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Highly efficient electrochemical recognition and quantification of amine enantiomers based on a guest-free homochiral MOF†

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We facilely synthesized a novel guest-free homochiral metal–organic framework, $(\text{Cu}_4\text{L}_4)_n$ [$\text{H}_2\text{L} = N$ -(2-hydroxybenzyl)-L-leucine] in space group *P1*. The $(\text{Cu}_4\text{L}_4)_n$ nanocrystals exhibit high electrochemical activity for rapidly discriminating chiral α -methylbenzylamine enantiomers and quantitatively determining the enantiomeric excess in the chiral amine mixture.

Many important molecules in the modern pharmaceutical and agrochemical industries, are chiral—that is, they are not superimposable on their mirror image, the pair of asymmetric molecules are known as enantiomers. The recognition and quantification of enantiomers is a major challenge particularly owing to their identical physical and chemical properties in an achiral environment.¹ In many cases, one of the enantiomers exhibits the desired responses while the other is inactive or even toxic.^{2–4} The present technology depends largely upon nuclear magnetic resonance (NMR), gas chromatography (GC) and high performance liquid chromatography (HPLC) based on homochiral stationary phases (CSPs), which is of particular importance for laboratories and industries in the discovery and development of enantiomeric substances and quality control of corresponding products.^{5–7} However, these methods require high concentrations of analytes, sophisticated operation, relatively expensive instrumentation, and GC and HPLC are also typically time-consuming.^{8,9} Molecularly imprinted polymers (MIPs) have more than 80 years of history,^{10,11} considerable advances have been made on the fundamental study.^{12,13} Although the approach behaves special desired selectivity because the MIPs create three-dimensional cross-linked polymers with tailor-made memory of the shape, size and functional groups for a template or target molecule,^{14,15} this sometimes also suffers from some problems, such as incomplete template removal, poor mass transfer, low binding capacity and slow binding kinetics, which restricts its applications in various aspects.¹⁶ In contrast, the electrochemical sensors can provide highly selective, low cost, fast speed, real time and on-line

operation. Unlike chromatographic instruments, the technology can be easily adapted for detecting a wide range and low concentrations of analytes, while remaining inexpensive.¹⁷ However, there are only a handful of progress at the present chiral sensing using this approach.

The key of the electrochemical chiral sensing is the material used to prepare the sensor's electroactive surface. The emerging chiral metal–organic framework (CMOFs) are the intriguing class of crystalline materials formed usually by the self-assembly of metal ions and chiral polydentate ligands. These materials are highly promising for electrochemical chiral sensors owing to ultrahigh surface area, precise network structures, fine-tuned chiral channels and pores, regularly ordered functionalities and host–guest interactions involved.^{18,19} At present, however, only a few achiral MOFs have been attempted as electrochemical sensors for detecting a few achiral analytes.^{20–22} Such as, Gassensmith *et al.* demonstrated that γ -cyclodextrin-derived CDMOF-2 is capable of measuring CO_2 concentrations quantitatively in the presence of ambient oxygen.²³ Wu *et al.* developed a water-stable Cu-MOF, $\{[\text{Cu}_2(\text{HL})_2(\mu_2\text{-OH})_2(\text{H}_2\text{O})_5] \cdot \text{H}_2\text{O}\}_n$ ($\text{H}_2\text{L} = 2,5$ -dicarboxylic acid-3,4-ethylene dioxathiophene) used for the simultaneous detection of ascorbic acid and L-tryptophan.²⁰ In addition, there are few studies to incorporate the functionality of achiral MOFs and its derivatives into enzyme biosensor applications.^{24–26} To the best of our knowledge, until now no reports have been found regarding CMOFs for selective chiral detection. On the other hand, aiming at practical applications, the CMOFs must retain their structural integrity in both cases, the aqueous environment and the removal of the guest molecules trapped in the channels for creating empty pores and accessible metal sites.²⁷ However, most CMOFs of current reports are susceptible to deterioration when exposed to water or even humid air, and during the conventional guest removal procedure of heating in vacuum.²⁸ It is particularly challenging to render networks with

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chiral topologies robust to guest loss.²⁹ The synthesis of guest-free CMOFs with highly water and air stability is also an extremely good choice.

Herein, we present the synthesis, structure of a novel guest-free CMOFs, that is, $(\text{Cu}_4\text{L}_4)_n$, constructed from (Cu_4L_4) clusters using an enantiopure reduced Schiff base, N -(2-hydroxybenzyl)- L -leucine] [L -H₂L) as a starting material. The CMOF exhibits high stability towards air and water. Remarkably, an electrochemical chiral sensor was easily obtained by a glass carbon electrode directly covered with $(\text{Cu}_4\text{L}_4)_n$ nanocrystals without desolvation and further post-modification. The sensor behaved high electroconducting and distinctive oxidation signals for each enantiomer of $R(+)$ - and $S(-)$ - α -methylbenzylamine (MBA). Not only is the sensor able to discriminate enantioselectively, but it also allows for the rapid quantitative determination of enantiomeric excess in the chiral amine mixture.

Single crystals of $(\text{Cu}_4\text{L}_4)_n$ were readily obtained by mixing $\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$ and the ligand L -H₂L (N -(2-hydroxybenzyl)- L -leucine), lithium hydroxide aqueous solution, acetonitrile and ethanol for 2 days at room temperature. To rapidly synthesize $(\text{Cu}_4\text{L}_4)_n$ nanocrystals, we also developed a 2 min, room-temperature method, which was achieved by adding $\text{Cu}(\text{OAc})_2$ aqueous solution into the mixture of ligand L -H₂L, lithium hydroxide aqueous solution, DMF and DMSO. The flaky nanocrystals were obtained in a high yield (73%) without the need for

any activation procedures. SEM images in Fig. 1a showed their micro-morphologies with a size of $\sim 500 \times 100 \times 90$ nm.

Single-crystal X-ray structural analysis revealed that $(\text{Cu}_4\text{L}_4)_n$ crystallizes in the chiral space group $P1$ and has guest-free chiral open channels (Fig. 1b). The asymmetric cluster of the framework consists of four penta-coordinated $\text{Cu}(\text{II})$ centers and four chiral $L(\text{II})$ ligands. Each $\text{Cu}(\text{II})$ center exhibits a square-pyramidal coordination geometry bridged by two phenolate O atoms (O1 and O4), two carboxylato O atoms (O3 and O8) and one secondary amine N atom (N1). Each $L(\text{II})$ ligand maintains a quadridentate bridging coordination mode. Each carboxylate group in $L(\text{II})$ ligands serves as a *syn-anti* bridge between $\text{Cu}(\text{II})$ ions to afford a 1D chain. Then the $\text{Cu}(\text{II})$ ions are further connected by the corresponding two L ligands through the phenolate O to generate a 2D sheet structure. The guest-free chiral channels are square-shaped, and phenyls on channels walls are perpendicularly with the coordination layers (Fig. 1c and d).

The ideal materials as the electrochemical chiral sensing should possess not only precise framework structures, but also homochirality, specific porous properties and high stability in heat, air and water. The solid-state circular dichroism (CD) spectra of $(\text{Cu}_4\text{L}_4)_n$ crystals and its enantiomer, $(\text{Cu}_4\text{L}_4)_n'$ exhibit strong opposite Cotton effect around 234, 292 and 387 nm, which confirmed the homochirality of $(\text{Cu}_4\text{L}_4)_n$ (Fig. 2a). The

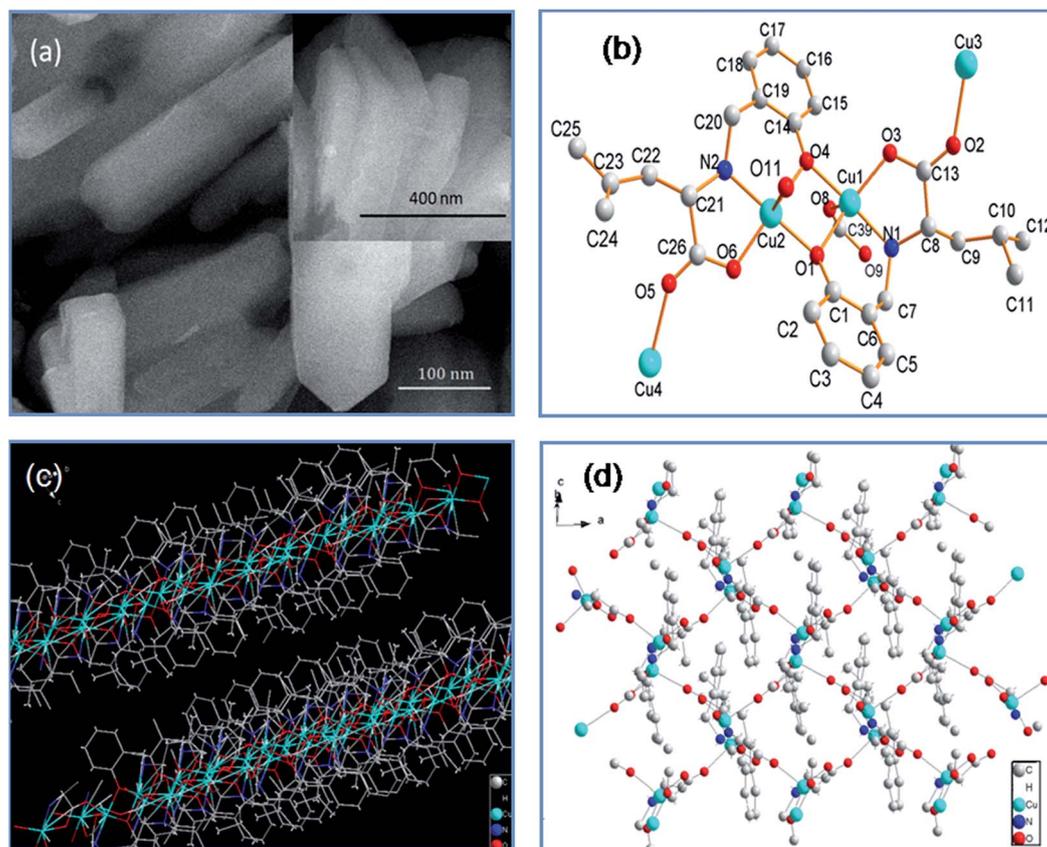


Fig. 1 SEM images of $(\text{Cu}_4\text{L}_4)_n$ crystals with different magnifications (a). Local coordination environments of the $\text{Cu}(\text{II})$ center and the chiral ligands in $(\text{Cu}_4\text{L}_4)_n$, hydrogen atoms are omitted for clarity (b). Side view of 2D sheet structure (c). Framework structure of $(\text{Cu}_4\text{L}_4)_n$ integrated, showing the microporous channels and the hydrophobic surface. Color codes: Cu red, N blue, C grey, H white (d).



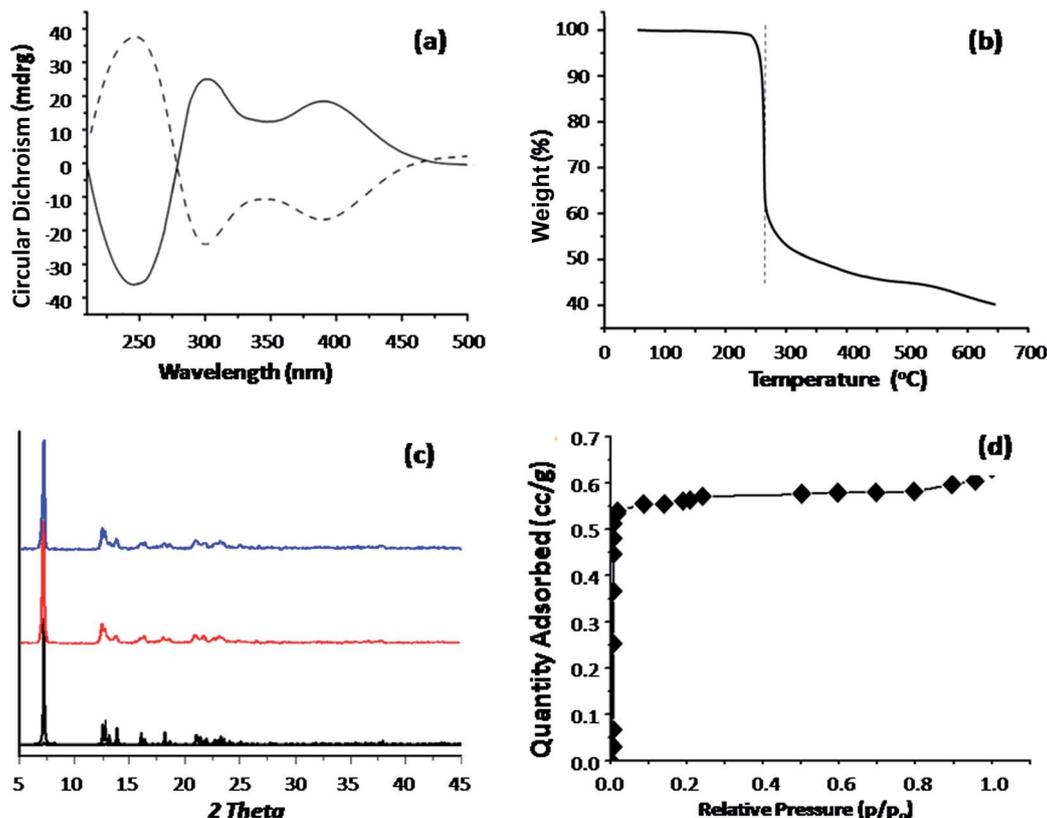


Fig. 2 $(\text{Cu}_4\text{L}_4)_n$ properties: (a) CD spectra of $(\text{Cu}_4\text{L}_4)_n$ (solid) in comparison with that of its enantiomer $(\text{Cu}_4\text{L}_4)_n'$ (dash); (b) TGA analysis; (c) PXRD patterns of simulated $(\text{Cu}_4\text{L}_4)_n$ from the single crystal data (black), as-synthesized product for 110 °C for 24 h (red) and after immersing into water for 2 weeks (blue); (d) nitrogen gas sorption isotherm at 77 K.

polymeric lattice of $(\text{Cu}_4\text{L}_4)_n$ is free-guest and can be retained upon heating at 280 °C, which is supported by the thermogravimetric (TGA) analysis (Fig. 2b). After being suspended in water for a week and exposed in air for one month, the PXRD patterns of these crystals were still in conformity with the simulated one from the single-crystal data (Fig. 2c), indicating an excellent water and air stability of this CMOF. The porosity of $(\text{Cu}_4\text{L}_4)_n$ was further probed by N_2 sorption, follow a typical type I isotherm corresponding to micropores, and its BET surface area is $56 \text{ m}^2 \text{ g}^{-1}$ (Fig. 2d).

Pure α -MBA enantiomers are key building blocks and intermediates in a number of pharmaceutical and agromedical compounds that possess a wide range of biological activities.³⁰ However, there is no report found concerning electrochemical detection $R(+)$ - and $S(-)$ - α -MBA enantiomers, which may be due to its poor electroactivity. To evaluate the ability of our CMOF discriminating α -MBA enantiomers, the electrochemical chiral sensor was assembled by coating $(\text{Cu}_4\text{L}_4)_n$ on the surfaces of GCE. In particular, present $(\text{Cu}_4\text{L}_4)_n$ is used without desolvation. In many instances, guest molecules filling in the pore of MOFs and solvent molecules coordinated at open-metal sites (OMS) of MOFs should be evacuated and decoordinated by an activation process prior to every use.³¹

The properties of the CMOF-functionalized electrodes were firstly tested by cyclic voltammetry (CV) in K_2HPO_4 - KH_2PO_4 buffer solution (0.1 mol L^{-1} PBS, pH 5). As shown in Fig. 3, the

bare GCE behaves low and overlapped currents for $R(+)$ - α -MBA and $S(-)$ - α -MBA. In contrast, the modified electrode, $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ causes a significantly increased oxidation current, and the degree of $R(+)$ - α -MBA is obviously larger than the case of $S(-)$ - α -MBA. Obvious difference is observed on anodic characteristic peaks for $R(+)$ - α -MBA at -34.8 mV and for $S(-)$ - α -MBA at 6.5 mV . No obvious reduction peak appears for the amine enantiomers, showing the irreversibility of their electrochemical reactions. These suggest that chiral $(\text{Cu}_4\text{L}_4)_n$ material has outstanding conductivity and different electrocatalytic

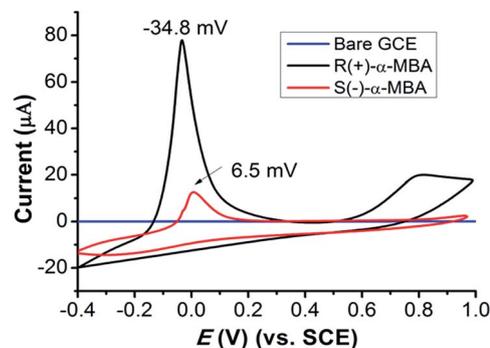


Fig. 3 CVs of chiral $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ electrode for the detection of 0.05 mmol L^{-1} $R(+)$ - and $S(-)$ - α -MBA in 0.1 mol L^{-1} PBS (vs. SCE) at pH 5, 50 mV s^{-1} of scan rate.



functions on the oxidation of $R(+)$ - and $S(-)$ - α -MBA. Based on the facts, the strategy for enantiodiscrimination of $R(+)$ - and $S(-)$ - α -MBA enantiomers with the CMOF material can be achieved.

Differential pulse voltammetry (DPV) is a powerful method in electrochemical analysis because of its higher sensitivity and better resolution than CV.³² Therefore, DPV was utilized to investigate the effect of pH value detected PBS solution containing the $R(+)$ - or $S(-)$ - α -MBA.

As displayed in Fig. 4, the electrochemical behavior of $R(+)$ - or $S(-)$ - α -MBA is strongly dependent on the pH values of the aqueous solutions. $R(+)$ - α -MBA at examined four pH values overall shows a pair of well-defined oxidation peaks in the potential range of -71 mV to $+735$ mV, and obviously higher than those of $S(-)$ - α -MBA. In comparison, a characteristic peak of $S(-)$ - α -MBA at pH 4 is splitting into two ones with increasing pH to 5, 6 and 7. In both peaks, these anodic peak potentials of $R(+)$ - and $S(-)$ - α -MBA at ~ 720 mV are not suitable for assaying the enantiomers owing to their weak peak currents. Therefore, we chosen pH 5 as further chiral measurements, based on the both peak shift and difference of the peak currents from $R(+)$ - α -MBA to $S(-)$ - α -MBA, although the peak shift from $R(+)$ - α -MBA to $S(-)$ - α -MBA is significantly larger for pH 6 than pH 5 and pH 7.

Under optimal pH 5, a series of α -MBA enantiomers with different concentrations were investigated using the $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ electrode. Fig. 5 shows the dependence of DPVs of α -MBA enantiomers on the concentrations. The two anodic peaks for $R(+)$ - α -MBA at -100.3 mV and for $S(-)$ - α -MBA at -84.2 mV are clearly to be distinguished, and shows successive enhancement on increasing the concentration of the enantiomers. And the oxidization degree of $R(+)$ - α -MBA is larger than that of $S(-)$ - α -MBA. The calibration curves are obtained in the range of 0.001 to 0.1 mmol L^{-1} with a detection limit of 1.3 $\mu\text{mol L}^{-1}$ ($S/N = 3$), and the linear regression equations of $R(+)$ - and $S(-)$ - α -MBA can be expressed as Ip^R (μA) = $882.1C + 19.6$ ($R^2 = 0.9967$) and Ip^S

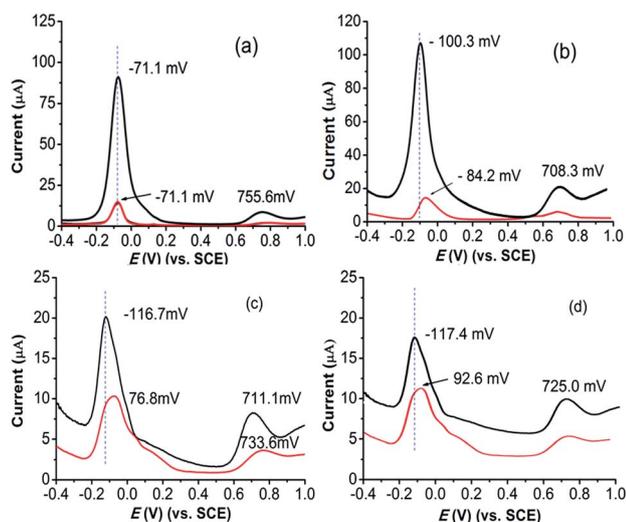


Fig. 4 DPV responses of chiral $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ electrode to different pH values in the presence of 0.1 mmol L^{-1} $R(+)$ - α -MBA (black) and $S(-)$ - α -MBA (red): (a) pH 4, (b) pH 5, (c) pH 6 and (d) pH 7. Other conditions as in Fig. 3.

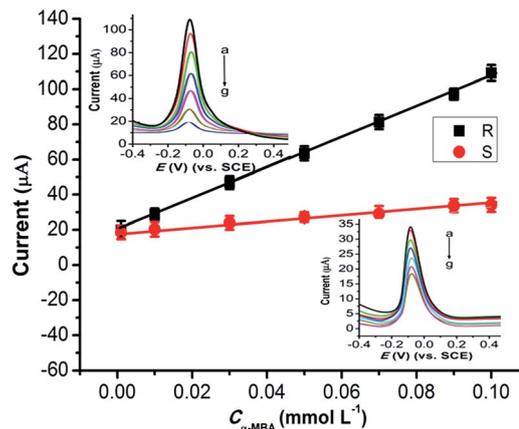


Fig. 5 DPV responses of chiral $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ electrode to different concentrations of $R(+)$ / $S(-)$ - α -MBA at pH 5: 0.01 , 0.03 , 0.05 , 0.07 , 0.09 , 0.10 mmol L^{-1} . Inset: plot of current intensity vs. logarithmic value of $R(+)$ / $S(-)$ - α -MBA concentration. Detection in 0.1 mol L^{-1} PBS (vs. SCE) at pH 5, 50 mV s^{-1} of scan rate.

(μA) = $159.5C + 19.8$ ($R^2 = 0.9921$), respectively. The results suggest the successful detection of $R(+)$ - and $S(-)$ - α -MBA enantiomers by $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ electrode in this concentration range. This attempt from only $(\text{Cu}_4\text{L}_4)_n$ nanocrystals of inexpensive and easy synthesis is a remarkable and potentially valuable feature of this system. By contrast, the most other materials for the enantiomeric recognition require multistep syntheses of the ligand and/or the complex.

The fast and convenient determination of the enantiomeric excess in development of single enantiomer pharmaceuticals and synthesis of chiral compounds has important application because a large number of samples must be screened.⁶ To characterize the ability of the $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ predicting EE value of α -MBA enantiomers in the mixture, here, the DPV analysis was performed for a series of solutions with the total concentration of $R(+)$ - and $S(-)$ - α -MBA being kept as 0.025 mM . As shown in Fig. 6, the oxidation peaks of $R(+)$ - α -MBA at ~ 100 mV and $S(-)$ - α -MBA at ~ 84 mV combine into a single one, which gradually shifts in positive direction as $S(-)$ - α -MBA concentration increased. More importantly, the oxidation currents against $S(-)$ - α -MBA% in the mixture of α -MBA enantiomers exhibits a linear dependence as $S(-)$ - α -MBA% increased with the correlation coefficient of 0.9903 . The results indicate that the proposed chiral sensor can rapidly and quantitatively determine enantiomeric excess in the mixture of α -MBA enantiomers.

Enantiomers should be distinguished in the presence of chiral centers on the electrode surface.³³ The fact that only chiral $(\text{Cu}_4\text{L}_4)_n$ introduced on GCE surface recognizes α -MBA enantiomers, suggests the formation of efficient chiral micro-environment. The oxidation of $R(+)$ - and $S(-)$ - α -MBA should begin their exchange electrons with electrode by passing through the CMOF layer on the electrode surface. As shown as in Fig. 5, the negative shift of oxidation potential and the significantly increasing of peak current for $R(+)$ - α -MBA than that of $S(-)$ - α -MBA behave that $R(+)$ - α -MBA is easier to lose its electron on the $(\text{Cu}_4\text{L}_4)_n/\text{GCE}$ surface and faster oxidation. The



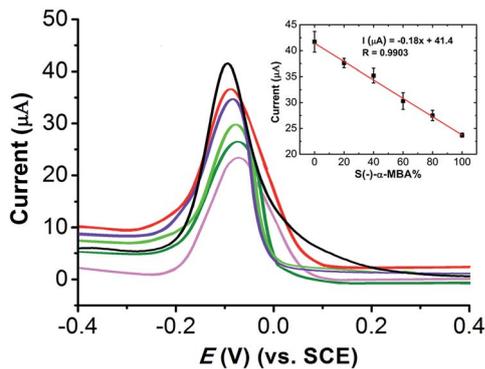


Fig. 6 The DPV for $S(-)$ - α -MBA% in the mixture of α -MBA enantiomers. Inset: linear relationship between $S(-)$ - α -MBA% and the oxidation currents. Detection in 0.1 mol L^{-1} PBS (vs. SCE) at pH 5, 50 mV s^{-1} of scan rate. Error bars represent standard deviation for three independent measurements.

present success must result from the interaction between the chiral host, $(\text{Cu}_4\text{L}_4)_n$ and α -MBA enantiomers favouring enantioselectivity during the adsorption process. The interactions can be explained by three-point interaction model between the host and analytes,³⁴ and at least one of these interactions must be stereoselective.¹ As described above, the structure of $(\text{Cu}_4\text{L}_4)_n$ crystal, possesses the guest-free chiral channels and rich-phenyls at wall perpendicularly with the coordination layers. A remarkable increase of $R(+)$ - α -MBA in the electrochemical response indicates that the more matching of the stereoscopic configuration between conformation of $R(+)$ - α -MBA molecule and chiral $(\text{Cu}_4\text{L}_4)_n$ nanocrystals than case of $S(-)$ - α -MBA, which should be the major interaction force in the recognition. Besides this, the weak interactions like π - π stacking, hydrogen bonding, hydrophobic interaction and dipole-dipole interactions may also play some role.

In summary, we have successfully synthesized a new 2D homochiral $(\text{Cu}_4\text{L}_4)_n$ with guest-free chiral channels. The chiral electrochemical sensor assembled with $(\text{Cu}_4\text{L}_4)_n$ nanocrystals shows excellent performance for selective oxidation recognition $R(+)$ - and $S(-)$ - α -MBA. This is the first attempt to directly apply a CMOF to discriminate α -MBA enantiomers. The features of this new approach are simplicity, rapidity and sensitivity. The CMOF-based sensor provides a promising platform for studying the application of CMOFs in area of chiral electrochemistry.

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References

1 Y. W. Peng, T. F. Gong, K. Zhang, X. C. Lin, Y. Liu, J. W. Jiang and Y. Cui, *Nat. Commun.*, 2014, **5**, 4406.

- 2 C. Wattanakit, Y. B. Saint Côme, V. Lapeyre, P. A. Bopp, M. Heim, S. Yadnum, S. Nokbin, C. Wattanakit, J. Limtrakul and A. Kuhn, *Nat. Commun.*, 2014, **5**, 3325.
- 3 X. Kuang, Y. Ma, H. Su, J. Zhang, Y. B. Dong and B. Tang, *Anal. Chem.*, 2014, **86**, 1277.
- 4 C. Liu, B. Wang, J. Diao and Z. Zhou, *RSC Adv.*, 2016, **6**, 59842.
- 5 M. Lämmerhofer, *J. Chromatogr. A.*, 2010, **1217**, 814.
- 6 T. J. Wenzel and J. D. Wilcox, *Chirality*, 2003, **15**, 256.
- 7 G. Mirri, S. D. Bull, P. N. Horton, T. D. James, L. Male and J. H. R. Tucker, *J. Am. Chem. Soc.*, 2010, **132**, 8903.
- 8 E. G. Shcherbakova, T. Minami, V. Brega and T. D. James, *Angew. Chem.*, 2015, **54**, 7236.
- 9 J. Tian and A. Rustum, *RSC Adv.*, 2016, **6**, 83020.
- 10 S. Sheybani, T. Hosseiniifar, M. Abdouss and S. Mazinani, *RSC Adv.*, 2015, **5**, 98880.
- 11 S. Farzaneh, E. Asadi, M. Abdouss, A. Barghi-Lish, S. Azodi-Deilami, H. A. Khonakdar and M. i. Gharghabi, *RSC Adv.*, 2015, **5**, 9154.
- 12 G. Vlatakis, L. I. Andersson, R. Muller and K. Mosbach, *Nature*, 1993, **361**, 645.
- 13 M. O. Ramstro and R. J. Ansell, *Chirality*, 1998, **10**, 195.
- 14 L. X. Chen, S. F. Xu and J. H. Li, *Chem. Soc. Rev.*, 2011, **40**, 2922.
- 15 T. Chang, Y. Liu, X. Yan, S. Liu and H. Zheng, *RSC Adv.*, 2016, **6**, 66297.
- 16 L. X. Chen, S. F. Xu and J. H. Li, *J. Mater. Chem.*, 2012, **22**, 3360.
- 17 D. W. Kimmel, G. LeBlanc, M. E. Meschievitz and D. E. Cliffel, *Electrochemical sensors and biosensors, Anal. Chem.*, 2012, **84**, 685.
- 18 K. Tanaka, T. Muraoka, Y. Otubo, H. Takahashi and A. Ohnishi, *RSC Adv.*, 2016, **6**, 21293.
- 19 M. Yoon, R. Srirambalaji and K. Kim, *Chem. Rev.*, 2012, **112**, 1196.
- 20 X. Q. Wu, J. G. Ma, H. Li, D. M. Chen, W. Gu, G. M. Yang and P. Cheng, *Chem. Commun.*, 2015, **51**, 9161.
- 21 L. Yang, S. Kinoshita, T. Yamada, S. Kanda and H. Kitagawa, *Angew. Chem., Int. Ed.*, 2010, **49**, 5348.
- 22 H. Ji, S. Hwang, K. Kim, C. Kim and N. C. Jeong, *ACS Appl. Mater. Interfaces*, 2016, **8**, 32414.
- 23 J. J. Gassensmith, J. Y. Kim, J. M. Holcroft, O. K. Farha, J. F. Stoddart, J. T. Hupp and N. C. Jeong, *J. Am. Chem. Soc.*, 2014, **136**, 8277.
- 24 P. H. Ling, J. P. Lei and H. X. Ju, *Biosens. Bioelectron.*, 2015, **71**, 373.
- 25 W. J. Ma, Q. Jiang, P. Yu, L. F. Yang and L. Q. Mao, *Anal. Chem.*, 2013, **85**, 7550.
- 26 Y. C. Fu, P. H. Li, L. J. Bu, T. Wang, Q. J. Xie, J. H. Chen and S. Z. Yao, *Anal. Chem.*, 2011, **83**, 6511.
- 27 O. K. Farha and J. T. Hupp, *Acc. Chem. Res.*, 2010, **43**, 1166.
- 28 J. H. Wang, M. Li and D. Li, *Chemistry*, 2014, **20**, 12004.
- 29 D. Bradshaw, T. J. Prior, E. J. Cussen, J. B. Claridge and M. J. Rosseinsky, *J. Am. Chem. Soc.*, 2004, **126**, 6106.
- 30 M. Breuer, K. Ditrach, T. Habicher, B. Hauer, M. Keßler, R. Sturmer and T. Zelinski, *Angew. Chem., Int. Ed.*, 2004, **43**, 788.



- 31 H. K. Kim, W. S. Yun, M.-B. Kim, J. Y. Kim, Y.-S. Bae, J. D. Lee and N. C. Jeong, *J. Am. Chem. Soc.*, 2015, **137**, 10009.
- 32 Y. Zhang, Y. Wang, H. Wang, J. H. Jiang, G. L. Shen, R. Q. Yu and J. Li, *Anal. Chem.*, 2009, **81**, 1982.
- 33 L. Chen, K. L. Li, H. Zhu, L. C. Meng, J. T. Chen, M. X. Li and Z. W. Zhu, *Talanta*, 2013, **105**, 250.
- 34 Y. Liu, W. Xuan and Y. Cui, *Adv. Mater.*, 2010, **22**, 4112.

