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# Porous 3D flower-like CoAl-LDH nanocomposite with excellent performance for NO<sub>2</sub> detection at room temperature†

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The 3D flower-like CoAl-layered double hydroxide (CoAl-LDH) was successfully prepared using the functional template agent of fluoride ions *via* a facile one-step hydrothermal route. Various techniques proved that all the samples presented 3D flower-like microstructural morphology. Representatively, the CA-2 sample, which was synthesized with the molar ratio of Co : Al of 3.65 : 1, had considerably abundant pores in its thin nanosheets. The average pore size was 2–4 nm, the specific surface area was equal to 49.45 m<sup>2</sup> g<sup>-1</sup>, and the thickness of nanosheets was approximately 3.068 nm. The CA-2 sample showed an excellent response to 0.01–100 ppm NO<sub>2</sub> with ultrafast response/recovery time at room temperature (RT). The detection limit of the sensor even reached 10 ppb. The superior gas sensing performance could be attributed to the synergistic effects of the functional template agent of fluoride ions and specific porous 3D flower-like nanostructure. The current study showed that the 3D flower-like CoAl-LDHs might be a promising material in practical detection of NO<sub>2</sub> at RT.

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## Introduction

As a notorious gas, nitrogen dioxide (NO<sub>2</sub>) plays a central role in the formation of photochemical smog, acid rain and PM<sub>2.5</sub> as well as ozone formation in the atmosphere.<sup>1–3</sup> According to the National Institute for Occupational Safety and Health, NO<sub>2</sub> may cause harm to life at concentrations higher than 20 ppm.<sup>4</sup> Faced with this, the sensitive monitoring of NO<sub>2</sub> is extremely important to public health and the environment. Many NO<sub>2</sub> sensing materials such as SnO<sub>2</sub>,<sup>5,6</sup> TiO<sub>2</sub>,<sup>7,8</sup> WO<sub>3</sub>,<sup>9,10</sup> and ZnO<sup>11</sup> have been investigated as solid-state gas sensor materials. Khuspe *et al.*<sup>6</sup> prepared nanostructured SnO<sub>2</sub> to detect NO<sub>2</sub> gas at 200 °C. 3D hierarchical flower-like WO<sub>3</sub> was applied by Wang *et al.*<sup>9</sup> to detect NO<sub>2</sub> at 100 °C. Using ZnO nanorods/TiO<sub>2</sub> materials, Zou *et al.*<sup>11</sup> studied their detection of NO<sub>2</sub> gas at 180 °C. Many efforts have been made to improve NO<sub>2</sub> gas detecting sensors,<sup>12–14</sup> but there are still some problems such as low sensitive response and high energy consumption.

Recently, two dimensional (2D) layered materials have become attractive in gas sensing because of being tunable and functional.<sup>15,16</sup> As typical 2D materials, LDHs have been

exploited for abundant applications ranging from catalysis<sup>17,18</sup> to electrochemistry<sup>19</sup> and gas sensing.<sup>20,21</sup> In LDHs, A<sup>n+</sup> anions are located in the interlayer gallery to balance positively charged brucite-like host layers of edge-sharing MO<sub>6</sub> octahedra;<sup>22–27</sup> notably, hydrogen bonds inside the interlayer space can provide an interacting site for external molecules,<sup>28</sup> therefore, LDHs lately has been used as gas sensing materials.<sup>29</sup> Table 1 showed a comparison of the gas sensing characteristic of LDHs reported in the literature and in present work. From this table, it was observed that Morandi *et al.*<sup>20</sup> reported Pt/Zn/Al-LDH as CH<sub>4</sub> gas sensor material operating at 450 °C, not only consumed too much energy, but also increased cost by using precious metal. (ZnAl-LDH/PAN)<sub>n</sub> as NH<sub>3</sub> sensor at RT was investigated by Xu *et al.*,<sup>30</sup> which faced a number of difficulties including complex preparation process and long response/recovery time. Our previous reported work,<sup>21</sup> reported an MgAl-LDH based gas sensor for NO<sub>x</sub> gas detection at RT, but there were still some shortcomings such as lower sensitivity and detection limit only at ppm level.

In this work, to further improve gas sensitivity, a 3D flower-like CoAl-LDHs nanocomposite with ultra-thin nanosheets (NSs) was synthesized *via* a simple hydrothermal method. Fluoride ion was utilized as a functional template agent under mild conditions. The functionalized composite was systematically characterized. One of sample showed excellent performance of high sensitivity, and ultra-fast response/recovery. Its detection limit was able to reach ppb level. Synthetic approach and related properties of newly-synthesized were discussed in detail.

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Table 1 Comparison of the gas sensing characteristics of LDHs reported in literature and in present work

Sensor material	Operating temperature	Gas	Gas concentration	Response	Response time	Recovery time	Ref.
Pt/ZnAl-LDH	450 °C	CH <sub>4</sub>	500 ppm	5	—	—	20
ZnAl-calcein-LDH	80 °C	Ethanol	100 ppm	5	—	—	30
Ag/MgAl-LDH	RT	Ethanol	200 ppm	7.8	10 s	50 s	31
ZnAl-LDH/PANI	RT	NH <sub>3</sub>	1000 ppm	16	150 s	240 s	32
NiAl-LDH	RT	O <sub>3</sub>	700 ppb	1.84	8 s	74 s	33
HPTS/NiFe-LDH	RT	CO <sub>2</sub>	—	—	—	—	34
MgAl-LDH	RT	NO <sub>x</sub>	100 ppm	76%	1.3 s	—	21
CoAl-LDH	RT	NO <sub>2</sub>	100 ppm	26.61	1.3 s	14.6 s	This work

## Experimental

### Materials and chemicals

All of the chemical reagents used in the experiment were analytical grade as purchased and be used directly without any further purification. Co(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O, Al(NO<sub>3</sub>)<sub>3</sub>·9H<sub>2</sub>O and urea were purchased from Xilong Chemical Co., Ltd., NH<sub>4</sub>F was purchased from Beijing Yili Fine Chemical Co., Ltd.

### Synthesis of functionalized CoAl-LDHs

Functionalized CoAl-LDHs were synthesized through a urea hydrolysis reaction without any surfactant. 0.8 mmol Co(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O (0.233 g), 0.4 mmol Al(NO<sub>3</sub>)<sub>3</sub>·9H<sub>2</sub>O (0.15 g), 4 mmol NH<sub>4</sub>F (0.15 g) and 28 mmol urea (1.7 g) were mixed in 40 mL deionized water and the mixed solution was continuously stirred for 0.5 h. The solution was then moved into a 50 mL Teflon-lined stainless steel autoclave. It was subsequently sealed, put into the electric blower drying box and heated at 90 °C for 6 h. Until being cooled naturally to room temperature, the obtained pink CoAl-LDHs was thoroughly washed with deionized water and ethanol. It was dried at 60 °C in air for 12 h and named as CA-2 (the molar ratio of Co : Al = 3.65 : 1). In addition, changed amount of Co(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O (0.4 mmol, 0.117 g) and (1.2 mmol, 0.349 g), and resulting samples were labelled as CA-1 (the molar ratio of Co : Al = 3.40 : 1), CA-3 (the molar ratio of Co : Al = 3.80 : 1). It should be noted that the synthetic condition was the same as CA-2. CAW was a sample without adding NH<sub>4</sub>F. By varying hydrothermal time *i.e.* from 1 to 5 hours, five different samples were obtained (see Fig. S2a†). The specific synthetic condition was listed in Table S1.†

### Materials characterizations

Crystalline structure and chemical composition of the samples were characterized through X-ray powder diffraction (XRD, Rigaku TTRIII X-ray diffractometer,  $\lambda = 1.5406 \text{ \AA}$ ) and Fourier transform infrared spectroscopy (FT-IR, Varian FTIR spectrophotometer), respectively. The morphology and size of the products were studied by scanning electron microscopy (SEM, HITACHI S-4800) and transmission electron microscopy (TEM, JEOL JEM-2100) at an accelerating voltage of 200 kV. Specific surface areas and pore size distribution of the samples were performed by using the Brunauer-Emmett-Teller (BET). Chemical state of elements in composite were studied by using

X-ray photoelectron spectroscopy (XPS). The charge transfer and carrier concentration were evaluated by electrochemical impedance spectroscopy (EIS) and Mott-Schottky (MS) plot using electrochemical working station (CHI660C, Shanghai, China) at room temperature. Temperature program desorption (TPD) studies was performed by AutoChem TP5080 chemisorption analyzer with a temperature ramp of 10 °C min<sup>-1</sup>, and using mass spectra (QIC-20, Hidden) to record TPD signals.

### Fabrication and testing of gas sensor

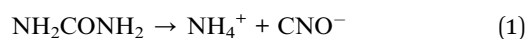
Manufacture method of CoAl-LDHs composites sensor film and test measurement were similar to our reported previously.<sup>21</sup> Sensor resistance test was performed at room temperature (RT = 25 °C) and a relative humidity (RH = 26%). After NO<sub>2</sub> reacted with the sensor to get a response, fresh air was injected and started to recover. The response was calculated as follows:

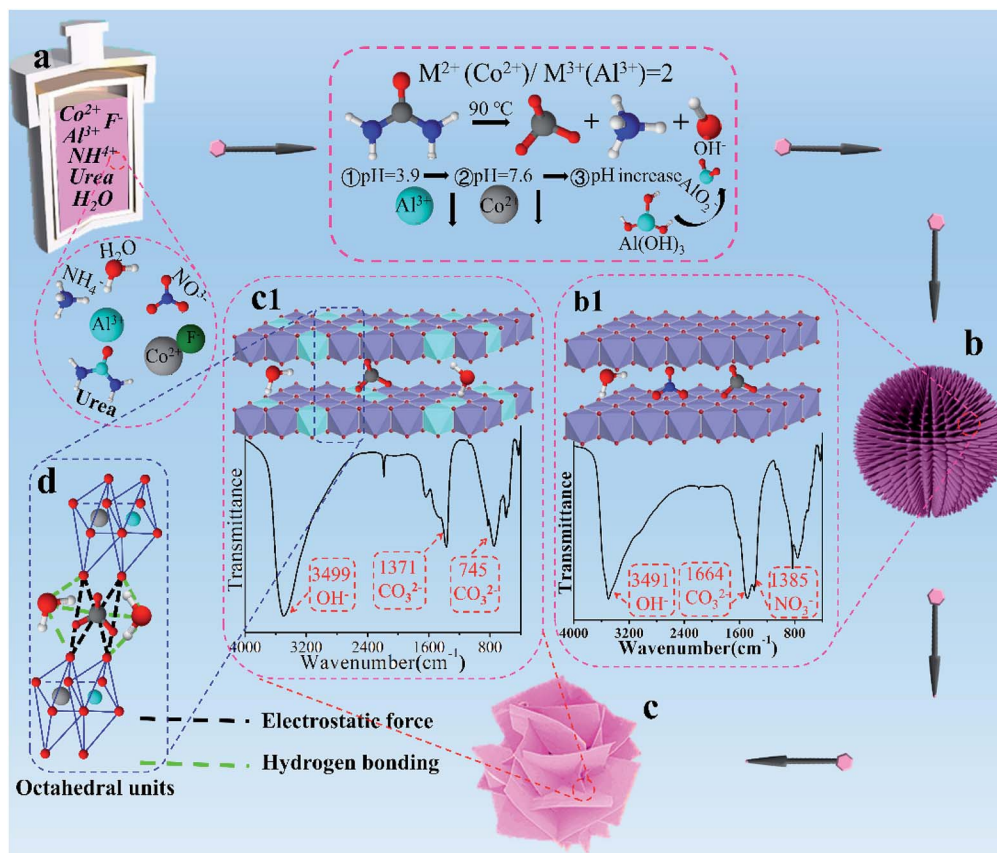
$$R = R_a/R_g$$

where  $R_a$  and  $R_g$  represent resistance in air and target gas respectively. The response and recovery time are defined as time taken to reach 90% of resistance change value. Repeatability was studied by using four sensors under each sensor detection.

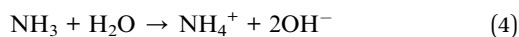
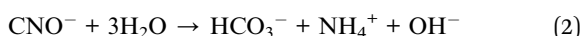
## Results and discussions

As demonstrated in Scheme 1a, urea, a common ammonia-releasing agent, was utilized as pH buffer to release OH<sup>-</sup> slowly and constantly, and provided carbonate ions simultaneously during the hydrolysis.<sup>35</sup> This offers the basic environment to precipitate Co<sup>2+</sup>, yet it might prevent oxidation of Co<sup>2+</sup> because of its reducibility. Moreover, F<sup>-</sup> played the role of functional template agent in formation of LDHs.<sup>36-38</sup> During the hydrothermal reaction, a large number of particles came into being and formed microspheres over time, which taken the juts on the surface of microspheres as crystal nucleus and further grew into nanofibers with F<sup>-</sup> as structure-orientation agent under the unreacted urea solution. The steric hindrance and magnetic interaction forced nanofibers to orient radially,<sup>39</sup> and the urchin-like grew up slowly (Scheme 1b), eventually formed in 3D flower-like structure (Scheme 1c). The main reactions in the precursor solution were described as follows:<sup>40,41</sup>





Scheme 1 The formation mechanism of functionalized 3D flower-like CoAl-LDH (a) precursor solution containing  $\text{Co}^{2+}$ ,  $\text{Al}^{3+}$ ,  $\text{F}^-$ ,  $\text{NH}_4^+$  and urea; (b and b1) C-2; (c and c1) CA-2; and (d) octahedral units of CA-2.



In the reaction process, pH gradually increased with slow hydrolysis of urea. At the beginning of crystallization, continuous hydrolysis was in favor of precipitation of  $\text{Al}(\text{OH})_3$  due to the pH value of  $\text{Al}^{3+}$  (pH = 3.9) was lower than that of  $\text{Co}^{2+}$  (pH = 7.6).<sup>42</sup> However, there were no diffraction peaks of aluminum hydroxide or hydrous oxide appearance (Fig. S2a†), that is,  $\text{Al}^{3+}$  first formed  $\text{AlO}_2^-$  in alkali aqueous. Then the complexation between  $\text{AlO}_2^-$  and  $\text{OH}^-$  occurred to produce  $[\text{Al}_{13}(\text{OH})_{32}(\text{H}_2\text{O})]^{7+}$ .<sup>43</sup> During this process,  $\alpha\text{-Co}(\text{OH})_2$  was formed in approximately 1 h (Fig. S2a†), which contained carbonate ions and nitrate ions in the interlayer gallery (Scheme 1b1). With reaction time increasing,  $\text{Al}^{3+}$  complex in  $[\text{Al}_{13}(\text{OH})_{32}(\text{H}_2\text{O})]^{7+}$  would promptly transport into the lattice of  $\alpha\text{-Co}(\text{OH})_2$  and substitute  $\text{Co}^{2+}$  to form CoAl-LDH nanosheet (Fig. S2c†).<sup>43,44</sup> This attributed to the ionic radius of  $\text{Al}^{3+}$  (0.053 nm) was lower than that of  $\text{Co}^{2+}$  (0.070 nm), eventually formed ultrathin nanosheets (Scheme 1c) with carbonate ions intercalated into 3D flower-like CoAl-LDH.

To investigate the crystal structure and purity of samples, XRD of CoAl-LDHs after hydrothermal reaction was depicted in

Fig. 1a. The study showed that all XRD peaks for CoAl-LDHs could be indexed in a hexagonal cell ( $a = 0.308$  nm,  $c = 2.301$  nm) with the rhombohedral symmetry  $R\bar{3}m$ . Interlayer spacing ( $d_{003} = 0.767$  nm) proved that carbonate anion intercalated in LDHs. The existence of carbonate anions was likely produced from urea hydrolysis in the hydrothermal process. There were four characteristic diffraction peaks with equal spacing at  $11.6^\circ$ ,  $23.32^\circ$ ,  $34.25^\circ$  and  $46.27^\circ$ , corresponding to (003), (006), (012) and (018) crystal planes (CoAl-LDHs JCPDS: 51-0045), respectively. The typical doublet (110) and (113) crystal planes of CoAl-LDHs could be clearly observed within the scope of  $2\theta = 60\text{--}62^\circ$ . The diffraction peaks of (003), (006) crystal planes were stronger compared with the other peaks indicating that CoAl-LDHs grew along an axial direction during the formation. Furthermore,

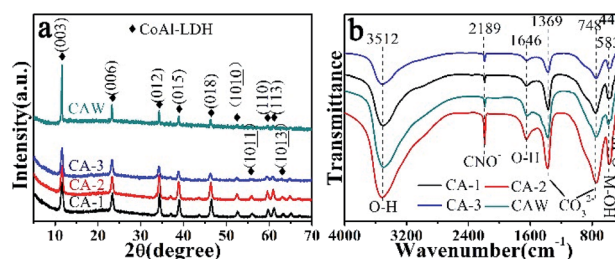


Fig. 1 XRD diffraction pattern (a) and FT-IR spectra (b) of CA-1, CA-2, CA-3 and CAW.



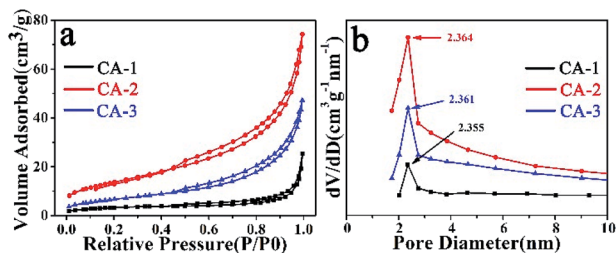


Fig. 2 BET (a) and BJH (b) curves of CA-1, CA-2 and CA-3.

there were two diffraction peaks ( $101\bar{1}$ ) and ( $101\bar{3}$ ) appeared after using  $F^-$  as functional template agent. This may be due to the effect of functional  $F^-$  controlling the crystal structure. On the other hand, no other peaks appeared, which clearly indicated that high purity of CoAl-LDHs were formed during preparation process.

FT-IR spectrum was carried out to investigate the chemical composition of CoAl-LDHs (Fig. 1b). There were four characteristic peaks associated with CoAl-LDH. The broad band at  $3512\text{ cm}^{-1}$  belonged to O–H stretching vibration from H-bonded OH group and water molecules in the interlayer.<sup>45</sup> The peak at  $1646\text{ cm}^{-1}$  could be assigned to the bending mode of interlayer water molecule. The peaks at  $1369$  and  $748\text{ cm}^{-1}$  corresponded to  $\nu_3$  bending vibration of  $\text{CO}_3^{2-}$  indicating that carbonate ions had been intercalated into LDHs. The low frequency peak at  $583\text{ cm}^{-1}$  was attributed to the M–OH bending mode of Co and Al, and  $445\text{ cm}^{-1}$  could be assigned to M–O stretching vibration.<sup>46</sup> Furthermore, the narrow peak appeared at  $2189\text{ cm}^{-1}$  was the typical stretching vibration of C≡N bonds in CNO anions indicating the incomplete decomposition of urea.<sup>47</sup>

The specific surface area and pore size distributions of the as-synthesized composites CA-1, CA-2 and CA-3 were investigated by using BET technique. The detailed information was

presented in Fig. 2 and Table S2.† All the three samples represented typical IV isotherms with H3 type hysteresis loop, which made clearly the presence of mesoporous structure. According to the pore size distribution curve, there was very slight change in the distribution centers. The pore size distribution was about 2–4 nm. Relative to other two samples, the CA-2 possessed the largest surface area ( $49.45\text{ m}^2\text{ g}^{-1}$ ) and its total pore volume was  $0.10\text{ cm}^3\text{ g}^{-1}$ . High pore volume could provide a delivery channel for electron easily transmitted to internal space, which was beneficial for  $\text{NO}_2$  gas sensing. This meant CA-2 would process higher  $\text{NO}_2$  sensitivities.

Morphology of the samples was examined by SEM and TEM. SEM images of the as-prepared CA-1, CA-2 and CA-2 samples were shown in Fig. 3. Fig. 3a–f distinctly revealed that the morphologies of CA-1, CA-2, and CA-3 not similar to each other. When the molar ratio of Co : Al was much lower, the morphology of CA-1 was insufficient to support formation of flawless nanosheets (Fig. 3b). When the molar ratio of Co : Al was much higher, excess primitives of CA-3 were attached to spaces between the neighboring nanosheets to form microscopic nanostructures (Fig. 3f). The CA-2 sample with the optimal ratio of Co : Al displayed a 3D flower-like structure (Fig. 3d), which was composed of a large number of well-dispersed ultrathin nanosheets, and there was a lot of inter-spaces between the layered nanopetals and thin nanosheets. The as-prepared CAW sample presented well-defined nanosheet structure was shown in Fig. S4.†

As shown in Fig. 4, low magnification TEM images of CA-2 demonstrated that 3D flower-like nanostructure was formed and the sample consisted of many ultrathin nanosheets with mesoporous size about 2–4 nm. The formation of mesoporous attributed to the release of  $\text{CO}_2$  during the urea hydrolysis process, and it was beneficial to improve the gas sensing properties.

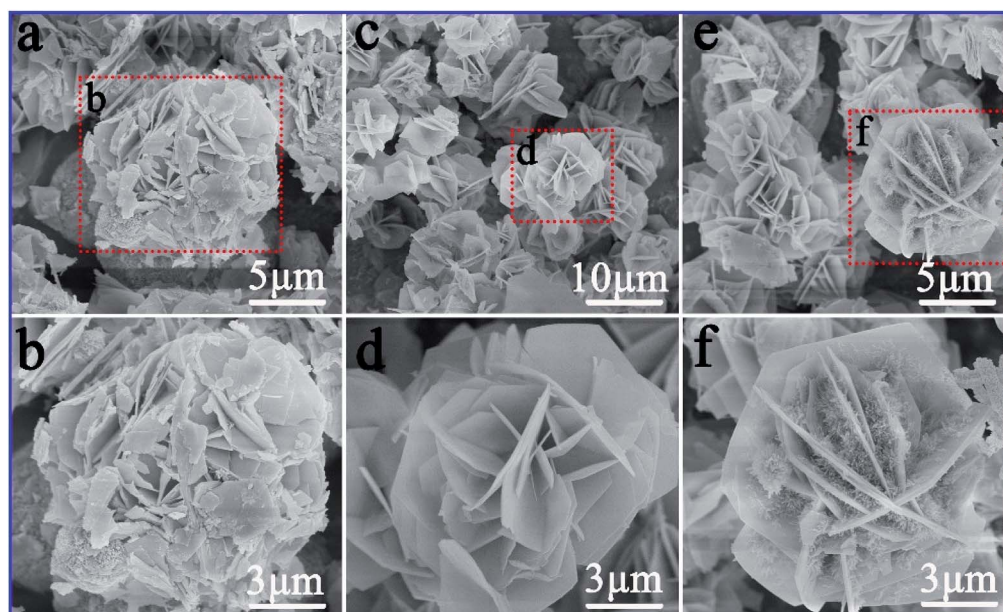


Fig. 3 SEM images of CA-1, CA-2 and CA-3.



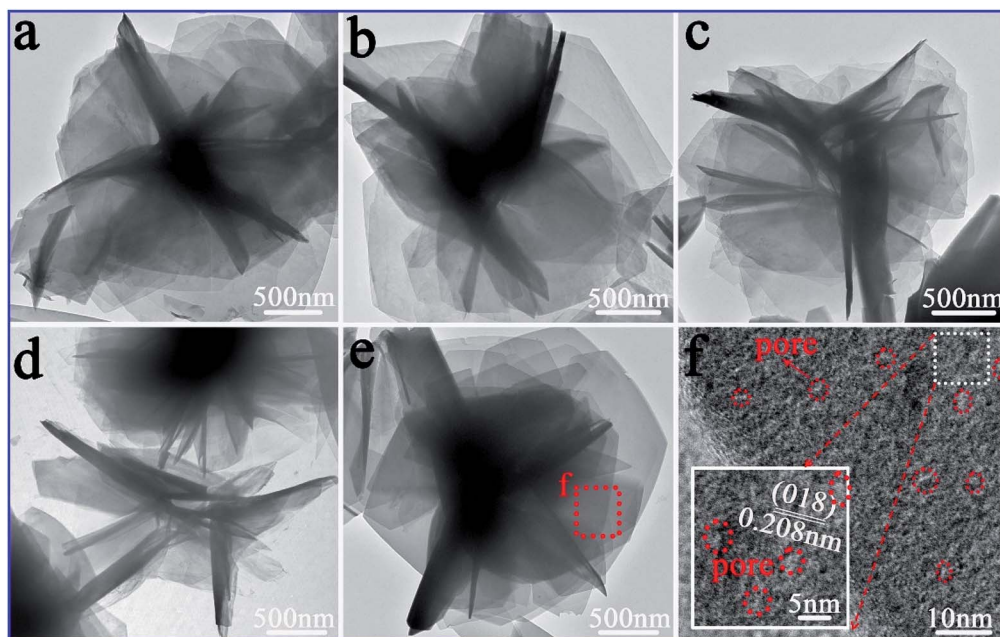


Fig. 4 TEM images of the CA-2 sample (a–e), together with a HRTEM image (f) from the (e) picture.

As illustrated in Fig. 5a, CA-2 sample was composed of many 3D flowers with the size of 2.4–3.4  $\mu\text{m}$ . The well-defined flower clearly illustrated that abundant nanosheets (Fig. 5b) were assembled to form a 3D flower-like architectures. Petals in Fig. 5c further confirmed the flowers consisted of uniform and ultrathin nanosheets with 4.0 layers and thickness of 3.068 nm. Moreover, as shown in HRTEM of Fig. 5d and e, the value of the interplanar spacing between the adjacent lattice fringes were 0.235 and 0.391 nm, which were consistent with (015) and (006) planes of CoAl-LDHs, respectively. The polycrystalline diffraction rings (Fig. 5f) from inside to outside were (015), (018) and (113) planes, respectively, which in accordance with the results of XRD (Fig. 1a). TEM and HRTEM images of CAW sample shown in Fig. S5<sup>†</sup> confirmed that the as-preparation sample with nanosheet structure has low crystallinity.

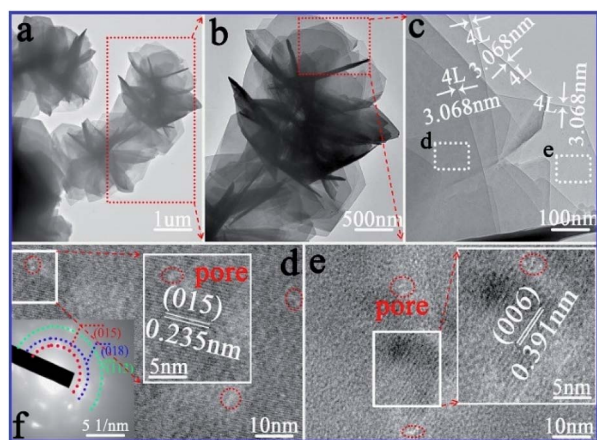


Fig. 5 Microstructure of CA-2: (a–c) gradually enlarged TEM images, (d and e) HRTEM images of different compositions, and (f) SAED pattern.

The element mapping and EDS spectra of CA-2 were used to verify the element distribution (Fig. 6). The bright image in Fig. 6b–e indicated that Co, Al, O and F elements were distributed uniformly. As Fig. 6f showed that Co atomic element contents was 26.49%, Al 7.24%, O 57.73%, and F 8.54%. The atomic ratio of Co/Al in CA-2 was calculated to be 3.65 : 1.

To reveal the surface chemical composition and the element electronic states of the flower-like CoAl-LDHs, the corresponding XPS results were presented in Fig. 7. Fig. 7a–c depicted the high-resolution Co 2p spectrum in CoAl-LDHs. For comparison, distributions of Co 2p XPS peaks were similar in three samples. The peaks of Co 2p<sub>3/2</sub> was at 781.9, 781.7 and 781.8 eV, and Co 2p<sub>1/2</sub> at 797.3, 797.2 and 797.2 eV, with the spin–orbit splitting value of 15.4, 15.5 and 15.4 eV, respectively. These all proved that Co ion existed in the form of Co(OH)<sub>2</sub>. The other three satellite bands demonstrated that Co ion existed in the form of divalent Co<sup>2+</sup> state.<sup>48,49</sup>

In the O 1s spectra (Fig. 7d–f), the spectra could divide into two peaks by using Gaussian fitting. Binding energy at 530.9 eV (peak 1) could be attributed to Co–OH and/or Al–OH groups,<sup>50</sup> indicating there was a large amount of OH groups inside the interlayer which could provide an interaction site for external molecules. The peak 2 (532.2–532.6 eV) could be attributed to chemisorbed oxygen.<sup>51</sup> The calculated chemisorbed oxygen of CA-1, CA-2 and CA-3 was about 18.64%, 24.91% and 20.19% (Table S3<sup>†</sup>), respectively, and CA-2 sample had the largest calculated chemisorbed oxygen. More chemisorbed oxygen participated in surface reactions, the higher response sensitivity might be produced.

Al 2p spectrum was presented in Fig. 7g–i. It could be concluded that there was only Al<sup>3+</sup> species peaks existed in CoAl-LDHs (BE = 72.9–73.0 eV).<sup>52</sup>

XPS spectrum of F<sup>−</sup> in 682–692 eV (Fig. 7j–l) clearly indicated that the substitution of OH<sup>−</sup> by F<sup>−</sup> did not cause a large lattice



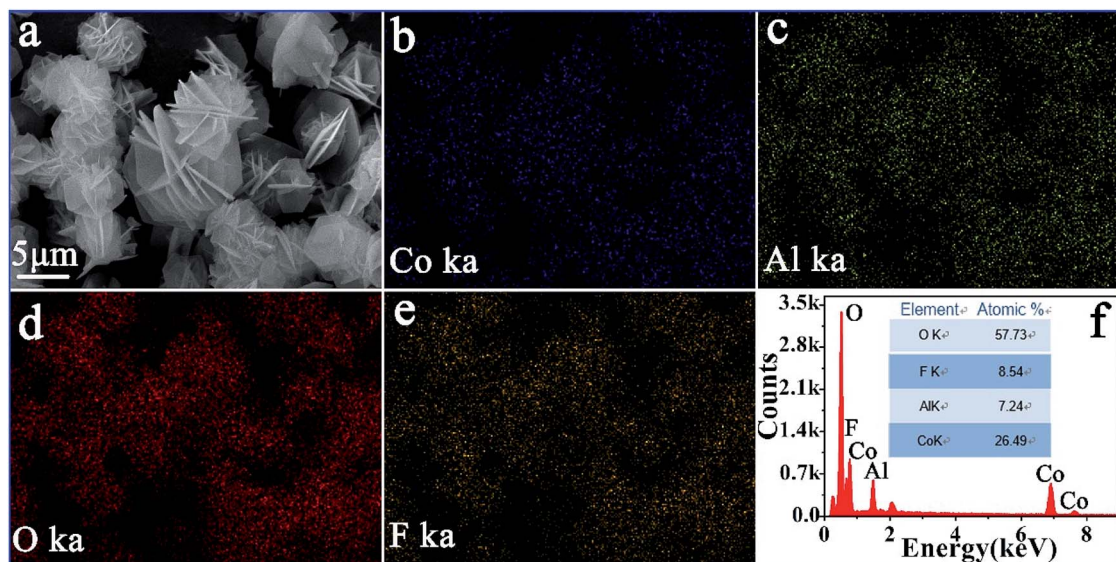


Fig. 6 SEM image/EDS mapping of CA-2: (a) bright field image, (b–e) Co, Al, O and F element mapping, and (f) EDS spectra.

distortion because ionic radius of  $F^-$  (0.117 nm) was similar to that of  $OH^-$ .<sup>53</sup> The existence of fluoride ions ( $F^-$ ) enhanced carrier concentration and decreased charge transfer resistance.<sup>54,55</sup> This result was consistent with those of MS (Fig. 9e) and EIS (Fig. 9f).

### Enhanced $NO_2$ gas sensing properties

For comparison, gas sensing properties to  $NO_2$  were investigated for CA-1, CA-2 and CA-3 based sensors at RT, as shown in Fig. 8.

As shown Fig. 8a, the CA-2 sensor shown the maximum response towards 100 ppm  $NO_2$  was 26.61, which was 5.03 and 1.3 times higher than that of CA-1 and CA-3 sensor, respectively. Their detection limit towards  $NO_2$  all could reach to 0.01 ppm, and the response of CA-2 was significantly higher than that of other two sensors, as Table S4† summarized. Fig. S6† presented that slope of the response curve was higher for low concentration than high concentration of  $NO_2$ . This further illustrated that CoAl-LDHs based sensor possessed a great ability to detect low concentration  $NO_2$ .

Fig. 8b and c showed response/recovery time of CA-1, CA-2 and CA-3 based sensors at RT under different  $NO_2$  concentration. The response time tended to increase while the recovery time was the opposite. It was worth noting that the response time of the CA-2 sensor was only 1.3 s (Table S4†) at 100 ppm, faster than that of the other two sensors. Even at the lowest detection limit (10 ppb), the response time of CA-2 sensor was still the shortest (4.6 s). Moreover, it could be clearly observed that the recovery time of CA-2 sensor was faster than that of other two sensors at every  $NO_2$  concentration (Table S4†). This was further verified by  $NO_2$ -TPD (Fig. 8d and S7†). Compared with the other two samples, the physical desorption peak of CA-2 sample shifted towards lower temperature with a larger area. This indicated that CA-2 possessed a high adsorption capacity for  $NO_2$  at RT (25 °C), which was conducive to the desorption process at lower temperature.<sup>56,57</sup>

To further investigated the gas sensing properties of CA-2 sensor, selectivity, repeatability, stability, MS and EIS tests were carried out at RT as presented in Fig. 9.

Fig. 9a depicted the dynamic response–recovery curve of CA-2 sensor at RT under different concentration of  $NO_2$ . When  $NO_2$  was injected and reacted with CA-2 sensor, its resistance dropped rapidly to a minimum value in a short time, and recovered back to its initial value after purging the gas with fresh air. The response gradually reduced with the injected concentration of  $NO_2$  decreasing and ultimately reached to a lower limit detection of 0.01 ppm.

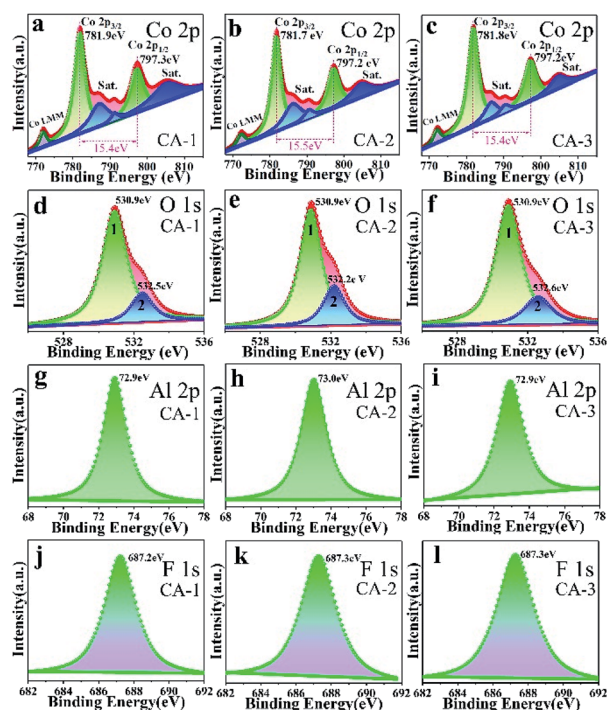


Fig. 7 XPS spectra of CA-1, CA-2 and CA-3 samples: (a–c) Co 2p, (d–f) O 1s, (g–i) Al 2p, and (j–l) F 1s.



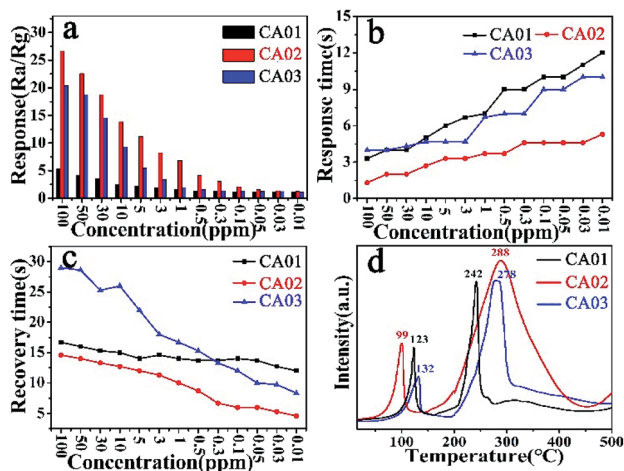


Fig. 8 (a) Response; (b) response time; (c) recovery time and (d)  $\text{NO}_2$ -TPD of the CA-1, CA-2 and CA-3 sensor.

Fig. 9b showed that CA-2 sensor almost had no response to five detection gases (*e.g.*  $\text{NH}_3$ ,  $\text{CO}$ ,  $\text{CH}_4$ ,  $\text{H}_2$  and  $\text{H}_2\text{S}$ ) at RT, but significantly higher response to  $\text{NO}_2$ , demonstrating that it had an excellent selectivity toward  $\text{NO}_2$ .

It was well-known that repeatability and long-term stability were vitally important parameters for sensor devices. As illustrated in Fig. 9c, response–recovery curves of CA-2 resistance to 10 ppm of  $\text{NO}_2$  at RT maintained the initial resistance upon 30 cycles without a clear attenuation, this revealed the sensor's stable and repeatable character. In addition, the measurement of the response values of CA-2 sensor to 10 ppm  $\text{NO}_2$  at RT and its initial resistance in air atmosphere ( $R_a$ ) for 54 days further verified its excellent long-term stability in Fig. 9d.

Generally, low resistance and superior electron transportation are critical to sensitive materials.<sup>58</sup> MS plot measurements were performed to investigate electric characteristics of CoAl-LDHs. As Fig. 9e showed, slope of CoAl-LDHs's

plot was negative, indicating CoAl-LDHs exhibited a p-type behavior. Based on the calculation,<sup>59</sup> carrier concentration of CA-1, CA-2, CA-3 and CAW were  $2.31 \times 10^{15}$ ,  $9.80 \times 10^{15}$ ,  $5.82 \times 10^{15}$  and  $3.08 \times 10^{15}$  (Table S7†), respectively, and CA-2 had the largest carrier concentration of  $9.80 \times 10^{15}$ .

EIS measurements were carried out to study charge-transfer of the samples. As shown in Fig. 9f, CA-2 sample had the smallest semicircle radii. The relevant equivalent circuit model and the impedance parameters' results were given in Fig. S9 and Table S7.† It was well known that radius of high-frequency region was thought to be associated with charge transfer, and the smaller radius means lower resistance. Consequently, EIS results indicated that CA-2 had a superior charge transport property, which could provide an ideal electron transmission.

### Gas sensing mechanism

LDHs commonly had a large surface/volume ratio and their structures proved to be able to absorb detection gases because of its interlayer spacing could use as a gallery pathway. What's more, the hydrogen bond between surface hydroxyl group and interlayer water molecules was equivalent to bridge connecting the adjacent the positively charge layers. This bridging effect could provide natural channels for the effective and fast transportation of carriers.

It was known that gas sensing mechanism of solid-state sensors could be attributed to modulation of electrical conductivity on the surface in the presence and absence of the target gas.<sup>60,61</sup> The mechanism of  $\text{NO}_2$  sensing for CoAl-LDHs was proposed in Scheme 2 which predominantly showed the involvement of three steps: adsorption, charge transfer and desorption process.

In air (Scheme 2a), surface electrons of the samples were captured by  $\text{O}_2$  from acceptor level of the sensor, leading to the formation of ionized  $\text{O}_2^-$ . As a result, decreased the conductivity and enhanced the resistance of the sensing material (eqn (5) and (6)).<sup>62</sup>

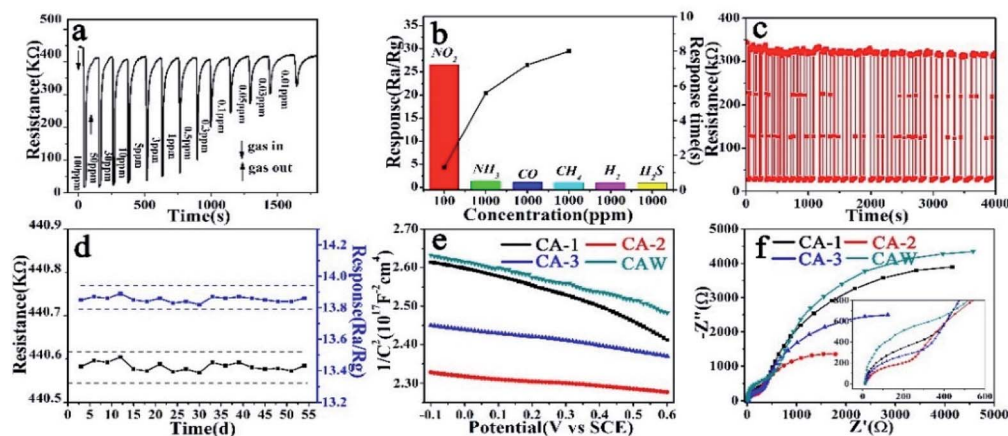
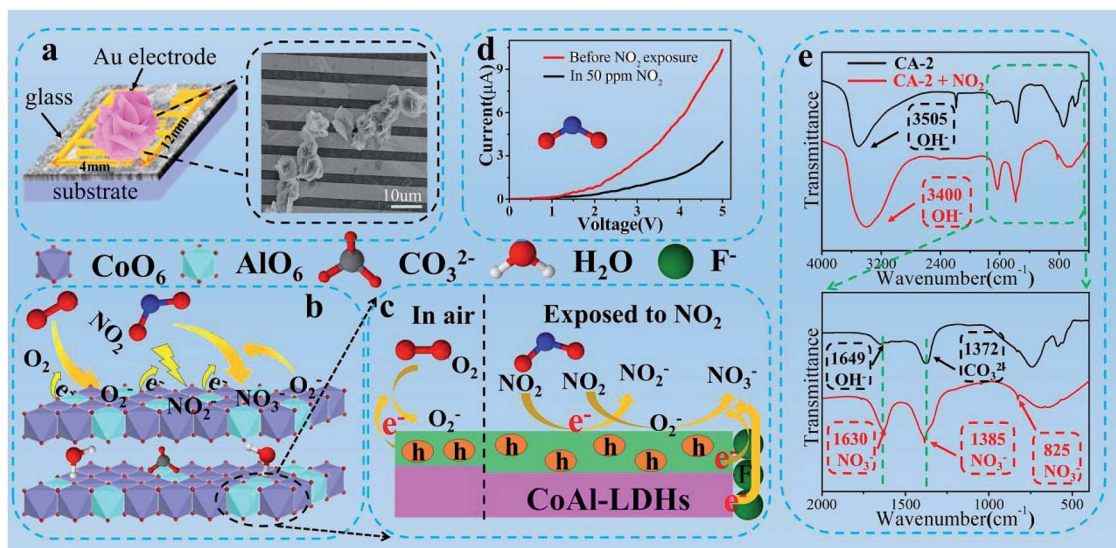
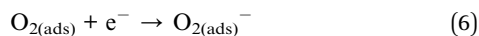


Fig. 9 Sensing properties (a) dynamic response–recovery curve of the CA-2 sensor for 100–0.01 ppm at RT; (b) response of CA-2 sensor for different gas; (c) thirty cycles of dynamic response curves of CA-2 sensor to 10 ppm  $\text{NO}_2$ ; (d) stability of CA-2 sensor to 10 ppm  $\text{NO}_2$  for 54 days at RT; (e) Mott–Schottky plots of CA-1, CA-2, CA-3 and CAW modified electrodes in 1 M KOH electrolyte measured at a frequency of 1 kHz; (f) EIS of CA-1, CA-2, CA-3 and CAW.



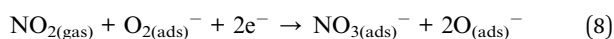
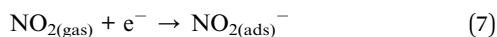


**Scheme 2** Schematic for the mechanism of NO<sub>2</sub> sensing for CoAl-LDHs. (a) SEM image of 3D flower-like CA-2 on Au electrode; (b and c) the model with response procedure of gas sensing; (d) *I*-*V* curve measured for CA-2 sensor in air and 50 ppm NO<sub>2</sub> exposure at RT; (e) IR spectra of CA-2 before and after NO<sub>2</sub> gas detection (CA-2 test NO<sub>2</sub> gas sensing for 2 h).



where “ads” means the “adsorbed”.

As shown in Scheme 2b and c, when the samples were exposed to NO<sub>2</sub>, the gas molecules not only captured free electrons from the acceptor electron level of CoAl-LDHs to produce NO<sub>2</sub><sup>-</sup>, but also reacted with O<sub>2</sub><sup>-</sup> forming NO<sub>3</sub><sup>-</sup>, which led to an increase in the hole density and ultimately a rapid decrease in the resistance (eqn (7) and (8)).



Scheme 2d showed the *I*-*V* plots for CA-2 sensor exposure to 50 ppm NO<sub>2</sub> gas. It was worth noting that the current was decreased when CA-2 sensor exposure to NO<sub>2</sub> gas. Because of the oxidation of NO<sub>2</sub>, it was easily adsorbed on the surface of CA-2 and withdrew electrons from CA-2. This led to the electron carrier concentration decreasing and significantly changed the resistance.

Therefore, 3D flower-like CA-2 sensor presented an outstanding NO<sub>2</sub> gas sensing performance at RT, especially high response, short response/recovery time. This could be ascribed to the following factors: first, the morphology of 3D flower-like CA-2 consisted of ultrathin nanosheets in its framework. Its thickness was 3.068 nm, which was in favored of the electrons transfer speed from internal to surface; second, there were a large number of mesoporous existed on the surface of ultrathin nanosheets which facilitated the gas diffusion/adsorption alleyway. NO<sub>2</sub> gas could be easily adsorbed at the outer/inner surfaces as well as the entire mesoporous wall. In addition, the largest specific surface area of CA-2 provided

a number of active sites, which provided more oxidative NO<sub>2</sub> molecules absorbed sites and finally led to the high response. In particularly, as functional template agent, the roles of fluoride ions (F<sup>-</sup>) (i) decreased the charge transfer resistance (Fig. 9f and Table S7<sup>†</sup>), *i.e.* the electron would transfer faster in adsorption and desorption process; (ii) enhanced the carrier concentration (Fig. 9e and Table S7<sup>†</sup>), which meant there were more electrons would react with oxygen to produce more O<sub>2</sub><sup>-</sup>. These were all beneficial for the high response of CA-2.

## Conclusions

In summary, a facile one-step hydrothermal method was developed to synthesize a 3D flower-like CoAl-LDHs architecture by using fluoride ion as a functional template agent. Specifically, the morphology of the functionalized 3D flower-like CA-2 sample consisted of ultrathin nanosheets with 3.068 nm thick in its framework, and a large number of mesoporous existed on the surface of ultrathin nanosheet. A systematic and comparative analysis indicated that the CA-2 based sensor showed a notable NO<sub>2</sub> gas sensing performance at RT, *i.e.*, high sensitivity, fast response/recovery time, and detection limit of ppb level. The superior performance was attributed to the 3D flower-like morphological and mesoporous structure of CoAl-LDHs as well as the fluoride ion as functional template agent. The existence of fluoride ion decreased the charge transfer resistance and enhanced the carrier concentration which led to much higher sensing response. Therefore, the functionalized CA-2 sample, displayed an outstanding sensing activity to super low-concentration NO<sub>2</sub> gas at RT, it would be a potential candidate to fabricate relevant sensor. And our facile synthetic method could be a promising method to prepare the basic gas sensor materials.



## Conflicts of interest

There are no conflicts to declare.

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