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Three CoS/CoO microspheres and their mixed matrix membranes for the highly efficient photocatalytic degradation of methyl blue†

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Photocatalytic degradation technology, as one of the most important advanced oxidation technologies for environmental remediation, has attracted great attention in recent years, but designing photocatalysts with excellent photocatalytic activity and good reusability remains a challenge. Herein, three CoS/CoO microspheres (CoS/CoO-M-1 (1), CoS/CoO-M-2 (2), and CoS/CoO-M-3 (3)) were prepared via a hydrothermal method using cobalt chloride hexahydrate, thiourea, deionized water and polyethylene glycol (PEG) with different polymerization degrees as raw materials, which have a uniform size distribution in the range of 5–24 μm and specific surface areas of 6.1924 $\text{m}^2 \text{g}^{-1}$ (1), 6.2870 $\text{m}^2 \text{g}^{-1}$ (2) and 6.6663 $\text{m}^2 \text{g}^{-1}$ (3). It is worth noting that all the three CoS/CoO microspheres showed a wide optical absorption range from the ultraviolet to the near-infrared (NIR) region and a high photocatalytic activity for degrading methyl blue under visible light (500 W metal halide lamp) irradiation. In order to improve the portability and recyclability of 3, a mixed matrix membrane (MMM) of 3 (3-MMM) was manufactured by coating a mixture dispersion of 3 and polyvinylidene fluoride (PVDF) on a glass substrate, which not only displayed excellent photocatalytic degradation performance, but also showed good portability and reusability (cycles > 12 times). Furthermore, adsorption and photocatalytic kinetics and possible mechanisms were studied.

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1 Introduction

Nowadays, with the rapid development of textile industry, a large quantity of wastewater containing organic dyes has been produced, which constitute 20% of the industrial water pollution in the world.^{1,2} These organic dyes in wastewater are chemically stable, highly toxic and difficult to directly biodegrade, which once released into the environment can cause various environmental problems.³ Thus far, although various physical and chemical treatment technologies have been developed to eliminate the toxicity of wastewater, these

technologies cannot completely decompose organic pollutants in industrial wastewater, and only transfer the pollutants from one state to the other, and even generate secondary pollutants.^{4–6} Recently, advanced oxidation technologies, such as photocatalytic degradation technology (PDT), have been considered as one of the most eco-friendly and efficient methods for purifying wastewater because of their strong oxidation ability, non-toxicity and low cost.^{7–9} However, a large number of photocatalysts with single components (such as metal oxides, metal sulfides, MOFs, and $\text{g-C}_3\text{N}_4$) or their composite components have been developed and reported successfully by esteemed investigators, but how to rationally design and fabricate of photocatalytic materials with excellent photocatalytic activity under simulated sunlight irradiation and good reusability is still a challenge, which severely limits the practical application of photocatalysis.^{10–13}

Thus far, compared with single-component photocatalysts, the construction of composite photocatalysts has been demonstrated to be one of the most efficient methods to enhance the photocatalytic activity under visible light irradiation.¹⁴ CoO has gained extensive interest in photocatalysis owing to its narrow band gap (2.4 eV), wide visible-light response, low cost and earth-abundance; however, it displays a weak photocatalytic performance due to rapid e^-/h^+ pair recombination.¹⁵ Similarly, CoS was also considered an

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attractive photocatalyst considering its excellent photoelectric properties, but it displays a large optical band gap (3.2 eV), poor visible-light absorption capacity, and rapid e^-/h^+ pair recombination.^{15,16} Hence, taking into consideration that CoO and CoS are both cobalt compounds and their special properties, a CoS/CoO composite photocatalyst with more excellent photocatalytic activity under visible light irradiation can be expected by modulating their structure, size or morphology, but the latter has little been considered. Meanwhile, the portability and recyclability are also required for practical application scenarios, but the powdery crystal photocatalysts are inconvenient to carry or recycle.¹⁷ It seems to be a good solution to fabricate a mixed matrix membrane (MMM) based on crystal photocatalysts and polymer matrixes.¹⁸ The flexible and self-supporting MMM can be easily tailored, carried, and conveniently removed from water, thus possessing many merits including processibility, portability and recyclability, which will facilitate the practical application of crystal photocatalysts. However, to the best of our knowledge, the MMM strategy has not been used in the synthesis of CoS/CoO/polymer membranes.¹⁹

Inspired by the above-mentioned factors, three CoS/CoO microspheres, namely, CoS/CoO-M-1 (1), CoS/CoO-M-2 (2), and CoS/CoO-M-3 (3) and 3-MMM were prepared and characterized. Interestingly, they displayed a wide optical absorption range from the ultraviolet, visible to the near-infrared region, and are full-solar range responsive photocatalysts, which could realize the photocatalytic degradation of methyl blue under visible light irradiation. In particular, 3-MMM showed not only excellent photocatalytic degradation performance, but also good portability and recyclability (cycles > 12 times). Furthermore, the adsorption and photocatalytic kinetics and possible mechanism were investigated.

2 Experiment section

2.1 Reagents and instruments

In this study, $\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$, thiourea, polyethylene glycol 200 (PEG-200), PEG-400, absolute ethanol, dimethylformamide (DMF), isopropyl alcohol (IPA), benzoquinone (BQ), ethylenediaminetetraacetic acid (EDTA) and polyvinylidene fluoride (PVDF) were used directly without purification. The crystal phase of the samples was detected using an XD-2 X-ray diffractometer (XRD). X-ray photo-electron spectroscopy (XPS) was performed using an Axis Ultra-DLD X-ray photoelectron spectrometer. Morphologies and elemental analyses were conducted using a JSM-6010 Plus scanning electron microscope equipped with EDS. The particle size of 1–3 was analysed using an HYL-1076 laser particle size distribution analyzer. The UV-vis-NIR absorption spectrum of the sample was recorded using a UV-3600 Plus UV-vis-NIR spectrophotometer.

2.1.1 Preparation of 1. First, 1.95 g $\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$ and 2.4 g thiourea were added to a mixture of 35 mL deionized water and polyethylene glycol 200 (PEG-200) ($V_{\text{(deionized water)}} : V_{\text{(PEG 200)}} = 4 : 1$) and stirred for more than 0.5 h. Then, it was subjected to a hydrothermal reaction at 180 °C for 16 h, and the precursor products were obtained by centrifugation and washed with

water and alcohol. The final products were then obtained by grinding and calcining at 700 °C in air for 1 h.

2.1.2 Preparation of 2. Sample 2 was synthesized in a manner analogous to that of 1 using PEG-200 and PEG-400 instead of PEG-200.

2.1.3 Preparation of 3. Sample 3 was synthesized in a manner analogous to that of 1 using PEG-400 instead of PEG-200.

2.1.4 Preparation of PVDF-M and 3-MMM. First, 160 mg PVDF was added into 2 mL DMF, dispersed ultrasonically for 0.5 h, stirred for more than 4 h, spread evenly onto a glass slide, and dried at 50 °C, and the film was removed to obtain PVDF-M. Then, 20 mg 3 and 160 mg PVDF were added into 2 mL DMF, dispersed ultrasonically for 0.5 h, stirred for more than 4 h, spread evenly on a glass slide, and dried at 50 °C, and the film was removed to obtain 3-MMM.

2.2 Photoelectric measurements and photocatalytic activity evaluation

Photoelectrochemical studies of the title samples were conducted using a CHI660E electrochemical workstation equipped with a three-electrode cell. Here, sample-modified ITO, a Pt wire and a Ag/AgCl electrode were used as the working, auxiliary and reference electrodes, respectively. To prepare the working electrode, 5 mg of microcrystalline powder was first added into a mixed solution containing 50 μL of Nafion and 420 μL of ethanol, which was then subjected to ultrasonic treatment for 30 min. Following this, the obtained suspensions were dropped onto a clean ITO substrate with an effective area of 1.0 cm^2 (1 $\text{cm} \times 1 \text{ cm}$). A 0.1 mol per L KCl aqueous solution (50 mL) was used as the supporting electrolyte solution, and a 300 W Xe lamp was used to simulate the sunlight illumination source. The Xe lamp irradiation interval was set at about 20 seconds. Electrochemical impedance spectroscopy (EIS) was performed under the same conditions.

The photocatalytic activities of 1, 2, 3 and 3-MMM were evaluated *via* the photodegradation of methyl blue dye using a multi-channel parallel photocatalytic reaction system with a 500 W metal halide lamp as the light source (NAI-GHY-DGNKW, Shanghai Nai Experimental Instrument Co., Ltd, China). Samples 1, 2, 3 and 3-MMM (containing 20 mg powder sample) were placed in a test tube, immersed in 20 mL aqueous solution containing contaminants (80 mg L^{-1}) of methyl blue, and stirred for 60 min in the dark to establish the adsorption-desorption equilibrium. At every 10 minutes of irradiation time, six parallel suspension solutions of 1, 2 or 3 were centrifuged and analyzed individually, while 3-MMM was tested directly without centrifugation using a UV-vis spectrophotometer.

3 Results and discussion

3.1 Structural characterization

The crystal structures and phase purity of as-synthesized 1, 2, 3 and 3-MMM were analyzed using X-ray diffraction (Fig. 1). As shown in Fig. 1a, the two characteristic diffraction peaks at 30.86° and 35.50° represent the Bragg reflection from the (100)



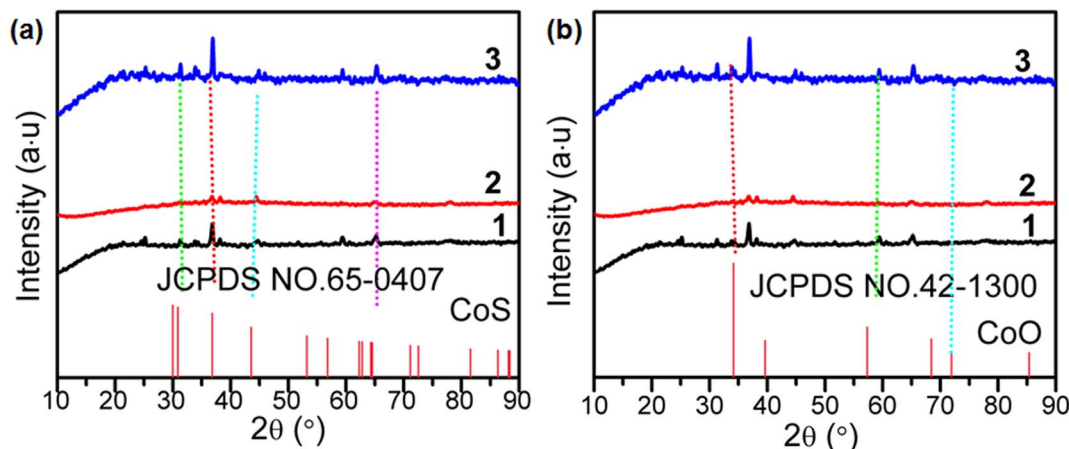


Fig. 1 (a) XRD patterns of CoS and 1–3. (b) XRD patterns of CoO and 1–3.

and (101) planes of the hexagonal phase of CoS (JCPDS card no. 65-0407), respectively.^{20,21} Both 43.56° and 64.55° are also consistent with JCPDS card no. 65-0407. The three characteristic diffraction peaks (2b) at 34.15° , 57.3° and 71.9° are also similar to the three characteristic peaks of CoO (JCPDS card no. 42-1300, Fig. 1b).^{22,23} The above results indicate the coexistence of CoS and CoO phases in as-synthesized 1, 2 and 3. Meanwhile, it can be seen from Fig. 1 that 1, 2 and 3 prepared from PEG solvents with different degrees of polymerization are not very similar,²⁴ and the crystallization degree of sample 3 is better than that of samples 1 and 2. It is obvious that the diffraction intensities of the primary peaks increase with the addition of PEG-400.²⁵ In addition, the XRD pattern of 3-**MMM** revealed the maintenance of the crystal structures and phase purity of 3 in the PVDF matrix (Fig. S1†).

The composition and elemental valence of 3 as a representative microsphere were further investigated by the XPS technique. It can be seen from Fig. 2a that the survey spectra of 3 clearly disclose the characteristic peaks of C 1s, Co 2p, O 1s and S 2p, which also indicates that these elements exist in 3. The fitted peaks of Co 2p and O 1s of 3 are shown in Fig. 2b and c. For the Co 2p spectrum (Fig. 2b), the peaks at 780.9 eV and 796.05 eV can be attributed to Co 2p_{3/2} and Co 2p_{1/2}, which also proves the presence of Co²⁺ in 3.²⁶ It is known that the oxidation state of cobalt oxide in XPS can be characterized by the gap between the Co 2p_{3/2} peak and the Co 2p_{1/2} peak and by intense shake-up peaks from the spin–orbit components.²⁷ The binding energy splitting between Co 2p_{3/2} and Co 2p_{1/2} was approximately 15.15 eV. This indicated that 3 consists of CoS and CoO, which matches well with the XRD results shown in Fig. 1. Fig. 2c shows the high-resolution XPS spectrum of O 1s in 3. The O 1s spectrum of 3 is deconvoluted into three peaks at 530.86 eV, 532.15 eV and 533.19 eV, the Co–O bond at 530.86 eV, and the chemical shift at 532.15 eV and 533.19 eV can be attributed to C–OH and adsorbed H₂O, respectively.^{28,29} It was also found that the binding energies of S 2p_{1/2} and S 2p_{3/2} are 162.84 eV and 161.65 eV respectively (Fig. 2d), indicating that S^{2–} exists in 3.^{26,30} Meanwhile, Fig. 2d shows the S 2p spectrum and the well-resolved peaks are in good agreement with the binding energies

of S 2p in cobalt sulfide.^{27,30} As shown in Fig. S2,† the survey spectra of 3-**MMM** also mainly contain four elements, namely C, O, S and Co, which is consistent with 3. However, the XPS peak strength of the Co element in 3-**MMM** is lower than that of the Co element in 3, which is attributed to the less content of 3 in the PVDF matrix. These results further confirmed that the crystal structure of 3 in the PVDF matrix is basically unchanged.

The morphological features of 1, 2 and 3 at 5 μm and 10 μm were analyzed by scanning electron microscopy (SEM). It can be seen from Fig. 3 that the morphology of 1, 2 and 3 with different solvents is not the same. The surface of 1 is rough with burrs, like the small burrs of “litchi”, and has uneven size distribution and serious aggregation.^{31,32} The surface of 2 is slightly rough, and has different sizes and relatively uniform distribution, among which the small number of microspheres are aggregated (Fig. 3b and b-1). The structures of 3 are like flowers with different sizes, uneven distribution and aggregation (Fig. 3c and c-1).^{33,34} However, the SEM images of 3-**MMM** reveals that there is little morphological difference between the sample 3 in 3-**MMM** and the as-prepared sample 3, which is probably due to the stirring for a long time in the film-making process (Fig. S3a†). In addition, the elemental compositions of 1, 2, 3 and 3-**MMM** were analyzed *via* EDS elemental mapping (Fig. 3a-2-a-4, b-2-b-4, c-2-c-4 and S3b†), indicating the homogeneous distribution of Co, O and S elements, and that no other elements are present.³⁵

Furthermore, the particle size and specific surface area of 1, 2 and 3 were analyzed using a laser particle size distribution analyzer. As shown in Fig. S4,† the particle size of 1, 2 and 3 are mainly distributed between 2 and 20 μm , and the average particle size is related to the polymerization degree of PEG ranging from 8.61 to 12.63 μm (ref. 36) (12.63 μm (1), 9.60 μm (2), and 8.61 μm (3), respectively). The specific surface areas of the three samples are 544.1 $\text{m}^2 \text{m}^{-3}$ (1), 728.76 $\text{m}^2 \text{m}^{-3}$ (2) and 811.32 $\text{m}^2 \text{m}^{-3}$ (3), respectively. Notably, with the increase in the polymerization degree of solvent PEG, the particle size of microsphere becomes smaller and smaller, and their specific surface area becomes higher and higher. Furthermore, N₂ adsorption and desorption experiments were carried out on

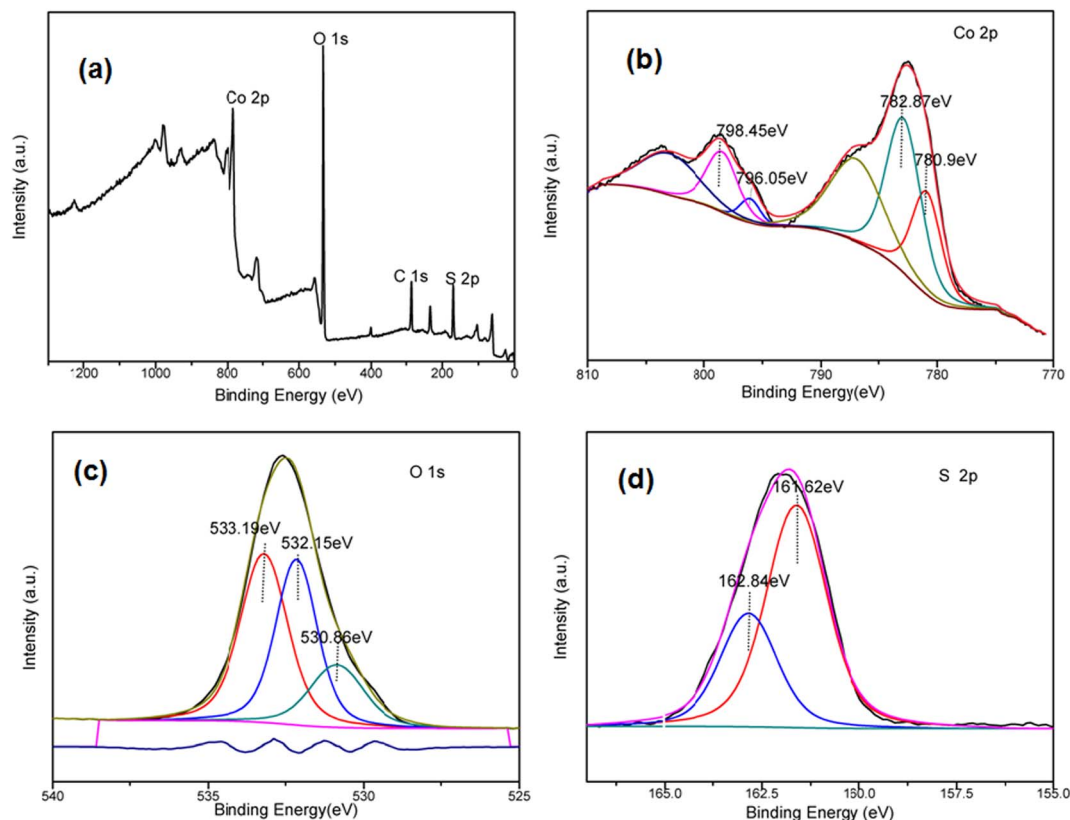


Fig. 2 XPS spectra of **3**: (a) survey spectra of **3**; (b) Co 2p XPS spectra; (c) O 1s XPS spectra; and (d) S 2p XPS spectra.

samples **1**, **2** and **3** at 77 K and the specific surface areas of **1**, **2** and **3** were tested (Fig. S5†), the resulting BET surface areas of three samples are $6.1924 \text{ m}^2 \text{ g}^{-1}$, $6.2870 \text{ m}^2 \text{ g}^{-1}$ and $6.6663 \text{ m}^2 \text{ g}^{-1}$, respectively, which is consistent with the law tested using a HYL-1076 laser particle size distribution analyzer (specific surface area: $3 > 2 > 1$). When the relative pressure p/p_0 varies between 0.4 and 1.0, the nitrogen isothermal adsorption–

desorption curve has a relatively obvious jump, and the samples also have hysteresis loops in the shape of small shuttles, and the curves of **1**, **2** and **3** are similar.

3.2 Optical characterization

The optical absorption properties of the as-synthesized black CoS/CoO microspheres were analyzed by UV-vis-NIR diffuse

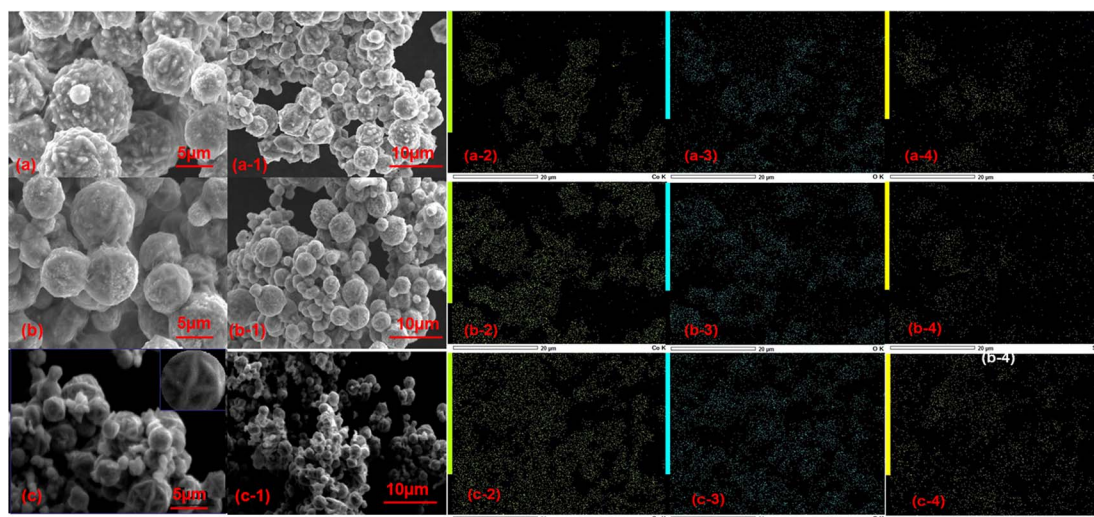


Fig. 3 ((a), (a-1), (a-2), (a-3) and (a-4)) SEM and EDS mappings of **1** respectively. ((b), (b-1), (b-2), (b-3) and (b-4)) SEM and EDS mappings of **2** respectively. ((c), (c-1), (c-2), (c-3) and (c-4)) SEM and EDS mappings of **3** respectively.



reflectance spectroscopy. From Fig. 4a, we can clearly see that **1**, **2** and **3** have a wide absorption range from the ultraviolet and visible regions to the NIR region, and an absorption edge up to about 1800 nm, which are in accordance with their dark appearance, and are favorable for full-solar-range-driven photocatalysis. According to the previous study, the band gaps (E_g) of the samples could be estimated by the plot of $(\alpha h\nu)^2$ versus photon energy ($h\nu$). As shown in Fig. 4b, the E_g values of **1**, **2** and **3** were estimated to be 0.39, 0.70, and 0.97 eV, respectively (Fig. 4b), which are lower than the most reported E_g values of CoO (2.4 eV), CoS (3.2 eV), other single components (such as metal oxides, metal sulfides, MOFs, and g-C₃N₄) or their composite components.^{37–39} The above-mentioned results show that the three CoS/CoO microspheres can be activated by visible light irradiation, and can greatly enhance their photocatalytic activity.

3.3 Photoelectric measurements

The optical bandgaps of title samples fall in the visible region, which encouraged us to further study their photocurrent response abilities that were generally used to evaluate the suitability as promising photocatalytic materials. The time-dependent photocurrent curves of title samples were recorded, as shown in Fig. 5a. As can be clearly seen, samples **1–3** exhibited the fast and obvious photoelectric switching behaviors at the beginning and end of illumination. Even after the multiple cycles, their on/off photocurrent response ratios did not decrease significantly, accounting for the good operation stabilities and performance reproducibilities. Under the visible light conditions, the photocurrent density of sample **3** was 37.5 nA cm^{−2}, which was better than those of samples **1** and **2** (25.0 and 12.5 nA cm^{−2}, respectively). In general, the high photocurrent densities indicated that the obtained materials have a better carrier transport efficiency, which have often appeared in the cases of some previous work.^{40,41} As shown in Fig. 5b, the electrochemical impedance spectroscopy (EIS) results indicate that **3** has a smallest radius of Nyquist plot, which results in a lower charge transfer resistance of **3** than those of **1** and **2**. In short, the photogenerated charge and migration efficiency of **3**

are higher than those of **1** and **2**, which is consistent with the results of photocurrent.

3.4 Photocatalytic activity for MB degradation

Inspired by the structural characteristics and the narrow band gap of the title samples, methyl blue (MB) was selected as a model pollutant to study their photocatalytic activities.

In order to easily describe the process of sample adsorption and photocatalytic degradation of methyl blue solution, the decontamination efficiency can be expressed as

$$\text{Decontamination efficiency (\%)} = \left(1 - \frac{C_t}{C_0}\right) \times 100\% \quad (1)$$

where C_0 and C_t (mg L^{−1}) represent the initial concentration and the concentration at time t of the methyl blue solution, respectively.

The reaction kinetics of the adsorption and photocatalytic degradation of methyl blue solution was found to follow a pseudo-first-order kinetics, as the plot of $\ln(C_0/C_t)$ against the time fits a linear relationship according to the following equation:⁴²

$$\ln\left(\frac{C_0}{C_t}\right) = kt \quad (2)$$

As clearly displayed in Fig. 6a and S6,[†] a methyl blue solution (80 mg L^{−1}, 20 mL) could be absorbed by the titled samples with the increase in stirring time, and then reached the equilibrium of adsorption–desorption in about 60 min,⁴³ indicating their relatively low adsorption capacity. For sample **1**, the adsorption rate of MB reached 46.11% after 60 min. For sample **2**, its adsorption rate reached 46.80% after 60 min. Compared with **1** and **2**, sample **3** shows a slightly faster adsorption capacity, which can reach 50.72% after 60 min, which may be attributed to the larger specific surface area of **3** than those of **1** and **2**.^{44,45} Fig. 6b shows the secondary reaction kinetics of the adsorption of **1–3**, respectively. Samples **1** and **2** do not have a good linear relationship, while **3** shows a good linear relationship, where

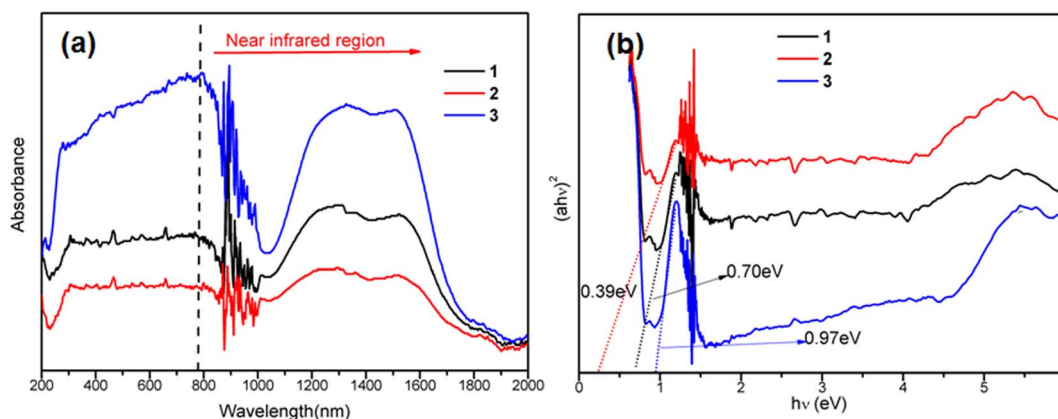


Fig. 4 (a) Solid-state UV-vis-NIR absorption spectra of **1**, **2** and **3**. (b) Plots of $(\alpha h\nu)^2$ versus photon energy ($h\nu$) for the bandgap energies of **1**, **2** and **3**.

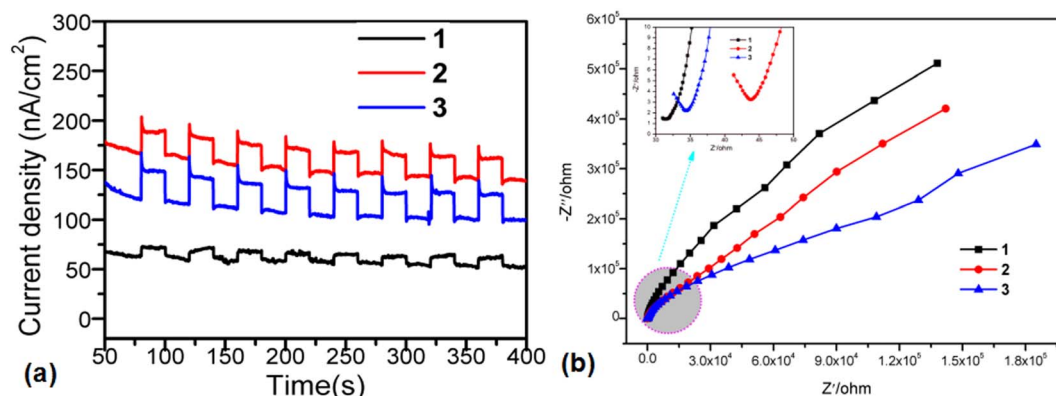


Fig. 5 (a) Photocurrent response behavior of 1, 2 and 3. (b) Electrochemical impedance spectra (EIS) of 1, 2 and 3.

the linear fitting constant is $R^2 = 0.9876$, indicating that the adsorption kinetics are stable compared with 1 and 2.

The above-mentioned adsorption results indicate that MB could not be completely adsorbed by the CoS/CoO microspheres, which can be further investigated for their photocatalytic activity.^{46,47} Samples 1–3 were put into an MB solution, which was kept in the darkness for 60 min to ensure the absorption/desorption equilibrium state prior to the irradiation. Subsequently, under visible light irradiation, the absorbance intensity of MB molecules in the UV-vis spectra decreased gradually as the irradiation time increases (Fig. S7†). As shown in Fig. 6a, the photocatalytic degradation rates of 1, 2 and 3 at 10 min reached 25.00%, 24.90% and 56.74% respectively. After light irradiation for 60 minutes, their photocatalytic degradation rates reached 96.00% for 1, 95.87% for 2, and 97.30% for 3, respectively. However, in the absence of the catalysts, the concentration of MB with irradiation time has hardly any reduction and the degradation is nearly negligible (Fig. S8†).

These results suggest that these CoS/CoO microspheres could be considered as good candidates for efficient visible-light-driven photocatalysts. As shown in Fig. 7, the photocatalytic degradation process also follows first-order reaction kinetics. Samples 1 and 2 showed similar rates ($k_1 = 0.0885 \text{ min}^{-1}$, $k_2 = 0.0898 \text{ min}^{-1}$), and the degradation rate of 3

($k_3 = 0.109 \text{ min}^{-1}$) is relatively fast, and the degradation is basically balanced in 30 min. Sample 3 performed with a slightly higher photocatalytic rate than 2 and 1, which may be attributed to its stronger photocurrent density, and more effective separation of photogenerated electron-hole pairs. In a word, *via* the adsorption and photocatalytic degradation process, the total decontamination efficiencies of 1, 2 and 3 reached 97.84%, 97.83% and 98.68% for MB respectively, indicating almost complete decomposition of organic pollutants (Fig. 6a and S9†).

The stability of a photocatalyst is an important index to evaluate its research value and actual utilization.⁴⁸ As sample 3 has a higher photocatalytic rate than that of 1 and 2, sample 3 is used to investigate the stability by adding 20 mg sample into 20 mL methyl blue solution. After adsorption and photocatalytic degradation of MB, the sample was centrifuged, filtered, washed, and dried for secondary use, as shown in Fig. 8.^{49,50} Unfortunately, the photocatalytic degradation rate of 3 dropped to 77% after 12 times of consecutive uses, which can arise from the mass loss of powdery crystal photocatalysts during the centrifugation and collection process. In order to improve the portability and recyclability, a mixed matrix membrane (MMM) of 3 (3-MMM) was produced by coating a mixture dispersion of 3 and a polyvinylidene fluoride membrane (PVDF-M) onto a glass

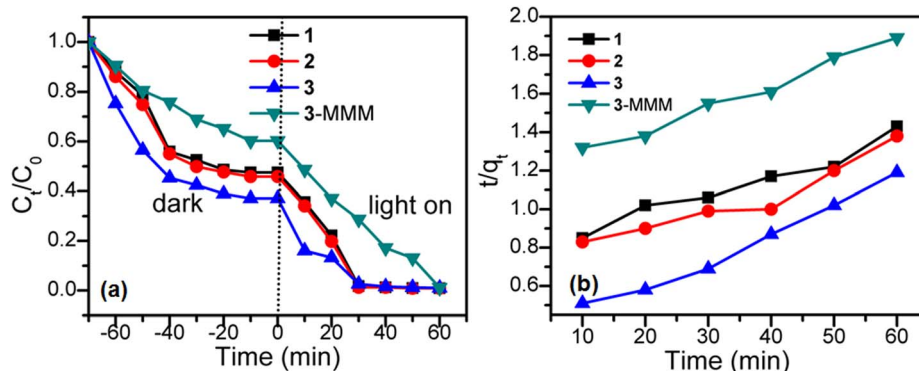


Fig. 6 (a) Adsorption and photocatalytic degradation of the methyl blue solution (80 mg L^{-1} , 20 mL) by 1, 2, 3 and 3-MMM. (b) Pseudo-second-order adsorption kinetic fitting of 1, 2, 3 and 3-MMM in the MB solution.



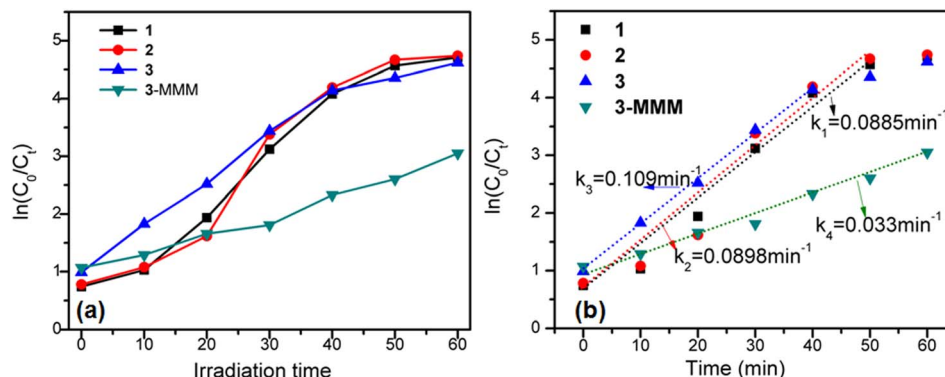


Fig. 7 (a) First-order kinetic fitting of photocatalytic degradation of 1, 2, 3 and 3-MMM in the MB solution. (b) Reaction rate of photocatalytic degradation of MB.

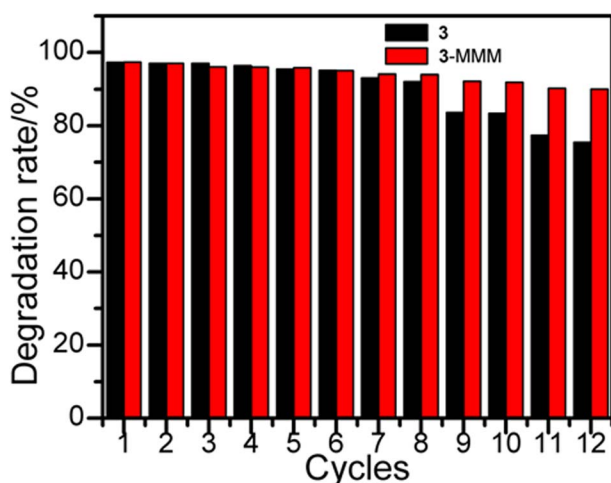


Fig. 8 Methyl blue was photocatalyzed for 12 cycles by 3 and 3-MMM.

substrate, and then dried at 50 °C for 12 h. For sample **3-MMM**, the adsorption rate of MB reached 33.45% at 60 min, and also shows low adsorption rate and a good linear relationship, where the linear fitting constant is $R^2 = 0.9901$ (Fig. 6a). Furthermore, using the above-mentioned experimental condition, the stability of **3-MMM** was tested directly without centrifugation, filtration, washing and drying, while it was taken out with tweezers. It is worth noting that the photocatalytic degradation rate of **3-MMM** reached 97.34% after light irradiation for 60 minutes, and the total decontamination efficiency reached 98.23% and remained basically unchanged even after the cycle of 12 times, which is better than the most reported photocatalysts (Table S1†). Meanwhile, as shown in Fig. S8 and S9,† it can be seen that **PVDF-M** has no obvious adsorption and photocatalytic effect on methyl blue. The above-mentioned result reveals that **3-MMM** not only has excellent photocatalytic degradation performance, but also possesses good portability and reusability.¹⁰ Moreover, the comparison of the XRD pattern before and after recycling confirms the maintenance of the structure of **3-MMM** (Fig. S1†), indicating that it possesses good stability.

3.5 Possible photocatalytic mechanism

In order to understand the possible mechanism of photocatalytic degradation, the radical trapping experiments were tested. In this study, isopropyl alcohol (IPA, capture% $\cdot\text{OH}$), ethylenediamine tetraacetic acid (EDTA-2Na, capture% h^+) and benzoquinone (BQ, capture% O_2^-) were used to determine the main active species that are generated in the photo-degradation process. Before light irradiation, 0.5 mmol IPA, EDTA-2Na or BQ were added to 20 mL MB 80 mg L^{-1} solution containing 20 mg of 3, respectively. Fig. 9 shows that the photocatalytic degradation rate of 3 was reduced from 97.30% to 5.30% and 22.37% after adding BQ and EDTA-2Na to the above solution respectively, illustrating that O_2^- and h^+ are the important active species in the photocatalytic reaction.⁵¹ By adding IPA, the photocatalytic degradation rate was only reduced to 79.92%, which illustrates that $\cdot\text{OH}$ is not the most critical reaction species.⁴⁹

On the basis of the above-mentioned experimental results and the similar reported work, the possible photocatalytic mechanism was proposed for the photodegradation of MB by 3. During the photocatalytic reaction, the electrons and holes of 3 containing CoS/CoO microspheres can first be separated under

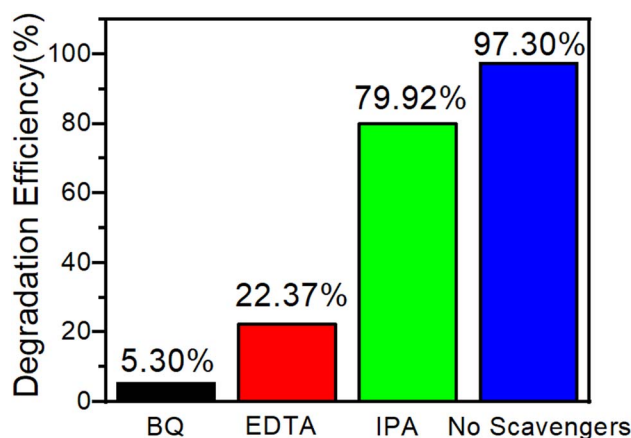


Fig. 9 Trapping experiment of active species during the photocatalytic degradation of MB over 3 under light irradiation.



light irradiation, in which the electrons are moved to the CB, departing from the holes in the VB.^{49,52} The electrons are further captured by O₂ in the solution to form superoxide ions ([•]O₂[−]) that will then be converted into less [•]OH, which collectively participate in the degradation of MB. At the same time, the remaining photoinduced holes also contribute to the decomposition of MB.

4 Conclusion

In summary, using cobalt chloride hexahydrate, thiourea, deionized water and polyethylene glycol (PEG) with different polymerization degrees as raw materials, three CoS/CoO microspheres have been prepared by a simple hydrothermal method. Notably, with the change in the polymerization degree of solvent PEG, the morphologies, particle sizes and specific surface areas of three microspheres can be effectively modulated, leading to different band gaps, photocurrent response abilities and photocatalytic degradation rates. In particular, the mixed matrix membrane of 3 (3-**MMM**) not only showed excellent photocatalytic degradation rate of MB, but also showed good portability and reusability (cycles > 12 times), which has potential prospects in the industrial application to treat organic pollutants in waste water. This result will provide guidance for the further rational design and synthesis of novel excellent visible-light-driven photocatalysts and their mixed matrix membranes based on the energy-band engineering and mixed matrix membrane strategy, and the elucidation of the structure–property relationship. Further studies by our group are in progress.

Data availability

All relevant data are available in the manuscript and its online ESI.†

Conflicts of interest

There is no conflict to declare.

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References

- 1 X. Linghu, Y. Shu, L. Liu, Y. Zhao, J. Zhang, Z. Chen, D. Shan and B. Wang, *J. Water Proc. Eng.*, 2023, **54**, 103994.

- 2 I. Ahmad, M. Muneer, A. S. Khder and S. A. Ahmed, *ACS Omega*, 2023, **8**, 22708–22720.
- 3 S. Selvinsimpson, S. Eva Gnana Dhana Rani, A. GaneshKumar, R. Rajaram, I. Sharmila Lydia and Y. Chen, *Environ. Res.*, 2021, **195**, 110787.
- 4 N. A. Alamrani and H. A. Al-Aoh, *Adsorpt. Sci. Technol.*, 2021, **2021**, 1–12.
- 5 T. Huang, M. He, Y. Zhou, S. Li, B. Ding, W. Pan, S. Huang and Y. Tong, *RSC Adv.*, 2016, **6**, 100392–100400.
- 6 S. He, Y. Chen, X. Li, L. Zeng and M. Zhu, *ACS ES&T Eng.*, 2022, **2**, 527–546.
- 7 K. Vellingiri, V. Choudhary, D. W. Boukhvalov and L. Philip, *ACS ES&T Water*, 2022, **2**, 1475–1499.
- 8 C. H. Mak, X. Han, M. Du, J. Kai, K. F. Tsang, G. Jia, K. C. Cheng, H. H. Shen and H. Y. Hsu, *J. Mater. Chem. A*, 2021, **9**, 4454–4504.
- 9 C. C. Wang, J. R. Li, X. L. Lv, Y. Q. Zhang and G. Guo, *Energy Environ. Sci.*, 2014, **7**, 2831–2867.
- 10 J. Yang, H. Song, Y. Zhang and X. Zhu, *Surf. Interfaces*, 2021, **24**, 101092.
- 11 F. T. Geldasa, M. A. Kebede, M. W. Shura and F. G. Hone, *RSC Adv.*, 2023, **13**, 18404–18442.
- 12 H. R. Dihom, M. M. Al-Shaibani, R. M. S. Radin Mohamed, A. A. Al-Gheethi, A. Sharma and M. H. B. Khamidun, *J. Water Proc. Eng.*, 2022, **47**, 102705.
- 13 Z. Saddique, M. Imran, A. Javaid, S. Latif, N. Hussain, P. Kowal and G. Boczkaj, *Water Resour. Ind.*, 2023, **29**, 100211.
- 14 H. Jiang, H. Chen, Y. Fu and C. Guo, *Appl. Surf. Sci.*, 2023, **611**, 155555.
- 15 H. R. Pouretedal and M. Kiyani, *J. Iran. Chem. Soc.*, 2014, **11**, 271–277.
- 16 D. Inoue, T. Yoshikawa, T. Okumura, Y. Yabuki and M. Ike, *J. Hazard. Mater.*, 2021, **414**, 125497.
- 17 Q. Fang, C. Chen, Z. Yang, X. A. Chen, X. Chen and T. Liu, *J. Alloys Compd.*, 2020, **826**, 154084.
- 18 A. Gulinigaer, A. Salamu, Y. Tuerxun, Y. Rouzhahong, M. Mamat, L. Xie, Z. Wu, F. Xiaerding and Y. Sun, *Russ. J. Phys. Chem. A*, 2020, **93**, 2852–2857.
- 19 W. Xiang, Q. Wang, Z. Li, J. Dong, J. Liu, L. Zhang, T. Xia, Y. He and D. Zhao, *Sep. Purif. Technol.*, 2024, **330**, 125268.
- 20 Y. Ohigashi, A. Higuchi, N. C. Rosero-Navarro, A. Miura and K. Tadanaga, *J. Sol-Gel Sci. Technol.*, 2022, **101**, 24–28.
- 21 Z. Shi and A. H. Jayatissa, *Prog. Nat. Sci.*, 2020, **30**, 352–359.
- 22 W. Shi, J. Gao, H. Sun, Z. Liu, F. Guo and L. Wang, *Chin. J. Chem. Eng.*, 2022, **49**, 213–223.
- 23 A. S. Alkorbi, H. M. Asif Javed, S. Hussain, S. Latif, M. Shabir Mahr, M. S. Mustafa, R. Alsaiari and N. Alhemiary, *Opt. Mater.*, 2022, **127**, 112259.
- 24 J. K. Kim, J. Y. Ju, S. K. Choi, S. Unithrattil, S. S. Lee, Y. Kang, Y. Kim, W. B. Im, S. Choi and J. Power, *Sources*, 2018, **378**, 562–570.
- 25 H. Kawasaki, T. Ohshima, Y. Yagyu, T. Ihara, K. Mitsuhashi, H. Nishiguchi and Y. Suda, *J. Appl. Phys.*, 2022, **61**, SA1019.
- 26 S. Zhang, X. Yu, F. Yan, C. Li, X. Zhang and Y. Chen, *J. Mater. Chem. A*, 2016, **4**, 12046–12053.



- 27 X. Liu, Q. Li, Y. Fan, W. Chen, L. Lin, W. Xu, H. Zhuang and Q. Li, *Mater. Lett.*, 2022, **315**, 132010.
- 28 L. Qi, J. Yu and M. Jaroniec, *Phys. Chem. Chem. Phys.*, 2011, **13**, 8915–8923.
- 29 M. He, Y. Zhou, T. Huang, S. Nie, Y. Wang, Z. Xu, Y. Huo, R. Xu, X. Chen and H. Peng, *Compos. Sci. Technol.*, 2020, **200**, 108403.
- 30 S. Dhiman and B. Gupta, *Environ. Technol. Innovation*, 2021, **23**, 101765.
- 31 K. Qi, J. Yu and K. Chen, *Cryst. Res. Technol.*, 2013, **48**, 1083–1086.
- 32 N. Hu, L. Huang, W. Gong and P. K. Shen, *ACS Sustain. Chem. Eng.*, 2018, **6**, 16933–16940.
- 33 X. Chen, N. Li, S. Xu, H. Wang and Y. Cai, *Materials*, 2018, **11**, 511.
- 34 G. Liu and J. Shao, *J. Mater. Chem. A*, 2017, **5**, 9801–9806.
- 35 J. C. Xing, Y. L. Zhu, M. Y. Li and Q. J. Jiao, *Electrochim. Acta*, 2014, **149**, 285–292.
- 36 Y. Mao, X. Shen, Z. Wu, L. Zhu and G. Liao, *J. Alloys Compd.*, 2020, **816**, 152604.
- 37 P. J. Wibawa, H. Saim, M. A. Agam and H. Nur, *Phys. Procedia*, 2011, **22**, 524–531.
- 38 M. Tian, J. Wang, R. Sun, M. Yao and L. Li, *Catalysts*, 2021, **11**, 515.
- 39 T. Luo, X. Sun, D. Ma, G. Wang, F. Yang, Y. Zhang, J. Huang, H. Zhang, J. Wang and F. Peng, *J. Phys. Chem. C*, 2023, **127**, 1372–1380.
- 40 J. Lu, J. Zhang, X. Yang, D. Jia and S. Zhao, *Inorg. Chim. Acta*, 2022, **537**, 120962.
- 41 B. Zhang, M. Pang, J. Li, Y. Yang, M. Z. Liu and X. Chen, *Mol. Cryst. Liq. Cryst.*, 2022, **757**, 85–94.
- 42 X. Bu, Y. g. Liu, C. A. Xie and B. Wang, *ACS Appl. Opt. Mater.*, 2023, **1**, 688–700.
- 43 G. Yang, Z. F. Yan and T. C. Xiao, *Appl. Surf. Sci.*, 2012, **258**, 8704–8712.
- 44 W. Sangchay, *Energy Procedia*, 2016, **89**, 170–176.
- 45 C. Zhang, Y. Li, M. Li, D. Shuai, X. Zhou, X. Xiong, C. Wang and Q. Hu, *J. Hazard. Mater.*, 2021, **420**, 126607.
- 46 S. Okeil, J. Krausmann, I. Dönges, S. Pflieger, J. Engstler and J. J. Schneider, *Dalton Trans.*, 2017, **46**, 5189–5201.
- 47 C. Shi, F. Tao and Y. Cui, *Int. J. Biol. Macromol.*, 2018, **114**, 400–407.
- 48 N. Cruz-González, O. Calzadilla, J. Roque, F. Chalé-Lara, J. K. Olarte and M. Meléndez-Lira, *Int. J. Photoenergy*, 2020, **2020**, 1–9.
- 49 X. Qiao, C. Wang and Y. Niu, *J. Hazard. Mater.*, 2020, **391**, 122121.
- 50 M. Guan, H. Li, X. Chen and S. Zhang, *Waste Manage.*, 2021, **120**, 635–641.
- 51 Q. Zhu, Y. Sun, F. Na, J. Wei, S. Xu, Y. Li and F. Guo, *Appl. Catal., B*, 2019, **254**, 541–550.
- 52 H. Xu, X. Wang, M. Dai, W. Wang, D. Lu, M. Zhang, Y. Chen and H. Song, *Mater. Chem. Phys.*, 2021, **260**, 124158.

