Supporting information

Synthesis of high-temperature CO₂ adsorbents from organolayered double hydroxide with markedly improved CO₂ capture capacity

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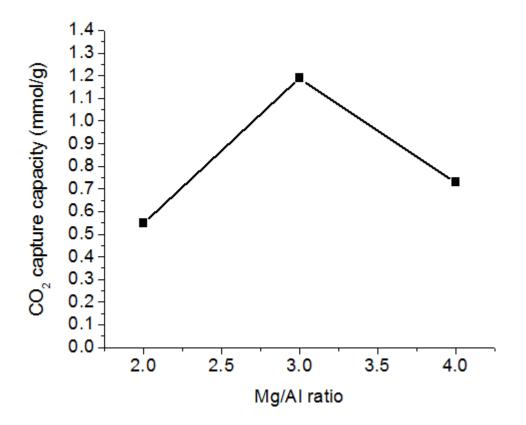


Figure S1. Effect of Mg/Al ratio on the CO₂ capture capacity of Mg_xAl₁-stearate (x = 2, 3 and 4) evaluated at 200 °C.

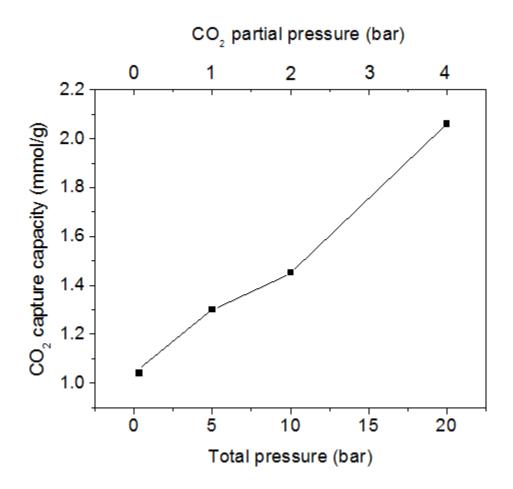


Figure S2. The effect of operating pressure (CO₂ partial pressure) on the CO₂ capture capacity of Mg_3Al_1 -stearate derived adsorbent. Testing condition: 1.5 g adsorbent, 20% CO₂, 20 % Ar in N₂ (total flow rate 100 ml/min), 400 °C.

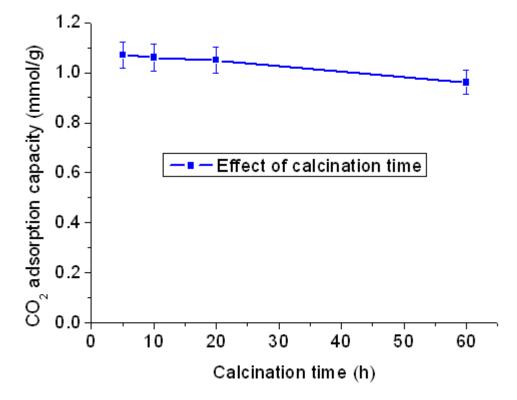


Figure S3. The effect of calcination time on the CO_2 capture capacity of the Mg_3Al_1 -stearate derived mixed oxide. Calcination temperature: 400 °C, adsorption temperature: 400 °C.

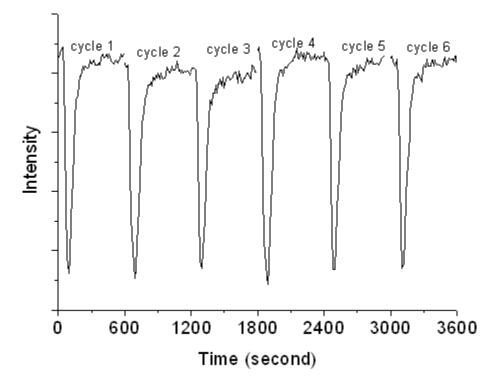


Figure S4. CO_2 adsorption/desorption cycling test for Mg₃Al₁-stearate derived mixed oxide. Adsorption temperature: 200 °C, desorption temperature 400 °C.

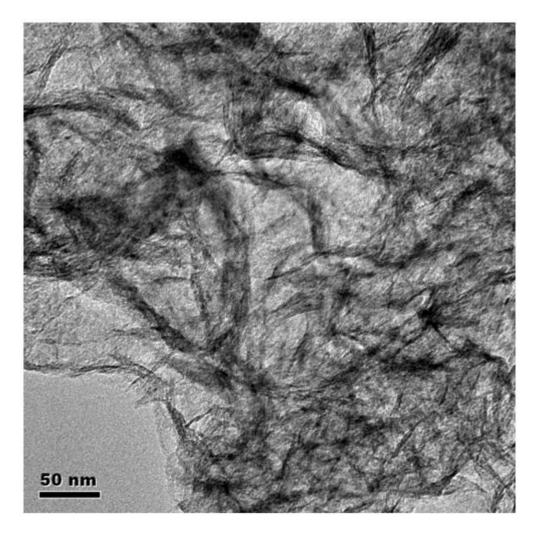


Figure S5. TEM image of the calcined Mg₃Al₁-stearate (400 °C, 5 h, in air).

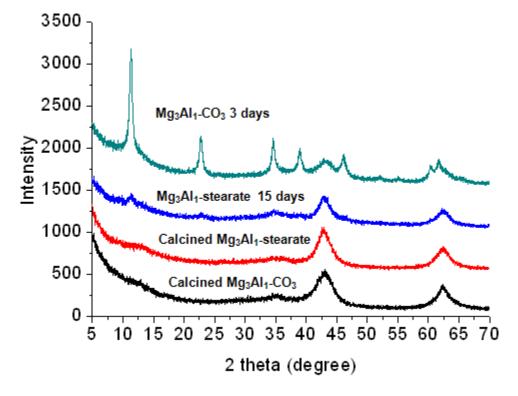


Figure S6. XRD patterns of the Mg_3Al_1 -CO₃ and Mg_3Al_1 -stearate after calcination at 400 °C and exposure to ambient atmosphere for several days.