

## EDGE ARTICLE



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Hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs as catalyst precursors for CO<sub>2</sub> hydrogenation to methanol†Tomasz Kondratowicz,<sup>a</sup> Marta Gajewska,<sup>b</sup> Jiangtong Li,<sup>c</sup> Molly Meng-Jung Li,<sup>c</sup> Zoë R. Turner,<sup>a</sup> Chunping Chen<sup>\*a</sup> and Dermot O'Hare<sup>\*a</sup>

We report a new synthetic strategy for preparing well-organised, spherical and mesoporous, mixed-metal, hollow-core@layered double hydroxides. Hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs ( $x + y + z = 2.32 \pm 0.06$ ) were prepared by exploiting a unique “memory effect” feature of LDH materials. The reconstruction with simultaneous incorporation of Cu<sup>2+</sup> and Zn<sup>2+</sup> into the LDH shell was achieved by exposing hollow-SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDO to an aqueous solution containing Cu<sup>2+</sup> and Zn<sup>2+</sup> cations. The effect of a single reconstruction step with various concentrations of Cu<sup>2+</sup> and Zn<sup>2+</sup> solutions (20–80 mM), as well as the implementation of five successive cycles of calcination–reconstruction on the chemical composition, morphology, texture and structure of the resulting materials are described. Hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs are precursors to active catalysts for CO<sub>2</sub> hydrogenation to methanol. The most active catalyst exhibits a space-time yield for methanol of 1.68 g<sub>MeOH</sub> g<sub>Cu</sub><sup>-1</sup> h<sup>-1</sup> at 270 °C (3 : 1 CO<sub>2</sub> : H<sub>2</sub>, 30 bar) which represents a 1.7-fold increase in space-time yield compared to commercial Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> catalyst under the same conditions.

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## Introduction

Over several decades, rapid development of civilisation and industrial expansion have sharply increased carbon dioxide (CO<sub>2</sub>) emissions, reaching approximately 35 billion tons in 2021.<sup>1</sup> This has led to serious environmental issues including global warming, acid rain, ocean acidification and rising sea levels. The primary challenge now is to mitigate anthropogenic CO<sub>2</sub> emissions by (i) improving energy efficiencies to reduce CO<sub>2</sub> production, and (ii) implementing carbon capture and storage (CCS) and/or carbon capture and utilisation (CCU) technologies to manage the emissions effectively.<sup>2,3</sup> CCS technology typically involves capturing CO<sub>2</sub> from gas streams (*e.g.*, flue gas) followed by storing it underground. However, this requires high investment and poses risks such as potential leakage of CO<sub>2</sub>.<sup>4</sup> Unlike CCS, CCU upgrades CO<sub>2</sub> waste into valuable products, such as polymers, chemicals and fuels.<sup>3–5</sup> In particular, methanol has been proposed as a promising product from CO<sub>2</sub> hydrogenation. Methanol has the potential to play a dual role in both the energy and chemicals sectors as highlighted by “methanol economy” concept.<sup>6</sup> It can be used directly

as fuel or blended with conventional fuels. It is also a key feedstock for the production of chemicals such as formaldehyde, dimethyl ether and acetic acid.<sup>7</sup> As one of the ten most important industrial chemicals, global methanol production exceeded 110 million tons in 2018, with expectation to double by 2030.<sup>8</sup> Since the 1960s, methanol has been produced on an industrial scale from CO-rich syn-gas (CO/CO<sub>2</sub>/H<sub>2</sub>) generated from fossil fuels, using a Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> catalyst at 3–5 MPa pressure and 200–300 °C.<sup>9</sup> This well-established process can be readily used for the CO<sub>2</sub> hydrogenation to produce green methanol, using captured CO<sub>2</sub> and green hydrogen. Such approach has already been demonstrated at pilot-scale plants in Europe and Asia.<sup>2,10,11</sup>

It is widely accepted that metallic copper (Cu<sup>0</sup>) produces the active sites for CO<sub>2</sub> hydrogenation to methanol, with a well-established relationship between the addressable specific Cu<sup>0</sup> surface area and catalyst performance.<sup>12,13</sup> The Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> combination remains the most widely used catalyst system for methanol production *via* CO<sub>2</sub> hydrogenation. ZnO acts as Cu promoter which can enhance and stabilise the Cu dispersion and adsorption of CO<sub>2</sub>, while Al<sub>2</sub>O<sub>3</sub> can enhance thermal and chemical stability. In addition to these core components, various promoters such as Cr, Ga, Zr, Mg, Sr, Ba, and others have been extensively studied to improve catalytic performance by enhancing Cu dispersion, reducing Cu<sup>0</sup> particle size, increasing Cu stability and reducibility.<sup>14–18</sup> Among these, Mg has garnered particular attention due to its ability to modify the basicity for CO<sub>2</sub> adsorption and stabilise the Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> interface. Recently, layered double hydroxides (LDHs) have

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been shown to offer great potential as precursors in making robust Cu/Zn<sub>x</sub>AlO<sub>y</sub> catalysts. Excellent metal dispersions are achieved by the incorporation of Cu, Zn, and Al into the LDH metal hydroxide layers.<sup>19–21</sup> LDHs are a large family of anionic layer materials, the most frequently observed formulation can be written as  $[M_{1-x}^{II}M_x^{III}(\text{OH})_2]^{x+}[(A^{n-})_{x/n}] \cdot m\text{H}_2\text{O}$ , consisting of positively-charged mixed metal hydroxide layers ( $[M_{1-x}^{II}M_x^{III}(\text{OH})_2]^{x+}$ ) intercalated with hydrated charge compensating anions ( $[(A^{n-})_{x/n} \cdot m\text{H}_2\text{O}]$ ). LDHs offer advantages such as compositional flexibility and atomic-scale dispersion of metal ions within the layers.<sup>22</sup> Upon calcination between 300–600 °C, LDHs transform into homogeneous mixed metal oxides, also known as layered double oxides (LDOs),<sup>23</sup> which exhibit an unique “memory effect” that may allow reconstruction of the original layered structure in the presence of water, either in the gas or liquid phase.<sup>24,25</sup> This behavior has allowed incorporation of new metal cations into the LDH matrix, simply *via* exposing the LDO to aqueous cations solutions.<sup>26–28</sup> Despite the well-established, cost-effective methods for LDH synthesis, conventional techniques often result in aggregated powders with poor morphology and irregular particle size (so-called stone-like morphology), limiting their porosity and practical applications.<sup>29</sup> One approach to mitigate these limitations is the decoration of LDH platelets on well-defined core inorganic or organic supports, *e.g.* SiO<sub>2</sub>,<sup>30</sup> Fe<sub>3</sub>O<sub>4</sub>,<sup>31</sup> TiO<sub>2</sub>,<sup>32</sup> Cu<sub>2</sub>O,<sup>33</sup> zeolites,<sup>34</sup> MOF<sup>35</sup> and carbon<sup>36</sup> to create core-shell hybrids with shapes including spheres,<sup>30</sup> cubes,<sup>33</sup> wires<sup>37</sup> and rods.<sup>38</sup> Notably, core-shell structures typically contain vertically oriented LDH platelets forming a three-dimensional (3D) honeycomb-like LDH shell that exhibits enhanced textural properties and active site accessibility compared to unsupported LDH materials.<sup>39,40</sup> Additionally, such hybrids can serve as templates for hollow structures,<sup>40–42</sup> which have demonstrated enhanced properties over the parent core-shell materials such as higher porosity and concentration of basic sites.<sup>42</sup>

Previously, we have reported the synthesis of silica@Cu<sub>x</sub>-ZnAl-LDH ( $x = 0.8\text{--}4.0$ ) core-shell hybrids, using commercially available ES757 silica, mesoporous MCM-48, and SBA-16 as a core, with various Cu<sub>x</sub>ZnAl-LDHs used as a shell.<sup>20</sup> Upon thermal activation, these core-shell materials exhibited significantly improved catalytic performance for CO<sub>2</sub> hydrogenation to methanol compared to their equivalent unsupported Cu<sub>x</sub>-ZnAl LDH precursors. Hollowed core@shell materials offer the advantage of greater catalytic activity per mass of catalyst as a result of reduction in inactive core, as well as potential for

improved gas diffusion through the catalyst. Herein, we report a new advance by developing a novel strategy for the synthesis of hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDH catalyst precursors for CO<sub>2</sub> hydrogenation to methanol.

## Results and discussion

### Synthetic strategy to obtain hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs

The concept for developing mixed-metal hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs is schematically shown in Fig. 1. In the first step, solid SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDH core-shell particles (S@MA-LDH) were synthesised by the pH controlled coprecipitation of Mg<sub>2</sub>Al-CO<sub>3</sub> LDH in the presence of non-porous spherical SiO<sub>2</sub> particles. The silica core from SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDH was then selectively etched under static conditions by treatment with a NaOH solution to yield hollow-SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDH spheres (H-S@MA-LDH), followed by calcination to produce hollow-SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDO spheres (H-S@MA-LDO), according to our previous protocol.<sup>42</sup> Afterwards, the hollow-SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDO spheres were added to aqueous Cu and Zn nitrate solutions, facilitating ion exchange between the transition metals and Mg<sup>2+</sup> cations during reconstruction of the Mg<sub>2</sub>Al-LDO into Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDH through the memory effect, to obtain hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDH (H-S@CZMA-LDH<sub>X</sub>, where X depends on applied reconstruction conditions). The impact of a single reconstruction step using various concentrations of Cu and Zn solutions, as well as the implementation of repeated calcination and reconstruction cycles on the chemical composition, morphology, structural and textural properties of the hollow sphere materials has been studied.

### Synthesis of hollow-SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDH spheres

Fig. 2 shows the transmission electron microscope (TEM) images of pristine solid S@MA-LDH core-shell and the H-S@MA-LDH hollow spheres obtained after the leaching of the SiO<sub>2</sub> core. It can be clearly observed that sample before etching (Fig. 2A) exhibits typical core-shell structure, with a dark solid core sphere surrounded by a bright shell composed of vertically grown hierarchical nanosheets of LDH. The average diameter of the core-shell particles is 562 ± 25 nm, of which the SiO<sub>2</sub> core diameter is 340 ± 9 nm and the LDH shell thickness is 112 ± 22 nm, consistent with S@MA-LDH prepared previously.<sup>37</sup> The EDX elemental maps in Fig. 2C clearly demonstrate that Si is mainly distributed in the core while Mg and Al are uniformly dispersed across the shell, confirming the formation of core-

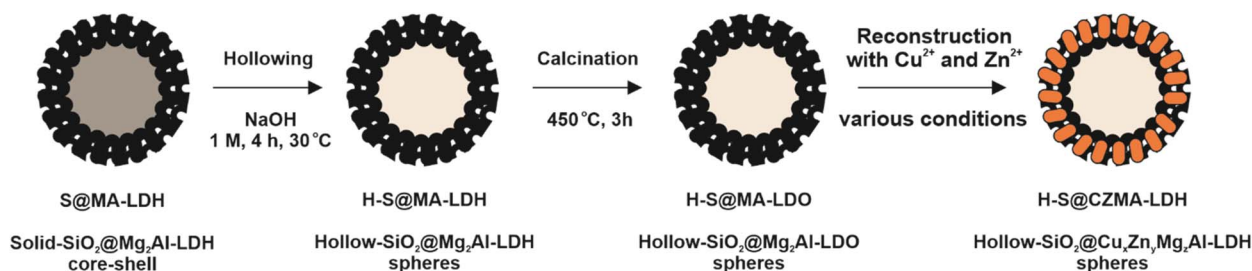


Fig. 1 Synthetic scheme used to obtain mixed-metal (Cu, Zn) hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDH spheres.

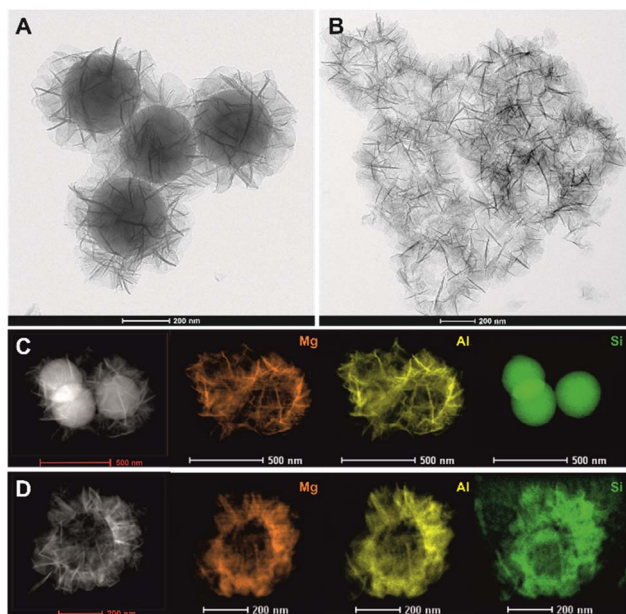


Fig. 2 TEM images of S@MA-LDH solid core-shell (A), H-S@MA-LDH hollow spheres (B) and Mg, Al, Si elemental EDX mapping of S@MA-LDH (C) and H-S@MA-LDH (D).

shell structure with SiO<sub>2</sub> as core and Mg<sub>2</sub>Al-LDH as shell. After 4 h of immersion in an alkali solution, the SiO<sub>2</sub> core was removed as evidenced in Fig. 2B, resulting in hollow shell structure with a clearly visible void inside the sphere while retaining the original particle size and spherical shape of the parent S@MA-LDH. The disappearance of contrast between the core and the shell confirms the high efficiency of the core removal. The elemental mapping analysis (Fig. 2D) further support this, as Si is absent in the core region but appears in the shell, likely due to partial deposition of Si species after leaching process.

The elemental compositions were further quantitatively analysed using elemental microanalysis (CHN), energy dispersive X-ray spectroscopy (EDX), inductively coupled plasma optical emission spectroscopy (ICP-OES) and thermogravimetric analysis (TGA). As shown in Table S1,† both EDX and ICP data reveal that Mg/Al molar ratio remained similar before and after the leaching process. However, the molar ratio of Si/Al significantly drops from 3.77 to 0.68, indicating that most SiO<sub>2</sub> has been removed. This is consistent with TGA analysis showing that the SiO<sub>2</sub> content decreases from 48.00 wt% to 17.07 wt%. The presence of SiO<sub>2</sub> in the hollow spheres, even at longer leaching times (up to 48 h) or at much harsh leaching conditions (2 M, 50 °C, 20 h),<sup>42</sup> is likely related to the formation of Si–O–Al (Mg) bonds.<sup>43</sup> Using the EDX, TGA and CHN data, we can estimate the chemical formula of S@MA-LDH and H-S@MA-LDH as [SiO<sub>2</sub>]<sub>0.81</sub>@[Mg<sub>2.18</sub>Al<sub>1.00</sub>(OH)<sub>6.36</sub>(CO<sub>3</sub>)<sub>0.13</sub>(-OH)<sub>0.74</sub>(H<sub>2</sub>O)<sub>3.64</sub>]<sub>0.19</sub> and [SiO<sub>2</sub>]<sub>0.44</sub>@[Mg<sub>2.16</sub>Al<sub>1.00</sub>(OH)<sub>6.32</sub>(CO<sub>3</sub>)<sub>0.25</sub>(OH)<sub>0.50</sub>(H<sub>2</sub>O)<sub>0.58</sub>]<sub>0.56</sub>, respectively.

The powder X-ray diffraction (XRD) patterns are shown in Fig. S1.† The virgin core-shell sample presents a combination of the characteristic features of both silica (amorphous

structure, broad feature at  $2\theta = 22^\circ$ ) and crystalline LDH (Bragg peaks at  $2\theta = 11.5^\circ, 23.2^\circ, 34.8^\circ, 39.3^\circ, 46.7^\circ, 60.8^\circ$ , and  $62.1^\circ$ ), which could be assigned to the (003), (006), (012), (015), (018), (110), and (113) reflections, typical of the double-layered structure in the trigonal  $R\bar{3}m$  space group.<sup>44</sup> After 4 h of leaching, the broad feature at  $2\theta = 22^\circ$  disappears, while the positions of the LDH Bragg reflections remain unchanged. However, the relative intensities slightly increase in H-S@MA-LDH, due to its higher proportion of crystalline LDH phase. The unit cell parameters  $a$ ,  $c$ , LDH basal spacing  $d(003)$ , and the crystallite domain size in the layer stacking direction,  $D_{(003)}$  and in the  $ab$ -plane  $D_{(110)}$  of H-S@MA-LDH are similar to that of S@MA-LDH (Table S2,†), which confirms that applied core leaching process presents minimal impact on the structural properties of the LDH phase.

The N<sub>2</sub> adsorption isotherms of S@MA-LDH and H-S@MA-LDH are shown in Fig. S2,† and the determined textural parameters are listed in Table S3.† The shape of both isotherms is similar to IVa type, according to the IUPAC classification, with H3 hysteresis loops characteristic for mesoporous materials with slit-shape pores.<sup>45</sup> Importantly, leaching of the predominant amount of the SiO<sub>2</sub> template from S@MA-LDH leads to an increase in the porosity for the H-S@MA-LDH spheres, the mesopore ( $V_{\text{meso}}$ ) and total pore ( $V_{\text{total}}$ ) volume increase from 0.38 to 0.66 cm<sup>3</sup> g<sup>-1</sup> and 0.50 to 0.89 cm<sup>3</sup> g<sup>-1</sup>, respectively. Furthermore, the N<sub>2</sub> BET surface area ( $S_{\text{BET}}$ ) doubles from 108 to 224 m<sup>2</sup> g<sup>-1</sup>.

### Synthesis of hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs through use of the “memory effect”

**Single immersion.** It is well-known that LDH materials possess significantly lower cation exchange ability compared to their associated LDO.<sup>46</sup> Therefore, the H-S@MA-LDH spheres were first calcined (properties of calcined H-S@MA-LDO are summarised in Fig. S3,†), and then the M<sup>2+</sup>:M<sup>3+</sup> brucite layer composition manipulated through use of the memory effect. The reconstruction and simultaneous incorporation of Cu<sup>2+</sup> and Zn<sup>2+</sup> into the brucite layers of the LDH are achieved by treating H-S@MA-LDO with aqueous solutions containing Cu<sup>2+</sup> and Zn<sup>2+</sup> cations at total concentrations of 20, 40, 60 and 80 mM (molar ratio of Cu to Zn = 1.30) at room temperature, which promotes a slow rehydration/reconstruction process. The reconstructed samples were designated as H-S@CZMA-LDH<sub>x</sub> mM, where  $x$  is the concentration (mM) of the solution used. The elemental composition of the samples before and after a single modification were analysed using the ICP-OES. The results, expressed as Mg/Al, (Cu + Zn + Mg)/Al, and (Cu + Zn)/Al molar ratios, are presented in Fig. 3. A clear trend shows that as the concentration of metal (Cu<sup>2+</sup> + Zn<sup>2+</sup>) ions increases, the (Cu + Zn)/Al molar ratio in the reconstructed samples gradually increases, reaching values of 0.86, 1.52, 1.82 and 1.84, for 20, 40, 60 and 80 mM, respectively. Conversely, the Mg/Al ratio decreases from 2.28 to 1.52, 0.85, 0.53 and 0.41, respectively. These changes strongly suggest a more extensive exchange of divalent metal ions with higher concentration solutions. Notably, the (Cu + Zn)/Al ratio appears to reach a plateau after modification with 60 mM, indicating that further



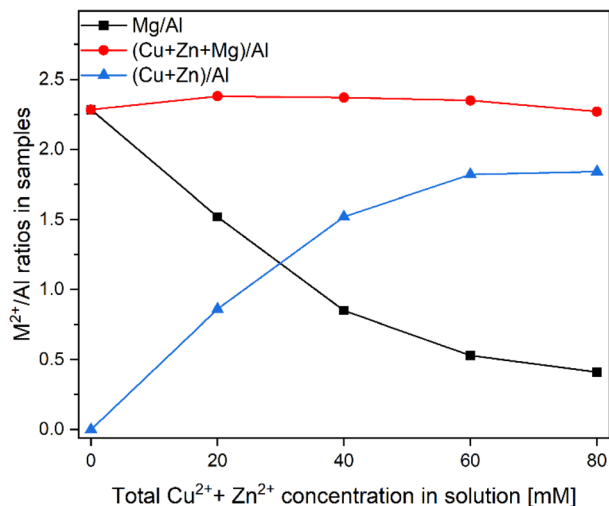


Fig. 3 Ratio of Mg/Al, (Cu + Zn + Mg)/Al and (Cu + Zn)/Al in H-S@CZMA-LDHs as a function of the initial aqueous  $[\text{Cu}^{2+}] + [\text{Zn}^{2+}]$ .

increasing the concentration of metal solution will not result in additional ion exchange as a maximum has been reached. Interestingly, the (Cu + Zn + Mg)/Al ratio, representing the total  $\text{M}^{2+}/\text{Al}^{3+}$  ratio, remains relatively constant (between 2.27 and 2.38), regardless of the metal concentration solutions. Moreover, these values are remarkably close to the initial Mg/Al ratio prior to the reconstruction process (Mg/Al = 2.28), suggesting that the exchange ratio between  $\text{Mg}^{2+}$  and  $(\text{Cu}^{2+} + \text{Zn}^{2+})$  is nearly equal to 1 : 1.

Importantly, after LDH reconstruction and simultaneous incorporation of  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  into the  $\text{Mg}_z\text{Al}$  layers, the particles still maintain their spherical morphology with a visible hollow core and hierarchical organised platelets on the thin  $\text{SiO}_2$  shell (Fig. 4A–D). A slight morphological degradation occurred when thicker and aggregated particles formed at high metal concentration (80 mM) (Fig. 4D). EDX elemental mapping has been carried out for H-S@CZMA-LDH\_20 mM (Fig. 4E); this sample represents an optimised balance between structural integrity with desired spherical morphology and elemental incorporation. A uniform elemental distribution (Mg, Al, Cu, Zn) is observed in the platelets across the spheres, confirming the incorporation of  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  ions into the LDH shell layer.

The XRD patterns (Fig. S4†) show that after reconstruction of H-S@MA-LDO in 20 mM solution, characteristic LDH Bragg diffraction features appear. However, when the concentration of the reconstruction solution increased beyond 40 mM, a less crystalline LDH phase becomes apparent, accompanied by a series of intense reflections corresponding to layered metal hydroxy salts (LHS), specifically  $\text{Cu}_2(\text{OH})_3\text{NO}_3$  (rouaite) with the monoclinic  $P2_1$  space group.<sup>47,48</sup> The formation of the LHS phase during reconstruction process is consistent with the findings of Wu *et al.*,<sup>46</sup> who studied the reconstruction of the bulky  $\text{Mg}_3\text{Al-LDO}$  (Mg/Al = 3) using Zn-nitrate solutions of different concentrations (12–108 mM). The presence of the LHS phase was also detected in samples when the concentration of reconstruction solution reaches 24 mM or higher. Moreover, the

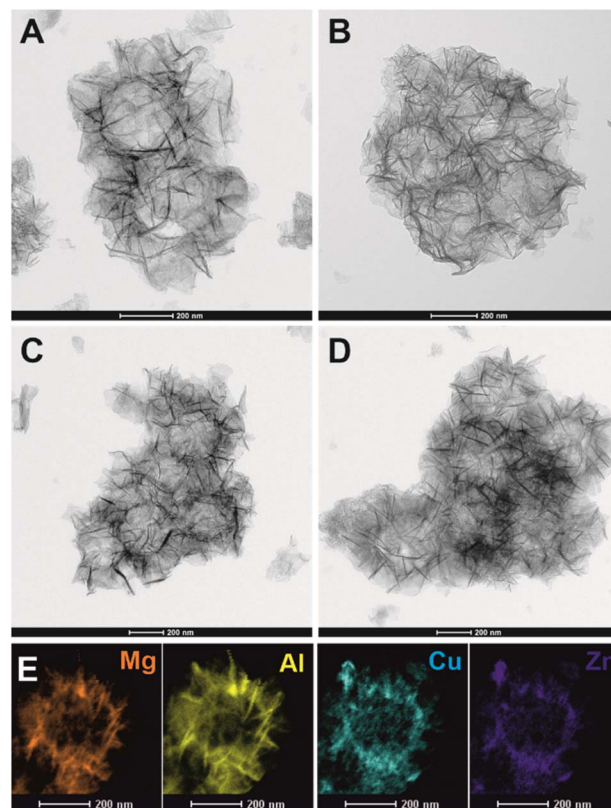


Fig. 4 TEM images of H-S@CZMA-LDHs obtained after immersing H-S@MA-LDO in solutions containing  $\text{Cu}^{2+} + \text{Zn}^{2+}$  at a concentration of 20 mM (A), 40 mM (B), 60 mM (C) and 80 mM (D) and Mg, Al, Cu, Zn elemental EDX mapping of H-S@CZMA-LDH\_20 mM (E).

intensity of reflections attributed to the LHS phase progressively increases with increasing the concentration of used solutions. A detailed phase analysis using LeBail refinement of the XRD data was performed using TOPAS software (Fig. S5, S6 and Table S4†). Good agreement was obtained using a multi-phase model containing  $\text{Cu}_x\text{Zn}_y\text{Mg}_z\text{Al-LDH}$ ,  $\text{Cu}_2(\text{OH})_3\text{NO}_3$  and silica. In general, a gradual increase in the content of the LHS phase is observed with the increase in the concentration of CuZn-containing solutions. The content of both LHS and LDH phases in the discussed H-S@CZMA-LDHs are 14.7, 23.9, 38.6 at% and 73.3, 62.5, 50.1 at%, respectively (Fig. S6†). Moreover, the silica content is around 11.3–13.6 at% which is lower than what we found from EDX result probably due to the existing of amorphous  $\text{SiO}_2$  phase.

The IR spectra of H-S@CZMA-LDHs as shown in Fig. S7† exhibit the characteristic bands of LDH: O–H stretching at  $3440\text{ cm}^{-1}$  from the hydroxide group and bending vibration from water molecules ( $1637\text{ cm}^{-1}$ ), indicating successful rehydration of LDO through the memory effect. It is worth pointing out that the pure LDH phase is reconstructed with the incorporation of  $\text{CO}_3^{2-}$  anions in the interlayer space at low CuZn concentration (20 mM), which is confirmed by the strong absorption band at  $1360\text{ cm}^{-1}$  (associated with  $\nu_3$  mode of  $\text{CO}_3^{2-}$ ).<sup>49</sup> However, at higher concentrations of Cu and Zn metal ion solutions (40–80 mM), new bands are observed at  $1336$  and  $1417\text{ cm}^{-1}$ , which correspond to the  $\nu_3$  mode of  $\text{NO}_3^-$ . Such

signals can be assigned to  $\text{Cu}_2(\text{OH})_3\text{NO}_3$  phase<sup>50,51</sup> as well as the anion exchanged nitrate in LDH phase. At the same time, the  $\text{CO}_3^{2-}$  signal at  $1360\text{ cm}^{-1}$  disappears or overlaps with the neighbouring bands, which does not exclude the presence of  $\text{CO}_3^{2-}$  ions in the samples.

After the reconstruction and incorporation with  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  cations, the adsorption–desorption isotherms (Fig. S8†) of the H-S@CZMA-LDHs remained type IVa with an H3 hysteresis loop according to IUPAC classification. However, the total pore volume and BET surface area (Table S3†) vary. The  $\text{N}_2$   $S_{\text{BET}}$  values for samples immersed in 20 mM and 40 mM solutions are comparable to those of the parent H-S@MA-LDH (225 and  $237\text{ vs. }224\text{ m}^2\text{ g}^{-1}$ ) while  $V_{\text{total}}$  slightly decreases ( $0.81$  and  $0.78\text{ vs. }0.89\text{ cm}^3\text{ g}^{-1}$ ). However, the samples modified in more concentrated solutions (60 and 80 mM) exhibit a significant reduction in  $S_{\text{BET}}$  and  $V_{\text{total}}$  ( $156$  and  $113\text{ m}^2\text{ g}^{-1}$ ,  $0.55$  and  $0.46\text{ cm}^3\text{ g}^{-1}$ , respectively). A similar trend is observed for the mesopore volume which gradually decreases from  $0.67$  to  $0.37\text{ cm}^3\text{ g}^{-1}$  with increasing metal concentration. This porosity reduction is likely due to the increasing presence of the highly crystalline  $\text{Cu}_2(\text{OH})_3\text{NO}_3$  phase (Fig. S4–S6†). In addition, gradually changing morphology of the samples (Fig. 4) may contribute to partial pore blockage.

**Calcination–reconstruction cycle.** As demonstrated above, a single immersion of H-S@MA-LDO spheres introduces significant amounts of  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  ions into LDH layers. However, this is accompanied by a gradual degradation of a hollow shell structure and the formation of undesirable phases at high concentration. To address this, we investigated the effect of repeated calcination–reconstruction cycles (five such repetitions were performed, and the samples were designated as H-S@CZMA-LDH<sub>Ry</sub>, where  $y = 1, 2, 3, 4$  or  $5$ , depending of cycle number) to obtain pure LDH material with high metal incorporation. A salt solution concentration of 15 mM was used and 1 g of the solid sample was mixed with 60 mL of the solution. After five cycles, the theoretical  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  content in the sample is similar to that of H-S@CZMA-LDH<sub>20 mM</sub> obtained from a single immersion approach in 20 mM solution ( $4.5\text{ vs. }4.0\text{ mmol}_{\text{Cu+Zn}}\text{ g}^{-1}$ ).

The elemental composition of the samples after each calcination–reconstruction cycle was analysed by ICP-OES as shown in Fig. S9.† The  $(\text{Cu} + \text{Zn})/\text{Al}$  molar ratio in the reconstructed samples steadily increases with each cycle, reaching 0.18, 0.36, 0.53, 0.74 and 0.96, while the  $\text{Mg}/\text{Al}$  ratio decreases from 2.28 to 2.11, 1.98, 1.78, 1.57 and 1.25. These trends suggest a stepwise exchange of divalent metal ions during the calcination–reconstruction cycle. In addition, the ratio of  $(\text{Cu} + \text{Zn} + \text{Mg})/\text{Al}$  remained relatively constant (2.21–2.34), regardless of the number of cycles. Given the initial  $\text{Mg}/\text{Al}$  of 2.28, the exchange ratio between  $\text{Mg}^{2+}$  and  $\text{Cu}^{2+} + \text{Zn}^{2+}$  appears to be 1 to 1, which is the same as that found in the single immersion approach. The  $\text{Cu}/\text{Zn}$  ratio across all sample was 1.42–1.44, closely matching the theoretical value of 1.30. Based on these results, the metal composition of the LDH in H-S@CZMA-LDHs after successive cycles is determined as follows:  $\text{Cu}_{0.10}\text{Zn}_{0.07}\text{Mg}_{2.11}\text{Al}$ ,  $\text{Cu}_{0.21}\text{Zn}_{0.15}\text{Mg}_{1.98}\text{Al}$ ,  $\text{Cu}_{0.31}\text{Zn}_{0.22}\text{Mg}_{1.78}\text{Al}$ ,  $\text{Cu}_{0.44}\text{Zn}_{0.31}\text{Mg}_{1.57}\text{Al}_{1.00}$ ,  $\text{Cu}_{0.56}\text{Zn}_{0.39}\text{Mg}_{1.25}\text{Al}$ , respectively.

The XRD patterns of the samples after each cycle are shown Fig. 5. In all cases, only distinct characteristic Bragg reflections for a crystalline LDH structure are observed. Notably, a systematic shift in the positions of (110) and (113) Bragg reflections toward lower  $2\theta$  values is observed as highlighted in the inset of Fig. 5. This shift is attributed to the differing ionic radii of  $\text{Mg}^{2+}$  compared to the  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  ions that replace it (octahedral ionic radii:  $\text{Mg}^{2+}$  0.72 Å,  $\text{Cu}^{2+}$  0.73 Å and  $\text{Zn}^{2+}$  0.74 Å).<sup>52</sup> The substitution affects the lattice parameters (Table 1). Particularly, the  $a$ -lattice parameter (the metal–metal distance in brucite-like layers) increases gradually from 0.3049 nm in the parent H-S@MA-LDH to 0.3073 nm after five cycles. This increase reflects the progressive replacement of  $\text{Mg}^{2+}$  ions with  $\text{Cu}^{2+}$  and  $\text{Zn}^{2+}$  ions, consistent with the ICP-OES analysis in Fig. S9.† The  $c$ -lattice parameter remains within 2.302–2.318 nm, corresponding to an LDH basal spacing of 7.67–7.73 Å with  $\text{CO}_3^{2-}$  ions in the interlayer space.<sup>53</sup> The presence of  $\text{CO}_3^{2-}$  anions in the interlayer space were further confirmed by IR spectroscopy (Fig. S10†). All reconstructed samples, regardless of the number of the calcination–reconstruction cycles, possess a similar spectral shape with a clearly distinct  $\nu(\text{C–O})$  stretching mode at  $1360\text{ cm}^{-1}$ .

The H-S@CZMA-LDHs exhibited similar porosity with nearly identical adsorption–desorption type IVa isotherms and H3 hysteresis loops (Fig. S11†). Their specific BET surface area and pore volumes are summarised in Table 1. The  $S_{\text{BET}}$  and  $V_{\text{total}}$  are in the range of  $158$ – $179\text{ m}^2\text{ g}^{-1}$  and  $0.66$ – $0.74\text{ cm}^3\text{ g}^{-1}$ , respectively. The mesopore volumes are between  $0.68$ – $0.75\text{ cm}^3\text{ g}^{-1}$ , with the highest volume obtained after the fourth

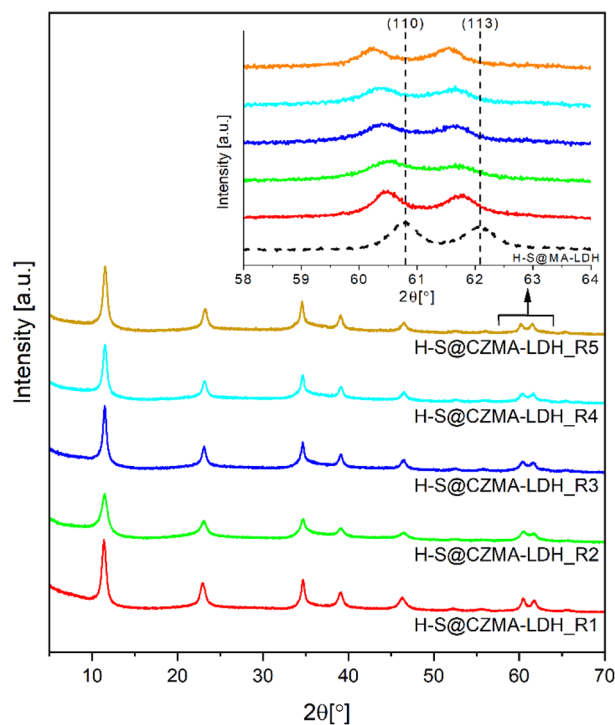


Fig. 5 XRD patterns for H-S@CZMA-LDHs after five successive iterations of calcination–reconstruction with a 15 mM CuZn-containing solution.

**Table 1** Crystal and textural properties of the H-S@MgAl-LDH and H-S@CZMA-LDHs after five successive iterations of calcination–reconstruction with a 15 mM CuZn-containing solution

Sample	LDH lattice parameters <sup>a</sup> [nm]		LDH <i>d</i> (003) [nm]	LDH crystallite size <sup>b</sup> [nm]		<i>S</i> <sub>BET</sub> [m <sup>2</sup> g <sup>-1</sup> ]	<i>V</i> <sub>micro</sub> [cm <sup>3</sup> g <sup>-1</sup> ]	<i>V</i> <sub>meso</sub> [cm <sup>3</sup> g <sup>-1</sup> ]	<i>V</i> <sub>total</sub> [cm <sup>3</sup> g <sup>-1</sup> ]
	<i>a</i>	<i>c</i>		<i>D</i> <sub>(003)</sub>	<i>D</i> <sub>(110)</sub>				
H-S@MA-LDH	0.3049	2.308	0.769	7.9	17.9	224	0.008	0.66	0.89
H-S@CZMA-LDH_R1	0.3062	2.318	0.773	13.9	15.0	163	0.009	0.68	0.67
H-S@CZMA-LDH_R2	0.3061	2.318	0.773	9.8	9.2	157	0.012	0.70	0.66
H-S@CZMA-LDH_R3	0.3065	2.312	0.771	15.8	9.9	158	0.012	0.74	0.73
H-S@CZMA-LDH_R4	0.3067	2.306	0.769	14.8	10.9	158	0.010	0.75	0.74
H-S@CZMA-LDH_R5	0.3073	2.302	0.767	14.2	13.5	179	0.002	0.66	0.74

<sup>a</sup>  $a = 2d_{110}$ ,  $c = 3d_{003}$ . <sup>b</sup> Crystallite size of LDH in stacking or plane direction ( $D_{(003)}$  and  $D_{(110)}$ , respectively), calculated according to Scherrer equation:  $D_{(hkl)} = 0.9 \times \lambda / (\beta \times \cos \theta)$ ,  $\lambda = 0.154$  nm,  $\theta$  is the position (deg.), and  $\beta$  is the FWHM (rad.) of the Bragg diffraction peaks.

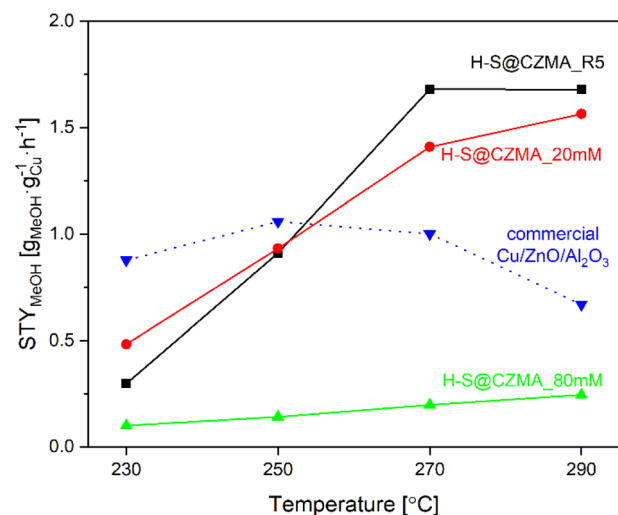
reconstruction cycle. Although, no clear correlation was found between porosity and the number of modifications, the calcination–reconstruction process had a minimal impact on the overall porosity of the materials.

### Spherical hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al as catalyst precursors

To deconvolute the relationship between catalyst structure/composition (including copper phase and size) and catalytic CO<sub>2</sub>-to-methanol activity, three samples (H-S@CZMA-LDH\_20 mM, H-S@CZMA-LDH\_80 mM and H-S@CZMA-LDH\_R5) were selected as promising catalyst precursors. As shown by elemental analysis (Table S5<sup>†</sup>), H-S@CZMA-LDH\_80 mM possesses two times the Cu + Zn content of H-S@CZMA-LDH\_20 mM but contains two phases of Cu<sub>2</sub>(OH)<sub>3</sub>-NO<sub>3</sub> and CuZnMgAl-LDH. In turn, H-S@CZMA-LDH\_R5, developed after five calcination–reconstruction cycles, retains a pure LDH structure with a similar metal content to H-S@CZMA-LDH\_20 mM. All samples were calcined in air at 330 °C and reduced in H<sub>2</sub> at 290 °C.

The relationship between CO<sub>2</sub> conversion, selectivity to methanol and CO, and reaction temperature is detailed in Fig. S12A–C.<sup>†</sup> All catalysts exhibit an increase in CO<sub>2</sub> conversion and CO selectivity and a decrease in methanol selectivity with increasing reaction temperature. This is consistent with thermodynamic control for CO<sub>2</sub> hydrogenation.<sup>5</sup> CO is the main by-product generated *via* a reverse water gas shift process (RWGS, CO<sub>2</sub> + H<sub>2</sub> ⇌ CO + H<sub>2</sub>O) which is an endothermic reaction ( $\Delta H^0 = 41.2$  kJ mol<sup>-1</sup>). When the temperature increases, the RWGS becomes more predominant, leading to an increased CO selectivity and a decreased methanol selectivity. The catalysts with comparable Cu + Zn loading (H-S@CZMA\_20 mM and H-S@CZMA\_R5) show comparable CO<sub>2</sub> conversion, reaching 16.5 and 17.5% at 290 °C, respectively. In contrast, the Cu-rich catalyst (H-S@CZMA\_80 mM) shows poor activity for CO<sub>2</sub> hydrogenation with a CO<sub>2</sub> conversion rate below 2.4%. As the temperature increases, MeOH selectivity drops from 70.6 to 21.3% and from 65.4 to 19.7% for H-S@CZMA\_20 mM and H-S@CZMA\_R5, respectively. However, H-S@CZMA\_80 mM exhibits a more moderate decrease in methanol selectivity (from 85.2 to 58.2%) over the same temperature range.

Given the wide range of reaction conditions used by different research groups, a direct comparison of the catalytic performance of our catalysts with those reported in the literature is challenging. However, we have summarised the performance of various Cu-containing catalysts with respect to the reaction conditions in Table S6.<sup>†</sup> We have been able to compare the catalytic efficiency of our catalysts with the commercial Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> catalyst, under the identical reaction conditions (Fig. S12D<sup>†</sup>). Although on a gram catalyst basis, the commercial Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> catalyst showed better CO<sub>2</sub> hydrogenation efficiency to methanol with CO<sub>2</sub> conversion of 9.4–22.2% and selectivity of 52.9–17.1% to MeOH in the temperature range of 230–290 °C, the situation changes dramatically when we normalise on a per Cu wt% basis. After normalisation, the space-time yield of methanol (STY<sub>MeOH</sub>) per gram Cu (g<sub>Cu</sub>) is shown in Fig. 6. At low temperature (230 °C), the commercial Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> catalyst is more effective than our catalysts. However, at higher temperatures, H-S@CZMA\_20 mM and H-S@CuZnMgAl\_R5 showed significantly higher STY<sub>MeOH</sub>. At



**Fig. 6** The space-time yield for methanol production (STY<sub>MeOH</sub>) of H-S@CZMA catalysts and commercial Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> as a function of temperature (CO<sub>2</sub>: H<sub>2</sub> = 3, 30 bar).

270 °C, the  $STY_{MeOH}$  for H-S@CZMA\_20 mM and H-S@CZMA\_R5 are 1.4 and 1.7 times higher than the commercial Cu/ZnO/Al<sub>2</sub>O<sub>3</sub> catalyst (1.41 and 1.68 *vs.* 1.00  $g_{MeOH} g_{Cu}^{-1} h^{-1}$ , respectively). This advantage became even more pronounced at the higher temperature, where determined values of  $STY_{MeOH}$  was 1.56 and 1.68 *versus* 0.67  $g_{MeOH} g_{Cu}^{-1} h^{-1}$ , respectively.

In order to understand the correlation between catalyst structure and catalytic CO<sub>2</sub>-to-methanol activity, comprehensive characterisation studies were conducted on the synthesised catalysts. XRD measurements were performed for thermally activated samples. After calcination at 330 °C (Fig. S13A†), the H-S@CZMA-LDH\_20 mM and H-S@CZMA-LDH\_R5 converted into equivalent amorphous LDO solids, with the crystalline LDH features disappearing due to dehydration, dehydroxylation and partially decarbonylation. Two broad scattering features are observed at *ca.*  $2\theta = 35.3$  and  $61.0^\circ$  which are attributed to the formation of poorly-crystalline ZnO phase.<sup>54</sup> In contrast, the H-S@CZMA-LDO\_80 mM showed highly the presence of a crystalline CuO phase, most likely due to its high Cu content (approx. 39 at%) of Cu<sub>2</sub>(OH)<sub>3</sub>NO<sub>3</sub> in H-S@CZMA-LDH\_80 mM, which facilitated CuO formation.<sup>47</sup> After reduction in H<sub>2</sub> at 290 °C (Fig. S13B†), an additional signal at  $2\theta = 43.3^\circ$  appears in both H-S@CZMA\_20 mM and H-S@CZMA\_R5. This weak scattering feature is assigned to the (111) Bragg reflection of metallic nanocopper (Cu<sup>0</sup>). H-S@CZMA\_80 mM presented two additional sharp Bragg reflections at  $2\theta = 43.4^\circ$  and  $50.6^\circ$ , which could be assigned to the (111) and (200) Bragg reflections of Cu<sup>0</sup>.<sup>55</sup> These differences in crystallinity of the metallic Cu phases in the catalysts were quantified using the Scherrer equation. For H-S@CZMA\_20 mM and H-S@CZMA\_R5, the mean Cu crystallite domain lengths are 3.2 and 2.7 nm, respectively. While the highly crystalline H-S@CZMA\_80 mM possesses Cu crystallites with much larger domain lengths of 33.9 nm.

TEM electron microscopy was used to evaluate the size distribution of metallic copper particles in our samples, as shown in Fig. S14.† The H-S@CZMA\_20 mM sample presents a narrow Cu particle size distribution (2–6 nm) with an average size of  $3.8 \pm 0.7$  nm. Similarly, the H-S@CZMA\_R5 catalyst showed Cu nanoparticles predominantly within the same range, with a slightly broader average size of  $3.6 \pm 1.2$  nm. In contrast, H-S@CZMA\_80 mM catalyst displays significantly larger Cu particles, with 66% of the particles in the 10–40 nm range and an average size of  $30.1 \pm 18.5$  nm.

These insights into the Cu particle size, supported by the results of XRD and TEM measurements, allow us to conclude that this parameter is strongly dependent on the conditions applied during the reconstruction process. Both of the more active methanol catalysts contains similarly small sized metallic copper nanoparticles incorporated into the catalyst precursor matrix (pure LDH) using less concentrated solutions, regardless of whether a single immersion or multiple calcination–reconstruction cycles were used. Conversely, the use of highly concentrated copper solution in single immersion process resulted in the formation of large particles/aggregates of metallic copper, leading to materials with the low catalytic efficiency.

## Conclusions

We have developed a systematic synthesis approach to hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs where LDH nanosheets are anchored perpendicularly to the surface of thin SiO<sub>2</sub> shells. Our approach leverages the memory effect of LDHs to regenerate hollow-SiO<sub>2</sub>@Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDHs within a pre-formed hollow-SiO<sub>2</sub>@Mg<sub>2</sub>Al-LDH template. We have demonstrated that the concentration of Cu<sup>2+</sup> and Zn<sup>2+</sup> solution significantly influences the composition, crystallinity, and porosity of the reconstructed LDH. A pure Cu<sub>x</sub>Zn<sub>y</sub>Mg<sub>z</sub>Al-LDH phase with desired morphology and porosity can be obtained in a single immersion approach using 20 mM. Higher concentration solutions greater than 40 mM result in the poorly crystalline LDH phase and a new phase with high crystallinity (Cu<sub>2</sub>(OH)<sub>3</sub>NO<sub>3</sub>), leading to lower porosity and poorer hollow morphology. Another approach using multiple calcination–reconstruction cycles allows gradual incorporation of Cu<sup>2+</sup> and Zn<sup>2+</sup> ions into the pure and crystalline LDH matrix, while maintaining good morphology and high porosity. Furthermore, our investigation underscored the significant influence of the reconstruction conditions on the formation of catalytically active phase. It was clearly demonstrated that a catalyst with a high content of the Cu<sup>0</sup> metallic phase, but in the form of large particles, shows a significantly lower efficiency in methanol production compared to those with a lower Cu content, but smaller and well-dispersed Cu<sup>0</sup> particle sizes. We believed that this research provides valuable insight into the design and synthesis of a new generation of hollow spherical material with promising properties and potential, particularly in fields related to climate protection and CO<sub>2</sub> management.

## Data availability

General experimental details, synthetic protocols and additional characterising data supporting this article have been included as part of the ESI.†

## Author contributions

T. Kondratowicz performed the synthetic and experimental work as well as conceptualised the research; M. Gajewska assisted with the electron microscopy studies; M. Li and J. Li conducted the catalytic CO<sub>2</sub> to MeOH experiments; C. Chen and Z. R. Turner supervised the work; D. O'Hare conceptualised and supervised the research and acquired the funding.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 Z. Liu, Z. Deng, S. J. Davis, C. Giron and P. Ciais, Monitoring global carbon emissions in 2021, *Nat. Rev. Earth Environ.*, 2022, **3**, 217–219.
- 2 I. Dimitriou, P. García-Gutiérrez, R. H. Elder, R. M. Cuéllar-Franca, A. Azapagic and R. W. K. Allen, Carbon dioxide utilisation for production of transport fuels: process and economic analysis, *Energy Environ. Sci.*, 2015, **8**, 1775–1789.
- 3 M. González-Castaño, B. Dorneanu and H. Arellano-García, The reverse water gas shift reaction: a process systems engineering perspective, *React. Chem. Eng.*, 2021, **6**, 954–976.
- 4 N. MacDowell, N. Florin, A. Buchard, J. Hallett, A. Galindo, G. Jackson, C. S. Adjiman, C. K. Williams, N. Shah and P. Fennell, An overview of CO<sub>2</sub> capture technologies, *Energy Environ. Sci.*, 2010, **3**, 1645–1669.
- 5 J. Zhong, X. Yang, Z. Wu, B. Liang, Y. Huang and T. Zhang, State of the art and perspectives in heterogeneous catalysis of CO<sub>2</sub> hydrogenation to methanol, *Chem. Soc. Rev.*, 2020, **49**, 1385–1413.
- 6 C. Filosa, X. Gong, A. Bavykina, A. D. Chowdhury, J. M. R. Gallo and J. Gascon, Enabling the Methanol Economy: Opportunities and Challenges for Heterogeneous Catalysis in the Production of Liquid Fuels via Methanol, *Acc. Chem. Res.*, 2023, **56**, 3492–3503.
- 7 A. García-Trenco, A. Regoutz, E. R. White, D. J. Payne, M. S. P. Shaffer and C. K. Williams, PdIn intermetallic nanoparticles for the Hydrogenation of CO<sub>2</sub> to Methanol, *Appl. Catal., B*, 2018, **220**, 9–18.
- 8 O. Bazaluk, V. Havrysh, V. Nitsenko, T. Baležentis, D. Streimikiene and E. A. Tarkhanova, Assessment of Green Methanol Production Potential and Related Economic and Environmental Benefits: The Case of China, *Energies*, 2020, **13**, 3113.
- 9 K. Wei, H. Guan, Q. Luo, J. He and S. Sun, Recent advances in CO<sub>2</sub> capture and reduction, *Nanoscale*, 2022, **14**, 11869–11891.
- 10 P. S. Murthy, W. Liang, Y. Jiang and J. Huang, Cu-Based Nanocatalysts for CO<sub>2</sub> Hydrogenation to Methanol, *Energy Fuels*, 2021, **35**, 8558–8584.
- 11 S.-T. Bai, G. De Smet, Y. Liao, R. Sun, C. Zhou, M. Beller, B. U. W. Maes and B. F. Sels, Homogeneous and heterogeneous catalysts for hydrogenation of CO<sub>2</sub> to methanol under mild conditions, *Chem. Soc. Rev.*, 2021, **50**, 4259–4298.
- 12 C. Baltes, S. Vukojević and F. Schüth, Correlations between synthesis, precursor, and catalyst structure and activity of a large set of CuO/ZnO/Al<sub>2</sub>O<sub>3</sub> catalysts for methanol synthesis, *J. Catal.*, 2008, **258**, 334–344.
- 13 H. Lei, Z. Hou and J. Xie, Hydrogenation of CO<sub>2</sub> to CH<sub>3</sub>OH over CuO/ZnO/Al<sub>2</sub>O<sub>3</sub> catalysts prepared via a solvent-free routine, *Fuel*, 2016, **164**, 191–198.
- 14 V. D. B. C. Dasireddy, S. Š. Neja and L. Blaž, Correlation between synthesis pH, structure and Cu/MgO/Al<sub>2</sub>O<sub>3</sub> heterogeneous catalyst activity and selectivity in CO<sub>2</sub> hydrogenation to methanol, *J. CO<sub>2</sub> Util.*, 2018, **28**, 189–199.
- 15 V. D. B. C. Dasireddy, N. S. Štefančič, M. Huš and B. Likozar, Effect of alkaline earth metal oxide (MO) Cu/MO/Al<sub>2</sub>O<sub>3</sub> catalysts on methanol synthesis activity and selectivity via CO<sub>2</sub> reduction, *Fuel*, 2018, **233**, 103–112.
- 16 H. Ren, C.-H. Xu, H.-Y. Zhao, Y.-X. Wang, J. Liu and J.-Y. Liu, Methanol synthesis from CO<sub>2</sub> hydrogenation over Cu/γ-Al<sub>2</sub>O<sub>3</sub> catalysts modified by ZnO, ZrO<sub>2</sub> and MgO, *J. Ind. Eng. Chem.*, 2015, **28**, 261–267.
- 17 C. S. Santana, L. F. Rasteiro, F. C. F. Marcos, E. M. Assaf, J. F. Gomes and J. M. Assaf, Influence of Al, Cr, Ga, or Zr as promoters on the performance of Cu/ZnO catalyst for CO<sub>2</sub> hydrogenation to methanol, *Mol. Catal.*, 2022, **528**, 112512.
- 18 K. Lee, H. Yan, Q. Sun, Z. Zhang and N. Yan, Mechanism-Guided Catalyst Design for CO<sub>2</sub> Hydrogenation to Formate and Methanol, *Acc. Mater. Res.*, 2023, **4**, 746–757.
- 19 M. M. J. Li, C. Chen, T. Ayvali, H. Suo, J. Zheng, I. F. Teixeira, L. Ye, H. Zou, D. O'Hare and S. C. E. Tsang, CO<sub>2</sub> Hydrogenation to Methanol over Catalysts Derived from Single Cationic Layer CuZnGa LDH Precursors, *ACS Catal.*, 2018, **8**, 4390–4401.
- 20 M. Lyu, J. Zheng, C. Coulthard, J. Ren, Y. Zhao, S. C. E. Tsang, C. Chen and D. O'Hare, Core-shell silica@CuZnAl LDH catalysts for efficient CO<sub>2</sub> hydrogenation to methanol, *Chem. Sci.*, 2023, **14**, 9814–9819.
- 21 A. M. H. Lim and H. C. Zeng, Controlling Nanosheet Spacing of ZnAl-Layered Double Hydroxide Assemblages for High-Efficiency Hydrogenation of CO<sub>2</sub> to Methanol, *Ind. Eng. Chem. Res.*, 2023, **62**, 1877–1890.
- 22 S. Li, D. Wang, X. Wu and Y. Chen, Recent advance on VOCs oxidation over layered double hydroxides derived mixed metal oxides, *Chin. J. Catal.*, 2020, **41**, 550–560.
- 23 M. Mokhtar, A. Inayat, J. Ofili and W. Schwieger, Thermal decomposition, gas phase hydration and liquid phase reconstruction in the system Mg/Al hydrotalcite/mixed oxide: A comparative study, *Appl. Clay Sci.*, 2010, **50**, 176–181.
- 24 S. Abelló, F. Medina, D. Tichit, J. Pérez-Ramírez, J. C. Groen, J. E. Sueiras, P. Salagre and Y. Cesteros, Aldol Condensations Over Reconstructed Mg–Al Hydrotalcites: Structure–Activity Relationships Related to the Rehydration Method, *Chem.–Eur. J.*, 2005, **11**, 728–739.
- 25 L. Dubnová, R. Daňhel, V. Meinhardová, V. Korolova, L. Smoláková, T. Kondratowicz, O. Kikhtyanin and L. Čapek, Reconstruction of the ZnAl Mixed Oxides Into the Layered Double Hydroxide Catalysts Active in the Aldol Condensation of Furfural: The Role of ZnO Particles, *Front. Chem.*, 2022, **9**, 803764.
- 26 P. Liu, M. Derchi and E. J. M. Hensen, Promotional effect of transition metal doping on the basicity and activity of calcined hydrotalcite catalysts for glycerol carbonate synthesis, *Appl. Catal., B*, 2014, **144**, 135–143.
- 27 S. K. Jana, Y. Kubota and T. Tatsumi, High activity of Mn-MgAl hydrotalcite in heterogeneously catalyzed liquid-phase selective oxidation of alkylaromatics to benzylic ketones with 1 atm of molecular oxygen, *J. Catal.*, 2007, **247**, 214–222.
- 28 P. Liu, Y. Guan, R. A. v. Santen, C. Li and E. J. M. Hensen, Aerobic oxidation of alcohols over hydrotalcite-supported



- gold nanoparticles: the promotional effect of transition metal cations, *Chem. Commun.*, 2011, **47**, 11540–11542.
- 29 H. Suo, C. Chen, J.-C. Buffet and D. O'Hare, Dendritic silica@aqueous miscible organic-layered double hydroxide hybrids, *Dalton Trans.*, 2018, **47**, 16413–16417.
- 30 S. Zhao, W.-C. Tseng, F. Hu, F. Zhong, H. Liu, S. Wen, G. Zheng, C. Qin and C. Gong, Layered double hydroxide-coated silica nanospheres with 3D architecture-modified composite anion exchange membranes for fuel cell applications, *J. Mater. Sci.*, 2020, **55**, 2967–2983.
- 31 M. Shao, F. Ning, J. Zhao, M. Wei, D. G. Evans and X. Duan, Preparation of Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>@Layered Double Hydroxide Core-Shell Microspheres for Magnetic Separation of Proteins, *J. Am. Chem. Soc.*, 2012, **134**, 1071–1077.
- 32 Y. Dou, S. Zhang, T. Pan, S. Xu, A. Zhou, M. Pu, H. Yan, J. Han, M. Wei, D. G. Evans and X. Duan, TiO<sub>2</sub>@Layered Double Hydroxide Core-Shell Nanospheres with Largely Enhanced Photocatalytic Activity Toward O<sub>2</sub> Generation, *Adv. Funct. Mater.*, 2015, **25**, 2243–2249.
- 33 C. Wang, B. Ma, S. Xu, D. Li, S. He, Y. Zhao, J. Han, M. Wei, D. G. Evans and X. Duan, Visible-light-driven overall water splitting with a largely-enhanced efficiency over a Cu<sub>2</sub>O@ZnCr-layered double hydroxide photocatalyst, *Nano Energy*, 2017, **32**, 463–469.
- 34 R. Li, T. Xue, R. Bingre, Y. Gao, B. Louis and Q. Wang, Microporous Zeolite@Vertically Aligned Mg-Al Layered Double Hydroxide Core@Shell Structures with Improved Hydrophobicity and Toluene Adsorption Capacity under Wet Conditions, *ACS Appl. Mater. Interfaces*, 2018, **10**, 34834–34839.
- 35 M. Lyu, C. Chen, J.-C. Buffet and D. O'Hare, A facile synthesis of layered double hydroxide based core@shell hybrid materials, *New J. Chem.*, 2020, **44**, 10095–10101.
- 36 Y. Ni, L. Yao, Y. Wang, B. Liu, M. Cao and C. Hu, Construction of hierarchically porous graphitized carbon-supported NiFe layered double hydroxides with a core-shell structure as an enhanced electrocatalyst for the oxygen evolution reaction, *Nanoscale*, 2017, **9**, 11596–11604.
- 37 L. Yu, H. Zhou, J. Sun, F. Qin, F. Yu, J. Bao, Y. Yu, S. Chen and Z. Ren, Cu nanowires shelled with NiFe layered double hydroxide nanosheets as bifunctional electrocatalysts for overall water splitting, *Energy Environ. Sci.*, 2017, **10**, 1820–1827.
- 38 Q. Guo, Q. Zhang, H. Wang, Z. Liu and Z. Zhao, Core-shell structured ZnO@Cu-Zn-Al layered double hydroxides with enhanced photocatalytic efficiency for CO<sub>2</sub> reduction, *Catal. Commun.*, 2016, **77**, 118–122.
- 39 C. Chen, R. Felton, J.-C. Buffet and D. O'Hare, Core-shell SiO<sub>2</sub>@LDHs with tuneable size, composition and morphology, *Chem. Commun.*, 2015, **51**, 3462–3465.
- 40 T. Kondratowicz, O. Horký, S. Slang, L. Dubnová, M. Gajewska, L. Chmielarz and L. Čapek, Hollow @CuMgAl double layered hydrotalcites and mixed oxides with tunable textural and structural properties, and thus enhanced NH<sub>3</sub>-NO<sub>x</sub>-SCR activity, *Nanoscale Adv.*, 2023, **5**, 3063–3074.
- 41 M. Li, P. Yuan, S. Guo, F. Liu and J. P. Cheng, Design and synthesis of Ni-Co and Ni-Mn layered double hydroxides hollow microspheres for supercapacitor, *Int. J. Hydrogen Energy*, 2017, **42**, 28797–28806.
- 42 T. Kondratowicz, S. Slang, L. Dubnová, O. Kikhtyanin, P. Bělina and L. Čapek, Controlled silica core removal from SiO<sub>2</sub>@MgAl core-shell system as a tool to prepare well-oriented and highly active catalysts, *Appl. Clay Sci.*, 2022, **216**, 106365.
- 43 P. Kenyon, S. Roberts, Z. R. Turner, N. H. Rees and D. O'Hare, Influence of the Alumina Interface layer in Silica@Layered Double Hydroxide Core-Shell Particles, *J. Phys. Chem. C*, 2024, **128**, 12249–12258.
- 44 G. Mitran, T. Cacciaguerra, S. Loridant, D. Tichit and I.-C. Marcu, Oxidative dehydrogenation of propane over cobalt-containing mixed oxides obtained from LDH precursors, *Appl. Catal., A*, 2012, **417–418**, 153–162.
- 45 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, Physisorption of gases, with special reference to the evaluation of surface area and pore size distribution (IUPAC Technical Report), *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 46 M. Wu, J. Zhang, Y. Peng, J. Zhou, X. Ruan, J. Liu, Q. Liu, Y. Xi, R. Frost and G. Qian, An investigation into mechanism of cation adsorption by reconstruction of calcined layered double hydroxide, *Microporous Mesoporous Mater.*, 2017, **242**, 182–189.
- 47 T. N. Ramesh and T. L. Madhu, Thermal Decomposition Studies of Layered Metal Hydroxynitrates (Metal: Cu, Zn, Cu/Co, and Zn/Co), *Int. J. Inorg. Chem.*, 2015, **2015**, 536470.
- 48 M. R. Jalal, H. Hojjati, J. R. Jalal, S. Ebrahimi and M. R. Z. Bighashi, Synthesis of Copper Hydroxide Nitrate (Cu<sub>2</sub>(OH)<sub>3</sub>NO<sub>3</sub>) micro-sheets by plasma electrolysis of Cu(NO<sub>3</sub>)<sub>2</sub> aqueous solution in atmospheric air, *JITL*, 2018, **2**, 109–112.
- 49 M. Smyrnioti, C. Tampaxis, T. Steriotis and T. Ioannides, Study of CO<sub>2</sub> adsorption on a commercial CuO/ZnO/Al<sub>2</sub>O<sub>3</sub> catalyst, *Catal. Today*, 2020, **357**, 495–502.
- 50 C. Henrist, K. Traina, C. Hubert, G. Toussaint, A. Rulmont and R. Cloots, Study of the morphology of copper hydroxynitrate nanoplatelets obtained by controlled double jet precipitation and urea hydrolysis, *J. Cryst. Growth*, 2003, **254**, 176–187.
- 51 E. A. Secco and G. G. Worth, Infrared spectra of unannealed and of annealed Cu<sub>4</sub>(OH)<sub>6</sub>(NO<sub>3</sub>)<sub>2</sub>, *Can. J. Chem.*, 1987, **65**, 2504–2508.
- 52 R. D. Shannon, Revised effective ionic radii and systematic studies of interatomic distances in halides and chalcogenides, *Acta Crystallogr.*, 1976, **A32**, 751–767.
- 53 S. Miyata, Anion-Exchange Properties of Hydrotalcite-Like Compounds, *Clays Clay Miner.*, 1983, **31**, 305–311.
- 54 X. Fang, Y. Men, F. Wu, Q. Zhao, R. Singh, P. Xiao, T. Du and P. A. Webley, Improved methanol yield and selectivity from CO<sub>2</sub> hydrogenation using a novel Cu-ZnO-ZrO<sub>2</sub> catalyst supported on Mg-Al layered double hydroxide (LDH), *J. CO<sub>2</sub> Util.*, 2019, **29**, 57–64.
- 55 M. Salavati-Niasari, F. Davar and N. Mir, Synthesis and characterization of metallic copper nanoparticles via thermal decomposition, *Polyhedron*, 2008, **27**, 3514–3518.