Sepiolite-based adsorbents for carbon dioxide capture

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Sepiolite and the sepiolite-based materials were studied in terms of carbon dioxide adsorption. The pore structure and the surface characterization of the obtained materials were specified based on adsorption-desorption isotherms of nitrogen measured at -196°C and carbon dioxide at 0°C . The specific surface area (SSA) was calculated according to the BET equation. The pore volume was estimated using the DFT method. Pristine sepiolite has shown the following value of SSA and CO_2 uptake at 0°C – $105\text{ m}^2/\text{g}$ and 0.27 mmol/g , respectively. The highest value of these parameters was found for material obtained by KOH activation of mixture sepiolite and molasses (MSEP2) – $676\text{ m}^2/\text{g}$ and 1.49 mmol/g . Sample MSEP2 also indicated the highest value of total pore volume and micropores volume with a diameter up to 0.8 nm .

Keywords: sepiolite, molasses, carbon-clay composite, CO_2 adsorption.

INTRODUCTION

Carbon dioxide makes up the vast majority of greenhouse gas emissions (81%) but smaller amounts of methane (10%), nitrous oxide (6%) and fluorinated gases (3%) are also emitted¹. Absorption of CO_2 from flue gas by amine scrubbing is a traditional method and currently in operation. The amine solution has some disadvantages such as high volatility, low thermal stability, and high oxidation degradation².

Adsorption is a promising alternative method for CO_2 capturing and could replace an amine-based absorption technique³. Zeolites^{4, 5}, carbon-based materials^{6–9} and metal-organic frameworks^{10–12} are the most critical class of adsorbents actively investigated for gas adsorption applications. Unfortunately, the moisture typically present in the flue gas stream affects zeolites and metal-organic frameworks structural stability¹³. Carbon materials, exhibit better stability towards moisture and also against corrosive gases¹⁴. Because of high adsorption capacity, easy availability and low cost activated carbons are most commonly tested as CO_2 adsorbents^{6, 15–17}. A variety of carbonaceous adsorbents were also considered for CO_2 capture: carbon nanotubes⁹, carbon spheres¹⁸, nitrogen-doped nanoporous carbon¹⁹, porous carbon nanosheets²⁰. There are also attempts to convert CO_2 into valuable products such as methane²¹, formaldehyde²², methanol²³.

Methane is emitted from energy, industry, agriculture, and waste management activities. CH_4 lifetime in the atmosphere is much shorter than CO_2 , but CH_4 is more effective at trapping radiation than CO_2 . The comparative impact of CH_4 is 25 times greater than CO_2 over a 100 years²⁴. The most significant industrial methane-based process is methane steam reforming to syngas – a mixture of CO and H_2 . The dry reforming of methane was also investigated^{25, 26}.

Converting methane to alternative forms sources of energy, such as methanol^{27, 28}, formaldehyde^{29, 30} and other derivatives^{31–34} is a preferred method for methane reduction. Catalytic methane decomposition led to the production of pure hydrogen and carbon nanomaterials^{35–38}. Methane can be applied as vehicle fuel^{39, 40}.

Carbonaceous materials are right sorbents for other gases and vapors, e.g. hydrogen^{41–43}. They were also described as catalysts and catalysts supports^{44–46}.

Clays are hydrated aluminum phyllosilicates naturally formed through hydrothermal alteration of rocks. Clay minerals have a sheet-like structure and are composed of mainly tetrahedral silicate and octahedral hydroxide groups⁴⁷.

Sepiolite, formerly known as Meerschaum (sea froth), is a naturally occurring phyllosilicate clay mineral, that is widespread due to its unusual fibrous morphology and intracrystalline tunnels. The main advantages of sepiolite are the porosity, the fibrous structure, the fine grain size, and the high surface area, which make it a valuable material for a wide range of applications (e.g. pesticide and catalyst carrier, decolourizing and anticaking agents or pharmaceutical and cosmetics applications)^{48, 49}.

Due to their low price, abundance, and porosity, clay materials (as montmorillonite, kaolinite, and bentonite) have been used as supports for CO_2 adsorption^{50–52}. However, to the best of our knowledge, the carbon dioxide adsorption of sepiolite-based materials (sepiolite-carbon composites from biomass) has not yet been reported.

EXPERIMENTAL

Preparation of sepiolite-based adsorbents

The sepiolite (SEP) was mixed with potassium hydroxide (KOH) in a ratio of 1:1 (KSEP), potassium hydroxide and molasses in a ratio of 1:1:1 (MSEP1) and in a ratio of 1:2:1 (MSEP2) and left in the room temperature for three hours. Afterward, samples KSEP, MSEP1, and MSEP2 were dried at 200°C for 19 h. In the next step, the materials were exposed to 800°C in a furnace for 1h under a constant nitrogen flow $18\text{ dm}^3/\text{h}$. After that time, obtained materials were washed with the deionized water to a pH value of 6.5 to remove KOH. Then, the materials were soaked in hydrochloric acid (HCl) in a concentration of 1 mol/dm^3 for 19 h, washed again with the deionized water until the pH of filtrate became 6.5 and dried at 200°C for 12 h.

The sepiolite (SEP) was also mixed with potassium hydroxide (KOH) in a ratio of 1:1 and left in the room temperature for three hours, but the heating step (to 800°C) was omitted (ISEP). The rest of the preparation stage was the same as for the samples mentioned above.

Characterization of sepiolite and sepiolite-based adsorbents

Sepiolite and sepiolite-based adsorbents were characterized by nitrogen adsorption-desorption isotherms measured at -196°C using Quantochrome Autosorb Instrument. Before analysis, all the samples were outgassed at 250°C under vacuum for 16 h in a vacuum environment. The specific surface area (SSA) was estimated from the adsorption isotherm using the standard Brunauer-Emmett-Teller (BET) equation based on adsorption data. The micropore volume (MPV_{N_2} , MPV_{CO_2}) was assessed by the Density Functional Theory (DFT) method used to the nitrogen and carbon dioxide adsorption. The total pore volume (TPV) was calculated based on the volume of N_2 adsorbed at a relative pressure $p/p_0 \sim 1$. The DFT was also used to establish the pore size distributions (PSD) from sorption isotherms of nitrogen at -196°C (PSD_{N_2}) and carbon dioxide at 0°C (PSD_{CO_2}).

The CO_2 adsorption capacity at 0°C and 25°C were determined using the Quantachrome Autosorb Instrument.

X-ray diffraction patterns of products were acquired by using PANalytical X'Pert PRO diffractometer. Data were collected over a 2θ range of $5\div 80^{\circ}$ using $\text{CuK}\alpha$ radiation.

The morphology and elemental composition of products were examined by scanning electron microscopy (SEM). SEM images were performed using SU8020 (Hitachi) microscope with an accelerating voltage of about 15 kV at 5000x magnification.

RESULTS AND DISCUSSION

Figure 1 presents nitrogen adsorption-desorption isotherms of obtained materials. Physisorption isotherms of all samples represent combination of type I and IV with a hysteresis loop of type H3 (for SEP, ISEP, and MSEP2) and H4 (for KEP, MSEP1), according to the IUPAC classification. The type I/IV isotherms showed that both, micropores and mesopores had been identified in samples. Capillary condensation in the pores occurred at a relative pressure (p/p_0) above 0.40, the most probably due to the presence of a large amount of narrow slit-like mesopores.

The textural properties (BET surface area, total pore volume, micropore volume) of sepiolite and sepiolite-based adsorbents are listed in Table 1. The specific surface

Figure 1. Nitrogen adsorption-desorption isotherms of pristine sepiolite and sepiolite-based materials

Table 1. Textural properties of pristine sepiolite and sepiolite-based materials

Sample	SSA [m ² /g]	MPV_{CO_2} [cm ³ /g]	MPV_{N_2} [cm ³ /g]	TPV [cm ³ /g]
SEP	105	0.017	0.008	0.285
ISEP	351	0.069	0.044	0.530
MSEP1	570	0.107	0.119	0.403
MSEP2	676	0.113	0.086	0.613
KSEP	426	0.101	0.003	0.269

area of the materials increased following the chemical modifications. The highest surface area of $676 \text{ m}^2/\text{g}$ was achieved by KOH activation of mixture sepiolite and molasses (MSEP2). The other samples had a smaller BET surface area compared to MSEP2, but larger than pristine sepiolite. As presented in Table 1 the pores of examined materials exhibited a considerable increase in volume in comparison with sepiolite. In the case of sepiolite, the KOH treatment improves the porosity of sepiolite itself. The highest total pore volume was also achieved on sample MSEP2 – $0.61 \text{ cm}^3/\text{g}$. This sample also had the highest MPV_{CO_2} value – $0.11 \text{ cm}^3/\text{g}$ (as well as sample MSEP1). However, the highest value of MPV_{N_2} was found for sample MSEP1 – $0.12 \text{ cm}^3/\text{g}$. These values are much higher than those obtained for the pristine sepiolite.

Figure 2. shows the pore size distribution determined from CO_2 adsorption isotherms at 0°C (PSD_{CO_2}), and Figure 3. reveals the results of PSD calculated from N_2 adsorption isotherms at -196°C (PSD_{N_2}). Both PSDs were calculated by the DFT method. To clarify the graph, the maximum value of the horizontal axis is 20 nm, for the reason that the obtained materials did not contain a high number of larger pores. All the samples exhibited contents of pores with a diameter range of 0.3–20 nm. The largest volume was occupied by pores with diameters ranging from 0.4 to 0.8 nm and 1.3 to 2.7 nm, which confirms

Figure 2. The pore size distributions of pristine sepiolite and obtained materials calculated from CO_2 adsorption isotherms (D – pores diameter)

Figure 3. The pore size distributions of pristine sepiolite and obtained materials calculated from N_2 adsorption isotherms (D – pores diameter)

the development of microporosity and the presence of smaller mesopores in obtained materials.

The carbon dioxide adsorption isotherms of the examined samples are presented in Figure 4 (at 0°C) and Figure 5 (at 25°C). The CO_2 uptake at 1 bar and 0°C is in the range of 1.18 to 1.49 mmol/g, while for pristine sepiolite is 0.27 mmol/g. In the case of CO_2 uptake at 1 bar and 25°C is between 0.63 and 0.89 mmol/g for sepiolite-based materials and 0.18 mmol/g for pristine sepiolite. The highest value of CO_2 capacity at 0°C and 25°C , 1.49 mmol/g, and 0.89 mmol/g was confirmed for MSEP2 material.

Figure 4. Carbon dioxide adsorption isotherms at 0°C of pristine sepiolite and sepiolite-based materials

Figure 6 demonstrates the relationship between specific surface area and carbon dioxide adsorption (at 0°C), while Figure 7 presents the correlation between MPV_{CO_2} value and CO_2 uptake (also at 0°C). It can be seen, that there is a excellent correlation between specific surface

Figure 5. Carbon dioxide adsorption isotherms at 25°C of pristine sepiolite and sepiolite-based materials

Figure 6. The correlation between SSA and CO_2 adsorption (at 0°C) for pristine sepiolite and obtained materials

Figure 7. The correlation between MPV_{CO_2} and CO_2 adsorption (at 0°C) for pristine sepiolite and obtained materials

area and micropore volume with CO_2 uptake. It is, equal with other findings for micro- or mesoporous materials. The coefficient of determination (R^2) in both cases was high and amounted to – 0.82 and 0.95, respectively.

The XRD patterns of the pristine sepiolite and sepiolite-based materials were shown in Figure 8. For SEP, a diffraction peak with high intensity at $2\theta = 7.2^\circ$ was

Figure 8. X-ray diffraction patterns of pristine sepiolite and obtained materials

ascribed to the (110) reflection of the sepiolite framework, which is the characteristic peak of sepiolite itself^{53, 54}. After impregnation with KOH, there is no characteristic peak in patterns of ISEP, KSEP, and MSEP1 samples, indicating that the KOH treatment entirely changes the crystalline structure of sepiolite. In the case of MSEP2 material, the crystal structure was slightly changed, and the characteristic peak of sepiolite has been preserved.

The morphology of the examined materials was investigated by SEM, and typical SEM images are shown in Figure 9. It is well known that pore structure development is influenced by many factors, including inorganic impurities and the initial structure. The pristine sepiolite has fibrous morphology in the form of aggregates and some single fibers (Fig. 9a). After KOH treatment and pyrolysis (KSEP), the morphology seems to be similar. However, some single fibers were gone, and fibrous clusters appeared, as shown in Figure 9b. In the case of samples with molasses (MSEP1, MSEP2), changes in the structure of sepiolite can be observed. The materials MSP1 and MSP2 show an irregular shape and larger cavities compared with the pristine sepiolite (Fig. 9c, 9d).

Figure 9. Scanning electron micrographs of pristine sepiolite and obtained materials

CONCLUSIONS

Sepiolite-based materials have been prepared by KOH impregnation (ISEP), KOH impregnation in combination with the pyrolysis (KSEP), and KOH impregnation with the pyrolysis of sepiolite/molasses mixture (MSEP1, MSEP2). The results of structure characterization results showed the increased specific surface area and pore volume of obtained materials in comparison with pristine sepiolite. The surface treatment changed the porous structure of the pristine sepiolite. For SEP, the SSA, TPV, and MPV_{CO_2} value was 105 m²/g, 0.285 cm³/g, and 0.017 cm³/g. The highest value of these parameters was obtained for MSEP2 – 676 m²/g, 0.613 cm³/g, and 0.113 cm³/g, respectively. Carbon dioxide adsorption tests showed that impregnating the pristine sepiolite with potassium hydroxide significantly improved CO₂ adsorption capacity at 0°C and 25°C. The highest value of CO₂ capacity at 0°C and 25°C was obtained for MSEP2 material – 1.49 mmol/g, and 0.89 mmol/g, respectively. The sepiolite as a clay mineral is an inexpensive material, with abundant availability around the world. Moreover, it is vulnerable to chemical modification. Thus, the sepiolite treatment by potassium hydroxide, causing the increase of both – the microporosity and the CO₂ adsorption capacity.

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