Effect of Phosphate Mine Tailings from Morocco on the Mechanical Properties of Ceramic Tiles

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Abstract—The objective of this work is to study a way of recovering phosphate mine tailings from the exploitation of phosphates in the manufacture of ceramic faience products. Five formulations are studied, the dry pressing process is used to make the tiles. The formulations studied are produced by mixing the waste rock with abundant local clay in percentages which vary from 0 to 100%. The materials produced are characterized by their physical properties (water absorption, shrinkage), mechanical properties (resistance to bending) and surface properties (observation by scanning electron microscopy).

The results show that the integration of phosphate mine tailings improves the mechanical properties of ceramic tiles but increases the percentage of fire shrinkage and water absorption. Nevertheless, industrial use of the waste studied with the percentages 20 and 50% in clay-based formulas could be envisaged to design ceramic faience products, in accordance with the applicable mechanical and technical requirements. This recovery method makes it possible to manage the large quantities of mine discharges and therefore their environmental impact as well as to preserve non-renewable natural resources.

Keywords: Moroccan phosphate, mine tailings, recovery, ceramic industry, mechanical properties.

I. INTRODUCTION

Taking advantage of a favorable geological context, the mining industry in Morocco constitutes an important vector of economic and social development. Mining is characterized by the preponderance of phosphate, of which Morocco is leading producer, with three-quarters of the world's reserves. The phosphate deposits are found in four zones: the Khouribga zone (Plateau of OuledAbdoun), the Maskala basin (Marrakech), the Benguerir and Youssoufia region (the Gantour basin) and the Layoune-BouCraa zone [1, 2, 3]. The OuledAbdoun basin of Khouribga is among the sediment richest in phosphates. It contains 44% of the country's reserves, or around 26.8 billion tonnes of phosphate [4-7]. Following operations such as moving the dead ground and the intermediate layers very poor in P₂O₅, mining inevitably results in mine tailings. The latter is dumped in the mine area, so as not to interfere with the mining activity or generate a risk. In the absence of a sustainable management strategy, the waste rock could have direct impacts on the natural environment and on the health of the neighboring population. Pollutants can be leached into groundwater following episodes of intense rain [3, 8]. Several solutions for the technological development of the phosphate mining waste rock are proposed in the literature (development material, filling of open voids, substrate for revegetation on mining sites, raw material for cement and concrete), but they are prove to be inappropriate and unprofitable in the current context [3, 8, 9, 10].

This work therefore offers an easy, economical and ecological solution for the manufacture of ceramic tiles from the waste rock from phosphate mines. Our strategy aims to partially replace the clays used in the manufacture of tiles by seeking the composition that generates the products with the best physico-chemical and mechanical properties. This manufacturing route then preserves non-renewable natural resources.

II. MATERIALS AND METHODS

A. Preparation of mixtures

The mine tailings of phosphate (S), which is the subject of recovery in this study, is a waste resulting from the exploitation of phosphate by the Office Chérifien du Phosphate (OCP) in the Béni-Idir area (the area of Khouribga in Morocco) (Fig 1).
For the preparation of mixtures, for the manufacture of ceramics, the mine tailings of phosphate was used in percentages ranging from 0 to 100% mixed with abundant clay from the El Gara quarry in Morocco. Five different formulations are thus prepared and studied (Table 1).

### Table 1. The Different Formulas Studied.

| Formulas | Clay (A) | Phosphate waste (S) |
|----------|----------|---------------------|
| F1       | 100      | 0                   |
| F2       | 95       | 5                   |
| F3       | 80       | 20                  |
| F4       | 50       | 50                  |
| F5       | 0        | 100                 |

### B. Tile manufacturing process

In a porcelain ball-grinding jar, with a capacity of 500 grams, the mixture is kneaded for 25 minutes, with approximately 4% water and 1% of deflocculant (Fluicer®). The slurry thus produced is placed in an industrial dryer for 30 minutes at 110 ° C until completely dry. The product obtained is ground, sieved, then rewetted by spraying. The wet powder thus obtained is introduced into a manual tableting press. Under a constraint of 200 bars, several pellets are generated (ten for each formula). These are dried at 170 ° C for 25 minutes in an industrial roller drier. This achieves a residual water content of less than 1%, a favorable condition to avoid cracks during cooking. A final stage of cooking is carried out, it lasts approximately 40 minutes, during this stage the temperature gradually reaches 1200 ° C.

### C. Chemical and particle size characterisation

The chemical analysis for the determination of the oxides is carried out by X-ray fluorescence spectrophotometry using a Tiger S8® Bruker spectrometer [11]. The particle size analysis is carried out by dry sieving for the fraction sandy and sedimentation for silty and clay fractions [12, 13]

### D. Mechanical characterization of the manufactured tiles

The mechanical resistance of the manufactured tiles is made by three-point bending tests by a Shimadzu® single-column tensile testing machine, EZ LX series (7 tests for each formula) [14, 15].

### E. Microscopic observation of tiles

The surface analysis was carried out by SEM scanning electron microscopy observations. This analysis makes it possible to assess the granular compactness and cohesion. Microscopic observations are made by a Hirox SH-4000M device after metalization of the surfaces.

### III. RESULTS AND DISCUSSION

#### A. Chemical analysis

Table 2 presents the results concerning the chemical characterization by X-ray fluorescence spectrophotometry of clay A and mining waste S.

### Table 2. Chemical Compositions by X-ray Fluorescence (in mass %) of Clays A and Mining Waste S.

|        | A  | S  |
|--------|----|----|
| SiO₂   | 64.00 | 7.41 |
| Al₂O₃  | 15.15 | 0.71 |
| Fe₂O₃  | 5.23  | 0.32 |
| CaO    | 2.23  | 46.45 |
| MgO    | 1.70  | 1.25 |
| K₂O    | 4.28  | 0.14 |
| Na₂O   | 0.80  | 0.57 |
| P₂O₅   | 0.10  | 24.13 |
| SO₃    | 0.12  | 1.06 |
| TiO₂   | 0.65  | 0.10 |
| MnO    | 0.09  | 0.02 |
| BaO    | 0.18  | 0.02 |
| FL*    | 6.27  | 17.06 |

* Fire loss at 1140°C.

The chemical analysis (Table II) of the mining waste shows high CaO (46.45%) and P₂O₅ (24.13%) contents, against a low percentage of silica SiO₂ (7.41%) and alumina Al₂O₃ (0.71%) The high CaO content reflects the carbonate nature of the waste, confirmed by a considerable fire loss (17.06%). According to the literature, the presence of carbonates stabilizes the acid-base environment and can lead to the formation of refractory monolith, essential for thermal resistance [3]. On the other hand, the presence of alumina in clay materials, generally helps to have a low shrinkage during drying, or even during baking, and allows the manufacture of refractory materials after the passage at high temperature. It is therefore essential to use the sterile mixture with clays rich in alumina to have a balanced clay mixture.
A relatively high value in SO$_3$ is noted for the phosphate mining waste, the presence of this element reflects the richness of the environment in plastic materials [17], which plays an important role in improving plasticity [18].

**B. Particle size analysis**

The particle size analysis makes it possible to define various classes of materials regardless of their chemical composition. Table 3 presents the particle size distributions of components S, and A.

![Image](F1.png)
![Image](F2.png)
![Image](F3.png)
![Image](F4.png)
![Image](F5.png)

**Fig 2.** Photo of the materials produced

**TABLE 3. GRANULOMETRY OF THE PHOSPHATE MINING WASTE (S) AND THE CLAY (A).**

|       | Sand 50-2000µm | Silt 2-50 µm | Clay <2µm |
|-------|----------------|--------------|-----------|
| **S** | 8%             | 17%          | 75%       |
| **A** | 28%            | 18%          | 54%       |

The particle size analysis (Table 3 and Fig 3.) allows to classify the mining waste and clay A, among the category of fine clays according to the conventional classification of the United States Department of Agriculture [12-13]. This type of soil is characterized by strong plasticity in the wet state or compactness in the dry state.

![Image](Fig 3.png)

**Fig 3.** Classification of the components A and S according to USDA soil texture triangle [12].

**Note:** Henceforth the appellations F1, F2, F3, F4 and F5 will designate the tiles produced according to the corresponding formulations.

**C. Firing shrinkage and water absorption of ceramic tiles**

Ceramic materials have the particularity of undergoing volume variations when they are found in dry or humid environments. Depending on their structure, these modifications are more or less important. Indeed, large increases in volume can be observed (swelling phenomenon) when the water content increases, and conversely, a retraction can occur (shrinking phenomenon). On an industrial scale, this property has a considerable impact on the progress of the glazing process and on the technical adaptation of the material to changes in the surrounding climate.

In addition, ceramic tiles are classified in the beginning according to several groups. The mention of these is based on two criteria: the manufacturing method and the percentage of water absorption.

Starting from the method of manufacturing the tiles, two methods are distinguished: Method A- tiles made by extrusion and Method B- tiles made by dry pressing. We retain for our case the letter "B", to designate that these are tiles manufactured by dry pressing. In the same way, according to the percentage of absorption, ceramic tiles are classified according to three groups:

- **Group I** - Tiles with low absorption percentage (≤ 3%). It includes for tiles produced by dry pressing: The B1a group (absorption percentage ≤ 0.5%) and the B1b group (absorption percentage> 0.5% and ≤ 3%)
- **Group II** - Tiles with medium water absorption percentage (> 3% and ≤ 10%). It includes for the tiles produced by dry pressing: The BIIa group (absorption percentage> 3% and ≤ 6%) and the BIIb group (absorption percentage> 6% and ≤ 10%)
Group III - This group includes tiles with a high percentage of absorption (> 10%).

On the other hand, during firing, a ceramic piece undergoes a shrinkage expressed by a percentage (%). Two types of shrinkage are evaluated: a mass shrinkage (a mass loss is caused by the release of certain volatile components); a diameter shrinkage (a loss in diameter caused by fusion of components, which tend to fill the voids existing between the particles).

| Table 4. Assessment of Shrinkage and Water Absorption for the Manufactured Ceramic Materials |
|------------------------------------------|------------------|------------------|------------------|
| Mass loss (%) | Diameter loss (%) | Ratio loss (%) | Water absorption (%) |
| F1 | 6.7±0.63 | 1.97±0.24 | 3.40 | 2.3±0.15 |
| F2 | 7.96±1.69 | 2.55±0.77 | 3.12 | 6.4±1.73 |
| F3 | 8.84±1.07 | 2.08±0.34 | 4.25 | 10.3±1.77 |
| F4 | 10.7±0.76 | 1.72±0.17 | 6.22 | 14.5±1.47 |
| F5 | 14±1.03 | 4.36±0.10 | 3.19 | 18.5±2.36 |

According to table 4, only F5 has both considerably high Mass and diameter loss. F3 and F4 have relatively high mass shrinkages, but reduced diameters loss (the ratios being the highest 4.25 and 6.22 respectively).

Mass shrinkage is caused by the loss of volatile matter contained in the composition of the part formula during firing, this loss leads to the formation of cavities. A significant reduction in diameter is expressed when these voids are filled with particles that have undergone fusion, this compensation leads to the compaction of the part [19]. These materials have relatively maintained their shapes (reduced shrinkage diameter) but have considerably lost in mass. The density of the corresponding parts (the density) will then decrease after cooking. Demonstration of the porous aspect by microscopic observations will confirm this ascertainment.

Firing shrinkage relating to materials F1, F2, can be considered as comparable and low.

The results concerning the percentage of absorption in table IV show a significant variability, which confirms the relationship which exists between the formulation and the quality of the ceramics obtained. Note that F5 (100% sterile) has a very high absorption percentage (18.5%).

Going from F1 to F5 the percentage of absorption knows an increase, this is probably due to the integration of the phosphate mining waste in CaO, and poor in SiO2 and Al2O3.

According to table IV, F2 and F3 have a percentage of water absorption of between 6% and 10% which makes it possible to consider them, among the group BIIb; F2 has a percentage between 0.5% and 3%, so it is part of the BIIb class. F4 and F5 have a percentage that exceeds 10%, they then fall into category B III [14, 20].

D. Three-point bending Mechanical tests

Fig. 4 shows results of the three-point bending Mechanical tests for the various fabricated materials.

According to the fig. 4, we note that the phosphate mining waste cannot be used alone, its use at 100% in a formula, produces an F5 material with poor mechanical properties (flexural strength of 2.5 N/mm²). For the F1, F2, F3 and F4 clay-based formulas, the increasing integration of phosphate mining waste makes it possible to improve the mechanical properties of the manufactured tiles more and more.

For group BIII, which contains the tiles which are not necessarily the first choice from a commercial point of view, a flexural strength greater than 15 N/mm2 is required (for a thickness <7.5 mm), so F4 respects this criterion and can be marketed under the name BIII.

For the BIIb group with an average absorption percentage between 6% and 10%, a flexural strength of at least 18 N/mm2 is required, so only F3 (20% of S) meets this requirement and may be marketed according to this range [14, 20].

For group BIIb, a flexural strength greater than 27 N/mm2 is required, F1 does not satisfy this criterion.

E. Surface analysis of ceramic materials

The observation of the cohesion state and the texture of the materials, carried out by scanning electron microscopy (SEM), Fig. 5 show the results.

This observation allows the detection of microcracks within the clay matrix and the decohesions at the interfaces with the aggregates.
The image concerning F4 shows porosity visible in SEM. This finding agrees well with the results of firing shrinkage. The firing operation causes decomposition of the volatile matter, which causes a reduction in the density of the material. On the other hand, F5 seems not very cohesive and less homogeneous, the image highlights the presence of several microcracks.

IV. CONCLUSION

The physic-chemical and mechanical characterization shows that the phosphate mining waste cannot be used alone to manufacture tiling materials. In addition, its integration according to the 20% and 50% percentages with abundant local clay, makes it possible to produce ceramic tiles, complying with the applicable mechanical requirements. The quality control results of the tiles produced show that successful formulations can be used to design earthenware products. The recycling of phosphate discharges in the ceramic field can be one of the means of management and recovery combining ecological and economic aspects.

The use of phosphate mining waste as an alternative raw material in the ceramic industry ultimately has the advantages of:

• Reduce the cost of the local product by compensating part of the construction materials used
• Reduce the losses attached to manufacturing by improving the mechanical properties of the tiles produced
• Conserve non-renewable natural resources
• Avoid pollution

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