Utilization of Phosphoric Acid and Lime for Stabilizing Laterite for Lateritic Bricks Production

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Abstract: The use of innovative technologies in building construction using locally available materials can lead to affordable housing and boost the socio-economic development of a nation. Laterite soils were stabilized with varying percentages of 1M phosphoric acid (H₃PO₄) and 5% lime to ascertain its suitability as a sustainable material for hollow lateritic bricks production. The bricks were cured for 7, 14 and 28 days under ambient air condition. The compressive strength (f_c), bulk density (ρ_b), dry density (ρ_d), modulus of rupture (f_r). The results were a maximum f_c of 0.93 and 0.87 N/mm² at 5% lime and 4% H₃PO₄. Maximum ρ_b and ρ_d of 15.2 and 14.9 kN/m³ respectively were obtained at 4% H₃PO₄ stabilization. Maximum f_r of 0.2 N/mm² was obtained at combined 4% H₃PO₄ and 5% lime. In conclusion, the f_c suggests that both lime and H₃PO₄ have great potential in stabilizing laterite soils.

Keywords: Phosphoric acid; lateritic bricks; hydrated lime; compressive strength; modulus of rupture; dry density; pH.

Introduction

Bricks made from plain soils for centuries have been used in building mud house [1]. The most consistent of these soil types are laterites. Laterite soils which are highly weathered tropical and subtropical residual soils are old and popular construction material that has served humanity for centuries now.

In Nigeria and other tropical regions, lateritic soils are common and are used in construction such as production of bricks, sub base for road pavement among others. Heavy rainfall and elevated or warm temperature which are characteristics of tropical and subtropical regions, makes rock weathering intensive. This usually results in the rapid breakdown of feldspars and ferromagnesian minerals, the removal of silica and bases (Na₂O, K₂O, MgO) [2], and the concentration of iron and aluminum oxides [3]. This process of breakdown of feldspars and removal of silica is termed laterization [4] which also involves the leaching of SiO₂ and deposition of Fe₂O₃ and Al₂O₃.

Laterites, when used in making bricks, such bricks are called lateritic bricks and they have many attractive features, some of which are locally available and cheap materials, bricks of good esthetics and recyclability.

However, there can be a severe structural failure if the bricks are not properly strengthened, hence the need and emphasis for stabilization [5]. Soil stabilization is the process of improving the physical and engineering properties of a soil to obtain some predetermined targets [6]. Among the various soil improvement methods (mechanical, and chemical stabilization), the use of chemicals, especially lime and cement for soil stabilization to increase soil strength and to reduce settlement/compressibility, seems to be a more popular choice, probably because of its convenience.

Lime stabilization, as one of the commonest processes of improving the engineering properties of soils chemically, has been used extensively in the past and recently [2,7–9]. Lime is particularly suitable for fine grained soils. The reactions between lime and clay minerals present in soil will result in changes in the plasticity properties and soil structure; therefore, higher load capacity and mechanical resistance are induced [9]. The addition of excessive lime is not desirable, as it can lead to strength reduction, especially for silica rich soil. The reduction in strength can be attributable to excessive formation of silica gel, a highly porous material [9]. Thus, the amount of lime can vary from 5 to 12% by weight of the soil.

There has been a proliferation of literature regarding the application of non-calcium-based stabilizers, such as coconut husk ash [10], natural rubber latex [11], recycled sugarcane fiber waste [12], sawdust and egg shell ash [13], for soil stabilization. This is to replace cement, especially because its production has been associated with the depletion of nature’s non-

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renewable resources, consumption of substantial energy and contribution to carbon dioxide emission into the atmosphere [5]. Phosphoric acid (H₃PO₄), which is a mineral acid, non-toxic, that poses no threat to the environment, has been shown to be one of the alternative stabilizers especially for laterite soils [6,14–16]. As stated earlier, lateritic soil is highly weathered natural material formed by the concentration of hydrated oxides of iron and aluminium [4]. Phosphoric acid can effectively leach out alumina and other metallic oxides from clay minerals [6]. Thus, phosphate from phosphoric acid, can react with free iron and aluminium oxides present in the lateritic soil environment to form cementitious, highly insoluble aluminium and iron phosphate compounds such as strengite (AlPO₄·2H₂O), variscite (FePO₄·2H₂O) among others [17]. A typical reaction between phosphate with soil mineral is presented in Equation 1.

\[
\text{Al}^{3+} + 2\text{OH}^- + \text{H}_2\text{PO}_4^- + x\text{H}_2\text{O} \rightarrow \text{Al(OH)}_2\text{H}_2\text{PO}_4 \cdot x\text{H}_2\text{O} \tag{1}
\]

The product Al(OH)₂H₂PO₄ is hard and highly insoluble. A similar reaction exists between Fe₂O₃ (goethite), and the phosphate source. The product is Fe(OH)₂H₂PO₄·xH₂O, which is also hard and highly insoluble. It has, however, been shown that these precipitates can be formed in an acidic environment [6,15,16], fortunately, laterite and lateritic soils are acidic in nature [4]. Phosphate can also be precipitated in highly alkaline environment by calcium. According to Rooselers and Van Loosdrecht [18], the most stable calcium salt is calcium phosphate which is formed as presented in Equation 2.

\[
5\text{Ca}^{2+} + 3\text{PO}_4^{2-} + \text{OH}^- \rightarrow \text{Ca}_5(\text{PO}_4)_3\text{OH} \tag{2}
\]

Thus, if the pH of laterite soil can be increased in the presence of calcium and phosphate, cementitious compound of calcium phosphate could be formed. Baldovino et al. [9] stated that the addition of lime can cause a sudden increase in soil pH value due to the partial dissolution of calcium hydroxide Ca(OH)₂.

This research was, therefore, carried out to investigate the efficacy of the combination of lime and phosphoric acid on the stabilization of a laterite soil and to possibly ascertain the mix proportion required to produce durable and sustainable lateritic bricks.

Materials and Method

Materials

The following materials were used for this work: A suitable lateritic soil collected at the borrow pit opposite Odudua University, Ile-Ife, Nigeria. The lateritic soil was primarily identified by its physical characteristics (reddish colour, sticky texture). The longitude and latitude of the location from which the laterite was collected is 7°29'52.812" N, 4°26'57.546" E.

Phosphoric acid, also known as orthophosphoric acid (H₃PO₄), was prepared to a molarity of 1 mol/dm³ by dilution with distilled water. Lime (hydrated) was purchased from a dealer. It is recommended that potable water, i.e water safe for human consumption, should be used for batching. Pipe borne water hence was used for batching.

Method

Determination of the Physical Properties of Lateritic Soil

Some geotechnical properties such as specific gravity, Atterberg’s limits (Liquid limit, plastic limit and plasticity index), particle size distribution and compaction properties of the laterite soil were determined. The Atterberg’s limits and particle size distribution results were used for the classification of the soil.

Stabilization of Lateritic Soil for Brick Production

For the evaluation of the physical properties of lateritic bricks, the lateritic soil was sieved with a sieving machine with a 2 mm aperture. The bricks were produced in batches per stabilizer variable. Various percentages of H₃PO₄ up to 6% at 2% interval, 5% lime and a combination of H₃PO₄ and lime in their respective percentages, resulting in 7 stabilizer variables and 1 unstabilized control brick. The mixing was done using a mixing machine. These variations were made in order to evaluate how the proportions of the chemicals affect the brick’s physical properties. The bricks were produced using a hydraulic brick making machine of mold size 300 mm x 150 mm x 80 mm as shown in Figure 1.

![Figure 1. Brick with Dimensions (all dimensions in mm)](image)

Determination of Physical Properties of the Stabilized Bricks

Physical properties such as bulk density, dry density, water absorption, compressive strength, pH and modulus of rupture of the bricks were determined.
Both bulk and dry density of lateritic bricks are important parameters, by which bricks may be specified [5] and they simply indicate the bulk and dry mass of the brick per unit volume, respectively. The mass and volume of the bricks after curing were measured. The bulk density was determined by dividing the mass by volume. The volume is determined by mathematically calculating the volume of the shape dimensions of the brick.

A portion of the brick was obtained after crushing and oven dried to determine the water content. The dry density of the brick was determined using Equation 3.

\[ \rho_d = \frac{\rho_b}{1 + w} \]  

(3)

Where: \( \rho_d \) = brick dry density, \( \rho_b \) = brick’s bulk density, \( w \) = water content.

Water absorption tests on bricks are conducted to determine durability property of bricks such as degree of burning, quality and behavior of bricks in weathering. The degree of compactness of bricks can be obtained by water absorption test, as water is absorbed by pores in bricks. The specimen was air-dried and its weight obtained. The specimen was completely immersed in clean water at a temperature of 27±2°C for 24 hours. The specimen was removed and traces of water wiped with damp cloth and weighed after removal from water. The rate of water absorption of the bricks was then determined.

Compressive strength tests on bricks are carried out to determine the load carrying capacity of bricks under compression using compression testing machine. Whole bricks were taken and dimensions measured to the nearest 1mm. The specimen was placed with flat face horizontal and the load was applied at a uniform rate of 14 N/mm² (140 kg/cm²) per minute until failure occurs and the maximum load noted at failure. The load at failure is maximum load at which the specimen fails to produce any further increase in the indicator reading on the testing machine. The compressive strength of the brick specimens was determined.

The California Bearing Ratio (CBR) machine was used to determine the modulus of rupture \( (f_r) \) of the bricks in accordance with ASTM [19]. Each brick was placed over two supports with a clear distance of 274.6 mm between the supports. The distance between the end of the brick and each support was 12.7 mm. A plain steel bar was placed at the center of the brick to serve as point liner load and the plunger of the CBR machine was lowered down until it touched the bar before the machine was switched on. The load was applied at a loading rate of 50 kN and the load at which each brick failed in flexure was recorded along with the distance from the mid span to the plane of failure. The procedure was repeated for other brick samples. Two samples were tested for each stabilizer variation and the average result calculated, Equation 4 gives the necessary applied parameters.

The modulus of rupture \( (f_r) \) is calculated using Equation 4.

\[ f_r = \frac{3F(0.5L-x)}{bd^2} \]  

(4)

Where: \( F \) = Load at failure (N); \( L \) = length of brick (mm); \( x \) = Distance from the plane of fracture to the midpoint of the brick (mm); \( d \) = Net width of the sample less hole (mm) and \( b \) = Depth of the sample (mm).

Results and Discussion

Physical Properties and Classification of Lateritic Soil

The physical properties and classification of the lateritic soil are presented in Table 1. The Atterberg’s limits, i.e Liquid limit (LL), Plastic limit (PL) and Plasticity index (PI) of the natural soil sample are 51.35%, 39.26%, and 12.09%, respectively. The PI of the soil suggests that the soil is of medium plasticity. According to Das [20], soils with PI ranging from 1 to 5 are described as slightly plastic, soils with PI ranging from 5 to 10 are described as low plasticity and soils with PI ranging from 10 to 20 are described as medium plasticity.

Table 1. Results of Preliminary Tests

| Properties                  | Result   |
|-----------------------------|----------|
| Natural moisture content (%) | 28.54    |
| Specific gravity            | 2.82     |
| pH                          | 6.2      |
| Liquid limit (%)            | 51.35    |
| Plastic limit (%)           | 39.26    |
| Plasticity index (%)        | 12.09    |
| Percent of sand (%)         | 48.32    |
| Percent of silt and clay (%)| 51.68    |
| Soil classification (USCS)   | MH (High plasticity silt) |
| Soil classification (AASHTO)| A-7-5 (Clayey soil) |
| Optimum moisture content (%)| 17.30    |
| Maximum dry density (g/cm³) | 1.57     |

The selected lateritic soil sample was classified according to AASHTO [21] and USCS [22] classification systems using Atterberg’s limits and particle size distribution results. The particle size distribution results show that the percentage of soil passing sieve No. 200 is 51.68 (>36 min), hence the soil falls within the Silt-Clay materials. LL is 51.35% and PI is 12.09%, hence the soil falls in (A-7, A-7-5a, A-7-6b) group, but PI ≤ LL-30, therefore soil is A-7-5. In
addition to this, using USCS, more than 50% of soil sample passed sieve No. 200, which implies the soil is fine grained. Liquid limit is more than 50 and according to the USCS the soil is silt or clay. From the plasticity chart and considering the LL and PI the plot is below the “A” line, therefore soil is MH (High plasticity silt).

**Effect of Phosphoric Acid and Lime on the Bulk and Dry Density of Stabilized Lateritic Brick**

The results of the bulk and dry density for the bricks produced from laterite stabilized with different percentages of H₃PO₄ and lime after 7, 14, and 28 days curing are shown in Figures 2 and 3. It has been shown by Fadele and Ata [5] that density is part of the properties by which bricks may be specified. Bricks with density exceeding 1000 kg/m³ (9.81 kN/m³) may be classified as high density (HD) bricks. The values of the densities presented in Figures 2 and 3 indicate that the bricks are HD bricks. The values of density obtained in this study is slightly lower than that obtained by Fadele and Ata [5], this could be due to the method of compaction used. It was observed that there was a decrease in bulk density with the passage of time; this is largely due to the loss of moisture through the air-drying process. The bulk density of the bricks stabilized with the combination of lime and H₃PO₄ decreased with increasing H₃PO₄ and 5% lime at the early stages of curing. The 4% H₃PO₄ stabilized bricks gave the highest bulk and dry density of 15.2 kN/m³ and 14.9 kN/m³ respectively after 28 days curing which was followed by 6% H₃PO₄ stabilized bricks after 14 days curing. Under desirable conditions the optimum percentage of 4% diluted H₃PO₄ is optimum.

Statistical analysis of the results using 2-way Analysis of Variance (ANOVA) shows that neither the percentage of H₃PO₄ used nor the curing period had any significant effects on both the bulk and dry densities at 5% confidence level. However, with the addition of lime, the curing period (p = 0.035) had significant effect on both the dry and bulk densities with p < 0.05.

In order to determine the axial loading capacity of the bricks, the compressive test was used. The compressive strength of the bricks produced from laterite stabilized with variation of H₃PO₄ and lime after curing for 7, 14, and 28 days are presented in Figure 4. It shows that the compressive strength of the bricks increased with the increase in percentage of stabilization with lime and the combination of both lime and H₃PO₄. It was also observed that the compressive strength of the bricks increased with the curing age. According to ASTM [19], curing is important before testing to allow for considerable amount for cementation to take place. For bricks stabilized with only H₃PO₄ after 7 and 14 days, there was an increase in strength compared to the unstabilized control at 2% and 6%. Only at 4% was a decrease observed at both the 7th and 14th day curing. However, after 28 days curing it possessed the highest compressive strength of the H₃PO₄ stabilized brick. It was only next to (in terms of strength) bricks stabilized with 5% lime and the combination of 5% lime and 6% H₃PO₄.

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H₃PO₄ and 5% lime respectively. It has been shown by Sharma et al. in 2015 [23] that the compressive strength of bricks may increase up to the 28 days curing period and that the compressive strength at 56 days and 90 days curing period are the same with the 28 days curing period but are higher than the 7, 14, and 21 days curing period.

It was also observed that the 5% lime-stabilized bricks experience a slow and steady increase in strength within the 28 days of curing compared to the 2% and 6% H₃PO₄-stabilized bricks and bricks stabilized with both lime and H₃PO₄ with achieved a rapid increase in the first 14 days and a slow increase afterwards. It can therefore be deduced that H₃PO₄ as a stabilizer increases the compressive strength of bricks more rapidly than lime, however, lime-stabilized bricks possessed greater strength with time than H₃PO₄ stabilized bricks. The Nigerian Building and Road Research Institute recommends that the minimum percentage of H₃PO₄ required is therefore 4% and lime stabilization exceeded 0.2 N/mm² [24]. This requirement was not satisfied in any of the stabilizations, however, the compressive strength of the stabilized brick increased by approximately 75%, 75.4%, and 86% after stabilization and 28 days curing at 4% H₃PO₄, 6% H₃PO₄ and 5% lime, which gave compressive strength of 0.874 N/mm², 0.879 N/mm² and 0.932 N/mm² respectively.

Statistical analysis of the result using 2-way ANOVA shows that neither the curing period nor the percentage of H₃PO₄ had significant effect on the compressive strength of H₃PO₄ only stabilized bricks. However, when lime was added, both the percentage of H₃PO₄ and curing periods are significant factors affecting the compressive strength of the stabilized bricks with p = 0.039 and p = 0.0029, respectively.

**Effect of Phosphoric Acid and Lime on the Water Absorption of Stabilized Lateritic**

The water absorption test was carried out to determine the quantity of water absorbed by the brick when immersed in water for a specified time. It is used to test the durability of the brick under wet conditions. The bricks were tested at each curing period. The bricks specimens produced from laterite stabilized with H₃PO₄ and lime deteriorate after immersion in water for the specified duration of time. This can be attributed to the binding of the soil particles together brought by stabilization with H₃PO₄ which do not make the brick less permeable. The absorption of water leads to the dissociation of the bond holding the soil particles together. Lime-stabilized bricks exhibited a lower rate of water absorption but still crumbled. However, stabilized lateritic bricks lasted longer in water than the unstabilized lateritic bricks, it is suggested that firing the bricks could help in reducing the water absorption rate.

**Effect of Phosphoric Acid and Lime on the Modulus of Rupture of Stabilized Lateritic Brick**

The results of the modulus of rupture test for the bricks produced from laterite stabilized with different percentages of H₃PO₄ after 28 days curing are presented in Figure 5. According to Baldovino et al. in 2019 [9], modulus of rupture depends on the materials composition and dimensions. The results show that only the modulus of rupture at 4% H₃PO₄ and 5% lime stabilization exceeded 0.2 N/mm². A maximum value of 0.203 N/mm² was obtained as the modulus of rupture for bricks produced from laterite stabilized with lime at 5% stabilization. The result presented in Figure 5 shows that the control (unstabilized) brick has the lowest modulus of rupture followed by 6% H₃PO₄-stabilized brick. It was observed that there was a small decline in the modulus of rupture values for bricks stabilized with the combination of lime and H₃PO₄ as the H₃PO₄ increased. It is therefore not advisable to stabilize by combining them when a high modulus of rupture is expected. It was recommended that the minimum value for the modulus of rupture for adobe bricks to be 0.241 N/mm² [24]. This requirement was not satisfied in any of the stabilization variations; however, after stabilizing with 4% H₃PO₄ and 5% lime after 28 days curing, the modulus of rupture compared to the control increased by 115% and 117% respectively. This increment shows that H₃PO₄ and lime are potential stabilizers. The optimum percentages of stabilizer required is therefore 4% H₃PO₄ and 5% lime.
also a comparable compressive strength was achieved obtained apressive strength of the stabilized lateritic bricks was with silicates from the laterite. The highest com
high pH (9), the calcium from lime probably reacted and not calcium phosphate within the laterite. For only to formati
of the laterite because of the low pH. As such, any bricks in the present study were probably leached out and iron oxides (as well as silicates) in the lateritic rich in iron and aluminium oxides cementation to occur. As earlier mentioned, laterite is Rogers pH must be high alkaline, infact, Glendinning and that for calcium to beneficially react with laterite t
ions were injected into the laterite. Both studies show compounds (and not calcium phosphates) could be also showed that at pH below 7, only aluminium on and iron phosphates cementitious compounds (and not calcium phosphates) could be formed in laterite when both calcium and phosphate ions were injected into the laterite. Both studies show that for calcium to beneficially react with laterite the pH must be high alkaline, infact, Glendinning and Rogers [26] specified a pH of greater than 12 for cementation to occur. As earlier mentioned, laterite is rich in iron and aluminium oxides [4]. The aluminium and iron oxides (as well as silicates) in the lateritic bricks in the present study were probably leached out of the laterite because of the low pH. As such, any improvement in compressive strength might be due only to formation of aluminium and iron phosphates and not calcium phosphate within the laterite. For high pH (9), the calcium from lime probably reacted with silicates from the laterite. The highest compressive strength of the stabilized laterite bricks was obtained at 5% lime stabilization at a pH value of 9.4, also a comparable compressive strength was achieved at a lower pH value of 5.2 at 4% stabilization with H$_3$PO$_4$. The combination of lime and H$_3$PO$_4$ neutralizes the alkalinity and reduces the pH hence the reason for the low strength of lime and H$_3$PO$_4$ stabilized laterite bricks. It might, thus, be necessary to increase the pH of lime and H$_3$PO$_4$ stabilized laterite bricks by alkaline activation if calcium will beneficially react with silicate and/or phosphate ions from H$_3$PO$_4$.

**Figure 5.** Modulus of Rupture Stabilized Lateritic Bricks

**Effect of Phosphoric Acid and Lime on the pH of Stabilized Lateritic Brick**

The pH of the control brick was 6.3 which is almost neutral. It was observed that an increase in the proportion of H$_3$PO$_4$ stabilization resulted in a decrease in the pH value down to 5.0 at 6% H$_3$PO$_4$ (as presented in Figure 6), which implies that the bricks became more acidic. At 5% lime stabilization, the brick’s pH value was 9.4 and alkaline. However, the pH of bricks stabilized with the combination of 5% lime and varying percentage of H$_3$PO$_4$ decreased (becoming less alkaline) with increasing pH value as shown in Figure 6, 5% lime still having the dominant effect. The effect of 5% lime was even more dominant on the pH value even when combined with 6% H$_3$PO$_4$ because H$_3$PO$_4$ is a weak acid and of low concentration. It has been shown that pH has effect on the formation of cementitious compounds which can lead to improved geotechnical properties of laterites. It was shown by Ghobadi et al. [25] that shear strength parameters for clays increased considerably if the pore fluid pH is as high as 9 or as low as 3. Ayodele and Agbede [16] also showed that at pH below 7, only aluminium or and iron phosphates cementitious compounds (and not calcium phosphates) could be formed in laterite when both calcium and phosphate ions were injected into the laterite. Both studies show that for calcium to beneficially react with laterite the pH must be high alkaline, infact, Glendinning and Rogers [26] specified a pH of greater than 12 for cementation to occur. As earlier mentioned, laterite is rich in iron and aluminium oxides [4]. The aluminium and iron oxides (as well as silicates) in the lateritic bricks in the present study were probably leached out of the laterite because of the low pH. As such, any improvement in compressive strength might be due only to formation of aluminium and iron phosphates and not calcium phosphate within the laterite. For high pH (9), the calcium from lime probably reacted with silicates from the laterite. The highest compressive strength of the stabilized laterite bricks was obtained at 5% lime stabilization at a pH value of 9.4, also a comparable compressive strength was achieved at a lower pH value of 5.2 at 4% stabilization with H$_3$PO$_4$. The combination of lime and H$_3$PO$_4$ neutralizes the alkalinity and reduces the pH hence the reason for the low strength of lime and H$_3$PO$_4$ stabilized laterite bricks. It might, thus, be necessary to increase the pH of lime and H$_3$PO$_4$ stabilized laterite bricks by alkaline activation if calcium will beneficially react with silicate and/or phosphate ions from H$_3$PO$_4$.

**Figure 6.** The pH of Stabilized Lateritic Bricks

**Determination of the Optimum Percentage of Stabilizer Required**

The optimum percentage of stabilizer required is the stabilizer proportion that best enhance the engineering properties of the lateritic brick. From the tests carried out, the highest compressive strength of 0.93 N/mm$^2$ and 0.87 N/mm$^2$ was obtained at 5% lime stabilization and at 4% H$_3$PO$_4$ stabilization respectively. The best modulus of rupture of 0.204 N/mm$^2$ and 0.202 N/mm$^2$ was also obtained at 4% H$_3$PO$_4$ stabilization and 5% lime stabilization respectively. The best bulk and dry density result of 15.21 kN/m$^3$ and 14.94 kN/m$^3$ respectively was obtained at 4% H$_3$PO$_4$ stabilization, while none of the stabilizations passed the water absorption test. Also, stabilizations done by combining lime and H$_3$PO$_4$ did not prove to enhance the physical properties better than 4% H$_3$PO$_4$ or 5% lime stabilization. In conclusion, under desirable conditions the optimum percentage of 4% 1M H$_3$PO$_4$ or 5% lime can be used for stabilization of lateritic hollow bricks.

**CONCLUSION**

Phosphoric acid and lime both improved the compressive strength of the bricks. The impact of H$_3$PO$_4$ being rapid and minimal in the first 14 days curing, while that of lime was slow but greater. At 28 days of curing, 4% H$_3$PO$_4$ of 1 molar concentration gave a significant increase in compressive strength compared to the unstabilized control specimen. It has a value comparable to the 5% lime-stabilized specimen, which possessed the highest compressive strength. However,
bulk density of lime stabilized specimen and specimen stabilized with the combination of lime and H₃PO₄ decreases with the passage of time while H₃PO₄ on the other hand increased. The combination of lime and H₃PO₄ as a stabilizer gave lesser compressive strength compared to lime (only) stabilized bricks, this is due to the decrease in the alkalinity, as pH value has been shown to have impact on the strength. Hence, alkaline activation will be necessary if H₃PO₄ and lime are to be combined when stabilizing lateritic bricks. The 4% H₃PO₄ stabilized specimen gave the closest and optimum results from all the tests; hence it is the optimum stabilizer proportion.

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