



Cite this: *RSC Adv.*, 2019, 9, 6054

Preparation of CaMgAl-LDHs and mesoporous silica sorbents derived from blast furnace slag for CO₂ capture†

Haojie Jiang,^a Hongwei Guo,^a  Peng Li,^{*a} Yang Li^b and Bingji Yan^a

High volume blast furnace slag (BFS) resulting from iron-making activities has long been considered a burden for the environment. Despite considerable research efforts, attempts to convert BFS into high value-added products for environmental remediation are still challenging. In this study, calcium–magnesium–aluminium layered double hydroxides (CaMgAl-LDHs) and ordered mesoporous silica material (MCM-41) sorbents were simultaneously synthesized from BFS, and their CO₂ adsorption performance was evaluated. Calcium (Ca), magnesium (Mg) and aluminium (Al) were selectively extracted from BFS using hydrochloric acid. Leaching conditions consisting of 2 mol L⁻¹ acid concentration, 100 °C leaching temperature, 90 min leaching time and a solid-to-liquid ratio of 40 g L⁻¹ achieved a high leaching ratio of Ca, Mg and Al at 88.08%, 88.59% and 82.27%, respectively. The silica-rich residue (SiO₂ > 98.6 wt%) generated from the leaching process could be used as a precursor for MCM-41 preparation. Chemical composition, surface chemical bonds, morphology and textural properties of the as-synthesized CaMgAl-LDHs and MCM-41 sorbents were determined. Both the CaMgAl-LDHs and MCM-41 sorbents were found to be thermally stable and exhibited comparable adsorption uptake and rates over 20 CO₂ adsorption/desorption cycles. This work demonstrated that a total solution for the utilisation of BFS can be achieved and the resulting valuable products, *i.e.* CaMgAl-LDHs and MCM-41 are promising sorbents for CO₂ capture.

Received 12th October 2018
 Accepted 18th December 2018

DOI: 10.1039/c8ra08458k

rsc.li/rsc-advances

1. Introduction

One of the most significant environmental challenges faced by humans in the 21st century is global warming caused by the increase in carbon dioxide levels.¹ In response to the issue of greenhouse gases, many countries have proposed a series of carbon emission reduction plans. Notably, research into options for the capture, utilisation and long-term storage of carbon dioxide has become a hot topic in the field of low-carbon and zero-carbon technologies, which include renewable energy, clean energy and carbon capture & storage (CCS).^{1–5} (CCS) is being investigated as a potential option to capture CO₂ at the source of generation from various industrial facilities, and to transport and separate the captured CO₂ in storage areas.^{5–9}

Iron and steel manufacturing are not only major energy-consuming industries but also major emitters of greenhouse gases. The generation of large amounts of solid waste including blast furnace slag (BFS) and steel slag (SS) has caused severe

environmental problems involving water and soil contamination, and local air pollution.¹⁰ To reduce or resolve these growing environmental concerns, many current research activities are focusing on the development of new waste management strategies such as the preparation of useful materials from waste. In the past ten years, mineral carbonation is the most dynamic and the fastest growing CO₂ sequestration technology. In mineral carbonation, CO₂ gas is stored by promoting the reaction of magnesium or calcium oxides in silicate minerals with carbon dioxide to form carbonates;⁴ BFS and SS from the steel industry contain a large number of alkaline oxides, such as CaO and MgO, which can be used for CO₂ capture.¹¹

In Finland, Eloneva *et al.*^{2–5} studied the possibility of utilizing SS for the production of pure calcium carbonate. They found that dilute acetic acid only selectively dissolved calcium from the steel slag. Tian *et al.*¹² proposed a process for simultaneously producing CaO-based CO₂ sorbents and high-quality iron ore from waste SS. The results showed that the extraction of calcium was always accompanied by the undesired dissolution of elemental impurities such as Al, Mg and Fe, resulting in decreased CO₂ adsorption capacity.

As well as the intensive research activity on SS, BFS is also considered as valuable raw material in CO₂ sequestration. Compared with SS, BFS is more greatly enriched in elemental Ca, Mg, Si and Al, which together account for around 90% of its total mass; this means that the aqueous extraction of BFS could

^aShagang School of Iron and Steel, Soochow University, Suzhou 215021, P. R. China. E-mail: guohongwei@suda.edu.cn; lipeng@suda.edu.cn; Fax: +86-0512-62334926; Tel: +86-0512-62334926

^bThe State Key Laboratory of Refractories and Metallurgy, School of Materials and Metallurgy, Wuhan University of Science and Technology, Wuhan 430081, China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ra08458k



potentially leads to less contamination. Seonhye Lee *et al.*¹ studied the indirect sequestration of CO₂ using BFS as the raw material and achieved 49.76% extraction efficiency of Ca and 95.69% carbonation efficiency using NH₄NO₃ under typical conditions (2 mol L⁻¹ solvent concentration, 30 °C reaction temperature, 60 min reaction time, 10 g L⁻¹ solid-to-liquid ratio).

In their early research, Yasutaka Kuwahara *et al.*¹³ successfully synthesised a Ca-based layered double hydroxide (LDH) compound from BFS, and applied it in CO₂ fixation. In their methodology, the hydrochloric acid leaching resulted in a precursor solution suited for LDH preparation and hydrated silica gel. Inspired by this result, in this paper, a facile method for the simultaneous preparation of (LDHs) and ordered mesoporous silica (MCM-41) materials from BFS is reported. The objectives of this research are to focus on: (i) clarifying the influence of acid leaching conditions on the dissolution behavior of BFS, and seeking optimised conditions for the selective and highly efficient dissolution of elemental calcium, magnesium and aluminium; (ii) making best use of the BFS through synthesising LDHs and MCM-41 sorbents by recovering the leachate and residue after solvent extraction and subsequent structural characterisation, (iii) the evaluation of CO₂ sequestration ability for both sorbents through adsorption–desorption measurements.

2. Experimental

2.1 Materials and test facilities

Water granulated blast furnace slag was received from the Shagang Steel Co., Ltd., China. The slag was crashed and ball

Table 1 Chemical composition of blast furnace slag A: measured by XRF analysis, B: by ICP-OES analysis

	Al ₂ O ₃	SiO ₂	CaO	MgO	K ₂ O	Fe ₂ O ₃	Na ₂ O	MnO
Composition^A								
wt%	14.47	34.27	36.84	10.14	0.52	0.53	0.43	0.52
Composition^B								
wt%	14.66	32.77	36.14	10.79	NA	0.26	NA	NA

milled to a particle size of less than 74 μm. The chemical composition of the BFS was analysed using X-ray fluorescence (XRF, SHIMADZU-EDX800, Japan), and the concentrations of Ca, Si, Mg and Al in the solution were measured using inductively coupled plasma-optic emission spectrometry (ICP-OES, Vista-MPX, Varian, USA). The composition^A results from XRF quantitative measurements are shown in Table 1, revealing that the BFS was mainly composed of 36.84 wt% CaO, 34.27 wt% SiO₂, 10.14 wt% MgO and 14.47 wt% Al₂O₃, which together accounted for 95% of the total mass. Results of the ICP-OES analysis gave the composition^B of the solution, with these values used for estimating the leaching ratio of metal ions from the BFS.

The mineralogical phases were identified by X-ray diffraction (XRD, Ultimal VX, Rigaku, Japan) with CuKα radiation, utilising a generator voltage of 40 kV and tube current of 40 mA. Small-angle X-ray scattering (SAXS) measurements were carried out at room temperature using a SAXSess mc² (Anton Paar) instrument.

Surface functional groups were determined with Fourier transform infrared spectroscopy (Jasco FT-IR-4100) with KBr in the 1800–400 cm⁻¹ region.

Morphologies were characterized by scanning electron microscope (SEM, SU5000, Hitachi, Japan) and a transmission electron microscope (TEM, TECNAI G2 F20, FEI, USA). Textural properties were investigated by N₂ adsorption–desorption measurements using an ASAP 2000 instrument (Micromeritics, USA). Thermogravimetric analysis (TGA) was performed to evaluate thermal stability using a TGA-50/50H analyser (Shimadzu, Kyoto, Japan).

2.2 Experimental methods

2.2.1 Hydrochloric acid dissolution experiments. As illustrated in Fig. 1, extraction experiments were carried out in a 500 mL flat-bottomed flask glass reactor with a working volume of 200 mL. A mechanical stirrer (Dragonlab® MS7-H550-S, Dragon Laboratory Instruments Co. Ltd., Beijing, China) was installed at the center of the reactor and used to mix the slurry at 500 rpm. The glass reactor was placed in a temperature-controlled water bath to maintain a consistent

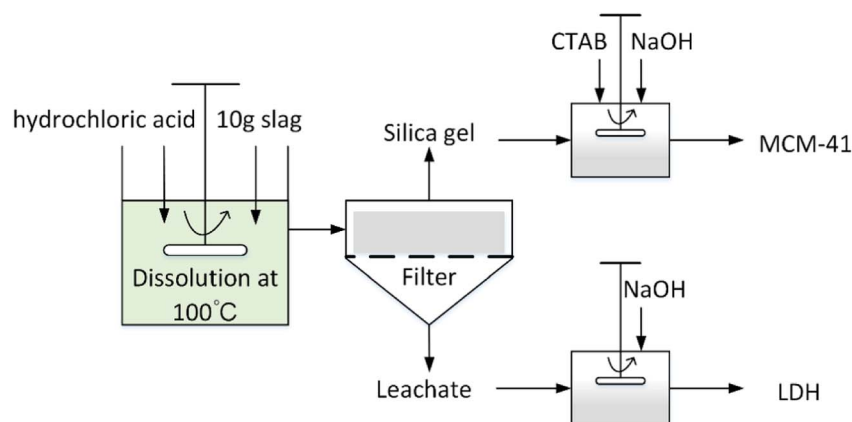


Fig. 1 Employed methodology for simultaneously producing LDHs and MCM-41 from blast furnace slag.



reaction temperature. To avoid evaporation losses, the reactor was equipped with a condenser cooled with tap water.

BFS (10 g) was added to the reactor containing a specific concentration (0.5 M, 1 M, 2 M or 3 M) of hydrochloric acid (11.74 M, Enox) at 100 °C. To monitor the progression of the dissolution, 5 mL of aqueous solution was sampled using quantitative pipette at 5, 10, 20, 40, 60, 80, 100, 120, 140, 160 and 180 min after the slag was added.

The leaching ratio of a specific element was calculated by the following equation:

$$\text{Leaching ratio (\%)} = \frac{C_x V}{M_s W_x} \times 100 \quad (1)$$

Here, C_x (mg L⁻¹) represents the concentration of a selected element in the extracted aqueous solution, V (L) is the volume of aqueous solution, M_s (mg) denotes the mass of the slag and W_x (wt%) is the elemental content in the slag.

2.2.2 Synthesis of LDHs. After the leaching process, the obtained leachate containing Ca²⁺, Mg²⁺, Al³⁺ can be used as precursor for LDHs synthesis. NaOH aqueous solution (2 M) was added dropwise into the leachate (150 mL) and the final pH adjusted to 12. The resulting mixture was then aged at 80 °C for 18 h to enable crystallisation. The aged mixture was filtered and washed with deionised water several times until the pH was 7. Finally, the obtained product was dried at 100 °C overnight.

2.2.3 Synthesis of MCM-41. The solid residue resulting from acid leaching was subjected to secondary acid leaching (2 M acid concentration, 1 h of leaching at 100 °C) to remove impurities. The XRF results in Table 2 show that the secondary acid leaching yielded solid residue containing around 98.6 wt% of silica and could be utilised as a precursor for mesoporous silica preparation.

In a typical procedure, *n*-cetyltrimethylammonium bromide (CTAB, 1.457 g) was dissolved in of deionised water (60 mL) at room temperature. Then, acid leaching residue (1.2 g) and NaOH (1 g) were added to the CTAB aqueous solution. The mixture was stirred until a clear solution was obtained, following which NaOH or HCl solution was added dropwise to adjust the pH to 11. The resulting solution was further stirred for another one hour, then placed in a Teflon-lined autoclave for hydrothermal treatment at 80 °C for 48 h. After that, the slurry was filtered, washed repeatedly using deionised water and air-dried overnight at 100 °C. Finally, the as-synthesized silica powder was calcined at 550 °C in air for 6 h.

2.2.4 Measurements of CO₂ adsorption capacity. CO₂ adsorption-desorption measurements of synthesized LDHs and MCM-41 materials were performed on a thermogravimetric analyser (CP-2000, Beijing). Precisely 10 mg sample was activated by heating it to a desired temperature and maintaining the temperature for 30 min under N₂ (50 mL min⁻¹) before CO₂ adsorption. The temperature was then adjusted to the adsorption temperature and the gas feed switched from N₂ to CO₂ with

a constant flow of 50 mL min⁻¹. After adsorption, the flow was switched from CO₂ to N₂ and the temperature adjusted to and maintained at the activated temperature for 30 min for completion of one desorption. This experiment was repeated twenty times to investigate the stability of CO₂ adsorption.

3. Results and discussion

3.1 Leaching of BFS

3.1.1 Effect of hydrochloric acid concentration. According to the results in Fig. 2, the dissolution of elemental Ca, Mg, Al and Si in BFS was strongly dependent on acid concentration. The leaching ratios of Ca, Mg, Al significantly improved when the acid concentration exceeded 1 M. Surprisingly the rate of dissolution of BFS in acid solution was extremely fast and the leaching ratios for these elements reached a within 5–10 min of initial contact; this suggested that the water-granulated BFS exhibits outstanding reactivity. In contrast, the dissolution behaviour of Si followed a different pattern. When higher concentrations of 2 M and 3 M were employed, the leaching ratio of Si was drastically reduced from 65% to 1.27% after 80 min and continuously decreased to nearly zero.

3.1.2 Effect of leaching temperature. Fig. 3 shows that increasing the temperature from 70 to 100 °C did not enhance the dissolution of Ca, Mg, Al and Si; on the contrary, the leaching ratios of these elements all decreased. In particular, the leaching ratio of Si dropped dramatically to 1.27% at 100 °C, which suggested that high temperature favours the formation of silica gel and consequently leads to decreased dissolution of Si. The newly forming silica gel with high surface area has the capability of caging metal ions, which may account for lower leaching ratios of Ca, Mg and Al at higher temperatures.

3.1.3 Effects of the solid-to-liquid ratio. To study the effect of the solid-to-liquid (S/L) ratio on the leaching behavior, the acid solution (concentration and volume) was kept constant and the amount of BFS doped in the solution was varied, *i.e.* 6, 8, 10, 12, 14 gram. As shown in Fig. 4, the general trend was that the leaching ratio of Ca, Mg, and Al increased as the S/L ratio decreased. At a S/L ratio of 40 g L⁻¹, the highest leaching ratio of Ca, Mg, Al was achieved at 88.08%, 88.59% and 82.27%, respectively. Meanwhile, the S/L ratio had little effect on the dissolution behaviour of Si.

3.1.4 Leaching selectivity. The degree of elemental selectivity was used to identify the optimised extraction conditions for subsequent synthesis of high purity LDHs and MCM-41 materials. As can be seen from Table 3, a leaching temperature of 100 °C, acid concentration of 2 M and S/L ratio of 40 g L⁻¹, yielded a satisfied elemental selectivity, that is the elemental selectivity for silicon was a minimum and high leaching ratio. Increase in the acid concentration resulted in the considerable decomposition of the amorphous silica phase within only a few minutes of leaching. Dissolved silicon presents as Si(OH)₄, which is monomeric silica in an acidic solution. Once the monomeric concentration exceeds the saturation level, the discrete silica particles start to polymerise to into a gel. The above experimental results indicated that high temperature and long leaching times can promote the polymerisation of Si according to eqn (2) and (3):¹⁴

Table 2 Chemical composition of the solid residue obtained from secondary acid leaching

Composition	SiO ₂	Al ₂ O ₃	MgO	TiO ₂	Fe ₂ O ₃	SO ₃	MnO	Cl
wt%	98.581	0.094	0.048	0.017	0.013	<0.01	<0.01	0.689



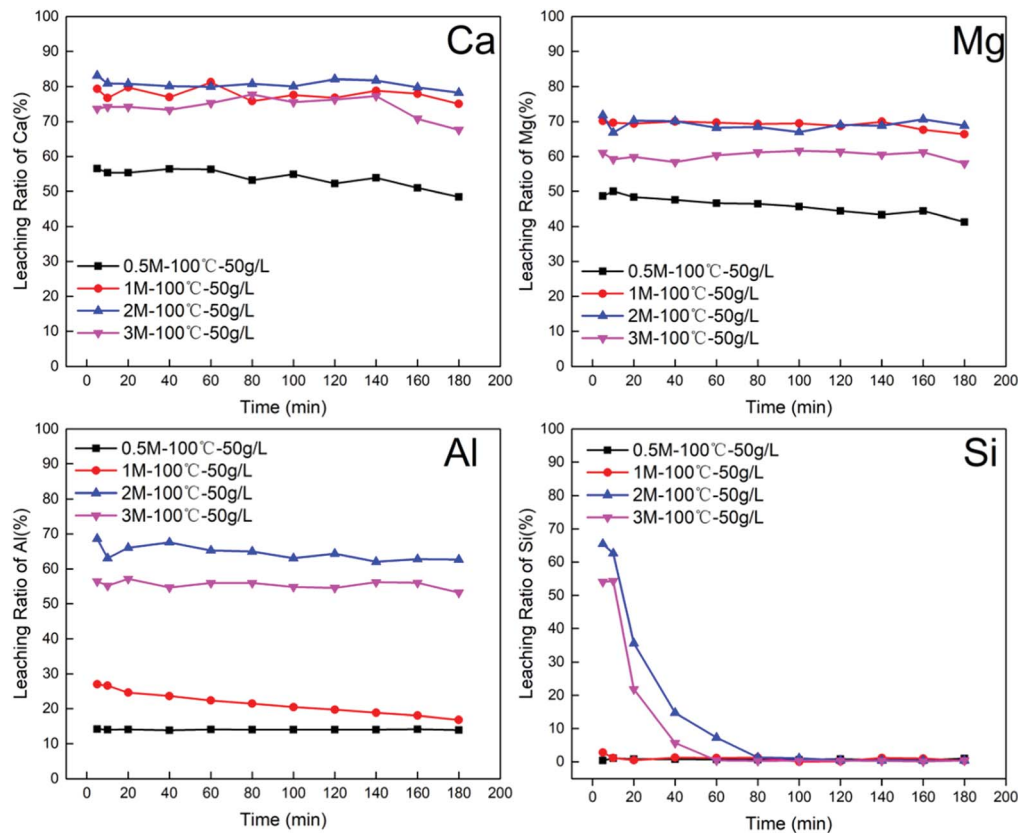
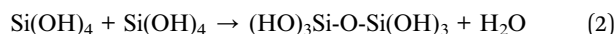


Fig. 2 Effect of the variation in the hydrochloric acid concentrations (0.5 M, 1 M, 2 M, 3 M) on the leaching ratio of Ca, Mg, Al, Si at condition: temperature of 100 °C, solid to liquid ratio of 50 g L⁻¹.



Another important index that also needs to be considered is the molar ratio of divalent cations to trivalent cations in the

leachate solution: the molar ratio should meet the general requirements (between 2 and 4)¹⁵ for the preparation of LDHs. The value of $n(\text{M}^{2+})/n(\text{M}^{3+})$ is strongly related to the hydrochloric acid concentration. With increased acid concentration, $n(\text{M}^{2+})/n(\text{M}^{3+})$ decreased, which suggests that the trivalent metal

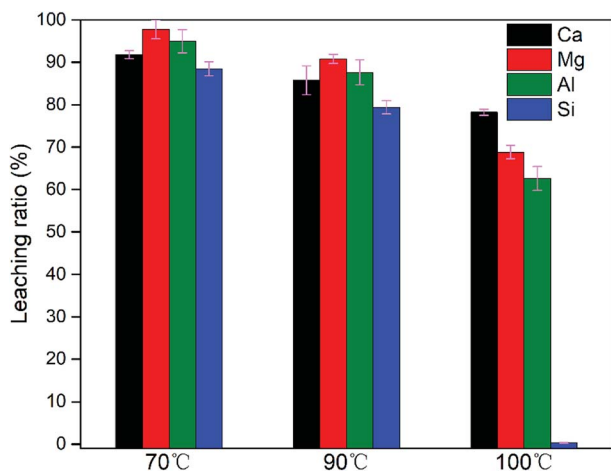


Fig. 3 Effect of leaching temperature (70 °C, 90 °C, 100 °C) on the leaching ratio of Ca, Mg, Al, Si at condition: acid concentration of 2 M, the solid to liquid ratio of 50 g L⁻¹, leaching time of 90 min.

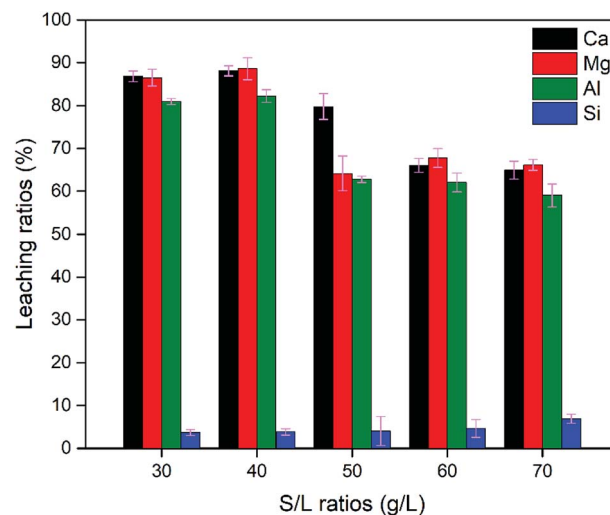


Fig. 4 Effect of the S/L ratio (30, 40, 50, 60, 70 g L⁻¹) on the leaching ratio of Ca, Mg, Al, Si at condition: temperature of 100 °C, acid concentration of 2 M, leaching time of 90 min.



Table 3 Elemental selectivity of four elements and $n(M^{2+})/n(M^{3+})$ in the leachate solution from different leaching conditions^a

Concentration of HCl (M)	Temperature (°C)	S/L ratios (g L ⁻¹)	Elemental selectivity (%)				$n(M^{2+})/n(M^{3+})$
			Ca ²⁺ (%)	Mg ²⁺ (%)	Al ³⁺ (%)	Si ⁴⁺ (%)	
0.5	100	50	67.94	22.36	8.52	1.17	10.60
1	100	50	69.36	23.59	6.81	0.24	13.65
2	100	50	58.99	20.05	20.73	0.23	3.81
3	100	50	59.55	19.67	20.47	0.31	3.87
2	70	50	35.90	15.45	17.13	31.52	3.00
2	90	50	36.45	15.63	17.16	30.77	3.04
2	100	30	52.70	21.01	24.11	2.17	3.06
2	100	40	50.89	20.38	22.91	5.82	3.11
2	100	60	50.54	20.96	23.47	5.03	3.05
2	100	70	52.33	22.08	24.10	1.49	3.09

^a Elemental selectivity (%) = $n_x/(n_{Ca} + n_{Mg} + n_{Al} + n_{Si}) \times 100\%$, x = Ca, Mg, Al, Si.

(Al³⁺) shows stronger resistance against acid attack compared with divalent alkali metals (Ca²⁺, Mg²⁺).

In summary, leaching conditions need to be rationally chosen for obtaining a high leaching ratio of elemental Ca, Mg and Al, low elemental selectivity for Si and a suitable $n(M^{2+})/n(M^{3+})$. Leaching conditions comprising 2 M acid concentration, 100 °C leaching temperature, 90 min leaching time and a S/L ratio of 40 g L⁻¹ allowed a high leaching ratio of Ca, Mg and Al to be achieved, at 88.08%, 88.59% and 82.27%, respectively, 5.82% of Si and a value of 3.11 for $n(M^{2+})/n(M^{3+})$.

3.2 Characterization of CaMgAl-LDHs sorbent

Fig. 5(a) shows the XRD pattern of as-synthesized CaMgAl-LDHs. All the sharp reflections with a high-degree of crystallinity were attributed to the successful formation of LDHs. The LDHs showed obvious crystallographic changes as a result of calcination at different temperatures in air. The sample maintained hydrotalcite structure when the calcination temperature was 200 °C. Once calcined at 400 °C, the sample showed weak reflections characteristic of hydrotalcite with a small amount of periclase, indicating decomposition of the layered structure. Calcination above 800 °C generated new crystalline phases assignable to CaO, MgO and mayenite (Ca₁₂Al₄O₃₃). The

thermal events of as-synthesized CaMgAl-LDHs from BFS was further studied by TGA analysis. The thermal decomposition of the CaMgAl-LDHs occurred in three main weight loss steps (ESI, Fig. S1†). In addition, this LDHs had a nominal structural formula of Ca_{1.89}Mg_{0.89}Al_{1.0}(OH)_{7.5}Cl_{0.83}(CO₃)_{0.11}· γ H₂O, and most of the intercalated anions were Cl⁻ (ESI, Table S1†).

The textural properties obtained from BET analysis are presented in Fig. 5(b). The specific surface area increased with increase in calcination temperature from 200 °C to 400 °C. The sample after calcination at 400 °C for 4 h had a specific surface area of 38.55 m² g⁻¹, which was twice as large as that of the LDHs. A slight increase in average pore diameter and pore volume were also observed. When the temperature further increased to 800 °C, the crystallisation of resulting oxides resulted in a decreased surface area and porosity.

SEM micrographs of as-synthesized LDHs calcined at different temperatures are shown in Fig. 6. Fig. 6(a) shows the formation of a typical plate-like structure with well-defined hexagonal shape.¹⁶ The particle size of LDHs was about 1 μm and the thickness was less than 40 nm. Fig. 6(b) shows that the plate-like particles stacked together after calcination at 200 °C. When the temperature further increased, calcination resulted in the disappearance of characteristic LDHs features with the formation of amorphous

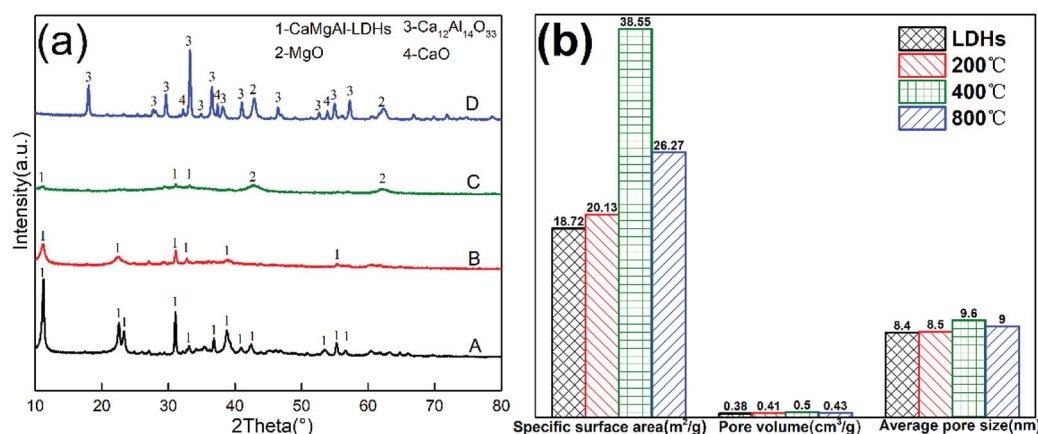


Fig. 5 (a) X-ray diffraction patterns of LDHs calcined at different temperatures for 4 h: (A) as-synthesized, (B) 200 °C, (C) 400 °C and (D) 800 °C (b) BET results of LDHs calcined at different temperatures.



mixed oxides (Fig. 6(c)), which further developed into mineral crystallites as a consequence of sintering events (Fig. 6(d)).¹⁷

3.3 Characterization of MCM-41 sorbent

The SEM image in Fig. 7(a) shows that the as-synthesized mesoporous silica nanoparticles on a scale of less than 200 nm had a good dispersity, with some coalescence into aggregates. The TEM micrograph in Fig. 7(b) reveals a highly ordered 2D hexagonal pore structure array of the MCM-41 type silica; this feature was also confirmed by the presence of (100), (110) and (200) plane reflections in Fig. 7(c).¹⁸ The diameter between the fringes of the pore in the mesoporous structure was estimated to be around 3 nm, which is in good agreement with the BJH pore size distribution analysis in Fig. 7(d), having a primary distribution in the range of 2.6–2.8 nm. The sorption isotherm in Fig. 7(d) shows a type IV isotherm according to the IUPAC classification with a sharp increase step in the relative pressure region of 0.31–0.35. This was attributed to capillary condensation within the narrow distribution of mesopores.¹⁹

Textural properties of MCM-41 including BET surface area (S_{BET}), total pore volume (V_{p}), average pore radius (D_{BJH}) and the wall thickness (b_{p}) are shown in Table 4. This material had a surface area of $1270 \text{ m}^2 \text{ g}^{-1}$, much higher than when synthesized from fly ash ($732 \text{ m}^2 \text{ g}^{-1}$),²⁰ iron ore tailings ($527 \text{ m}^2 \text{ g}^{-1}$),²¹ sodium silicate ($874 \text{ m}^2 \text{ g}^{-1}$)²² and pure TEOS ($1100 \text{ m}^2 \text{ g}^{-1}$).²⁰ The thickness of pore walls was comparable at 1.04 nm with results in the previous literature.²³

3.4 Evaluation of CO₂ capture performance

The CO₂ adsorption behaviour of as-synthesized Ca_{1.89}Mg_{0.89}-Al_{1.0}-LDHs and after calcination at 200 °C, 400 °C, 800 °C,

denoted as CaMgAl-LDHs-200, CaMgAl-LDHs-400, CaMgAl-LDHs-800, respectively, was measured at 400 °C. As seen from Fig. 8(a), after calcination at 400 °C for 4 h, CaMgAl-LDHs-400 exhibited a first contact capacity of 1.58 mmol g^{-1} , almost 2 times greater than that of CaMgAl-LDHs without thermal activation. However, calcination at 800 °C caused a great decline in the CO₂ adsorption. These results were in agreement with those of Ram Reddy *et al.*,²⁵ indicating that there exists an optimised thermal activation temperature for CaMgAl-LDHs. The events of dehydroxylation, decomposition of intercalated anions, followed by crystallization upon thermal calcination (confirmed by the XRD and TGA analyses) gave rise to the variation in surface area and a number of active basic sites, thus having a great influence on the CO₂ uptake.

The thermal stability of CaMgAl-LDHs-400 was evaluated through continuous CO₂ adsorption/desorption cycles. As shown in Fig. 8(b), CaMgAl-LDHs-400 demonstrates good reusability and cyclic performance throughout the 20 adsorption/desorption cycles. After 20 times cycles, the CaMgAl-LDHs-400 sorbent still adsorbs around 1.2 mmol g^{-1} of CO₂ at 400 °C and 1 atm, CO₂ adsorption capacity reduces 24%.

As depicted in Fig. 9(a), for MCM-41, the highest CO₂ adsorption capacity increases from 0.95 to 1.24 mmol g^{-1} when the adsorption temperature increases from 25 to 75 °C, which is a characteristic feature of physisorption processes.²⁶ However, the adsorption capacity decreased slightly to 1.13 mmol g^{-1} when the temperature exceeded 100 °C, which could be related to the dominant CO₂ desorption reaction at high temperature. The adsorption/desorption cycles for MCM-41 were conducted at 75 °C. As shown in Fig. 9(b), there was about 20% deterioration in the adsorption capacity after 20 adsorption/desorption cycles, indicating that the MCM-41 sorbents had a good thermal stability.

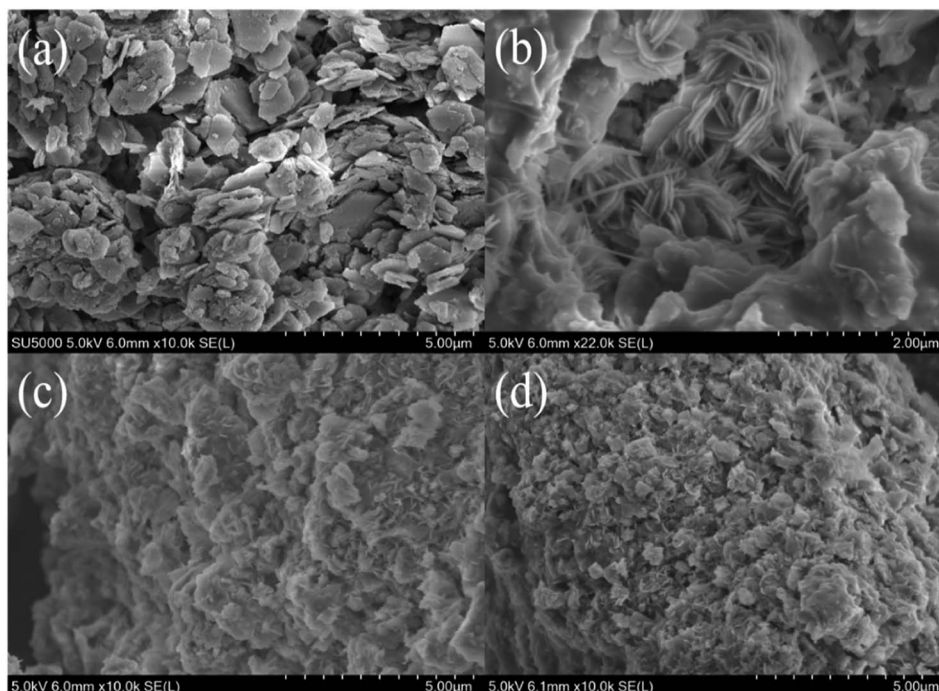


Fig. 6 SEM images of LDHs sorbents calcined at different temperatures: (a) as-synthesized, (b) 200 °C, (c) 400 °C and (d) 800 °C.



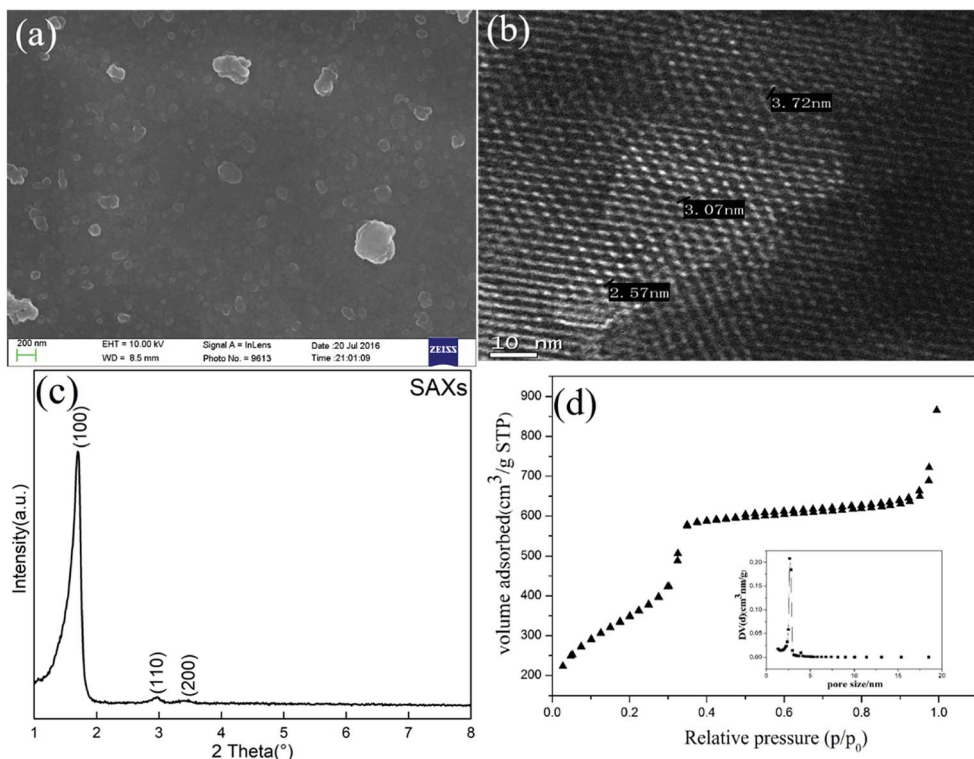


Fig. 7 (a) SEM image and (b) TEM image of MCM-41 sorbent prepared from BFS; (c) SAXs scattering pattern; (d) N₂ adsorption–desorption isotherm and pore size distribution.

Table 4 Textural properties of MCM-41 derived from BFS^a

Sample	S_{BET} (m ² g ⁻¹)	V_{p} (cm ³ g ⁻¹)	D_{BJH} (nm)	d_{100} (nm)	b_{p} (nm)
MCM-41	1270	1.342	3.034	3.531	1.04

^a S_{BET} , specific surface area calculated using BET equation. V_{p} , the total pore volumes were determined at a P/P_0 value of 0.995. D_{BJH} , the mean pore size distributions were determined by BJH mode applied to the desorption branches of the isotherms. b_{p} , the pore wall thickness, calculated from equation,²⁴ $b_{\text{p}} = 2d_{100}/3^{1/2} - D_{\text{BJH}}$, where d_{100} refers to the interplanar spacing of the (100) reflection by Bragg's equation ($2d_{100} \sin \theta = \lambda$).

3.5 Adsorptive mechanism for the two prepared composites for CO₂ capture

The composite stability for the prepared two composites was studied using the thermogravimetric analysis with results shown in Fig. 10. LDHs-400 sample was heated to 400 °C and maintained for a while, the weight loss was small. Similarly, MCM-41 sample was heated to 700 °C and maintained for a while, the weight loss was due to the adsorbed gas and water at the beginning and did not change as the temperature rose to 700 °C. This confirmed that both composites had good stability.

As depicted Fig. 8(a), the LDHs calcined at 400 °C has a maximum adsorption capacity, which was most likely due to

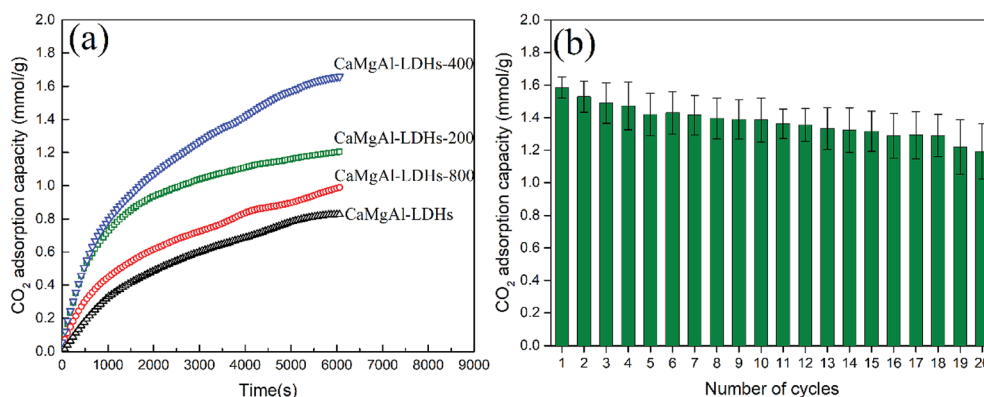


Fig. 8 (a) The effect of calcination temperature on the first CO₂ contact capacities of CaMgAl-LDHs sorbents adsorption temperature was set at 400 °C; (b) adsorption/desorption cycles for CaMgAl-LDHs-400 at 400 °C.



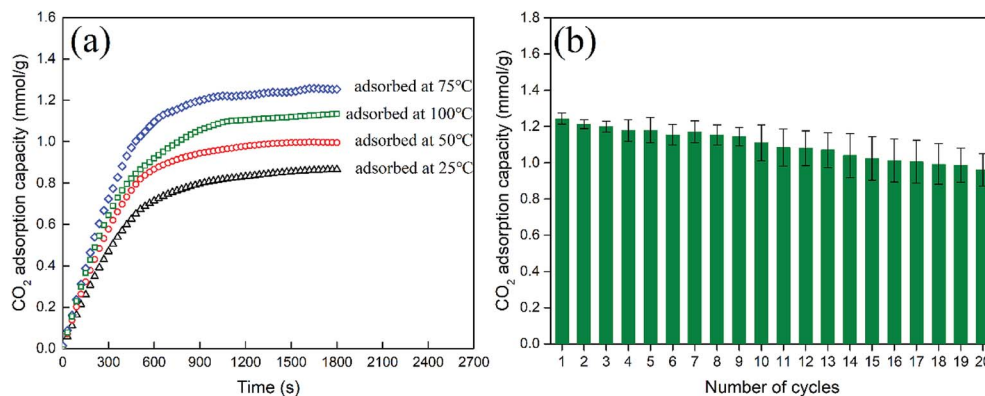


Fig. 9 (a) The first CO₂ contact capacities of MCM-41 sorbent at different temperatures (b) adsorption/desorption cycles for MCM-41(adsorbed at 75 °C, desorbed at 105 °C).

the optimum balance between the surface area and enough basic sites.²⁵ The BET analysis for LDHs calcined at different temperatures in Fig. 5(b) supported this theory. The LDHs-400 sample, which is an amorphous mixed oxide phase, had the maximum BET surface area compared to the other samples at 38.55 m² g⁻¹. The LDHs-200 sample had the minimum BET surface area because its layered structure was maintained, with CO₃²⁻ ions occupying the basic sites. Calcination at higher temperature (800 °C) decomposed most CO₃²⁻ ions to release some basic sites for CO₂ adsorption but resulted in a low BET surface area due to the phase transformation of amorphous mixed oxide into crystalline spinel, CaO and MgO phases. So, the LDHs-400 sample had not only the highest surface area but also enough basic sites for improving the CO₂ adsorption performance.

FTIR spectra patterns of CaMgAl-LDHs-400 and MCM-41 before and after adsorption are shown in Fig. 11. For LDHs-400, the reflection at 1633 cm⁻¹ for LDHs material was the δH₂O mode of interlayer water molecules.²⁷ The relatively weak CO₃²⁻ reflection at 1425–1475 cm⁻¹ indicated a low level of contamination with CO₂ before adsorption,²⁸ while after CO₂ adsorption the corresponding band located at around 1417 cm⁻¹ was strong. The AlO₆ stretching mode with the indicative reflection around 457 cm⁻¹ was observed after calcination.²⁶ The new reflection at 846 cm⁻¹ indicated the presence of Ca–O bonding, which was consistent with previous

results reported in the synthesis of hydroxalite with varied (Ca + Mg)/Al molar ratios.²⁹ After CO₂ adsorption, the samples exhibited extremely weak vibration peaks at 1633 cm⁻¹ and the disappearance of other vibration peaks. The disappearance of peaks related to metal–oxygen vibrations at low wavenumber of around 500 cm⁻¹ was due to the chemical reaction between basic sites and CO₂.

For MCM-41, the band around 3450 cm⁻¹ attributed to hydrogen bonded surface silanols (Si–OH) and adsorbed water molecules were due to O–H stretching and the band at 1633 cm⁻¹ was due to bending vibrations of adsorbed water molecules. The peaks centred at 1093 cm⁻¹, 804 cm⁻¹ and 460 cm⁻¹ corresponded to the asymmetric stretching, symmetric stretching and rocking vibrations of the Si–O–Si bonds, respectively. The band appearing at 956 cm⁻¹ was attributed to Si–O stretching in Si–OH. Silanol groups (Si–OH) on the surface of MCM-41 maybe the adsorption sites for capturing CO₂ molecules.³⁰ After CO₂ adsorption, the band at 956 cm⁻¹ disappeared and a very weak band at 1384 cm⁻¹ was observed for the symmetric stretching mode of linear CO₂ which verified that physisorbed CO₂ on mesoporous silica was detected.³¹ The intensity of band corresponded to OH stretching of free silanol groups around 3450 cm⁻¹ decreased after CO₂ adsorption and a weakly bonded adduct between silanol groups and CO₂: –OH···O=C=O formed which can easily break down as the temperature increases. As depicted Fig. 9(a), the CO₂

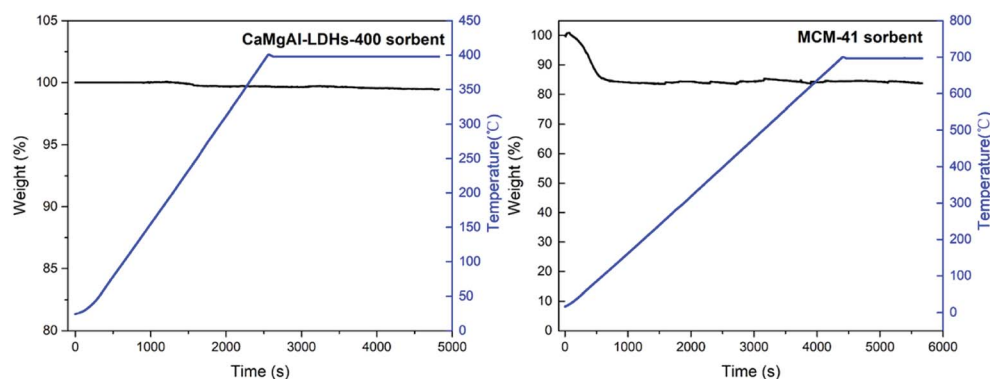


Fig. 10 Thermogravimetric analysis of CaMgAl-LDHs-400 and MCM-41.



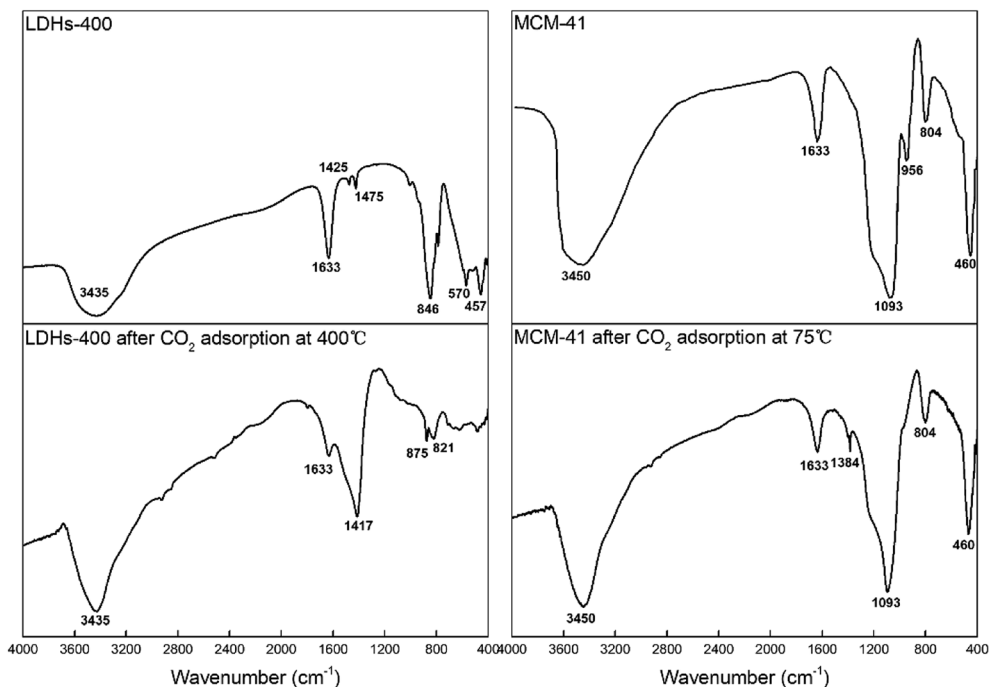


Fig. 11 FTIR spectra patterns of CaMgAl-LDHs-400 and MCM-41 before and after adsorption.

Table 5 Comparison of the CO₂ adsorption capacities of various materials

Sorbents	Materials	CO ₂ uptake	Ref.
LDHs	Mg ₃ Al-stearate-LDH-400	1.30 mmol g ⁻¹ at 300 °C, 1 bar	33
	Ca ₃ Al-LDH-550	0.87 mmol g ⁻¹ at 350 °C, 1 bar	34
	Mg ₃ Al-LDH-550	1.18 mmol g ⁻¹ at 350 °C, 1 bar	34
	CaMg ₂ Al-LDH-550	1.28 mmol g ⁻¹ at 350 °C, 1 bar	34
	Nano-sized 20 wt% K ₂ CO ₃ impregnated Mg ₃ Al-LDH-400	1.21 mmol g ⁻¹ at 200 °C, 1 bar	35
	Ca _{1.89} Mg _{0.89} Al _{1.0} -LDHs-400	1.66 mmol g ⁻¹ at 400 °C, 1 bar	In this study
Mesoporous silica	MCM-41/CPA	1.5 mmol g ⁻¹ at 25 °C, 1 bar	36
	MCM-48/CPA	1.1 mmol g ⁻¹ at 25 °C, 1 bar	36
	MCM-41-NH ₂	1.15 mmol g ⁻¹ at 30 °C, 2.1 bar	37
	TRI-PE-MCM-41	2.21 mmol g ⁻¹ at 55 °C, 1 bar	38
	MCM-41	1.25 mmol g ⁻¹ at 75 °C, 1 bar	In this study

adsorption capacity at higher temperature (100 °C) decreased because interaction between CO₂ and silanol groups is van der Waals force, which does not favor the adsorption at higher temperatures.³²

3.6 Comparison of the CO₂ adsorption capacities of various materials

Different kinds of LDHs and mesoporous silica and their CO₂ adsorption performances are compared in Table 5. The maximum CO₂ adsorption capacity of CaMgAl-LDHs derived from BFS is 1.66 mmol g⁻¹, which is higher than that of other LDHs synthesized using pure reagents. The maximum CO₂ adsorption capacity of MCM-41 derived from BFS was 1.25 mmol g⁻¹, which is comparable to other mesoporous silica synthesized from other materials. Comparing the results of this study with those reported in literature, BFS was found to be a promising raw material for CaMgAl-LDHs and MCM-41

production and their further use as efficient sorbents for capturing CO₂.

4. Conclusions

In this work, the conversion of blast furnace slag into two different kinds of promising sorbents of CO₂ has been presented. The calcium, magnesium, aluminium in BFS were selectively extracted by hydrochloric acid and used for preparation of layered double hydroxides (LDHs). The leaching residue which contained a high amount of silica was utilised for MCM-41 material synthesis. Several operating variables including initial hydrochloric acid concentration, leaching temperature, solid-to-liquid ratio, and leaching time were optimised. It was found that higher hydrochloric acid concentration, higher leaching temperature, and longer leaching time could promote silica gel formation; this provided a great



advantage for the selective leaching out of calcium, magnesium, aluminium with high efficiency. Characterisation of the as-synthesized CaMgAl-LDHs and MCM-41 sorbents was performed to determine chemical composition, morphology and textural properties. The structural formula of CaMgAl-LDHs was determined as $\text{Ca}_{1.89}\text{Mg}_{0.89}\text{Al}_{1.0}(\text{OH})_{7.5}\text{Cl}_{0.83}(\text{CO}_3)_{0.11} \cdot \gamma\text{H}_2\text{O}$, with Cl^- as the predominate intercalated anion. Thermal calcination at 400 °C enhanced the specific surface area, thus leading to improved CO_2 uptake and a maximum CO_2 adsorption capacity of 1.58 mmol g^{-1} at 400 °C. The MCM-41 sorbents derived from BFS exhibited a high surface area of 1270 $\text{m}^2 \text{g}^{-1}$ and showed comparable CO_2 adsorption capacity with a value of 1.24 mmol g^{-1} at 75 °C. In the CO_2 adsorption/desorption cycles, both sorbents showed good reusability and cyclic performance. With all these results, the CaMgAl-LDHs and MCM-41 sorbents derived from blast furnace slag are promising sorbents for practical applications in the field of CO_2 capture.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The National Natural Science Foundation of China (No. 51574169, No. 51604178, No. 51704202, No. 51604202, No. 51774209), the Natural Science Foundation of the Jiangsu Higher Education Institutions of China (No. 17KJB450002), are duly acknowledged for their financial support. The authors also are grateful for the analysis support from Soochow University.

References

- 1 S. Lee, J. W. Kim, S. Chae, J. H. Bang and S. W. Lee, *J. CO₂ Util.*, 2016, **16**, 336–345.
- 2 S. Teir, S. Eloneva, C. J. Fogelholm and R. Zevenhoven, *Energy*, 2007, **32**, 528–539.
- 3 S. Eloneva, S. Teir, J. Salminen, C. J. Fogelholm and R. Zevenhoven, *Energy*, 2008, **33**, 1461–1467.
- 4 S. Eloneva, S. Teir, J. Salminen, C. J. Fogelholm and R. Zevenhoven, *Ind. Eng. Chem. Res.*, 2008, **47**, 7104–7111.
- 5 S. Eloneva, S. Teir, J. Savolahti, C. Fogelholm and R. Zevenhoven, *Proceeding ECOS 2007*, Padova, Italy, 2007, vol. II, pp. 1389–1396.
- 6 C. Hall, D. J. Large, B. Adderley and H. M. West, *Miner. Eng.*, 2014, **65**, 156–162.
- 7 A. jun Xu, H. ning Zhang, Y. Yang, J. Cui, D. feng He and N. yuan Tian, *J. Iron Steel Res. Int.*, 2012, **19**, 34–38.
- 8 J. H. Bang, S. W. Lee, C. Jeon, S. Park, K. Song, W. J. Jo and S. Chae, *Energies*, 2016, **9**(12), 996.
- 9 W. Bao, H. Li and Z. Yi, *Ind. Eng. Chem. Res.*, 2010, **49**, 2055–2063.
- 10 J. Wang, L. Huang, R. Yang, Z. Zhang, J. Wu, Y. Gao, Q. Wang, D. O'Hare and Z. Zhong, *Energy Environ. Sci.*, 2014, **7**, 3478–3518.
- 11 S. H. Wang, Y. Bin Wang, Y. M. Dai and J. M. Jehng, *Appl. Catal., A*, 2012, **439–440**, 135–141.
- 12 S. Tian, J. Jiang, F. Yan, K. Li, X. Chen and V. Manovic, *Green Chem.*, 2016, **18**, 4022–4031.
- 13 Y. Kuwahara and H. Yamashita, *ISIJ Int.*, 2015, **55**, 1531–1537.
- 14 K. Song, S. Park, W. Kim, C. Jeon and J.-W. Ahn, *Metals*, 2017, **7**, 199.
- 15 Q. Wang and D. Ohare, *Chem. Rev.*, 2012, **112**, 4124–4155.
- 16 A. P. Tathod and O. M. Gazit, *Cryst. Growth Des.*, 2016, **16**, 6709–6713.
- 17 A. C. Vieira, R. L. Moreira and A. Dias, *J. Phys. Chem. C*, 2009, **113**, 13358–13368.
- 18 K. Flodström and V. Alfredsson, *Microporous Mesoporous Mater.*, 2003, **59**, 167–176.
- 19 P. Selvam, S. K. Bhatia and C. G. Sonwane, *Ind. Eng. Chem. Res.*, 2001, **40**, 3237–3261.
- 20 H. Misran, R. Singh, S. Begum and M. A. Yarmo, *J. Mater. Process. Technol.*, 2007, **186**, 8–13.
- 21 H. Yu, X. Xue and D. Huang, *Mater. Res. Bull.*, 2009, **44**, 2112–2115.
- 22 Y. Y. Zhou, X. xuan Li and Z. xing Chen, *Powder Technol.*, 2012, **226**, 239–245.
- 23 P. Kumar, N. Mal, Y. Oumi, K. Yamana and T. Sano, *J. Mater. Chem.*, 2001, **11**, 3285–3290.
- 24 D. Zhao, Q. Huo, J. Feng, B. F. Chmelka and G. D. Stucky, *J. Am. Chem. Soc.*, 1998, **120**, 6024–6036.
- 25 M. K. Ram Reddy, Z. P. Xu, G. Q. Lu and J. C. D. Da Costa, *Ind. Eng. Chem. Res.*, 2006, **45**, 7504–7509.
- 26 R. L. Frost, H. J. Spratt and S. J. Palmer, *Spectrochim. Acta, Part A*, 2009, **72**, 984–988.
- 27 R. Galindo, A. López-Delgado, I. Padilla and M. Yates, *Appl. Clay Sci.*, 2014, **95**, 41–49.
- 28 A. Boumaza, L. Favaro, J. Lédion, G. Sattonnay, J. B. Brubach, P. Berthet, A. M. Huntz, P. Roy and R. Tétot, *J. Solid State Chem.*, 2009, **182**, 1171–1176.
- 29 J. Zhou, Z. P. Xu, S. Qiao, Q. Liu, Y. Xu and G. Qian, *J. Hazard. Mater.*, 2011, **189**, 586–594.
- 30 W. Wang, J. Li, X. Wei, J. Ding, H. Feng, J. Yan and J. Yang, *Appl. Energy*, 2015, **142**, 221–228.
- 31 Z. Bacsik, N. Ahlsten, A. Ziadi, G. Zhao, A. E. Garcia-Bennett, B. Martín-Matute and N. Hedin, *Langmuir*, 2011, **27**, 11118–11128.
- 32 T. Witoon and M. Chareonpanich, *Songklanakarin J. Sci. Technol.*, 2012, **34**, 403–407.
- 33 Q. Wang, H. H. Tay, Z. Zhong, J. Luo and A. Borgna, *Energy Environ. Sci.*, 2012, **5**, 7526–7530.
- 34 P. W. Xiao, J. Y. Jun, J. Cheng, P. H. Zheng and P. X. Zhi, *Environ. Sci. Technol.*, 2008, **42**, 614–618.
- 35 Q. Wang, Y. Gao, J. Luo, Z. Zhong, A. Borgna, Z. Guo and D. O'Hare, *RSC Adv.*, 2013, **3**, 3414–3420.
- 36 M. Bhagiyalakshmi, L. J. Yun, R. Anuradha and H. T. Jang, *J. Porous Mater.*, 2010, **17**, 475–484.
- 37 M. R. Mello, D. Phanon, G. Q. Silveira, P. L. Llewellyn and C. M. Ronconi, *Microporous Mesoporous Mater.*, 2011, **143**, 174–179.
- 38 Y. Belmabkhout, R. Serna-Guerrero and A. Sayari, *Adsorption*, 2011, **17**, 395–401.

