Effects of Acid Treatment on the Clay Palygorskite: XRD, Surface Area, Morphological and Chemical Composition

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The palygorskite is an aluminum-magnesium silicate that has a fibrous morphology. Their physicochemical characteristics are the result of high surface area, porosity and thermal resistance which make it an attractive adsorbent. Its adsorption capacity can be increased through chemical reactions and/or heat treatments. The objective of this work is to verify the effects of acid activation on the palygorskite, treated with HCl at 90 °C at concentrations of 2, 4 and 6 mol L\textsuperscript{-1} in 2 and 4 hours, with clay/acid solution ratio 1 g 10 mL\textsuperscript{-1} and characterized by techniques: XRF, XRD and surface area. A significant increase in specific surface area was observed in the sample treated with HCl at the concentration 6 mol L\textsuperscript{-1}. The changes were more pronounced at stricter concentrations of acidity, with decreasing intensity of reflection of the clay indicated in the XRD. These changes were confirmed in the XRF with the leaching of some oxides and with increasing concentration of SiO\textsubscript{2}.

Keywords: Palygorskite clay, acid-activation, characterization.

1. Introduction

Palygorskite is a crystalline hydrated magnesium aluminium silicate that exists in nature as a fibrous mineral with large surface area, excellent chemical stability and strong adsorption properties. It consists of two layers of tetrahedral silica linked by magnesium ions in an octahedral configuration, connected by a continuous plane of tetrahedral basal oxygen atoms\textsuperscript{4}. The tetrahedral basal oxygen atoms invert apical direction at regular intervals coordinating talc like ribbons. Each talc-like ribbon alternates with channels along the fiber axis\textsuperscript{5}.

The palygorskite has the theoretical formula: Si\textsubscript{8}Mg\textsubscript{2}O\textsubscript{20} (OH\textsubscript{2})\textsubscript{4}(OH\textsubscript{4})\textsubscript{2}4H\textsubscript{2}O, which was first proposed by Bradley\textsuperscript{1,5}. However, the chemical composition of these mineral clays varies within certain limits according to their origin. Considerable amounts of Al\textsuperscript{3+} can replace Mg\textsuperscript{2+} in the octahedral sheet. Thus, depending on the degree of isomorphous substitution, the ideal palygorskite rich in magnesium or aluminum would be: Si\textsubscript{8}(Mg\textsubscript{5.3}Al\textsubscript{2.7})O\textsubscript{20}(OH\textsubscript{2})\textsubscript{4}(OH\textsubscript{4})\textsubscript{2}4H\textsubscript{2}O, whose structure contains three forms of water in its structure: zeolitic water, coordinated and structural water\textsuperscript{6}.

In addition to aluminum, other cations may replace magnesium, albeit of a less extensive form. The presence of trivalent cations into octahedral positions creates an excess of positive charge which is compensated by empty spaces caused by two trivalent ions, whose chemical composition of the palygorskite varies according to the origin of the sample\textsuperscript{7}. The differences in the ratio of divalent/trivalent cation, which alter the chemical composition of the palygorskite, will affect the proportion of trivalent ions located at the ends and within the octahedral sheet, and carry the differences in its behavior and composition, particularly in response to acid leaching\textsuperscript{8}.

Appropriate acid treatments of clay minerals modify their surface area and/or the number of acid sites by disaggregation of clay particles, elimination of mineral impurities, removal of metal-exchange cations and proton exchange. For palygorskite, acid activation can eliminate impurities, change its composition, generate more active OH groups on its surface, increase its area and pore volume as well as the number of adsorption sites by disaggregation of palygorskite particles\textsuperscript{9-11}. Thus, acid treatment have been used for increasing the surface area of clay minerals and for obtaining from these minerals solids with a high numbers of acid centers, properties that influence the application of these materials in descolourization, adsorption and catalysis in reactions in which catalysis with acidic properties are required\textsuperscript{10}.

The objective of this research is to analyze the structural changes developed in the palygorskite after treatment by

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hydrochloric acid at three different concentrations with variations in time and with temperature of 90 °C through the use of the techniques: XRF, XRD and specific surface area (BET).

2. Experimental

2.1. Materials

It was used a sample of Palygorskite, from Guadalupe, PI, provided in the form of agglomerate by the Itoaeoste company, which has a cation exchange capacity (CEC) of 40 meq/100g, and with structural formula: (Si_{40}Mg_{8.19}Al_{1.82}Fe_{0.37}Ti_{0.084}Mn_{0.02})(K_{0.052}O_{20}(OH)_{3.4}nH_{2}O[3]). The natural clay dried at 25°, and ground was sieved entirely on 200 mesh sieve. The hydrochloric acid (HCl) 35% PA (VETEC) and distilled water was utilized in this work.

2.2. Acid activation

For the activation assays, it was treated, under magnetic stirring, an amount of palygorskite in hydrochloric acid at concentrations of 2, 4, 6 mol L⁻¹ at 90 °C with reaction times of 2 or 4 hours[8,11]. The mixture was stirred in a beaker with a cover in order to maintain the constant volume of solution used, with the mass ratio of the clay/solution volume, 1 g : 10 mL. Initially, aequous solution was prepared in the requerid concentration with hydrochloric acid, which was added to the sample (10 g) and crushed, which was contained in a glass container. After completion of the reaction, the mixtures obtained were vacuum filtered (Buchner funnel and conical flask), and washed with distilled water until the pH of the filtrates remained at around 7. The materials obtained were dried in an oven at 60 ± 5 °C and after 24 hours were disaggregated in a mortar. The activated samples were characterized by X-ray fluorescence, X-ray diffraction and surface area.

2.3. Characterization of the palygorskite

The XRF analysis was performed on spectrometer for X-ray fluorescence - (WDS) model BOL - FRX - 030. The sample was prepared by fusion at a dilution of 1/10 and used as a flux, a mixture of lithium borate (Li₂BO₂·LiBO₂). The X-Ray Diffraction (XRD) was performed on equipment LabX - XRD 600, Shimadzu, with Cu-Kα radiation (λ = 1.5406 Å) with 2θ in the range of 5° to 75°, with scan rate of 2°/min and total exposure time of 40 minutes. The surface area was determined in an apparatus for physical adsorption of nitrogen Quantachrome NOVA 4200, using BET method for processing the data. Secondary electron images were acquired with a JEOL JSM 6360LV scanning electron microscope, operating at 20 kV.

3. Results and Discussion

3.1. X-ray diffraction

The diffractograms of the natural clay and acid treated are shown in Figure 1, of which the main indication of the presence of palygorskite occurs by the appearance of peak at 8.5° (1.05 nm). Besides this, other peaks at 14.0° (0.62 nm) and 28.1° (0.32 nm) are also observed in the sample of the natural. Besides the palygorskite there is also the presence of quartz, evidenced by the peak around 26.6° (0.34 nm). Table 1 shows a comparison between data palygorskite used in this work and data from crystallographic card (JCPDS-ICCD 00-031-0783) of Clay, revealing that there are no differences between them.

The intensity of the peaks characteristic at 8.5° (1.051 nm), 14.0° (0.636 nm) and 28.1° (0.32 nm) decreased progressively in intensity with increasing acid concentration in the experimental conditions established, but does not disappear. This is explained because when the acid concentration is increased to treat a decrease in intensity of the characteristic peak of palygorskite (8.5°), a fact caused by the change of the crystalline structure and followed by leaching in octahedral and tetrahedral sheets[1,12,13].

The basal distances of the activated samples are approximately equal to the natural sample; however there are small displacements, as for example in the HCl sample - 4 mol L⁻¹/4H/90 °C, which occurs a displacement of 1.03 to 1.00 nm.

In the activated samples under more intense conditions of temperature of 90 °C and a time of 4 hours it is observed that the peaks of palygorskite become broader, indicating the loss of crystallinity. At the same time, it was observed through the difftragrams, an increase in insoluble impurities in the sample, and the main reflections correspond to quartz which were found and become more intense with increasing acid attack, progressing in both sets of samples, indicating that the acid attack occurs only on the palygorskite, with the decrease in the intensity peaks relating to clay, with a consequent decrease in crystallinity, the only observations verified[14].

![Figure 1. Diffractogram of natural palygorskite and activated with HCl.](image)

| Peaks  | Crystallographic Card | Palygorskite (This work) |
|--------|------------------------|--------------------------|
| 8.5    | 1.04 nm                | 1.05 nm                  |
| 14.0   | 0.63 nm                | 0.62 nm                  |
| 28.1   | 0.31 nm                | 0.32 nm                  |
| 26.6   | 0.34 nm                | 0.34 nm                  |

Table 1. Comparison of crystallographic planes of palygorskite.
3.2. XRF analysis

The results of XRF are shown in Table 2, which shows the chemical compositions of natural and activated palygorskites, where it is observed that Al$_2$O$_3$, Fe$_2$O$_3$, MgO, and K$_2$O are the major constituents of the octahedral sheets of the clay.

The palygorskite is a magnesium silicate, which can be substituted for aluminum in the structure. The natural sample has 11.48% aluminum and 8.28% Mg. This indicates that the substitution of magnesium by aluminum is larger.

The amounts of Mg, Al and Fe were reduced with increasing acid concentration, confirming that the ions Mg$^{2+}$, Al$^{3+}$ and Fe$^{3+}$ were leached in all activated samples. After acid treatment for a long period of time (96 h and 50°C), the hydrochloric acid solution can convert part of the crystals of palygorskite into amorphous silica. However, in this study this result was not observed, it is observed the distinction between the plans and the similarity with the initial crystalline sample, as observed in Figure 1, unlike a diffraction pattern of an amorphous material, which appears only a very broad peak.

Some explanations can be found in studies which affirm that magnesium rich clays are more easily attacked by acids and the octahedral sheets easily undergo leaching, this because its fibers have a smaller size and thus offers a greater specific surface area and a greater number of corners and edges, points which are more easily attacked by acid.

However, samples rich in aluminum, as is the case with clay of this research, have greater fiber length, as well as a larger incidence of trivalent cations that limit the progress of the attack. Another explanation is found in researches that consider that the Al occupies the center and the Mg, exclusively, the edges of the octahedral ribbons of the palygorskite which facilitates the leaching of the cations.

When the palygorskite was treated in the condition HCl - 6 mol L$^{-1}$/2 h/90°C, 53% of the Al$_2$O$_3$, 70% Fe$_2$O$_3$ and

![Figure 2. Measurements of surface area of the samples of palygorskite natural and activated.](image)

Table 2. Chemical Composition of the clays natural and activated with hydrochloric acid, under different conditions.

| Treatment                | MgO   | Al$_2$O$_3$ | SiO$_2$ | K$_2$O | TiO$_2$ | MnO   | Fe$_3$O$_4$ | Loss on Fire |
|-------------------------|-------|-------------|---------|--------|---------|-------|-------------|--------------|
| Natural                 | 8.28  | 11.48       | 63.67   | 0.32   | 0.87    | 0.21  | 3.27        | 11.90        |
| 2 mol L$^{-1}$/2H/90°C  | 8.17  | 11.26       | 63.93   | 0.27   | 1.03    | 0.17  | 3.25        | 11.92        |
| 4 mol L$^{-1}$/2H/90°C  | 5.01  | 7.26        | 71.09   | 0.11   | 0.89    | ND    | 1.66        | 13.98        |
| 6 mol L$^{-1}$/2H/90°C  | 3.29  | 5.42        | 78.97   | 0.18   | 0.90    | ND    | 0.98        | 10.26        |
| 2 mol L$^{-1}$/4H/90°C  | 5.72  | 8.09        | 72.10   | 0.14   | 1.11    | ND    | 2.04        | 10.80        |
| 4 mol L$^{-1}$/4H/90°C  | 5.00  | 7.61        | 73.84   | 0.14   | 0.93    | 0.04  | 1.80        | 10.64        |
| 6 mol L$^{-1}$/4H/90°C  | 4.79  | 6.97        | 75.65   | 0.15   | 0.97    | ND    | 1.71        | 9.76         |

ND = Non-detectable.

![Figure 3. SEM micrographs of natural palygorskite.](image)
Figure 4. SEM micrographs of samples of activated palygorskites with HCl at 90 °C.
57% of MgO were removed from the sample of palygorskite. In this sample was found 78.97% SiO₂. The percentage of the content of SiO₂ increases with the exiting of Mg, Al, Fe and others, besides silicon being the dominant constituent in the tetrahedral sheets and being relatively resistant to acid attack.1,20.

3.3. Surface area

The surface area measurements of natural and activated palygorskite are shown in Figure 2, showing the increased value of the area as the acid treatment progresses. The natural palygorskite has a surface of 113 m² g⁻¹. The maximum is 310 m² g⁻¹ in the sample 6M/2H/90 °C. The surface area of the sample treated at 6M/4H/90 °C decreases slightly to a value of 290 m² g⁻¹, as shown in Figure 2, being it possible when the treatment time was 2 hours, but there was variation in the surface area when the material was treated in a high concentration (6 mol L⁻¹). And treatments using time of 4 hours is observed that as the concentration increases, there is also an increase in surface area due to the strength of the acid.

The increased surface area of the natural palygorskite to acid treated samples is related mainly to the elimination of exchangeable cations and increase of SiO₂. The increased surface area to a maximum value and additional decrease were described in the literature for activation with acid on the palygorskite21,22 and clays such as: halloysites23, and bentonites24.

The surface area is one of the most important physicochemical properties of palygorskite, especially when it is used as an adsorbent to remove heavy metals and dyes ions. The specific surface area of the acid activated palygorskite has great influence on the adsorption of Cu (II)25.

3.4. Morphological analysis

In order to check the morphology of natural and activated samples with hydrochloric acid are shown in Figures 3 and 4.

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