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Regulating photocatalysis by external-stimuli manipulation of the microenvironment in europium–organic frameworks†

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Although catalysis is highly dependent on the catalytically active site, accurately regulating the microenvironment around the active site is an important way to improve the overall performance of catalysts. Herein, an Eu-based MOF with a rod-shaped secondary building unit (SBU) was obtained through a self-assembly strategy, and named Eu-MOF-T (T = temperature). Eu-MOF-T can absolutely and kinetically transform into another structure, Eu-MOF-S (S = solvent), under a stimulus in different solvents. Interestingly, Eu-MOF-S can convert into Eu-MOF-T absolutely and thermodynamically upon temperature stimulation. These transformations are mainly caused by microenvironmental changes, including the ligand torsion angle and cavity volume in the structure, and take place in the form of single-crystal-to-single-crystal. Incredibly, Eu-MOF-S exhibits a halved cavity volume due to the bending of the ligand, yet demonstrates a superior photocatalytic CO₂RR capacity of 7886.1 μmol g⁻¹ h⁻¹, while Eu-MOF-T, which possesses a larger cavity porosity, has a capacity of only 599.7 μmol g⁻¹ h⁻¹. Theoretical calculations further reveal that Eu-MOF-S is more favorable for the formation of COOH* and promotes its further conversion into CO during the CO₂RR.

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Introduction

Metal–organic frameworks (MOFs), as excellent model catalysts, have been increasingly used in small molecule catalysis in recent years, such as water oxidation, CO₂ reduction, organic small molecule transformations, *etc.*^{1–4} which is attributed to their well-defined catalytic sites, tunable structures, and suitable material transport channels, but their unstable feature makes their application in catalysis very limited.^{5–8} The design and preparation of framework-stabilized MOFs are essential prerequisites for the development of their catalytic properties. Currently known frameworks, which can stabilize above 400 °C, such as UiO-66,⁹ NU-1000,¹⁰ MIL-101,¹¹ and MOF-808,¹² are too rigid and only have a single type of nodal metal. This makes them unable to

reflect the advantages in the field of catalysis, especially in the field of photocatalysis. Designing MOFs with high thermal stability, good framework flexibility and node diversity is one of the next tasks for MOF designers.^{13–15}

Although the catalytic performance is highly dependent on the active sites of the catalyst, including the number and distribution of the sites, the impact of the microenvironment surrounding the catalytic site on the catalytic outcome is typically significant yet often disregarded.^{16,17} Certain pivotal microenvironments intricately maintain a multifaceted and delicate equilibrium within living systems, actively participate in the regulation of diverse physiological processes, and assume an indispensable role in upholding the robust development of organisms.¹⁸ Natural enzymes, as the catalysts with the highest catalytic efficiency in nature, have abundant amino acid residues around their centers to act as microenvironments to enhance the catalytic efficiency of the catalytic sites.¹⁹ The microenvironment is an extremely important factor in determining the physical or chemical behavior of the surrounding molecules as a powerful complement to the catalytic site, the main types of which include the ligand situation, spatial accessibility, residual groups, conjugated environments, and guest molecules, and designing and making full use of these types can significantly enhance catalytic effects.^{20–22}

The crystalline nature of MOF materials provides a good basis for studying microenvironmental changes induced by

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external stimuli at the atomic level.^{23–26} The microstructure of MOFs with a certain framework flexibility can obviously respond to changes in external stimuli, and the magnitude of such changes can be determined by means of single-crystal testing.^{25,27,28} Single-crystal-to-single-crystal (SCSC) conversion has been a hot topic in the field of MOF research in recent years, and the relationship between structure and performance in applying SCSC to the catalytic process is rarely reported at present. By introducing PDMAEMA-functionalized pollen as “smart” anchors, Lee’s group was able to “turn on” and “turn off” the catalytic activity of MOF@P-pollen with tunable dispersion, environmentally responsive activity, and significantly enhanced liquid-phase photocatalytic performance.²⁹ However, among the thousands of MOFs reported in the past decades, only a few were able to respond to external stimuli, and this practical necessity motivates us to continue researching this type of material.^{30,31}

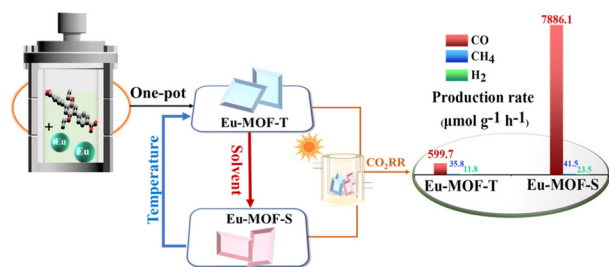
In order to deeply investigate the mechanism of the influence of the microenvironment around the active site on the photocatalytic performance to understand the microenvironment–activity relationship, structurally flexible MOFs with well-defined microenvironments were constructed in this study using the rare-earth metal Eu and flexibly branched bicarboxylic acid ligands (Eu-MOF-T and Eu-MOF-S, Scheme 1). The reversible single-crystal-to-single-crystal (SCSC) conversion of Eu-MOFs under external stimuli was performed by a simple strategy, in which Eu-MOF-T can be converted into Eu-MOF-S using a solvent under 20 min immersion, and conversely, Eu-MOF-S can be fully converted into Eu-MOF-T again at 220 °C. The photocatalytic results show a larger production of CO₂ to CO for Eu-MOF-S.

Materials and equipment: All chemicals were commercially sourced and used without further purification.

Results and discussion

The Experimental section is given in the ESI.† The detailed information for Eu-MOF is summarized in Tables S1 and S2.†

Two Eu-MOFs with different coordination environments were prepared by a mild and convenient method using the



Scheme 1 Brief schematic diagrams of the construction, structural transformation and photocatalytic properties of Eu-MOFs (in the experiment, the crystals were all colorless and transparent, but here the blue crystal represents Eu-MOF-T and the red crystal represents Eu-MOF-S).

same Eu metal salt and the flexible ligand DTDA as precursors. First, Eu-MOF-T was obtained by self-assembly using the classical one-pot synthesis method. Eu-MOF-T was immersed in different solvents to form Eu-MOF-S, which occurred in a SCSC manner. Single crystal X-ray diffraction studies show that the crystal structures belonging to the triclinic system of both Eu-MOFs were constructed from the same Eu metal ion and the DTDA ligand, in the $P\bar{1}$ space group. The asymmetric units all contain an Eu(III) cation, one and a half DTDA²⁻ anions, and two coordinating water molecules (Fig. 1a and b). The difference is that the guest molecules of Eu-MOF-T contain three DMF and eight water molecules, while Eu-MOF-S contains only one DMF and six water molecules. And the torsion angles of the DTDA ligands in the two MOFs are very different (Fig. 1c, d, Fig. S3a and b†). The DTDA ligand simultaneously coordinates with four Eu(III) ions in a saturated coordination mode, and each Eu(III) ion coordinates with six DTDA ligands and two water molecules in an eight-coordinate mode. For two structures, the Eu(III) ions are connected by four carboxyl groups from four DTDA ligands to form a twisted paddlewheel unit (Eu₂(CO₂)₄). These units further extend indefinitely to form rod-shaped SBUs (Fig. S3g and h†), and ultimately form an infinite three-dimensional structure by DTDA cross-linking (Fig. 1e and f). From a topological point of view, the spatial structure of the entire complex can be simplified as a new 5,5,6 connected three-node network with the point symbol {3².4⁴.6².7.8} {3².4⁶.5.6} {3².4⁶.5².6⁵}. The pore structures of the two Eu-MOFs are clearly different (Fig. S3b–f†). The void

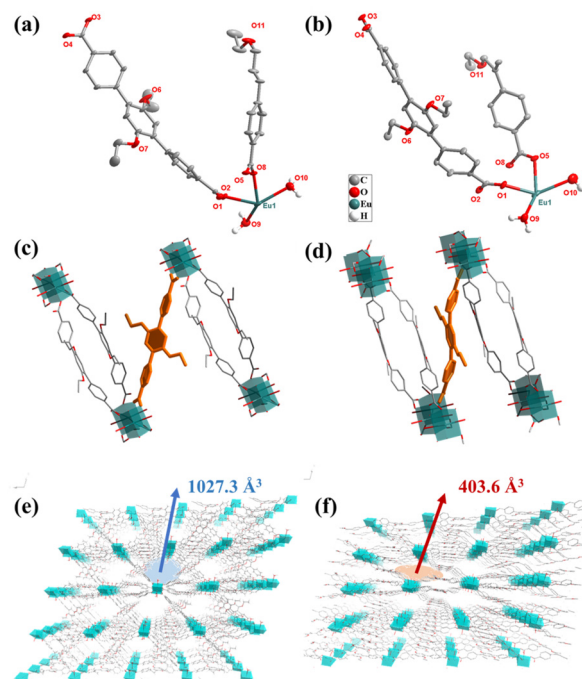


Fig. 1 Asymmetric units of Eu-MOF-T (a) and Eu-MOF-S (b); partial structural diagrams of Eu-MOF-T (c) and Eu-MOF-S (d); 3D structural diagrams of Eu-MOF-T (e) and Eu-MOF-S (f); the labeling in the figure is the pore volume.

volume of Eu-MOF-T was 1027.3 \AA^3 (41.4% of the unit cell volume), whereas the pore volume of Eu-MOF-S was reduced to roughly half, *i.e.*, 403.6 \AA^3 (21.2% of unit cell volume) (Fig. 1e, f and S3e and f†).

Powder X-ray diffraction (PXRD) was conducted to determine the structural transformation processes that regulate solvent- and temperature-mediated polycrystalline transformations, with the kinetic product Eu-MOF-S interconverting with the thermodynamic product Eu-MOF-T. The PXRD patterns of the synthesized Eu-MOF-T were very similar to the crystal structure simulations (Fig. S4a†), but changed significantly under different solvents submerged for different times. Specifically, the initial peak of Eu-MOF-T ($2\theta = 6.1^\circ$) disappeared rapidly after 20 min of immersion, while a new peak ($2\theta = 8.2^\circ$) appeared, and after 24 h of immersion, it was found to be completely transformed into Eu-MOF-S in all solvents without the appearance of any other new structures (Fig. 2a). Interestingly, this conversion was achieved through an SCSC manner in a variety of solvents. Amazingly, Eu-MOF-S is gradually transformed into Eu-MOF-T during the warm-up process (Fig. 2b). As known from Fig. 2f, this process can be divided into three stages. The first stage is the induction period; the PXRD spectra do not change significantly until 190°C . Both Eu-MOF-T and Eu-MOF-S have guest molecules and their conversion has to overcome a barrier for activation. The transformation starts from the second stage ($190\text{--}220^\circ\text{C}$), during which the 2θ peak at 8.2° for Eu-MOF-S gradually disappears and a new peak at 6.1° subsequently increases. In the third stage, above 220°C , the peak at 8.2° almost disappears, while the intensity of the peak at 6.1° continues to increase, and the final PXRD is in good agreement with the simulated Eu-MOF-T. This indicates that the polycrystalline transformation of Eu-MOF-S and Eu-MOF-T is successful and reaches the thermodynamic minimum.

Although the Eu-MOF-T and Eu-MOF-S transformations show low signal-to-noise ratios in the infrared spectra, the characteristic infrared variation bands of both structures can

still be observed by testing at different temperatures (Fig. S4c and d†). The characteristic peaks for the asymmetric stretching vibration of Eu–O (595 cm^{-1}), as well as the characteristic peaks of C–O and C–H of the DTDA ligand, and the peaks of DMF and H_2O , can be clearly identified in the IR spectrum. With the increase of temperature, the infrared bands of Eu-MOFs changed significantly, and the different frequencies of the C–H ($\sim 2850\text{--}3000 \text{ cm}^{-1}$) and C–O ($\sim 1630 \text{ cm}^{-1}$) stretching vibrations led to an increase in the peak bandwidth and a blue shift, which is in agreement with the results of the structural transformations (Fig. 2c and d). The vibrations of DMF disappeared, whereas the benzene rings of DTDA ligands remained unchanged, and the Eu–O bonds were unchanged. From the thermogravimetric analysis (TGA), it can be seen that Eu-MOF-T and Eu-MOF-S underwent a strong pyrolysis at 410°C (Fig. S5a and b†), with the final thermal weight loss reaching 50.9% and 58.5%, respectively. In the N_2 adsorption and desorption experiments at 77 K (Fig. S6a and b†), Eu-MOF-T adsorbed $128 \text{ cm}^3 \text{ g}^{-1}$ with a pore size distribution of $7.3\text{--}12.7 \text{ \AA}$, whereas Eu-MOF-S did not adsorb or desorb N_2 , and the pore size distribution obtained by CO_2 adsorption and desorption experiments was $6.9\text{--}12.7 \text{ \AA}$, which suggests that Eu-MOF-T has a larger pore size. In CO_2 adsorption and desorption experiments (Fig. S6c and d†), Eu-MOF-T outperforms Eu-MOF-S at both temperatures, but the presence of a large hysteresis loop in the isothermal adsorption and desorption curves of Eu-MOF-S indicates a strong CO_2 adsorption capacity, which can capture and immobilize CO_2 molecules more efficiently during the catalytic process, which is conducive to the photocatalytic CO_2 RR.

Further spectroscopic measurements were performed to determine the actual effect of the ligand-twisted microenvironment on the formation of Eu-MOFs. In general, PL signals are generated by photogenerated electron–hole complexation, and the peak intensity correspondingly indicates the degree of complexation. As shown in Fig. S5c,† both Eu-MOF-T and Eu-MOF-S show four PL emission peaks, which can be attributed to the typical three emission peaks of metal Eu and one emission peak of ligand DTDA. Among them, Eu-MOF-S has the lowest PL signal, indicating that it has the highest separation efficiency of photoelectrons and holes. At various temperatures, however, the fluorescence emission showed a significant redshift from low to high temperature tests, which was caused by molecular motion (Fig. 3a). In addition, the time-resolved photoluminescence (TRPL) decay spectra show a smaller average fitted lifetime of 31 ns for Eu-MOF-S relative to the 37 ns for Eu-MOF-T, suggesting that its singlet excited state decays faster and is capable of rapid electron transfer (Fig. S5e and f†). The UV-vis spectra showed that Eu-MOF-T and Eu-MOF-S possessed similar light absorption properties, with two distinct absorption peaks at 286 nm and 355 nm. Compared with the DTDA ligand, the enhancement of absorption in the visible region of both MOFs is not negligible (Fig. S5d,† $\geq 400 \text{ nm}$), and both have competitive potential for the photocatalytic reduction of CO_2 . The band gaps of Eu-MOF-T and Eu-MOF-S are calculated to be 2.99 and 2.84 eV, respectively,

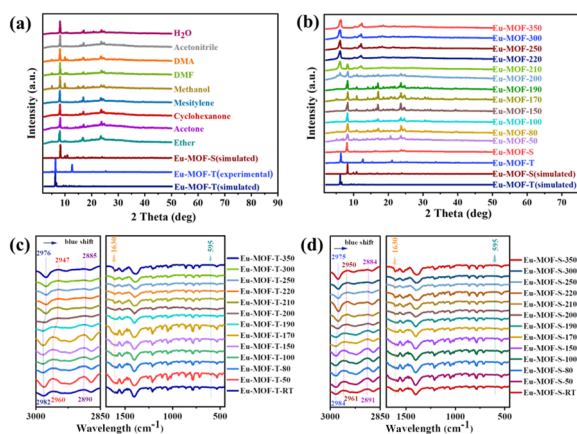


Fig. 2 (a) PXRD spectra of two Eu-MOFs in different solvents; (b) PXRD spectra of two Eu-MOFs at different temperatures; enlarged details of IR spectra of (c) Eu-MOF-T and (d) Eu-MOF-S at different temperatures.

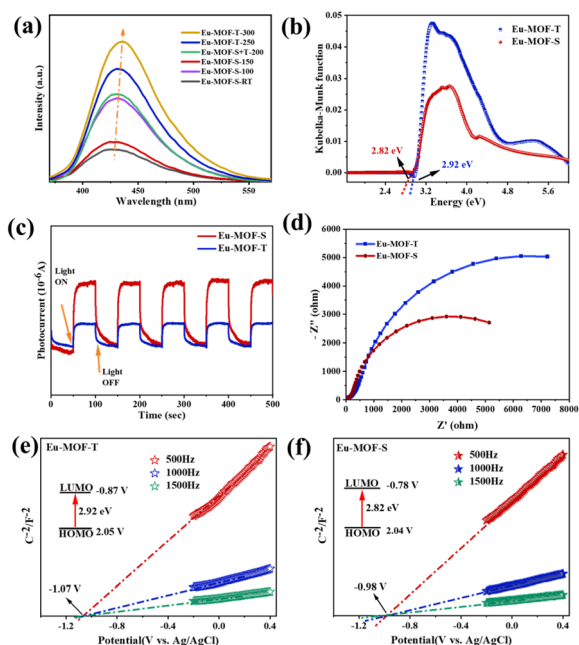


Fig. 3 (a) Fluorescence emission plots of Eu-MOFs at different temperatures; (b) Kubelka–Munk energy function plots of Eu-MOFs; (c) the corresponding plots of transient photocurrents of Eu-MOFs; (d) ESI impedance plots of Eu-MOFs; (e) Mott–Schottky plots of Eu-MOF-T (e) and Eu-MOF-S (f).

by the Kubelka–Munk function (Fig. 3b). As can be seen from the band gap structure, the prepared photocatalysts fully satisfy the favorable conditions for the conversion of CO₂ to CO. The ligand-twisted Eu-MOF-S photocatalyst with efficient photogenerated carrier separation efficiency, excellent light absorption properties and suitable bandgap structures can enhance the photocatalytic CO₂ reduction performance.

The transient photocurrent response was tested under visible light using an electrochemical workstation. When the light source is turned on, the catalysts absorb photons with energies equal to or greater than the bandgap energy, leading to the rapid generation of electron–hole pairs on a picosecond time scale, resulting in an instantaneous increase in the photocurrent. Over time, the process of the trapping and re-emission of light-generated carriers on the trap gradually takes effect, leading to a gradual decay of the current to the equilibrium current. The slow decay of the current when the light is turned off can be understood as being a result of the gradual re-emission and recombination of charge carriers from the trap. The photoresponsive capability of Eu-MOF-S is much stronger than that of Eu-MOF-T and boosts more photogenerated electron–hole pairs (Fig. 3c). In order to better define the carrier transport type of Eu-MOFs, Mott–Schottky experiments were performed. As shown in Fig. 3e and f, Eu-MOF-T and Eu-MOF-S have positive slopes and are both typical n-type semiconductors with flat band potentials (E_{fb}) of -1.07 V (V vs. Ag/AgCl) and -0.98 V (V vs. Ag/AgCl), respectively. Based on the optical bandgap values, the HOMO and LUMO positions of

the Eu-MOFs are roughly calculated, where the HOMO and LUMO positions of the Eu-MOF-T are at 2.12 V and -0.87 V, respectively, while those of the Eu-MOF-S are at 2.06 V and -0.78 V, respectively. The electron transfer ability of the two MOFs was investigated using EIS. In general, the smaller the R_{ct} value of the catalyst, the smaller the surface charge transfer impedance and the stronger the electron transfer ability. The conductivity of Eu-MOF-S was significantly increased after undergoing solvent stimulation, suggesting that ligand twisting significantly improves the electron transfer capacity of Eu-MOFs (Fig. 3d). The results show that the effective capture of photogenerated carriers by the twisted microenvironment of Eu-MOF-S can promote the separation of photogenerated electron–hole pairs. The captured photogenerated electrons rapidly migrate to the catalyst surface, which in turn promotes the photocatalytic CO₂ reduction.

The CO₂ photoreduction performance of two Eu-MOFs was tested using a visible 400 nm cartridge xenon lamp as a light source, acetonitrile as a solution, Ru(bpy)₃Cl₂ as a photosensitizer, and triethylamine (TEA) as a sacrificial agent, to investigate the effects of microenvironmental changes induced by external stimuli on the photocatalytic activity. The collected gaseous products were detected using gas chromatography and the liquid products were analyzed using a mass spectrometer. The results show that the photocatalytic CO₂ reduction products for Eu-MOF-T and Eu-MOF-S are indeed H₂, CO and CH₄.

Firstly, the reduction performance of Eu-MOFs at different activation temperatures was investigated, as shown in Fig. 4a. With the prolongation of the irradiation time, the CO production rate increased rapidly in the initial steps, then decreased, and increased slowly in the last stage. This may be attributed to the faster decomposition of the photosensitizer during the experimental process, and led to a sudden decrease in the reactivity. Due to the high stability of the catalyst, the

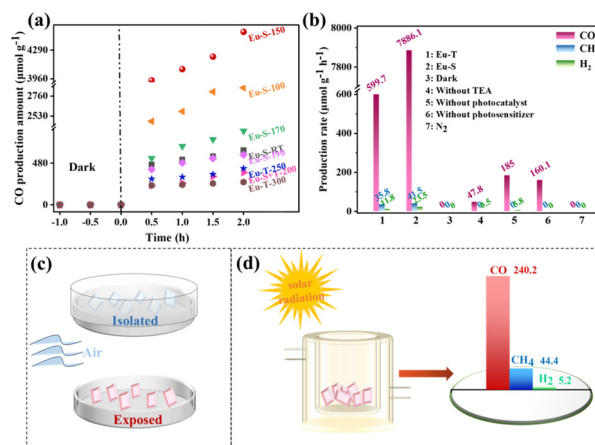


Fig. 4 (a) CO yield plots of photocatalytic CO₂ reduction by Eu-MOFs at different temperatures; (b) photocatalytic CO₂ reduction yield plots of Eu-MOFs under different conditions; (c) graphical representation of the glass-surface vessel opening/closing reaction; (d) graphical representation of photocatalytic CO₂ reduction by sunlight.

CO production amount continued to increase. After irradiation for 2 h, Eu-MOF-S showed the best catalytic effect at an activation temperature of 150 °C, and Eu-MOF-S was superior to Eu-MOF-T at any temperature. Next, blank and control experiments were carried out under the same experimental conditions (Fig. 4b) to investigate the yields and selectivities. It can be seen that Eu-MOF-S is optimal in terms of photocatalytic efficiency and selectivity (99.2%), and its catalytic activity of $7886.1 \mu\text{mol g}^{-1} \text{h}^{-1}$ is about 13 times better than that of Eu-MOF-T, reflecting the necessity of microenvironmental studies in stimulus-responsive MOFs. No CO_2 reduction products were detected under both N_2 and dark conditions, suggesting that the products originate from a light-driven catalytic reduction reaction of dissolved CO_2 , and confirming that the CO_2 feed is indeed the carbon source. In the absence of the sacrificial agent triethylamine, the catalytic activity was reduced to $47.8 \mu\text{mol g}^{-1} \text{h}^{-1}$ CO due to severe electron-hole complexation. The catalytic activity of Eu-MOF-S without the photosensitizer was significantly reduced to $160.1 \mu\text{mol g}^{-1} \text{h}^{-1}$. However, $[\text{Ru}(\text{bpy})_3]\text{Cl}_2$ itself produced $185 \mu\text{mol g}^{-1} \text{h}^{-1}$ CO and $5.8 \mu\text{mol g}^{-1} \text{h}^{-1}$ H_2 , which was subtracted from the catalytic reduction results of Eu-MOFs. Controlled experiments demonstrated that the Eu-MOF photocatalyst, light, photosensitizer, sacrificial agent and CO_2 were indispensable. In order to evaluate the stability of the Eu-MOF catalysts, cycling experiments were carried out, and only a slight decrease in the CO yield was observed in five test cycles (Fig. S7a†). The XRD spectra of Eu-MOFs before and after the reaction were almost the same (Fig. S7b†), indicating quality framework structure stability during the photocatalytic testing.

It is extremely interesting that Eu-MOF-T remains unchanged when placed in an isolated glass-surface vessel for one week after synthesis, while it can be converted into Eu-MOF-S in 3 days when placed in an open glass-surface vessel, *i.e.*, when exposed to air, indicating that a very weak stimulus can elicit a response from Eu-MOF-T (Fig. 4c). The solar photocatalytic CO_2 RR was carried out, and the experimental procedure was consistent with the “Photocatalytic reduction of CO_2 test” described in the ESI,† except that the xenon lamp was replaced by sunlight. The CO_2 to CO production rate was $240.2 \mu\text{mol g}^{-1} \text{h}^{-1}$ (Fig. 4d), which indicates that Eu-MOF-S is a new stimulus-responsive MOF material with potential for CO_2 reduction using solar energy.

Density Functional Theory (DFT) calculations were performed on the above reaction steps to understand the photoreductive activities of Eu-MOF-T and Eu-MOF-S at the atomic level, especially the key role of the microenvironment around the active site in CO_2 conversion. Fig. S6a† shows that the Gibbs free energy (ΔG) of Eu-MOF-T is lower than that of Eu-MOF-S. The theoretical bandgap values of Eu-MOF-T and Eu-MOF-S are 2.91 and 2.83 eV, respectively, which are in agreement with the experimental test results (Fig. S8b†). The ΔG of the photocatalytic CO_2 reduction process was calculated for Eu-MOFs according to two reaction pathways, where the negative ΔG value of the adsorbed CO_2 represents the effective capture and activation of CO_2 molecules. Fig. 5a shows the

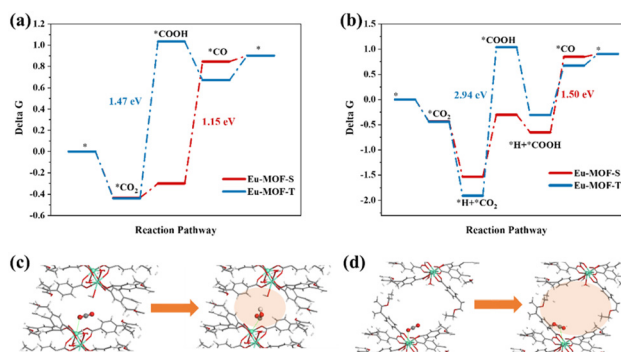


Fig. 5 (a and b) Gibbs free energy diagram of the CO_2 reduction pathway of Eu-MOFs, where in the a-figure path the proton originates from solution and in the b-figure path the proton originates from the adsorbed state H^+ near the catalytic site; (c) state diagram of CO_2 to *COOH near metal sites in the pores of Eu-MOF-S; (d) state diagram of CO_2 to *COOH near metal sites in the pores of Eu-MOF-T.

path of protons originating from the solution, and the first proton-coupled electron transfer (PCET) on Eu-MOF-T forms adsorbed COOH^* with a ΔG of 1.47 eV. In contrast, the lower energy barrier of ΔG (-0.3 eV) for Eu-MOF-S obviously favors the catalytic process. The other pathway is the proton originating from the adsorbed state *H near the catalytic site (Fig. 5b); the ΔG values of the process ($\text{*CO}_2 + \text{H}^+ + \text{e}^- \rightarrow \text{*COOH}$) are 2.94 eV and 1.23 eV for Eu-MOF-T and Eu-MOF-S, respectively. It follows that *COOH is the rate-determining step for CO_2 -to-CO and Eu-MOF-S is superior to Eu-MOF-T under both pathways. Detailed information on PCET throughout the photoreduction process and the corresponding structures of the reaction intermediates are provided in Fig. S9.† In the further PCET process, dehydration occurs to form *CO which then desorbs from the active site.

Further computational simulations show that the presence of the HOMO on the metal site and total density of states (DOS) originate from the Eu-4f atomic orbital for the Eu-MOF-S configuration, as seen in Fig. S10,† which facilitates electron transfer from the substrate to CO_2 during CO_2 adsorption. The next two hydrogenation steps of CO_2 to *COOH and *COOH to CO are also accompanied by the transfer of electrons, and the better electrical conductivity of Eu-MOF-S may result in the lowering of *COOH energy barriers, which creates ordered and fast electron transfer conditions for photocatalysis. It is noteworthy that *COOH and Eu sites are positively charged and do not approach each other easily, whereas the apparently small pore of Eu-MOF-S favors the confinement of *COOH , making its distance to the Eu site 2.9 Å, which is smaller than that of 3.76 Å for Eu-MOF-T. *COOH is forced to be close to the Eu site and appears to interact with O around Eu in the presence of H stabilization (differential charge density plot), reducing the overall energy (Fig. 5c and S9†). It can be seen that the modulation of the catalytic properties of crystalline MOF materials is ultimately achieved by changes in the microenvironment of the active site manipulated by external stimuli such as temperature and solvent, including the

ligand torsion angle, cavity volume, and electronic effects. This catalytic property enhancement accompanying the SCSC conversion process is a major breakthrough in the application field of single MOF materials, and also provides experimental and theoretical support for the study of stimulated crystalline materials.

Conclusions

In order to deeply investigate the mechanism of the microenvironment around the active site on the photocatalytic performance and to understand the microenvironment–activity relationship, in this paper, structurally flexible MOFs with a good microenvironment were constructed using rare-earth metal Eu and flexibly branched dicarboxylic acid ligands. They have flexibly branched dicarboxylic acid ligands, DTDA, and the two carboxyl groups are attached to two metal Eu(III) ions, respectively. The effects of different microenvironments around the metal active sites on the photocatalytic CO₂ reduction performance and the subtlety of the stimulus-responsive structural modulation were investigated by adjusting the torsion angle of DTDA in the model system. Interestingly, reversible SCSC conversion of the Eu-MOF can be achieved under external stimulation using a simple strategy, in which Eu-MOF-T can be converted into Eu-MOF-S in a solvent with immersion for 20 min, and in contrast, Eu-MOF-S can be completely converted into Eu-MOF-T again under an Ar atmosphere at 220 °C. Photocatalytic CO₂ reduction results showed that Eu-MOF-S with a larger ligand torsion angle significantly improved the photocatalytic activity and reached the optimum catalytic effect at an activation temperature of 150 °C. Utilizing the fundamental physical properties of the stimuli-responsive framework with host–guest interactions and rationally designing the ligand torsion, not only was a better photocatalytic performance obtained, but also the microenvironment–activity relationship was elucidated to achieve the performance tuning of the microenvironmental aspect of stimuli-responsive MOF catalysis.

Data availability

All relevant data are within the manuscript and ESI is available. CCDC 1923215 and 2354711.†

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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