	

TABLE 5 (Cont'd)

Description of Soil-Stabilizer Specimens	Percent of Stabilizer	Compressive Strength in P.S.I	Remarks
III Bitumen and Admixtures			
(a) cut-back MC-3 + 10% gasoline -			
dry cured	8	-- }	Slaked and failed
moist cured	8	17 }	Plastic failure
(b) 8% cut-back MC-3 + 10% kerosene + 0.8% aniline-furfural (2 l) -			
dry cured	-	0	
moist cured	-	35 }	Crumbly shear failure
(c) 8% cut-back MC-3 + 10% gasoline + 0.24% aniline-furfural 2 l + 0.08% pentachlorophenol -			
dry cured	-	12 }	Specimens slaked considerably in both cases
moist cured	-	44 }	
(d) 8% tar RT-6 + 0.8% aniline-furfural 2 l -			
dry cured	-	0 }	Specimens slaked completely and failed
moist cured	-	30 }	Plastic failure

forward.

Silicate of soda was a complete failure in immersion for both dry and moist cured specimens. Both abietic acid and abietic salt (Resin 321) showed very poor strength values. The addition of gasoline as a dispersing aid for the MC-3 treatment gave no improvement. However, definite improvement resulted from the addition of a small percentage of aniline-furfural to the MC-3. Further addition of pentachlorophenol in the combination gives even better results. These observations are in line with previous findings on the influence of these activators. The addition of aniline-furfural to the RT-6, however, did not show any marked change in the strength values. In view of the excellent compatibility of aniline-furfural with coke-oven pitch, the observed behavior with the RT-6 may be due to the character of the cutting oil.

The results obtained with soil of the Matanzas series show a fairly good

susceptibility to stabilization with Portland cement, and a rather satisfactory one with aniline-furfural; the general results are, however, quite below the level that would normally be expected of this well developed laterite soil.

HAVANA SOIL (TABLE 6)

The specimens failed almost completely in the wetting and drying tests with Portland cement. In the other exposures, the specimens showed relatively poor strength values considering the high percentages of cement employed.

The stabilization with the MC-3 was a total failure for both moist cured and dry cured specimens. The fact that moist cured samples with 6 percent of cut-back were relatively more stable than those with 8 and 10 percent is noteworthy. The soil contained a considerable amount of lime.

Only the moist cured samples withstood the exposure tests, with aniline-furfural showing better strength values than the cut-back MC-3. Optimum performance possible with this resin was not achieved, however, since the calcium carbonate in the soil neutralized the acid catalyst and slowed down the setting and hardening time of the resin. This is illustrated by the fact that the lesser percentage of aniline-furfural, viz., 3 percent gave better results than 4 percent. In the meantime, methodology has been developed to successfully stabilize calcium carbonate-containing soils with aniline-furfural resin.

The Havana soil with its illitic and montmorillonitic clay mineral and high calcium content presents extreme

difficulty in stabilization with the conventional stabilizers.

CATALINA SOIL (TABLE 7)

The specimens failed completely in wetting and drying in the Portland cement test and gave extremely poor values in freezing and thawing. They did better, though still poorly, in the immersion test. It is significant that in the supposedly semi-rigid or rigid system of soil-cement, the specimens with as much as 16 percent of cement exhibited plastic failures. This brings home the fact, observed separately on pure soil-water specimens, that partially developed lateritic soils may have a greater water affinity and consequently more pronounced swelling

TABLE 6

RESULTS OF STABILIZATION TESTS WITH HAVANA SOIL

Description of Soil-Stabilizer Specimens	Percent of Stabilizer	Compressive Strength in P S I	Remarks	
I Portland cement - moist cured	8	114	Shear failures	
	F-T 12	131		
	16	164		
	W-D	8	0	Failed on second wetting
		12	0	Failed on second wetting
		16	70	Considerable slaking was noticed
	IMM	8	128	Shear failures
		12	175	
		16	182	
	II Cut-back MC-3 - dry cured	6	0	Swelled, cracked and failed within one day of immersion
		IMM 8	0	
		10	0	
Cut-back MC-3 - moist cured	6	5	The 6 percent samples were definitely better than the 8 and 10 percent ones by both appearance and strength	
	IMM 8	0		
	10	0		
III Aniline-furfural (2 1) - dry cured	3	0	Cracked and failed after 1 day	
	IMM 4	0		
	5	0		Cracked and failed after three days
Aniline-furfural (2 1) - moist cured	3	30	Plastic failure	
	IMM 4	19		
	5	20		Crumbly shear failure

TABLE 7
RESULTS OF STABILIZATION TESTS WITH CATALINA SOIL

Description of Soil-Stabilizer Specimens	Percent of Stabilizer	Compressive Strength in P S I.	Remarks			
I Portland cement - moist cured	F-T	8	0 } Plastic failures			
		12		3 }		
		16		25 }		
	W-D	8	0	} Slaked and failed on second wetting		
			12		0 }	
		16	0	} Slaked and failed on second wetting but less fast		
			0		} Slaked and failed on third wetting	
		IMM	8	31	} Plastic failures	
			12	30		
	16		80			
	II Cut-back MC-3 - dry cured	IMM	6	0	} Swelled, cracked and failed but retained the shape, showed no strength	
			8	0		
10			0			
Cut-back MC-3 - moist cured		6	33	} Plastic failures, cracks commenced with a pressure of 10 to 20 psi		
		8	33			
		10	40			
III Aniline-furfural (2 l) - dry cured	IMM	3	97	} Crumbly shear failures		
		4	68			
		5	143			
	Aniline-furfural (2 l) - moist cured	3	78		} Plastic failure	
		4	70			} Plastic to crumbly shear
		5	90			

and shrinkage characteristics than normally expected in view of the kaolinite type of clay contained in such soils.

The dry-cured specimens failed completely with MC-3 cut-back. The moist-cured specimens showed fairly good compressive strength for such plastic systems.

The results obtained with aniline-furfural are in sharp contrast to those obtained with Portland cement and MC-3. Excellent compressive strength was exhibited by both the dry- and moist-cured specimens. The dry-cured specimens, for the first time, showed better strength characteristics

than the moist cured ones for any soil with any organic stabilizer. The nature of failure of the moist-cured specimens was crumbly shear. The strength and water-resistance obtained with 3 percent of stabilizer were sufficiently good to suggest lowering of the stabilizer quantity to 2 percent. The good values obtained here have an added significance because of the poor showing with this soil of the other stabilizers, even at high percentages.

GUINEA SOIL (TABLE 8)

The size composition as well as the lateritic character of the soil under

investigation would indicate that it could be easily stabilized by means of any of the conventional materials, Portland cement, asphalt and tar.

The low cohesion of this soil makes it imperative that the organic stabilizers possess cementing in addition to waterproofing properties. The proto-

TABLE 8

RESULTS OF STABILIZATION TESTS ON GUINEA SOIL (23)

Percent of Portland Cement	Without additive	Compressive Strength of the Soil-Cement Specimens		
		10% CaCl ₂ ^a	2% Hydrous AlCl ₃ ^a	2% Hydrous Al ₂ (SO ₄) ₃ ^a
A' After 7 days curing				
4	22	--	--	--
6	40	6.4	0	19
8	25	9.1	0	12.7
10	29	12.7	0	19
B After 14 days curing				
6	--	--	9.6	27
8	--	--	8	24
10	--	--	12.8	30

^aPercentage based on weight of Portland cement

TABLE 9

RESULTS OF STABILIZATION TESTS ON NIPE SOIL

Description of Soil-Stabilizer Specimen	Moisture Content During Molding	Compressive Strength in P S I	Remarks
Portland cement used 8% - Moist cured			
F-T	23.4 ^a	27.5	
W-D	23.4	77	
IMM	23.4	53.3	
F-T	27.0	--	Samples were bruised during molding
W-D	27.0	77.0	
IMM	27.0	--	Samples were bruised during molding
F-T	30.0	33.3	
W-D	30.0	43.0	
IMM	30.0	43.3	

^aOptimum moisture content as of Proctor

The results obtained with Portland cement were quite disappointing because the cement failed to set and harden. This condition was not helped by the admixture of calcium chloride as suggested by the Portland Cement Association, nor by the admixture of aluminum chloride or aluminum sulfate.

type of a stabilized system to be achieved with this type of soil is "sand-bitumen" (8).

The real problem in the stabilization of this specific soil was the presence of organic matter ranging in size from easily recognizable fractions of peanut shells and roots to colloidal dimensions

and viable matter ranging from seeds to microbes.

The seeds sprouted lustily in the soil-cement specimens; the microbes proved to be especially active in the immersion test and provided visual and kakodylic evidence of their existence by reducing the red, iron-oxide color of the soil to a dirty bluish-green and by polluting the atmosphere. Subsequently, some oil-treated specimens were immersed in water contained in an iron bath well covered with rust. The microbes not only changed the color of the soil but, in addition, so thoroughly reduced and dissolved the rust of the bath that the latter showed a clean and shiny metal surface after the seven day immersion test.

The problem of the organic matter could be overcome by chemically sterilizing makro- and microbial agents present in the soil. The relative proportion of organic matter in the fraction passing the No. 200 sieve can be gleaned from a comparison of the thermal curves obtained on the unoxidized and oxidized soil fraction.

NIPE SOIL (TABLE 9)

The tests with this soil were concerned mainly with the influence of mixing moisture on the properties of soil-cement. The results are not conclusive; but there is a tendency for a decrease in strength with increase in moisture content at constant compactive effort. The data do not permit decision as to whether this apparent effect is a primary one of the moisture, or a secondary one either caused by easier destruction of structure during mixing at higher moisture contents or by a resulting change of the density of the specimens. See reference (30).

CONCLUSIONS

The work described and documented in this paper was concerned with a limited number of tests on a limited number of soil systems. Despite these limitations, the following conclusions of general importance could be drawn:

(1) The susceptibility to stabilization

of laterite and lateritic soils may vary over a wide range from excellent to poor. As a general rule, this susceptibility increases with increasing degree of laterization as evidenced by the silica-sesquioxide ratio. This confirms conclusions drawn from previous work performed on temperate zone soils in which iron and aluminum ions had been substituted for the exchange ions of the natural soils.

(2) In the absence of the analytical data required for calculation of the $\text{SiO}_2/\text{R}_2\text{O}_3$ ratio, clay-type determination by differential thermal analysis may be used as an index to the degree of laterization. Highly laterized samples contain aluminum and iron oxides, possessing typical thermal curves, in addition to the Kaolinite mineral which possesses a silica-sesquioxide ratio of 2.

(3) Soils possessing a low degree of laterization may possess more pronounced swell and shrinkage characteristics than normally expected of red lateritic clay soils.

(4) Fairly high quantities of organic matter may be present in red tropical and subtropical soils. This organic matter, especially if biologically active, may prevent a soil from properly responding to stabilizing treatment, even though the soil be granulometrically well suited for such treatment. In such cases, chemical sterilization preceding or simultaneous with the stabilization is indicated.

It may be appropriate to append a few considerations to this paper. Depending upon environment, parent rock and time, soil forming processes may, in the tropical and subtropical zones, produce materials ranging from hard laterite "rock" to non-laterite soils similar to those found in temperate regions.

From the point of view of road use, either as an aggregate, or as a binder, or as a material to be treated with cementing or waterproofing agents, the material becomes more favorable the farther it has progressed on the way to laterization. This indicates that the greatest problem in the road use of laterite soils arises from the effect on

the soil systems of the severity of climate, especially the desiccation followed by the monsoon, rather than from the inherent soil properties which at the worst come to resemble more and more those of soils from the temperate regions.

It stands to reason that granular soil stabilization has been more widely employed in laterite areas than the other stabilization methods. Granular stabilized soils suffer most from climatic severity especially from long desiccation periods. This is true not only for tropical climate but also for our own gravel roads in the great plains and for the more frigid climate of Argentine Patagonia, as judged from the wind erosion in these areas.

Considering these general facts, and also the specific properties of most soils in the tropic regions, i.e., cementation and structure which are best left undisturbed, it would appear that the most promising method for low-cost road construction is "earth oiling" as described in the Current Road Problems Bulletin on "Soil Bituminous Roads." Of course, the oil to be employed - and it may be any type of oil, asphaltic, pyrogenous from coal, vegetable or animal origin, or any other, must be well fortified with anti-oxidant and with bactericidal additives.

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