

Cite this: *Chem. Sci.*, 2021, 12, 938

All publication charges for this article have been paid for by the Royal Society of Chemistry

Integrating hydrogen production with anodic selective oxidation of sulfides over a CoFe layered double hydroxide electrode†

Lina Ma,[‡] Hua Zhou,[‡] Ming Xu,^a Peipei Hao,^a Xianggui Kong^a and Haohong Duan^{*,b}

Replacing the sluggish oxygen evolution reaction (OER) with oxidation reactions for the synthesis of complex pharmaceutical molecules coupled with enhanced hydrogen evolution reaction (HER) is highly attractive, but it is rarely explored. Here, we report an electrochemical protocol for selective oxidation of sulfides to sulfoxides over a CoFe layered double hydroxide (CoFe-LDH) anode in an aqueous-MeCN electrolyte, coupled with 2-fold promoted cathodic H₂ productivity. This protocol displays high activity (85–96% yields), catalyst stability (10 cycles), and generality (12 examples) in selective sulfide oxidation. We demonstrate its applicability in the synthesis of four important pharmaceutical related sulfoxide compounds with scalability (up to 1.79 g). X-ray spectroscopy investigations reveal that the CoFe-LDH material evolved into amorphous CoFe-oxyhydroxide under catalytic conditions. This work may pave the way towards sustainable organic synthesis of valuable pharmaceuticals coupled with H₂ production.

Received 5th October 2020
Accepted 11th November 2020

DOI: 10.1039/d0sc05499b

rsc.li/chemical-science

Introduction

Electrochemical water splitting is considered to be a promising hydrogen production approach to deal with the increasing global energy demand and environmental problems associated with fossil fuel utilization.¹ Water electrolysis involves two half-reactions, including H₂ and O₂ evolution reactions (HER and OER).² The overall reaction rate is often restricted by the anodic OER because of its more sluggish kinetics.^{2c} As a result, much higher overpotential is needed for the OER to match the H₂ production rate, thereby undermining the overall energy conversion efficiency.³ Although advanced non-noble-metal OER electrocatalysts have been developed, accomplishing an OER efficiency comparable with that of the HER still remains a challenge.⁴ In addition, the economic value of O₂ is far inferior to that of H₂. Recently, electrooxidation of low-cost organic agents has emerged as an alternative strategy to replace the OER, achieving lower overpotential for total water splitting and at the same time producing high-value chemicals.⁵ Electrooxidation has been exploited for converting biomass-derived platform chemicals, *e.g.*, ethanol, glycerol, and 5-hydroxymethylfurfural (HMF).⁶ Despite the advanced concept of co-

production of valuable chemicals and H₂, only a handful of simple molecules were studied.^{6f,g} The anodic synthesis of functional and complex organic molecules, especially pharmaceuticals with medicinal significance, has rarely been explored.

In the family of pharmaceuticals, organosulfur compounds play essential roles in many biological processes linked with human disease therapy.⁷ For example, sulforaphane offers efficient chemoprotection against various cancers (*e.g.* prostate, lung, breast, and colon cancer).^{7f} Direct oxidation of their sulfide precursors is a straightforward and atom-economical synthetic method, and thus has been widely applied for the synthesis of sulfoxides.⁸ However, current synthetic methods often require homogeneous catalysts (*e.g.* Mn- and Fe-based catalysts) and strong oxidizing agents such as H₂O₂.⁹ Additional oxidants such as peroxy acids, prevalent iodine reagents or oxone are also needed.¹⁰ In 2012, He and co-workers reported a catalyst-free protocol for the selective oxidation of sulfides with an inorganic oxidant oxone in ethanol.¹¹ However, the utilization of oxone may result in purification issues and environmental contamination. More importantly, over-oxidation of sulfoxide to sulfone or co-oxidation of other functional groups in the sulfoxide molecule would occur and generate medically useless products. Recently, Xu and co-workers reported an efficient heterogeneous polyoxovanadate catalyst for sulfide oxidation; nevertheless the reaction was assisted by a strong oxidant (*tert*-butylhydroperoxide (TBHP)) at elevated temperatures.¹² Jiang's group reported a photocatalysis strategy to achieve sulfide selective oxidation driven by visible light under mild conditions, but the use of homogeneous catalysts (UO₂(OAc)₂·2H₂O) would make separation difficult.¹² Therefore, the

^aState Key Laboratory of Chemical Resource Engineering, Beijing University of Chemical Technology, Beijing 100029, China

^bDepartment of Chemistry, Tsinghua University, Beijing 100084, China. E-mail: hhduan@mail.tsinghua.edu.cn

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0sc05499b

‡ These authors contributed equally.



Table 1 Optimization of reaction conditions

Entry	Deviation from standard conditions	Conv. ^a (%)	Yield 2a ^a (%)	Yield 3a ^a (%)
1	None	>99	85(81)	Trace
2	H ₂ O as solvent	39	35(33)	2
3	MeCN as solvent	12	Trace	nd
4	N ₂ atmosphere	>99	81(81)	Trace
5	CC as the anode	39	11(10)	Trace
6	No electric current	<5	nd	nd

^a Conversion and yields were determined by ¹H NMR using CH₂Br₂ as an internal standard. nd = not detected. The isolated yields are given in parentheses. Standard reaction conditions and variations on the electrochemical oxidation of diphenyl sulfide: **1a** (0.5 mmol), *n*Bu₄NH₂PO₄ (0.5 mmol), MeCN/H₂O (2.0 mL, 1 : 1 v/v), RT, 2 h, CoFe-LDH anode (working area: 1 cm²), Pt cathode, applied potential: 2.0 V vs. Ag/AgCl (passing charge of about 96C), under air, unless otherwise noted.

which represents a high performance compared to existing homo- and heterogeneous catalysts (ESI, Table S3†). Specifically, H₂O was initially tested as the solvent for its wide use in the OER and anodic oxidations, but moderate conversion and yield of **2a** were obtained, accompanied by overoxidation (entry 2, Table 1). Replacing it by MeCN or other organic solvents showed even lower activities (entry 3 in Table 1, entries 1–4 in the ESI, Table S2†). We found that using a mixed solvent of MeCN and H₂O could furnish significantly higher efficacy, which is likely due to MeCN showing higher conductivity and capacity to dissolve the electrolyte, substrate, and reactants compared with other employed solvents, which may lead to catalytic enhancement (entries 5–6 in the ESI, Table S2†).¹⁹ The use of electrolytes other than *n*Bu₄NH₂PO₄ showed lower activity (entries 7–9 in the ESI, Table S2†), which is due to the promotion effect of *n*Bu₄NH₂PO₄ for the oxidation reaction.^{12b} In addition, the yield of **2a** didn't show obvious change when the reaction atmosphere was varied from air to N₂ (entry 4 in Table 1), indicating that H₂O in the electrolyte could likely be the source of oxygen in **2a**. It should be noted that under our reaction conditions in air, it remains possible that oxygen in the product can be from the air. A blank experiment using CC as the anode showed much lower conversion for sulfide oxidation, suggesting CoFe-LDH/CC was essential for catalytic activity of sulfide oxidation (entry 5 in Table 1). Finally, no reaction occurred in the absence of electric current, demonstrating the nature of electrocatalysis (entry 6 in Table 1). Pt ions were not observed in the electrolyte after electrooxidation by using an Inductive Coupled Plasma Emission Spectrometer (ICP), indicating that Pt was not leached into the electrolyte. According to the conversion of **1a** and total yields of products **2a** (or with **3a**), there remained other byproducts while their structures can't be recognized as no other signals were observed by crude NMR analysis.

In the following catalytic studies, the catalytic reactions were examined over the as-synthesized CoFe-LDH/CC under the optimized conditions. Fig. 3a shows the linear sweep voltammetry (LSV) curves in the presence and absence of **1a** over the

CoFe-LDH/CC anode. After adding **1a**, the current density significantly increases, and the required potential for gaining current density of 5 mA cm⁻² dramatically decreased from 1.90 V (without **1a**) to 1.39 V (with **1a**). The desired product **2a** increased with the time, and to our delight, the overoxidized product (sulfone **3a**) was rarely observed (<5% yield, Fig. 3b). More importantly, **2a** can be obtained on a gram scale with 83% yield (up to 1.63 g) (for details, see Experimental procedures in the ESI†). We then assembled a two-electrode setup in which CoFe-LDH/CC was used as the anode for **1a** oxidation and Pt as the cathode for the HER. Impressively, it shows two-fold higher

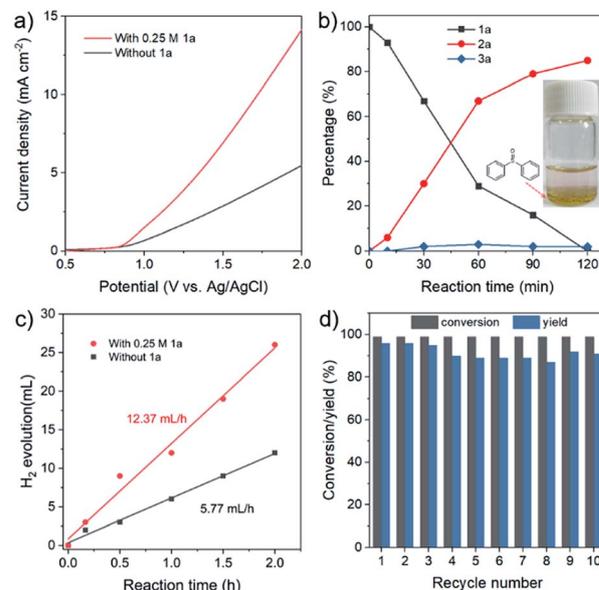


Fig. 3 (a) LSV curves of the CoFe-LDH/CC anode at a scan rate of 50 mV s⁻¹ in MeCN/H₂O (2.0 mL, 1 : 1 v/v) with and without 0.5 mmol **1a**. (b) Tracking experiments of the sulfoxidation procedure. (c) H₂ production with and without **1a**. (a–c) CoFe-LDH/CC anode and Pt cathode. (d) Cycle-dependent conversions for (methylsulfinyl) benzene.



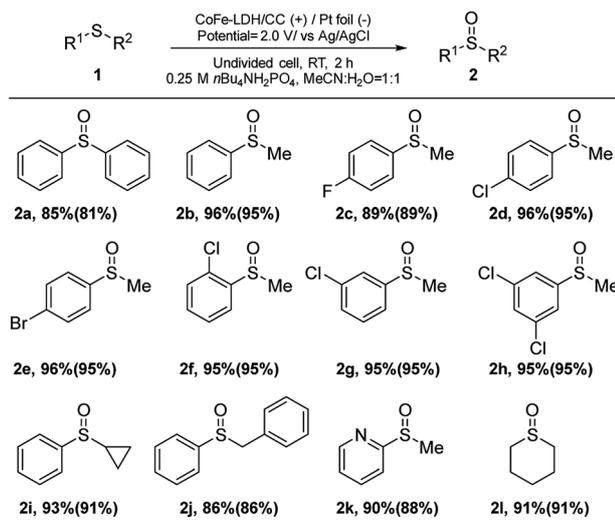


Fig. 4 Substrate scope for the formation of sulfoxides. Yields were determined by ¹H NMR using CH₂Br₂ as an internal standard. The isolated yields were given in parentheses.

activity for H₂ production compared with the one without **1a** that follows the OER process (Fig. 3c), indicative of enhanced energy efficiency. To a certain extent, the organic solvent MeCN has some slight effect on the faradaic efficiency of H₂ (90%), lower than the faradaic efficiency of 96% in aqueous electrolyte. After ten-cycle runs, no apparent decrease of activity was observed (Fig. 3d), indicative of promising catalytic stability. The structural stability of the used CoFe-LDH/CC anode was also demonstrated (ESI, Fig. S2†).

Subsequently, the generality of the electrocatalytic selective oxidation using CoFe-LDH/CC was studied by using 12 examples (Fig. 4). In general, the sulfides and their derivatives with functional groups including -H, -F, -Cl and -Br worked very well to produce sulfoxides (2a–2g) under the optimized conditions. The reaction efficiency was not affected by steric hindrance (2d, 2f and 2g). A multi-substituted sulfoxide (2h) was also produced efficiently. Aliphatic and benzyl groups could be tolerated, and target products 2i and 2j were furnished in 91% and 86% yields, respectively. Notably, nitrogen-containing hetero-aromatics can be selectively oxidized to form the corresponding sulfoxide (2k) without *N*-oxide formation. Finally, alkyl sulfoxide (2l) was well tolerated under our reaction conditions, given that it is often sensitive to oxygenation conditions.¹³

The late-stage oxygenation of complex molecules to construct sulfoxides is often accompanied by the existence of various functional groups, and maintaining these functional groups is very important but remains challenging. By using our selective method, we expect that complex pharmaceuticals can be precisely modified at the last stage, in which sulfide is oxidized to sulfoxide while the side reactions on other functional groups are avoided. As a proof-of-concept, we focused on four important drug molecules with sulfoxide moieties (Fig. 5). Specifically, the corresponding sulfides were synthesized first by the reported procedures (for Experimental procedures see the

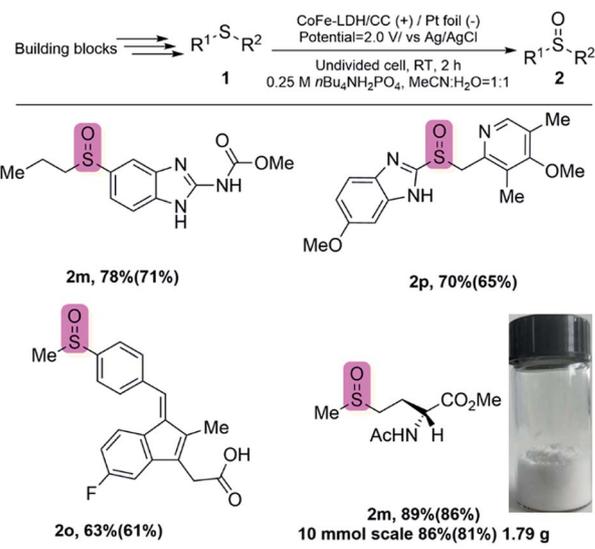


Fig. 5 Synthesis of existing pharmaceuticals: showcasing the applicability of this protocol by integrating with current procedures. Yields were determined by ¹H NMR using CH₂Br₂ as an internal standard. The isolated yields are given in parentheses. 2m, ricobendazole. 2n, omeprazole. 2o, sulindac. 2p, amino acid methionine. The sulfoxide moieties in the pharmaceuticals are highlighted in pink color.

ESI†), which was followed by electrochemical oxidation to the targeted sulfoxides by using CoFe-LDH/CC as the anode under the optimized conditions. The catalytic results show that, ricobendazole²⁰ (2m), an effective drug for a potential anticancer agent, was obtained by selective oxidation in 78% yield. Omeprazole²⁰ (2n), a well-known drug for treating gastroesophageal reflux, was also obtained in 70% yield. Sulindac²⁰ (2o), another anticancer reagent, was generated with 63% yield even in the presence of oxygen-sensitive sequential conjugated alkenes. Amino acid methionine (2p),²⁰ a biologically relevant compound, could be efficiently synthesized in good yield. Moreover, this electrochemical protocol could be used in the gram-scale synthesis of sulfoxides (2p) with 1.79 g and high yield of 86% was still obtained. Therefore, this protocol potentially enables access to electrochemical synthesis of valuable pharmaceutical related sulfoxides without external oxidants. In addition, the CoFe-LDH/CC electrocatalyst potentially overcomes disadvantages, such as catalyst reusability, which are often encountered by using homogeneous catalysts in conventional methods.

In order to understand the reaction mechanism, a set of control experiments were carried out (Fig. 6). When the standard reaction was performed in the presence of 2,2,6,6-tetramethyl-1-piperidinyloxy (TEMPO) or butylated hydroxytoluene (BHT) as the radical scavengers,²¹ the reaction was completely inhibited (Fig. 6a). According to a study on oxygenation of sulfides,^{12b,21} the sulfide radical (Ph₂S[•]) or persulfoxide radical (Ph₂SOO[•]) was generally present in the process. The EPR experiments were designed to detect radical intermediates by adding the radical trapping agent DMPO (5,5-dimethyl-1-pyrroline *N*-oxide). No obvious signals could be observed in



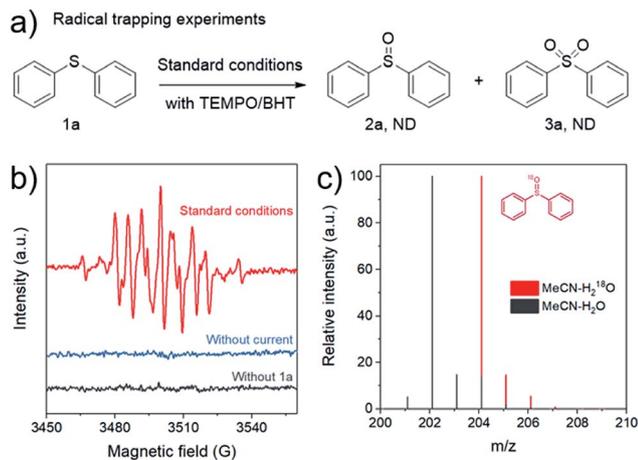


Fig. 6 (a) Radical trapping experiments, (b) EPR spectra of the DMPO-adduct under different conditions, (c) mass spectra of **2a** from isotope labelling experiments with H_2O or H_2^{18}O , respectively.

the absence of **1a** (Fig. 6b, black line). Under the standard conditions, a typical signal of the DMPO-radical adduct was determined by the EPR during reaction (Fig. 6b, red line). We speculated that it might be a sulfide radical ($\text{Ph}_2\text{S}^\bullet$) or persulfide radical ($\text{Ph}_2\text{SOO}^\bullet$) generated in the reaction with the DMPO adduct.²² These findings suggest that the electrooxidation reaction over CoFe-LDH/CC may involve a radical process. To confirm the source of the oxygen atom in the formed product, oxidation of **1a** was carried out in $\text{MeCN-H}_2^{18}\text{O}$ electrolyte, and the GC mass spectrum analysis shows that one ^{18}O atom was introduced into **2a** (Fig. 6c). This result suggests that water could likely supply oxygen for the sulfoxide oxidation under our reaction conditions.

To gain an insight into the active species in this catalytic system, the structural evolution of the CoFe-LDH/CC anode was in-depth investigated using XRD, TEM, SAED, X-ray photoelectron spectroscopy (XPS), and X-ray absorption fine structure spectroscopy (XAFS).²³ As shown in Fig. 7a, the XRD pattern of

the used CoFe-LDH/CC (1st recycled CoFe-LDH/CC) shows only two broad diffraction peaks at 24.5° and 44.0° , which are attributed to carbon cloth, indicating that CoFe-LDH undergoes dramatic reconstruction to form an amorphous structure under reaction conditions. Consistently, the SAED further confirmed the amorphous nature of the used CoFe-LDH (ESI, Fig. S3†).

We then investigated the structure of CoFe-LDH before and after reaction using XPS and XAFS techniques. As shown in Fig. 7b, the XPS signal of $\text{Co}2\text{p}_{3/2}$ in the starting material shows a typical satellite peak at 787.7 eV and a peak at 783.0 eV attributed to Co^{II} in LDHs. After the reaction, the intensity of the peak of Co^{2+} decreased with the appearance of a new peak at 780.1 eV which could be attributed to Co^{3+} , suggesting the oxidation of Co^{2+} to Co^{3+} in the electrooxidation process.²⁴ Concomitantly, the deprotonation of hydroxyl on CoFe-LDH occurred, giving rise to a peak at 529.1 eV in the $\text{O}1\text{s}$ XPS spectrum of the used CoFe-LDH that can be attributed to the typical M-O-M species in metal oxyhydroxides (Fig. 7c).^{24,25} Meanwhile, the $\text{Fe}2\text{p}$ XPS spectra reveal a stable Fe^{III} state in the fresh and the used CoFe-LDH (ESI, Fig. S4†). In addition, X-ray absorption near-edge structure (XANES) analysis in Fig. 7d shows that the Co K edge profile of the used CoFe-LDH/CC moved to a higher energy and next to that of the crystalline CoFe-oxyhydroxide reference (*c*-CoFeOOH, obtained from CoFe-LDH/CC by anodization in 1 M KOH, ESI, Fig. S5†), indicating that some Co^{II} ions in starting CoFe-LDH/CC were oxidized to Co^{III} during the reaction, which is consistent with the XPS data. In addition, the extended XAFS spectra reveal that the length of the first coordination shell C-O (2.03 Å) and the second coordination shell Co-Co(Fe) (3.13 Å) in the initial CoFe-LDH was decreased to 1.88 Å and 2.83 Å, respectively (Fig. 7e), which are identical to those of *c*-CoFeOOH, reflecting its metal oxyhydroxide nature. This structural transformation during electrooxidation was induced by oxidation of Co^{II} ions to Co^{III} .²⁶ Notably, the intensities of both Co-O and Co-Co(Fe) in the used CoFe-LDH are lower than those of fresh CoFe-LDH and *c*-CoFeOOH in the spectra of wavelet transformed EXAFS (ESI, Fig. S6†), owing to its amorphous structure. Moreover, the EXAFS analysis show that the as-formed oxyhydroxide structure was stable after four cycle reactions (Fig. 7f). We can conclude that the starting CoFe-LDH/CC material evolved into the corresponding amorphous oxyhydroxide during electrooxidation as the real active species. In addition, SCN^- , often adopted to poison metal sites,²⁷ was used as an indicator for active sites. As shown in Fig. S7,† a remarkable decrease of **2a** yield was observed by adding SCN^- , thus indicating that the metals in CoFe-LDH contribute to the catalytic activity. As an electrochemical reaction takes place at the interface between the catalyst and electrolyte, we evaluated the stability of CoFe-LDH for 7 h using chronoamperometry (CA) at 2.0 V versus the Ag/AgCl electrode (Fig. S8†). A current density of $\sim 11 \text{ mA cm}^{-2}$ was retained, indicating the stability of the catalyst over long reaction times.

On the basis of our results and the previous literature,^{6d,28} a reaction mechanism of this catalytic system is proposed in Fig. 8. In the neutral aqueous-organic electrolyte (MeCN as the organic phase), water molecules undergo Volmer and Heyrovsky

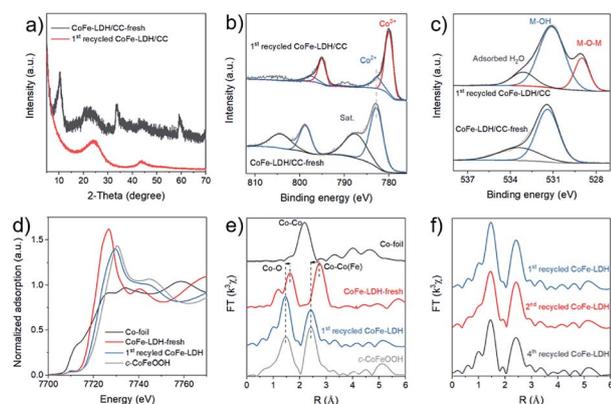


Fig. 7 (a) XRD patterns, (b) Co 2p XPS spectra, and (c) O1s XPS spectra of fresh and 1st recycled CoFe-LDH/CC. (d) Normalized Co K-edge XANES spectra and (e and f) Fourier-transform Co K-edge EXAFS spectra of reference samples, fresh and used CoFe-LDH/CC.



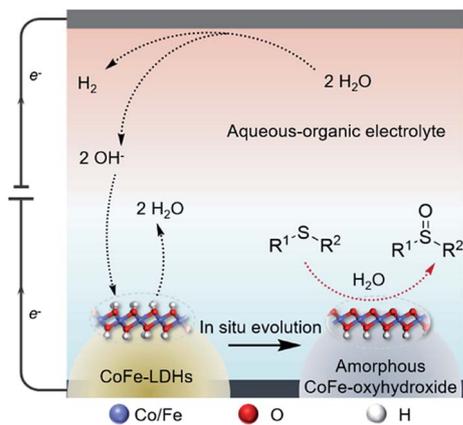


Fig. 8 Proposed reaction mechanism.

reactions to produce H_2 and hydroxide ions (OH^-) over the Pt cathode under work potential bias. Meanwhile, the starting CoFe-LDH was oxidized and evolved into amorphous CoFe-oxyhydroxide *via* the deprotonation process with the assistance of OH^- very fast in a few minutes (<5 minutes), and it may complete as soon as the reaction starts, as demonstrated by the anode color change and XRD and XPS results (Fig. S9†), ensuring the maintenance of its activity during the reaction time course. We synthesized reference materials ($Co(OH)_2/CC$ and $Fe(OH)_3/CC$) and compared their catalytic activity with that of CoFe-LDH/CC. As shown in Fig. S10a,† $Co(OH)_2/CC$ showed much higher activity than $Fe(OH)_3/CC$, suggesting that the $Co(OH)_2$ -derived structure is the main active phase. The CoFe-LDH/CC exhibits the highest performance among the three catalysts, indicating that a synergistic effect exists between Co and Fe in this reaction. The $Co2p_{3/2}$ XPS spectra of the used CoFe-LDH/CC, the derived CoFeOOH structure, exhibits a 0.43 eV shift to higher binding energy compared with that of the used $Co(OH)_2/CC$ (Fig. S10b†), indicating the modulation of the local electronic structure of Co cations in the presence of Fe species. Because of the modulated electronic structure, the redox behavior of Co cations could probably be manipulated by Fe incorporation, leading to the active phases in sulfide electrooxidation with higher activity, which is in agreement with recent reports.²⁹ This amorphous oxyhydroxide might act as the active phase for selective sulfide oxidation using water as the oxygen source.

Conclusions

In summary, we demonstrate an efficient electrochemical method for selective electrooxidation of sulfides with promoted HER over a CoFe-LDH/CC electrode. A variety of sulfide model compounds were selectively electrooxidized to sulfoxides in high yields (85–96%) at the anode under ambient conditions, showing high catalytic performance compared to the existing homo- and heterogeneous catalysts. Importantly, this method can be applied to the synthesis of four complex pharmaceuticals with sulfoxide moieties from their sulfide precursors in good yields, including ricobendazole (78%), omeprazole (70%),

sulindac (63%) and amino acid methionine (89%). Moreover, CoFe-LDH/CC was used in the gram-scale synthesis of diphenyl sulfoxide with 83% yield (up to 1.63 g) and amino acid methionine with 86% yield (up to 1.79 g). In addition, the CoFe-LDH/CC electrocatalyst could be reused, maintaining the yield for more than ten cycles. Mechanistic studies indicate that the reaction pathway proceeds through a radical process, and the *in situ* formed CoFe-oxyhydroxide may serve as the active species for the sulfide oxidation. The efficiency of this reaction system may pave the way for the electrochemical synthesis of valuable organic molecules by using heterogeneous catalysts without external oxidants under ambient conditions.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The work was financially supported by the National Natural Science Foundation of China (21978147, 21935001), and the China Postdoctoral Science Foundation (2019M660421). We greatly thank Profs. Duan Xue and Yadong Li for long-term support.

Notes and references

- (a) H. Jin, C. Guo, X. Liu, J. Liu, A. Vasileff, Y. Jiao, Y. Zheng and S. Z. Qiao, *Chem. Rev.*, 2018, **118**, 6337–6408; (b) S. Chu and A. Majumdar, *Nature*, 2012, **488**, 294–303; (c) Y. Yang, H. L. Fei, G. D. Ruan and J. M. Tour, *Adv. Mater.*, 2015, **27**, 3175–3180; (d) Y. Yu, Y. Shi and B. Zhang, *Acc. Chem. Res.*, 2018, **51**, 1711–1721; (e) J. X. Feng, S. Y. Tong, Y. X. Tong and G. R. Li, *J. Am. Chem. Soc.*, 2018, **140**, 5118–5126; (f) X. H. Gao, H. X. Zhang, Q. G. Li, X. G. Yu, Z. L. Hong, X. W. Zhang, C. D. Liang and Z. Lin, *Angew. Chem., Int. Ed.*, 2016, **55**, 6290–6294; (g) C. C. L. McCrory, S. Jung, I. M. Ferrer, S. M. Chatman, J. C. Peters and T. F. Jaramillo, *J. Am. Chem. Soc.*, 2015, **137**, 4347–4357; (h) Y. Wang, D. Yan, S. E. Hankari, Y. Zou and S. Wang, *Adv. Sci.*, 2018, **5**, 1800064.
- (a) Y. Jiao, Y. Zheng, M. Jaroniec and S. Z. Qiao, *Chem. Soc. Rev.*, 2015, **44**, 2060–2086; (b) W. J. Zhou, J. Jia, J. Lu, L. J. Yang, D. M. Hou, G. Q. Li and S. W. Chen, *Nano Energy*, 2016, **28**, 29–43; (c) J. Luo, J. H. Im, M. T. Mayer, M. Schreier, M. K. Nazeeruddin, N. G. Park, S. D. Tilley, H. J. Fan and M. Grätzel, *Science*, 2014, **345**, 1593–1596; (d) M. Gong, W. Zhou, M. C. Tsai, J. Zhou, M. Guan, M. C. Lin, B. Zhang, Y. Hu, D. Y. Wang and J. Yang, *Nat. Commun.*, 2014, **5**, 4695–4700; (e) H. Yin, S. Zhao, K. Zhao, A. Muqsit, H. Tang, L. Chang, H. Zhao, Y. Gao and Z. Tang, *Nat. Commun.*, 2015, **6**, 6430–6437; (f) M. W. Kanan and D. G. Nocera, *Nature*, 2008, **321**, 1072–1075; (g) P. Zhou, Y. Wang, C. Xie, C. Chen, H. Liu, R. Chen, J. Huo and S. Wang, *Chem. Commun.*, 2013, **53**, 11778–11781.



- 3 (a) F. Song, L. Bai, A. Moysiadou, S. Lee, C. Hu, L. Liardet and X. Hu, *J. Am. Chem. Soc.*, 2018, **140**, 7748–7759; (b) D. S. He, D. P. He, J. Wang, Y. Lin, P. Q. Yin, X. Hong, Y. Wu and Y. D. Li, *J. Am. Chem. Soc.*, 2016, **138**, 1494–1497; (c) H. Sun, X. Xu, Z. Yan, X. Chen, F. Cheng, P. S. Weiss and J. Chen, *Chem. Mater.*, 2017, **29**, 8539–8547; (d) P. Zhang, X. Sheng, X. Chen, Z. Fang, J. Jiang, M. Wang, F. Li, L. Fan, Y. Ren, B. Zhang, B. Timmer, M. Ahlquist and L. Sun, *Angew. Chem., Int. Ed.*, 2019, **58**, 9155–9159; (e) J. Hou, Y. Wu, B. Zhang, S. Cao, Z. Li and L. Sun, *Adv. Funct. Mater.*, 2019, **29**, 1908367.
- 4 (a) T. Binninger, R. Mohamed, A. Patru, K. Waltar, E. Gericke, X. Tuaeov, E. Fabbri, P. Levecque, A. Hoell and T. J. Schmidt, *Chem. Mater.*, 2017, **29**, 2831–2843; (b) G. Y. Chen, K. A. Kuttiyiel, D. Su, M. Li, C. H. Wang, D. Buceta, C. Y. Du, Y. Z. Gao, G. P. Yin, K. Sasaki, M. B. Vukmirovic and R. R. Adzic, *Chem. Mater.*, 2016, **28**, 5274–5281; (c) K. Xu, P. Z. Chen, X. L. Li, Y. Tong, H. Ding, X. J. Wu, W. S. Chu, Z. M. Peng, C. Z. Wu and Y. Xie, *J. Am. Chem. Soc.*, 2015, **137**, 4119–4125; (d) P. W. Cai, J. H. Huang, J. X. Chen and Z. H. Wen, *Angew. Chem., Int. Ed.*, 2017, **56**, 4858–4861; (e) X. Zou, Y. Liu, G. D. Li, Y. Y. Wu, D. P. Liu, W. Li, H. W. Li, D. J. Wang, Y. Zhang and X. X. Zou, *Adv. Mater.*, 2017, **29**, 1700404; (f) T. T. Liu, D. N. Liu, F. L. Qu, D. X. Wang, L. Zhang, R. X. Ge, S. Hao, Y. J. Ma, G. Du and A. M. Asiri, *Adv. Mater.*, 2017, **7**, 1700020; (g) T. T. Liu, X. Ma, D. N. Liu, S. Hao, G. Du, Y. J. Ma, A. M. Asiri, X. P. Sun and L. Chen, *ACS Catal.*, 2017, **7**, 98–102.
- 5 (a) B. You, X. Liu, N. Jiang and Y. Sun, *J. Am. Chem. Soc.*, 2016, **138**, 13639–13646; (b) H. G. Cha and K.-S. Choi, *Nat. Chem.*, 2015, **7**, 328–333; (c) Y. Zhou, Y. Gao, X. Zhong, W. Jiang, Y. Liang, P. Niu, M. Li, G. Zhuang, X. Li and J. Wang, *Adv. Funct. Mater.*, 2019, **29**, 1807651; (d) S. Li, X. Sun, Z. Yao, X. Zhong, Y. Cao, Y. Liang, Z. Wei, S. Deng, G. Zhuang, X. Li and J. Wang, *Adv. Funct. Mater.*, 2019, **42**, 1904780; (e) Q. Zhang, Y. Cao, Y. Yan, B. Yuan, H. Zheng, Y. Gu, X. Zhong and J. Wang, *J. Mater. Chem. A*, 2020, **8**, 2336–2342.
- 6 (a) X. Jin, M. Zhao, C. Zeng, W. Yan, Z. Song, P. S. Thapa, B. Subramaniam and R. V. Chaudhari, *ACS Catal.*, 2016, **6**, 4576–4583; (b) J. S. Luterbacher, J. M. Rand, D. M. Alonso, J. Han, J. T. Youngquist, C. T. Maravelias, B. F. Pflieger and J. A. Dumesic, *Science*, 2014, **343**, 277–280; (c) Y. Huang, X. Chong, C. Liu, Y. Liang and B. Zhang, *Angew. Chem. Int., Ed.*, 2018, **57**, 13163–13166; (d) C. Huang, Y. Huang, C. Liu, Y. Yu and B. Zhang, *Angew. Chem. Int., Ed.*, 2019, **58**, 12014–12017; (e) A. Corma, S. Iborra and A. Velty, *Chem. Rev.*, 2007, **107**, 2411–2502; (f) Y. Tong, P. Chen, M. Zhang, T. Zhou, L. Zhang, W. Chu, C. Wu and Y. Xie, *ACS Catal.*, 2017, **8**, 1–7; (g) J. Zhang, H. Wang, Y. Tian, Y. Yan, Q. Xue, T. He, H. Liu, C. Wang, Y. Chen and B. Y. Xia, *Angew. Chem., Int. Ed.*, 2018, **57**, 7649–7653.
- 7 (a) S. Otocka, M. Kwiatkowska, L. Madalińska and P. Kielbasiński, *Chem. Rev.*, 2017, **117**, 4147–4181; (b) I. Fernandez and N. Khiar, *Chem. Rev.*, 2003, **103**, 3651–3706; (c) B. M. Trost and M. Rao, *Angew. Chem., Int. Ed.*, 2015, **54**, 5026–5043; (d) J. A. Ellman, T. D. Owens and T. P. Tang, *Acc. Chem. Res.*, 2002, **35**, 984–995; (e) D. Kaldre, I. Klose and N. Maulide, *Science*, 2018, **361**, 664–667; (f) R. R. Merchant, J. T. Edwards, T. Qin, M. M. Kruszyk, C. Bi, G. Che, D.-H. Bao, W. Qiao, L. Sun, M. R. Collins, O. O. Fadeyi, G. M. Gallego, J. J. Mousseau, P. Nuhant and P. S. Baran, *Science*, 2018, **360**, 75–80; (g) H. Yehuda, S. Khatib, I. Sussan, R. Musa, J. Vaya and S. Tamir, *Biofactors*, 2009, **35**, 295–305.
- 8 (a) Y. Li, M. Wang and X. Jiang, *ACS Catal.*, 2017, **7**, 7587–7592; (b) J. Legros, J. R. Dehli and C. Bolm, *Adv. Synth. Catal.*, 2005, **347**, 19–31.
- 9 (a) R. Trivedi and P. Lalitha, *Synth. Commun.*, 2006, **36**, 3777–3782; (b) M. Matteucci, G. Bhalay and M. Bradley, *Org. Lett.*, 2003, **5**, 235–237; (c) F. Hosseinpour and H. Golchoubian, *Tetrahedron Lett.*, 2006, **47**, 5195–5197; (d) M. D. A. Maria, P. Mastroilli and C. F. Nobile, *J. Mol. Catal. A: Chem.*, 1996, **108**, 57–62; (e) J. Legros and C. Bolm, *Angew. Chem., Int. Ed.*, 2004, **43**, 4225–4228; (f) J. Legros and C. Bolm, *Angew. Chem., Int. Ed.*, 2003, **42**, 5487–5489; (g) H. Egami and T. Katsuki, *J. Am. Chem. Soc.*, 2007, **129**, 8940–8941; (h) B. Li, A.-H. Liu, L.-N. He, Z.-Z. Yang, J. Gao and K.-H. Chen, *Green Chem.*, 2012, **14**, 130–135.
- 10 (a) B. Yu, A.-H. Liu, L.-N. He, B. Li, Z.-F. Diao and Y.-N. Li, *Green Chem.*, 2012, **14**, 957–962; (b) E. S. Priestley, D. L. Cheney, I. DeLucca, A. Wei, J. M. Luetgen, A. R. Rendina, P. C. Wong and R. R. Wexler, *J. Med. Chem.*, 2015, **58**, 6225–6236; (c) D. P. de Sousa, C. Wegeberg, M. S. Vad, S. Mørup, C. Frandsen, W. A. Donald and C. J. McKenzie, *Chem.–Eur. J.*, 2016, **22**, 3810–3820; (d) M. V. Gómez, R. Caballero, E. Vázquez, A. Moreno, A. de la Hoz and A. Díaz-Ortiz, *Green Chem.*, 2007, **9**, 331–336; (e) Y. Liu, A. J. Howarth, J. T. Hupp and O. K. Farha, *Angew. Chem., Int. Ed.*, 2015, **54**, 9001–9005.
- 11 B. Yu, A.-H. Liu, L.-N. He, B. Li, Z.-F. Diao and Y.-N. Li, *Green Chem.*, 2012, **14**, 957–962.
- 12 (a) J.-P. Cao, Y.-S. Xue, N.-F. Li, J.-J. Gong, R.-K. Kang and Y. Xu, *J. Am. Chem. Soc.*, 2019, **141**, 19487–19497; (b) Y. Li, S. A.-e.-A. Rizvi, D. Hu, D. Sun, A. Gao, Y. Zhou, J. Li and X. Jiang, *Angew. Chem., Int. Ed.*, 2019, **58**, 13499–13506.
- 13 G. Laudadio, N. J. W. Straathof, M. D. Lanting, B. Knoops, V. Hessel and T. Noël, *Green Chem.*, 2017, **19**, 4061–4066.
- 14 (a) L. Zhou, M. F. Shao, C. Zhang, J. W. Zhao, S. He, D. M. Rao, M. Wei, D. G. Evans and X. Duan, *Adv. Mater.*, 2017, **29**, 1604080; (b) X. Long, G. X. Li, Z. L. Wang, H. Y. Zhu, T. Zhang, S. Xiao, W. Y. Guo and S. H. Yang, *J. Am. Chem. Soc.*, 2015, **137**, 11900–11903; (c) C. Zhang, M. F. Shao, L. Zhou, Z. H. Li, K. M. Xiao and M. Wei, *ACS Appl. Mater. Interfaces*, 2016, **8**, 33697–33703; (d) Q. Wang and D. O'Hare, *Chem. Rev.*, 2012, **112**, 4124–4155; (e) M. Shao, F. Ning, J. Zhao, M. Wei, D. G. Evans and X. Duan, *J. Am. Chem. Soc.*, 2012, **134**, 1071–1077; (f) Y. Zhao, B. Li, Q. Wang, W. Gao, C. J. Wang, M. Wei, D. G. Evans, X. Duan and D. O'Hare, *Chem. Sci.*, 2014, **5**, 951–958; (g) L. Yin, S. Li, X. Liu and T. Yan, *Sci. China Mater.*, 2019, **62**, 1537–1555.
- 15 (a) F. Song and X. Hu, *Nat. Commun.*, 2014, **5**, 4477–4485; (b) X. Zou, A. Goswami and T. Asefa, *J. Am. Chem. Soc.*, 2013,



- 135, 17242–17245; (c) B. Hunter, J. Blakemore, M. Deimund, H. Gray and J. Winkler, *J. Am. Chem. Soc.*, 2014, **136**, 13118–13121; (d) F. Li, J. Du, X. Li, J. Shen, Y. Wang, Y. Zhu and L. Sun, *Adv. Energy Mater.*, 2018, **8**, 1702598; (e) X. Wu, Y. Zhao, T. Xing, P. Zhang, F. Li, H. Lee, F. Li and L. Sun, *ChemSusChem*, 2018, **11**, 1761–1767; (f) L. Huang, Z. He, J. Guo, S. Pei, H. Shao and J. Wang, *Nano Res.*, 2020, **13**, 246–254; (g) H. Yang, H. Wang, Y. Zhang and Q. Wang, *Sci. China Mater.*, 2019, **62**, 681–689.
- 16 (a) H. Zhang, H. Li, B. Akram and X. Wang, *Nano Res.*, 2019, **12**, 1327–1331; (b) B. You, N. Jiang, X. Liu and Y. Sun, *Angew. Chem., Int. Ed.*, 2016, **55**, 9913–9917; (c) B. You, X. Liu, N. Jiang and Y. Sun, *J. Am. Chem. Soc.*, 2016, **138**, 13639–13646.
- 17 J. Zhao, Z. Lu, M. Shao, D. Yan, M. Wei, D. G. Evans and X. Duan, *RSC Adv.*, 2013, **3**, 1045–1049.
- 18 R. Yang, Y. Zhou, Y. Xing, D. Li, D. Jiang, M. Chen and S. Yuan, *ACS Appl. Energy Mater.*, 2018, **1**, 1200–1209.
- 19 C. Kingston, M. D. Palkowitz, Y. Takahira, J. C. Vantourout, B. K. Peters, Y. Kawamata and P. S. Baran, *Acc. Chem. Res.*, 2020, **53**, 72–83.
- 20 For detailed references. See the ESI. Section 2.†
- 21 Z. Cheng, P. Sun, A. Tang, W. Jin and C. Liu, *Org. Lett.*, 2019, **22**, 8925–8929.
- 22 (a) R. I. Sayler, B. M. Hunter, W. Fu, H. B. Gray and R. D. Britt, *J. Am. Chem. Soc.*, 2020, **142**, 1838–1845; (b) Z. Guan, S. Zhu, S. Wang, H. Wang, X. Zhong, F. Bu, H. Cong and A. Lei, *Angew. Chem., Int. Ed.*, DOI: 10.1002/anie.202011329.
- 23 (a) X. Lu, L. Gu, J. Wang, J. Wu, P.-Q. Liao and G.-R. Li, *Adv. Mater.*, 2017, **29**, 1604437; (b) J. A. Koza, C. M. Hull, Y.-C. Liu and J. A. Switzer, *Chem. Mater.*, 2013, **25**, 1922–1926; (c) M. Ludvigsson, J. Lindgren and J. Tegenfeldt, *J. Mater. Chem.*, 2001, **11**, 1269–1276; (d) T. Pauporte, L. Mendoza, M. Cassir, M. C. Bernard and J. Chivot, *J. Electrochem. Soc.*, 2005, **152**, 49–53.
- 24 (a) Z.-F. Huang, J. Song, Y. Du, S. Xi, S. Dou, J. M. V. Nsanzimana, C. Wang, Z. J. Xu and X. Wang, *Nat. Energy*, 2019, **4**, 328–338; (b) Z. Li, H. Duan, M. Shao, J. Li, D. O'Hare, M. Wei and Z. L. Wang, *Chem*, 2018, **4**, 1–12.
- 25 (a) Z. Li, K. Liu, K. Fan, Y. Yang, M. Shao, M. Wei and X. Duan, *Angew. Chem., Int. Ed.*, 2019, **58**, 3962–3965; (b) F. Dionigi, Z. Zeng, I. Sinev, *et al.*, *Nat. Commun.*, 2020, **11**, 2522.
- 26 (a) J. Huang, J. Chen, T. Yao, J. He, S. Jiang, Z. Sun, Q. Liu, W. Cheng, F. Hu, Y. Jiang, Z. Pan and S. Wei, *Angew. Chem., Int. Ed.*, 2015, **54**, 8722–8727; (b) A. Bergmann, T. E. Jones, E. Martinez Moreno, D. Teschner, P. Chernev, M. Gliech, T. Reier, H. Dau and P. Strasser, *Nat. Catal.*, 2018, **1**, 711–719.
- 27 (a) E. Zhang, T. Wang, K. Yu, J. Liu, W. Chen, A. Li, H. Rong, R. Lin, S. Ji, X. Zheng, Y. Wang, L. Zheng, C. Chen, D. Wang, J. Zhang and Y. Li, *J. Am. Chem. Soc.*, 2019, **141**, 16569–16573; (b) X. Li, W. Bi, M. Chen, Y. Sun, H. Ju, W. Yan, J. Zhu, X. Wu, W. Chu, C. Wu and Y. Xie, *J. Am. Chem. Soc.*, 2017, **139**, 14889–14892.
- 28 C. Huang, Y. Huang, C. Liu, Y. Yu and B. Zhang, *Angew. Chem., Int. Ed.*, 2019, **58**, 12014–12017.
- 29 X. Han, C. Yu, S. Zhou, C. Zhao, H. Huang, J. Yang, Z. Liu, C. Zhao and J. Qiu, *Adv. Energy Mater.*, 2017, **7**, 1602148.

