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Djubaedah, Euis
Department of Mechanical Engineering, Universitas Indonesia

Wulandari, Arum, Dyah
Department of Mechanical Engineering, Universitas Indonesia

Nasruddin
Department of Mechanical Engineering, G.L. Bajaj Institute of Technology and Management

Krisnandi, K., Yuni
Department of Chemistry, Universitas Indoensia

https://doi.org/10.5109/2740938
Surface Area Modification of Natural Zeolite Through NaCl Counterbalanced Treatment to Apply in Adsorption Heat Storage System

Euis Djubaedah¹, Dyah Arum Wulandari¹, Nasruddin¹*, Yuni K. Krisnandi²

¹Department of Mechanical Engineering, Universitas Indonesia, Indonesia
²Department of Chemistry, Universitas Indonesia, Indonesia

*Author to whom correspondence should be addressed:
E-mail: nasruddin@eng.ui.ac.id

(Received October 31, 2019; Revised January 23, 2020; accepted February 24, 2020).

Abstract: The natural zeolite from Blitar deposits, a region of Indonesia were studied by simultaneous surface area and porosity measurement used BET analysis. XRD analysis was performed and the result was the natural zeolite contained mordenite and clinoptilolite and small quantities of quartz. From the pattern is also evident that the treatment doesn’t affect the mineralogical structure, patterns are identical. Thermophysical analysis, morphology and Si/Al ratio were measured. The surface area increases significantly for natural zeolite after NaCl counterbalance treatment.

Keywords: adsorption; natural zeolite; NaCl counterbalanced; surface area

1. Introduction

Zeolites are microporous aluminosilicate mineral with three-dimensional crystal structure which have large open cavities in a regular arrangement as a cage and channels. The cavities occurred from the structural composition of zeolite¹,². Their frameworks are consisting of (SiO₂)₄⁻ and (AlO₂)₅⁻ tetrahedral, where both can build a secondary building unit consisting of single rings 4-, 6- and 8-, or double rings 4-4, 6-6 and 8-8 or branched rings 4-1, 5-1, etc.³. The type of framework structure will determine surface area, pore size, and porosity⁴. Zeolite has several advantages compared to other minerals especially its function as ion exchangers, catalysts and adsorbents. Indonesia's position, which is surrounded by oceans and traversed by a series of volcanic rings, makes Indonesia rich in the potential of natural zeolite minerals⁵,⁶. Zeolite can be used as a catalyst, ion exchange and as adsorbent⁶. In generally, zeolite minerals have the following chemical formula as follows⁷:

\[ M_x / n \left[(\text{AlO}_2) \times (\text{SiO}_2) \ w \ \text{H}_2\text{O}\right] \]  (1)

Where:
- \( M \) = the cation of valence \( n \)
- \( w \) = number of water molecules per unit cell zeolite
- \( x \) and \( y \) = total tetrahedral number per unit cell

To overcome the problems of the energy crisis and environmental issues, currently, many researchers developed advanced technologies that are more energy-efficient, environmentally friendly and used for renewable energy. Utilization of solar radiation to replace fossil energy plays an important role in efforts to preserve primary energy, prevents pollution and global warming⁸. One of the technologies that utilize solar radiation is Thermal Energy Storage (TES). Thermal energy storage applications are widely used for various purposes, for example in some industries there is often a lot of waste heat leftover from the process that has not been utilized even though the heat energy can be stored as a backup or reused for other process needs. Another example is the use of hot water that comes from solar thermal energy, excess heat during the day can be stored for use at night⁹,¹⁰. In recent years, research about sorption thermal energy storage systems are significantly increase become promise option for energy-efficient technology¹²,¹³. The advantage of adsorption heat storage systems is it has high storage density and can store heat energy for a long time in ambient temperature¹⁵,¹⁶. In the adsorption process, thermal energy can be stored through three steps¹²,¹³: (1) Regeneration/Charging; (2) Storing/Discharging; (3) Adsorption.

Extensive research has been performed in the literature to study the feasibility of various adsorbent-adsorbate pairs to perform adsorption thermal storage¹⁴,¹⁵. The known molecular sieves zeolite and silica gel are promising candidates for sorbents with the necessary
physical properties. In theory, silica gel has energy densities of 220 kWh/m³, and zeolite has energy density 188 kWh/m³. From several of studies that have been carried out, synthetic zeolite is more widely discussed than natural zeolite, perhaps because natural zeolite has many disadvantages including high impurities. The presence of impurities in natural zeolite can cause clogging of the pores of zeolites so it will reduce the adsorption performance.

Although the surface area is not the only parameter to determine the performance of the adsorbent, at least the surface area can be one of the parameters that should be considered in the selection of adsorbents for adsorption heat storage system.

In this study, surface area modification by NaCl counterbalanced treatment of Indonesian natural zeolite was investigated. The objectives of this paper are determining surface area, pore volume, pore size and the Si/Al ratio of natural zeolite after NaCl counterbalanced treatment.

2. Materials and Methodology

2.1 Material Preparation

The natural zeolites (NZ) were obtained from Blitar-East Java, a region of Indonesia. All zeolites were first milled to particle size in a range of 80 mesh and washing by deionized water (1:3 w/v) under stirring for 3 hours. After washing, zeolites dried in the oven at 150 °C for 3 hours. The process as mentioned above called activated zeolite. After the activation process, 10 g of natural zeolite was mixed with 0.5M NaCl solution (10g zeolite/100 ml solution) at 80 °C under stirring for 2 x 8 hours. After stirring complete, the mixture decantation and washing by deionized water and then filtered. Keep the zeolites at room temperature until dry. The end of the process is drying the zeolite treated at 80 °C – 100 °C in the oven to make sure there is no water content in the zeolite.

2.2 Material Characterization

The surface area and porosity of prepared samples were evaluated from the N2 adsorption-desorption isotherms, as calculated at −196 °C, in the Micromeritics ASAP 2020 system. Before the study, the samples (0.10 g) were outgassed at 350 °C for 12 h. Samples total surface area was determined using the Brunauer-Emmett-Teller (BET) equation, while the t-plot approach has been used to measure the volume and surface area of micropores. Total pore volume was determined from the adsorbed volume at the maximum relative pressure obtained by the adsorption isotherm (P/Po=0.99).

X-ray diffraction (XRD) pattern was performed by PANalytical diffractometer using Cu Kα radiation (λ=1.54060 Å), from 5° to 90° in 20 degrees. The PANalytical X’pert HighScore Pro software was used for the processing of diffractograms and phase identification in combination with the International Center for Diffraction Data (ICDD) files.

X-Ray Fluorescence (XRF): This is for analyzing the elemental composition of the starting, intermediate and final product. The method employed in this research is the energy-dispersive analysis (ED-XRF).

Scanning Electron Microscopy (SEM) and energy dispersive X-Rays spectroscopy (EDS) study were performed using the JEOL JSM-7100F electron microscope. Prior to the study, in order to improve the electrical conductivity, the samples were coated with gold in a sputter coating method.

Fourier Transform Infrared Spectra (FTIR): The infrared spectra deal with complex organic molecules. Data on the nature of a compound can be derived not only from the frequencies that are present but also by peak shape and strength.

Differential Thermal Analysis/ Thermogravimetrics Analysis (DTA/TGA): The thermal analysis is used to evaluate the phase diagram, measurement of the thermal transition and desorption in different atmospheres. It provides thermal and mass loss information. The thermal analysis (DTA/TGA) was carried out with PAN analytical.

3. Result and Discussion

3.1 BET Result

The surface area, isotherm adsorption and pore distribution of natural zeolite were determined by the Brunauer- Emmett-Teller (BET) method. Surface area modification of natural zeolite with NaCl counterbalanced is a very significant change in surface area and pore diameter. Table 1 shows the enhancement of the surface area of the zeolite from 202 m²/g becomes 297 m²/g, but pore size and pore volume decrease.

<table>
<thead>
<tr>
<th>Zeolite</th>
<th>Surface Area (m²/g)</th>
<th>Pore Size (nm)</th>
<th>Pore Volume (Cm³/g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Natural Activated</td>
<td>202</td>
<td>3.7</td>
<td>0.094</td>
</tr>
<tr>
<td>Natural Zeolite – NaCl Counterbalanced</td>
<td>297.5</td>
<td>2.92</td>
<td>0.070</td>
</tr>
</tbody>
</table>

The presence and form of the mesoporous can be inferred by studying the structure of the N₂ adsorption and the desorption of the isotherm. Nitrogen adsorption involves the physical interaction of gas molecules with the surface of the sample through the force of van der Waals. N₂ adsorption isotherms of Indonesian natural zeolite. Fig. 1 is compatible with type IV isotherms in the IUPAC classification system for mesoporous materials.
3.2 XRD Result

Mineralogical analysis of the natural zeolite sample (before and after NaCl counterbalance) was carried out. The results show in Fig.2 and Fig.3 that the natural zeolite formed from mordenite and clinoptilolite and small quantities of quartz. It is also clear from the sequence that the treatment does not change the mineralogical structure, the patterns are the same.

3.3 XRF Result

The chemical composition of the zeolite reported by the manufacturer (Zeocem, a.s.) and the findings of the X-ray fluorescence analysis are shown in Table 2.

<table>
<thead>
<tr>
<th>Element/</th>
<th>NZ-activated Concentration (%)</th>
<th>NZ-NaCl counterbalanced Concentration (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>MgO</td>
<td>7.21</td>
<td>7.04</td>
</tr>
<tr>
<td>Al₂O₃</td>
<td>8.36</td>
<td>8.51</td>
</tr>
<tr>
<td>SiO₂</td>
<td>64.77</td>
<td>69.56</td>
</tr>
<tr>
<td>P₂O₅</td>
<td>0.77</td>
<td>0.79</td>
</tr>
<tr>
<td>K₂O</td>
<td>3.58</td>
<td>3.60</td>
</tr>
<tr>
<td>CaO</td>
<td>6.64</td>
<td>2.75</td>
</tr>
<tr>
<td>TiO₂</td>
<td>0.77</td>
<td>0.72</td>
</tr>
<tr>
<td>MnO</td>
<td>0.17</td>
<td>0.14</td>
</tr>
<tr>
<td>Fe₂O₃</td>
<td>7.23</td>
<td>6.42</td>
</tr>
</tbody>
</table>

3.4 FTIR Result

The IR spectrum expresses the amount of infrared radiation transmitted through the frequency of the waves. The specific identity of the zeolite structure is usually drawn from the peaks that appear in wavenumber 1250 cm⁻¹ – 350 cm⁻¹, and the wave number 4000 cm⁻¹ – 1250 cm⁻¹ is an absorption group that is not a specific identity of the zeolite structure.

FTIR spectrum is shown in Fig. 4 that the presence of water appears at the peaks that occur in the range of 3400 cm⁻¹ – 3600 cm⁻¹ as a result of the stretching of OH vibrations. The area of the presence of water formed in the natural zeolite counterbalanced by NaCl is greater compared to natural zeolite activated and raw material. The absorption area is seen in the range of 1600 cm⁻¹ – 1700 cm⁻¹. The peak of 1638,15 cm⁻¹ is the presence of Si-O vibrations for natural zeolite counterbalanced by NaCl. It has slightly higher the natural zeolite activated and raw. And areas of 1052 cm⁻¹ is the presence of Al-O vibrations.
3.5 SEM Result

The morphologies of natural zeolite samples obtained from SEM analysis are given in Fig 5. And analysis of element composition of natural zeolite was observed by EDS analysis is shown in Table 3.

Table 3. EDS result of natural zeolite before and after treatment

<table>
<thead>
<tr>
<th>Element</th>
<th>Natural Zeolite activated Mass %</th>
<th>Natural Zeolite – NaCl Counterbalanced Mass %</th>
</tr>
</thead>
<tbody>
<tr>
<td>C</td>
<td>13.68</td>
<td></td>
</tr>
<tr>
<td>O</td>
<td>39.78</td>
<td>48.77</td>
</tr>
<tr>
<td>Na</td>
<td>1.16</td>
<td>4.19</td>
</tr>
<tr>
<td>Mg</td>
<td>0.07</td>
<td></td>
</tr>
<tr>
<td>Al</td>
<td>4.5</td>
<td>6.73</td>
</tr>
<tr>
<td>Si</td>
<td>26.41</td>
<td>37.62</td>
</tr>
<tr>
<td>Cl</td>
<td>0.16</td>
<td>0.17</td>
</tr>
<tr>
<td>K</td>
<td>2.87</td>
<td>0.51</td>
</tr>
<tr>
<td>Ca</td>
<td>4.93</td>
<td>0.85</td>
</tr>
<tr>
<td>Fe</td>
<td>6.44</td>
<td>1.15</td>
</tr>
<tr>
<td>Total</td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>Si/Al</td>
<td>5.87</td>
<td>5.59</td>
</tr>
</tbody>
</table>

The result of EDS analysis showed that the exchangeable cations in natural zeolite structure were Na - K - Ca. According to the ion-exchange background of the process, it can be observed the concentration of exchangeable cations (Na, K and Ca) after NaCl counterbalance treatment. Na cation is increased in concentration after ion exchange, Na⁺ is later expected to be able to carry H₂O groups, which is interpreted as the potential for H₂O absorption.

3.6 TGA/DTA Result

In the adsorption of thermal energy storage, the regeneration or desorption process is carried out starting from 25 °C - 200 °C, so it is necessary to analyze the thermal characteristics that occur at that temperature range. Fig. 6 below represents the corresponding sample TGA thermal analysis. The curve in red color stands for the natural zeolite counterbalanced by NaCl, while the one in color blue stands for natural zeolite activated.

Thermal examination reveals the presence of endothermic effects, common for natural zeolite, which does not cause structural changes and thermal dehydration, under which the natural zeolite under investigation loosens physically and chemically bound water. The
thermal characteristics of natural zeolite before and after counterbalanced NaCl have almost the same properties.

4. Conclusion

Natural zeolites were successively treated with a NaCl counterbalance. The effect of NaCl on the structure has been studied. There are no changes in a mineralogical structure after NaCl treatment. The Si / Al ratio of the zeolite sample decreased slightly. The NaCl treatment changes the pore structure of the zeolite sample and increases the specific surface area of the sample. The specific surface area of natural zeolites increased from 202 m²/g to 297 m²/g after being treated with NaCl. Heat treatment at low temperatures has a negligible effect on the pore structure of zeolite samples. Natural zeolite treated with NaCl has the better potential of water absorption ability compared to other samples, this can be seen from the results of FTIR and EDS analysis.

Acknowledgements

The authors acknowledge the financing support from Ministry of Research, Technology and Higher Education of the (KEMENRISTEKDIKTI) Republic of Indonesia for Grant Research Doctors (PDD) No. 1/E1/KP.PTNBH/2019 - 234/PKS/R/UI/2019 - NKB-1847/UN2.R3.1/HKP.05.00/2019.

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