

CO₂ Adsorption on Activated Carbon

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Editorial

Climate change is one of the major environmental, social and economic problems affecting the world. Carbon dioxide (CO_2) is considered to be the principal source of greenhouse gases that is responsible for global warming [1]. The IPCC believes that CO₂ concentration increases to 570 ppm by the year 2100, provoking a global average temperature to rise by about 1.9°C [2]. Various techniques have been used to reduce the CO₂ emissions among which the use of renewable energy, the energy control and the development of CO₂ capture and storage (CCS) technology [3]. The CCS technique has been considered as the best option for reducing carbon dioxide emissions from large point sources. The carbon dioxide (CO2) captured from large stationary sources can be safely injected and stored in appropriate geological formations, such as deep saline formations and depleted oil and gas reservoirs [4].

Adsorption process has become a potential attractive alternative for CO_2 storage in the context of carbon capture and sequestration technologies (CCS) because of its low energy requirements, cost effectiveness, and ease of use at a variety of pressures and temperatures [5]. Many different adsorbents can be used for this purpose such as clay, activated carbon (AC), zeolites, mesoporous silica, and organo-metallic structures [5].

Based on the above-mentioned studies, activated carbon (AC) is used as adsorbent with CO_2 [6]. The AC was characterized by SEM (Figure 1), wich present a roughness structure resembling a series of parallel lines. The AC gave rougher textures with heterogeneous surfaces and a greater variety of randomly distributed pores size. The N₂ isotherm (Figure 2) shows that the studied AC can be classified as a type I isotherm [6]. According to the IUPAC classification, the adsorbent is essentially formed by micropores with a very narrow microporosity. The textural parameters show that

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the AC has a significant porosity, with high values of total pore volume Vtotal (0.676 cm₃ g⁻¹), surface area S_{BET} (1565 m² g⁻¹) and micropore volume (0.649 cm³ g⁻¹).



Figure 1: The AC was characterized by SEM.



The CO_2 adsorption isotherms in formulated AC were measured volumetrically at various temperatures (298, 310 and 320 K), and the experimental data were modeled with the Langmuir and Freundlich methods [6]. The Langmuir isotherm was employed to describe the monolayer

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adsorption into the homogeneous surface. While the Freundlich isotherm was used for multilayer adsorption and surface heterogeneity (Figure 3). These parameters proved

that the CO_2 adsorption is spontaneous, exothermic and physisorption in nature.



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