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A single-atom Cu–N₂ catalyst eliminates oxygen interference for electrochemical sensing of hydrogen peroxide in a living animal brain†

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Hydrogen peroxide (H₂O₂) plays essential roles in various physiological and pathological processes. The electrochemical hydrogen peroxide reduction reaction (HPRR) has been recognized as an efficient approach to H₂O₂ sensing; however, the HPRR has always suffered from low tolerance against the oxygen reduction reaction (ORR), resulting in poor selectivity of the HPRR-based sensing platform. In this study, we find that the electrochemical HPRR occurs preferentially compared to the ORR when isolated Cu atoms anchored on carbon nitride (Cu₁/C₃N₄) are used as a single-atom electrocatalyst, which is theoretically attributed to the lower energy barrier of the HPRR than that of the ORR on a Cu₁/C₃N₄ single-atom catalyst (SAC). With the Cu₁/C₃N₄ SAC as the electrocatalyst, we fabricated microsensors that have a good response to H₂O₂, but not to O₂ or other electroactive neurochemicals. When implanted into a living rat brain, the microsensor shows excellent *in vivo* sensing performance, enabling its application in real-time quantitative investigation of the dynamics of H₂O₂ production induced by mercaptosuccinate and glutathione monoethyl ester in a living animal brain.

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Introduction

In physiological and pathological processes, H₂O₂ plays important roles, for example, as a biochemical mediator and signaling pathway modulator.^{1–7} In aerobic organisms, O₂ is the sole source for H₂O₂ production which is accomplished by one-electron reduction of O₂ to form superoxide ([•]O₂[–]) followed by the disproportionation of [•]O₂[–] mediated by superoxide dismutase (SOD).^{6–8} As a representative of reactive oxygen species (ROS), the level of H₂O₂ can be regulated by antioxidants such as glutathione peroxidase, catalase, thioredoxins and thiols to maintain the cellular redox homeostasis.^{9,10} In addition, H₂O₂

with a low physiological level can serve as an intracellular signaling molecule to regulate metabolism, enabling cellular adaption to the changes of stress and the environment.^{11,12} In contrast, overproduction of H₂O₂ can break the redox balance and thus cause oxidative stress, which is closely related with a variety of pathological events such as aging and progressive neurodegenerative disorders.^{13–15} Moreover, H₂O₂ is produced in enzymatic reactions with O₂ as an electron receptor, which can thus serve as an indicator of the enzymatic activity and the level of related substrates.^{16,17} Therefore, the information on the kinetics of H₂O₂ would largely advance these studies.^{18–20}

Electrochemical oxidation or reduction of H₂O₂ has been used to build a platform for H₂O₂ sensing;²¹ however, the oxidation of H₂O₂ often bears interference from some reductive biomolecules,²² and the reduction process always suffers from interference from O₂ with almost all electrocatalysts.^{23–26} Although the HPRR is more favorable than the ORR thermodynamically (eqn (1) and (2)), it often proceeds more slowly than the ORR under practical conditions due to the sluggish kinetics of the HPRR and low mass transport efficiency of H₂O₂,^{23,24} resulting in the occurrence of the HPRR at more negative potentials than the ORR, which makes it difficult to selectively sense H₂O₂ free from the interference from O₂ when potential-controlled amperometry is used. Therefore, the design and fabrication of high-performance electrocatalysts where the HPRR occurs prior to the ORR are very essential for selective H₂O₂ sensing *in vivo*.

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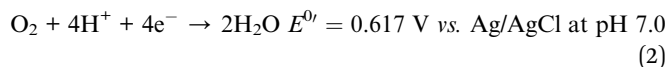
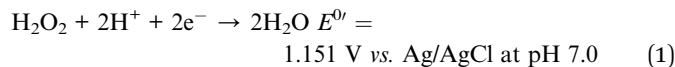
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Single-atom catalysts (SACs)^{27–32} with almost 100% atom utilization efficiency and uniform catalytic centers have recently received considerable attention in industrial chemical processing^{33–35} and electro/photochemical energy conversion^{36–39} such as the ORR,^{40–43} hydrogen evolution reaction (HER),^{44–46} oxygen evolution reaction (OER),^{47–49} and carbon dioxide reduction reaction (CO₂RR).^{50–54} SACs exhibit unique selectivity because of the homogeneous and well-defined structure compared with nanoparticle-based catalysts.^{55–57} In addition, the structure of SACs can be flexibly regulated by changing the active atom species and/or the support and by tuning the coordination environment to achieve a high catalytic performance.^{58–61} Such unique characteristics and the inner-sphere electron-transfer features of the HPRR and ORR suggest that the use of SACs may offer priority to one of these two reactions because transition metal single atoms with different coordination environments embedded in a well-defined carbon support have shown high capacity in tuning the binding strength with O species,^{62–64} providing a great opportunity for the design and fabrication of selective electrocatalysts for prioritizing the HPRR over the ORR and thus enabling amperometric sensing of H₂O₂ free from interference from O₂.

In this work, we report the design and synthesis of a SAC with atomically dispersed Cu sites anchored on mesoporous graphitic carbon nitride (Cu₁/C₃N₄) to realize selective electrochemical HPRR against the ORR. The Cu₁/C₃N₄ SAC synthesized by an impregnation method with a mesoporous structure exhibits a high electrocatalytic performance toward the HPRR with an onset potential at about 0.20 V (vs. Ag/AgCl) in neutral media. In contrast, the ORR commences at a more negative potential at around 0.0 V (vs. Ag/AgCl), demonstrating that the occurrence of the electrochemical HPRR prior to the ORR can be successfully achieved through rational design and fabrication of SACs. Density functional theory calculation (DFT) reveals that the priority given to the HPRR over the ORR on the Cu₁/C₃N₄ SAC is due to the synergetic effect of the Cu–N₂ site and its adjacent carbon site on the adsorption state of the *OH intermediate, resulting in a lower energy barrier of the HPRR on the SAC. This finding not only offers a new electrocatalyst for promoting the HPRR over the ORR, but also provides a novel avenue for selective *in vivo* monitoring of molecular events in a living rat brain with SACs as the electrocatalysts.

Results and discussion

To synthesize SACs, C₃N₄ with a mesoporous structure was prepared by the thermally induced self-condensation of cyanamide with colloidal silica as a hard template and was used as the support, enabling a large surface area with accessible active sites.⁶⁵ In order to find an electrocatalyst with selectivity for the HPRR over the ORR, a series of transition metal SACs (denoted

as M₁/C₃N₄, M = Fe, Co, Cu, Mn) were prepared by calcining C₃N₄ absorbed with metal ions (Fig. 1a). Fig. 1b and S1† show the transmission electron microscopy (TEM) images of the as-synthesized samples, retaining the mesoporous structure with ca. 10 nm pores of C₃N₄ (Fig. S2†), which is beneficial to the mass transport of H₂O₂. No clusters or nanoparticles were observed in the TEM images, showing that metal atoms exist as single atoms dispersed on the support in all the prepared catalysts. X-ray photoelectron spectra (XPS) suggest a similar structure of the four samples and the existence of M–N_x species positioned at around 398.9 eV (Fig. S3 and S4†).⁶⁶ Among the synthetic SACs, Cu₁/C₃N₄ possesses striking electrocatalytic performance toward H₂O₂ reduction in terms of onset potential and current intensity (Fig. S5†), demonstrating the vital role of the single-atom metal centers in tuning the reaction activity. Fourier transform infrared (FT-IR) spectra of the as-prepared C₃N₄ and Cu₁/C₃N₄ demonstrate the formation of extended

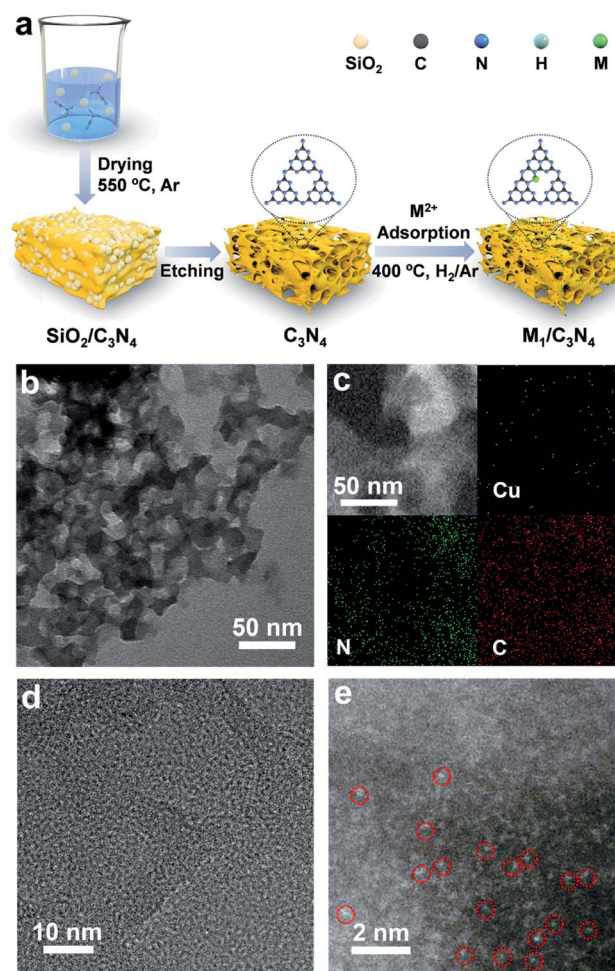


Fig. 1 Synthesis and characterization of the Cu₁/C₃N₄ SAC. (a) Illustration of the synthetic route to M₁/C₃N₄. (b) TEM image of Cu₁/C₃N₄. (c) HAADF-STEM and corresponding elemental mapping images of Cu₁/C₃N₄ showing the uniform distribution of Cu (yellow), N (green) and C (red). (d and e) High-resolution TEM image (d) and AC HAADF STEM image (e) of Cu₁/C₃N₄ (single Cu atoms are marked with red circles).



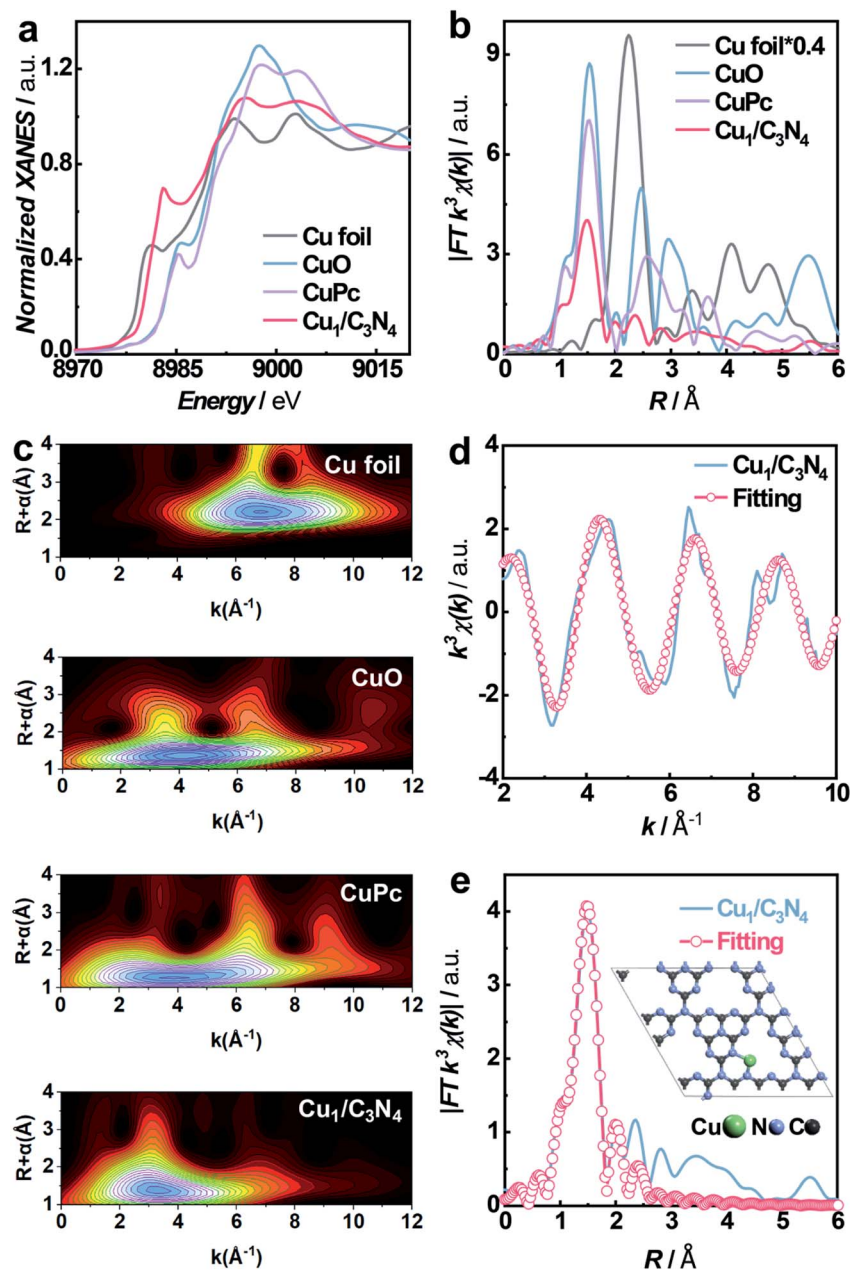


Fig. 2 XAS analysis of $\text{Cu}_1/\text{C}_3\text{N}_4$. (a and b) Normalized XANES spectra (a) and FT-EXAFS spectra (b) at the Cu K-edge of $\text{Cu}_1/\text{C}_3\text{N}_4$ (red line), Cu foil (black line), CuO (blue line) and CuPc (purple line). (c) Wavelet transforms of the Cu K-edge EXAFS spectra for Cu foil, CuO, CuPc and $\text{Cu}_1/\text{C}_3\text{N}_4$. (d and e) Corresponding fitting curves of the EXAFS spectra of $\text{Cu}_1/\text{C}_3\text{N}_4$ in k space (d) and R space (e). The inset in (e) is the schematic structure of $\text{Cu}_1/\text{C}_3\text{N}_4$.

CN heterocycles (adsorption bands between 1200 cm^{-1} and 1650 cm^{-1}) and the introduction of Cu atoms does not change the support structure obviously (Fig. S6†).⁶⁷ Besides, the corresponding energy-dispersive X-ray spectroscopy (EDS) demonstrates the existence of Cu, N and C elements in $\text{Cu}_1/\text{C}_3\text{N}_4$ (Fig. S7†) and the corresponding mapping confirms that Cu, N, and C elements are uniformly distributed over the entire C_3N_4 skeleton (Fig. 1c). The N_2 adsorption/desorption isotherms and pore size distribution plots of C_3N_4 and $\text{Cu}_1/\text{C}_3\text{N}_4$ show a mesoporous architecture with a large BET surface area (Fig. S8†), which means that more active sites are available to reactant

molecules (Table S1†). X-ray powder diffraction (XRD) patterns indicate that the crystal structure of C_3N_4 is maintained after the introduction of Cu (Fig. S9†). In addition, $\text{Cu}_1/\text{C}_3\text{N}_4$ exhibits no characteristic peaks related with Cu, demonstrating that no Cu clusters or nanoparticles exist in the sample. To further verify the isolated single-atom form of Cu, high-resolution TEM and aberration corrected high-angle annular dark-field scanning TEM (AC HAADF-STEM) were performed (Fig. 1d and e). The bright dots with atomic size distributed on the C_3N_4 support were recognized to be single Cu atoms. Different areas were examined carefully to confirm the absence of Cu clusters



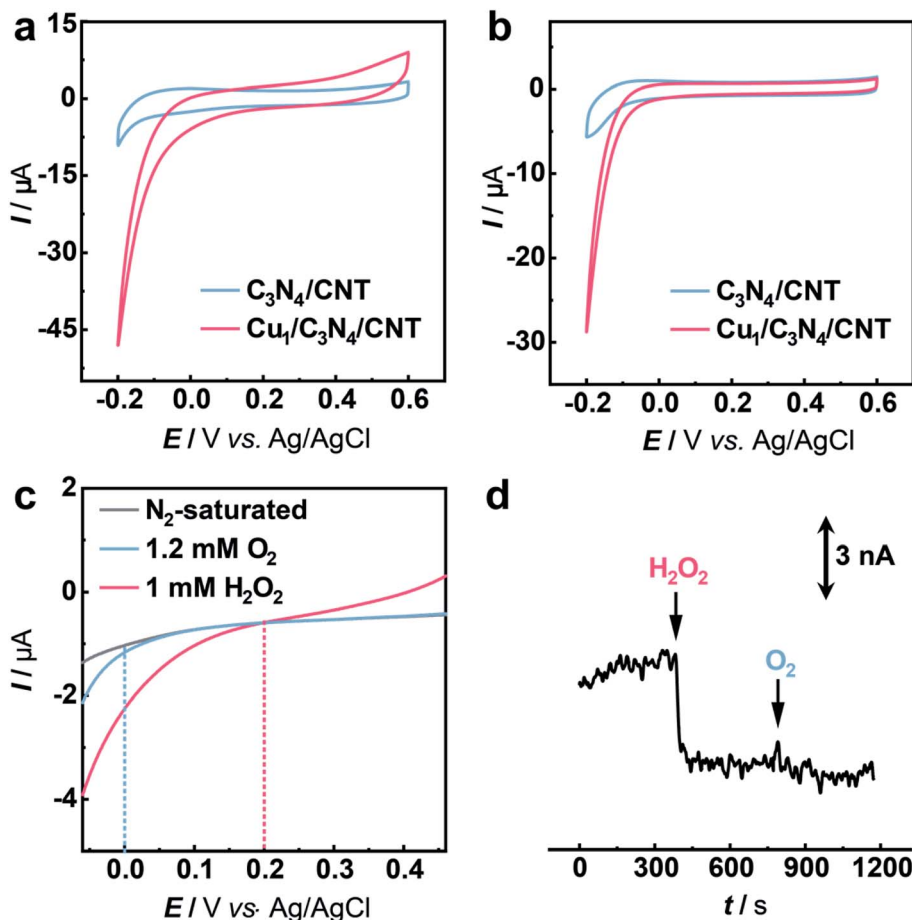


Fig. 3 Electrochemical study of the HPRR over the ORR. (a and b) CVs obtained with glassy carbon (GC) electrodes modified with $\text{C}_3\text{N}_4/\text{CNT}$ (blue curve) and $\text{Cu}_1/\text{C}_3\text{N}_4/\text{CNT}$ (red curve) in aCSF (pH 7.4) in the presence of 5 mM H_2O_2 (a) or under an O_2 atmosphere (ca. 1.2 mM O_2) (b). Scan rate: 50 mV s^{-1} . (c) LSVs obtained with the CuSAC-based electrode in aCSF without (gray curve) and with 1.2 mM O_2 (blue curve) or 1 mM H_2O_2 (red curve) in aCSF under a N_2 atmosphere. Scan rate: 50 mV s^{-1} . (d) Amperometric response recorded at the SAC-based electrode with the addition of $5 \mu\text{M H}_2\text{O}_2$ and $50 \mu\text{M O}_2$ in aCSF. Applied potential: 0.0 V vs. Ag/AgCl.

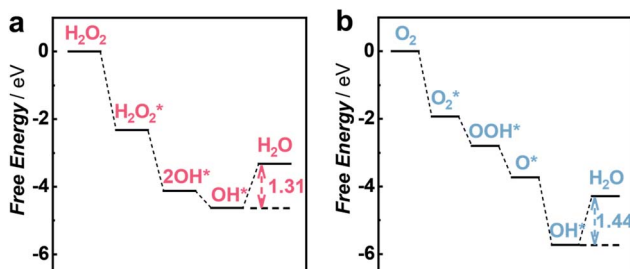


Fig. 4 DFT calculations for the HPRR and ORR on the $\text{Cu}_1/\text{C}_3\text{N}_4$ SAC. (a and b) Free energy diagrams of the HPRR (a) and ORR (b) at pH 7.0 on the active site of the $\text{Cu}_1/\text{C}_3\text{N}_4$ SAC.

or nanoparticles (Fig. S10[†]), revealing that only individual Cu single atoms are present in $\text{Cu}_1/\text{C}_3\text{N}_4$.

X-ray absorption spectroscopy (XAS) measurements at the Cu K-edge were performed to explore the electronic structure and coordination configuration of Cu species in $\text{Cu}_1/\text{C}_3\text{N}_4$. The X-ray absorption near-edge structure (XANES) spectrum of the as-prepared $\text{Cu}_1/\text{C}_3\text{N}_4$ is illustrated in Fig. 2a, with Cu foil and

CuO as references. The adsorption-edge of $\text{Cu}_1/\text{C}_3\text{N}_4$ is located between those of Cu foil and CuO, showing that the valence of the Cu atom is between 0 and +2, which originates from the strong interaction between Cu and C_3N_4 . Furthermore, the high-resolution Cu 2p XPS spectrum of $\text{Cu}_1/\text{C}_3\text{N}_4$ (Fig. S11[†]) reveals the existence of only one kind of Cu species. The main peaks located at 932.2 eV and 952.1 eV are characteristic peaks of Cu^+ , consistent with the XANES analysis.⁶⁶ No Cu^0 signals were observed from the Cu 2p spectrum, indicating that no Cu particles or clusters exist. Fig. 2b shows the Fourier transformed (FT) extended X-ray absorption fine structure (EXAFS) spectra of $\text{Cu}_1/\text{C}_3\text{N}_4$ and the references. Only one prominent peak at 1.5 Å similar to that of CuPc was observed for $\text{Cu}_1/\text{C}_3\text{N}_4$, which was ascribed to the first coordination shell of Cu–N. No Cu–Cu coordination peak at 2.2 Å like for Cu foil was observed, suggesting that Cu species are atomically dispersed in $\text{Cu}_1/\text{C}_3\text{N}_4$. In order to further explore the atomic distribution of Cu in $\text{Cu}_1/\text{C}_3\text{N}_4$, wavelet transform (WT) analysis of Cu K-edge EXAFS was performed. As shown in Fig. 2c, only one intensity maximum at 3.3 \AA^{-1} related to Cu–N coordination was observed from the WT plots of $\text{Cu}_1/\text{C}_3\text{N}_4$, which is different from the WT plot of Cu foil



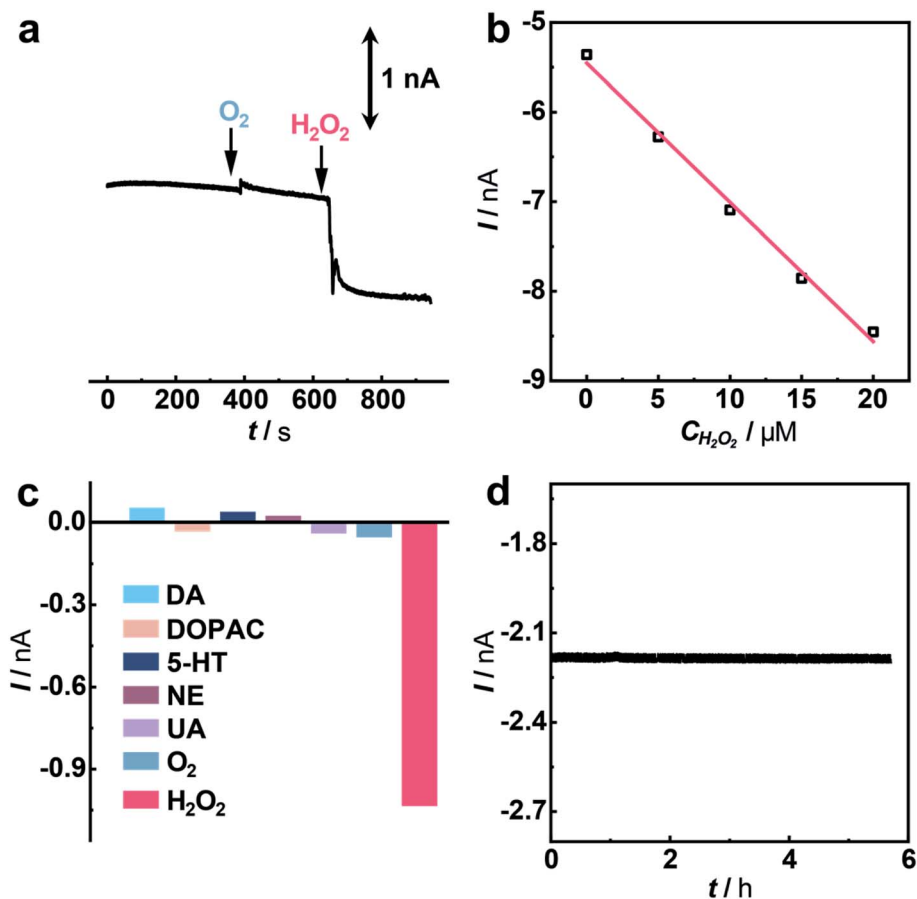


Fig. 5 Selective sensing of H₂O₂ with the CuSAC-based microsensor. (a) Amperometric responses recorded with the microsensor in aCSF toward the addition of 50 μM O₂ and 5 μM H₂O₂. Applied potential: 0.0 V vs. Ag/AgCl. (b) Calibration curve for amperometric H₂O₂ detection obtained at the microsensor in aCSF toward successive addition of 5 μM H₂O₂. Applied potential: 0.0 V vs. Ag/AgCl. (c) Amperometric responses recorded with the microsensor in aCSF toward 10 μM DA, 10 μM DOPAC, 10 μM 5-HT, 10 μM NE, 10 μM UA, 50 μM O₂ or 5 μM H₂O₂. Applied potential: 0.0 V vs. Ag/AgCl. (d) Long-time amperometric *i*-*t* curve recorded with the microsensor in aCSF toward 5 μM H₂O₂. Applied potential: 0.0 V vs. Ag/AgCl.

with one intensity maximum corresponding to the Cu-Cu contribution at 6.8 Å⁻¹, showing that no Cu-Cu bond is present in Cu₁/C₃N₄, but isolated Cu atoms exist. EXAFS fitting was carried out to obtain the atomic structure parameters of Cu in Cu₁/C₃N₄. According to the fitting curves (Fig. 2d and e, S12 and S13[†]) and the fitting parameters in Table S2,[†] the coordination number of Cu in Cu₁/C₃N₄ was calculated to be 2.2 and the mean bond length of Cu-N was 1.97 Å, suggesting that one Cu atom is coordinated by two N atoms to form a Cu-N₂ structure. The atomic structure model of Cu₁/C₃N₄ was constructed and is illustrated as the inset in Fig. 2e.

The electrocatalytic performance of C₃N₄ and Cu₁/C₃N₄ towards the HPRR in artificial cerebrospinal fluid (aCSF) was investigated by cyclic voltammetry (CV). In order to improve the electrical conductivity of Cu₁/C₃N₄ used for the HPRR, carbon nanotubes (CNTs) were incorporated with Cu₁/C₃N₄ because of the semiconducting properties of C₃N₄. Electrochemical impedance spectra (EIS) (Fig. S14[†]) reveal a much lower charge transfer resistance after mixing Cu₁/C₃N₄ with CNTs, which facilitates electron transfer to the active Cu sites. Fig. 3a and b show the CV curves of the Cu₁/C₃N₄/CNT catalyst (CuSAC)

towards the HPRR and ORR, respectively. Obviously, the CuSAC shows a high performance toward the HPRR with a high onset potential of ca. 0.20 V vs. Ag/AgCl, at which C₃N₄, CNTs and the C₃N₄/CNT composite exhibit no obvious current change (Fig. 3a and S15-S17[†]), implying that the single Cu atoms on the C₃N₄ substrate serve as active sites for the HPRR. In contrast, the ORR starts more negatively at ca. 0.0 V vs. Ag/AgCl (Fig. 3b). The activities of the catalyst toward the HPRR and ORR were further evaluated by linear sweep voltammetry (LSV) with comparable concentrations of H₂O₂ and O₂ (Fig. 3c). Similarly, the HPRR commences at ca. 0.20 V vs. Ag/AgCl (red curve), which is obviously more positive than the onset potential of the ORR (0.0 V vs. Ag/AgCl) (blue curve), again revealing that the HPRR occurs prior to and is thus discriminated from the ORR at the CuSAC-based electrode.

To further demonstrate the CuSAC-enabled priority of the HPRR over the ORR, potential-controlled amperometry was conducted with the CuSAC-based electrode. As displayed in Fig. 3d, the reduction current was observed after adding 5 μM H₂O₂ into the solution, while almost no current change was recorded with the addition of 50 μM O₂, indicating that H₂O₂ is



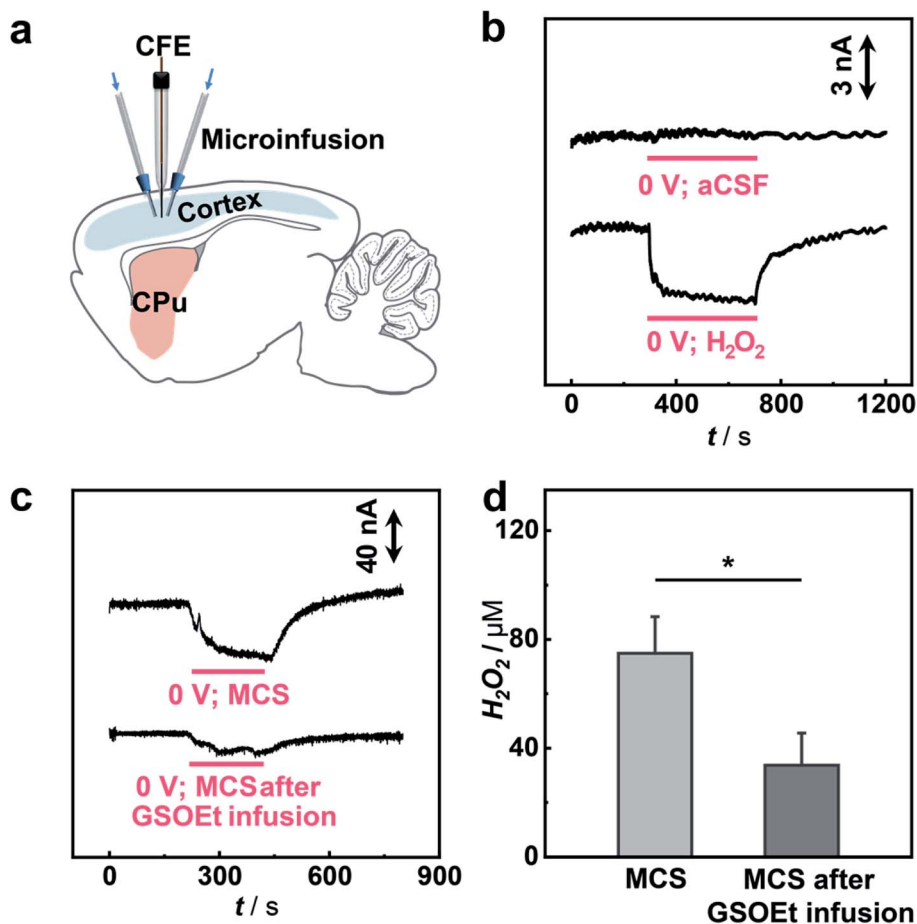


Fig. 6 *In vivo* sensing of H_2O_2 with the SAC-based microsensors. (a) Scheme of the sensing platform for *in vivo* monitoring of H_2O_2 . (b) Typical current responses at the microsensors implanted in the cortex of anesthetized guinea pigs upon local infusion (4 min, indicated by a red line) of aCSF without (top) and with (bottom) $100 \mu\text{M}$ H_2O_2 at 0.0 V vs. Ag/AgCl . (c) Typical current responses upon the local injection of 10 mM MCS (4 min, $1.0 \mu\text{L min}^{-1}$ indicated by a red line) without (top) and with (bottom) the pre-infusion of 10 mM GSOEt (15 min, $1.0 \mu\text{L min}^{-1}$). (d) Histogram of the H_2O_2 change calculated from (c). Error bars indicate the standard error of mean of 3 independent tests. Paired Student's *t*-test, *, $p < 0.05$.

preferentially reduced compared with O_2 . This finding is remarkable because, in almost all electrochemical systems, H_2O_2 reduction normally occurs at about the same or an even more negative potential than that of O_2 reduction, in spite of the more positive formal potential of the HPRR than that of the ORR in neutral pH. The use of SACs as the electrocatalyst successfully accelerates the HPRR and enables its occurrence prior to the ORR. Furthermore, after the HPRR test, the used $\text{Cu}_1/\text{C}_3\text{N}_4$ SAC retained the mesoporous structure (Fig. S18a†) with Cu, N, and C elements uniformly dispersed on the substrate (Fig. S18b†), without obvious clusters or nanoparticles observed in the high-resolution TEM images (Fig. S18c and S18d†), showing that the Cu species were still atomically dispersed. Meanwhile, the Cu 2p XPS spectrum of the used $\text{Cu}_1/\text{C}_3\text{N}_4$ also exhibits characteristic peaks of Cu^+ at 932.5 eV and 952.3 eV (Fig. S19†), verifying the negligible valence changes of Cu atoms. All the results demonstrate a high stability of $\text{Cu}_1/\text{C}_3\text{N}_4$.

To gain insights into the priority of the HPRR over the ORR, we carried out DFT calculations to investigate the reaction mechanism on the $\text{Cu}_1/\text{C}_3\text{N}_4$ SAC. Fig. 4a and b show the

calculated free energy diagrams of the two reactions on the $\text{Cu}_1/\text{C}_3\text{N}_4$ SAC, and the atomic configurations of the intermediates for the two reactions are illustrated in Fig. S20 and S21.† For H_2O_2 reduction, the H_2O_2^* dissociates into two OH^* spontaneously without electron transfer, which subsequently adsorb on the Cu atom and its adjacent C atom, respectively. The last OH^* desorption step on the C atom was regarded as the rate-determining step (RDS) owing to the endothermic nature with a free energy of 1.31 eV (Fig. 4a). For O_2 reduction, only OH^* desorption on the Cu atom is thermodynamically uphill with a free energy of 1.44 eV (Fig. 4b), which was determined to be the RDS. Due to the different adsorption sites of OH^* , the last OH^* desorption (*i.e.*, RDS) of the two reactions gives an energy difference of 0.13 eV , indicating that the HPRR occurs prior to the ORR. Therefore, the Cu– N_2 site and the surrounding C atom play important roles in intermediate adsorption and charge transfer for the HPRR. In contrast, only the Cu– N_2 site acts as an active center participating in the ORR. These results suggest that the synergism between the Cu– N_2 site and its adjacent C atom on C_3N_4 leads to a different pathway for the HPRR,



enabling priority of the HP RR over the ORR on the Cu₁/C₃N₄ SAC.

Having demonstrated the excellent electrocatalytic performance of the Cu₁/C₃N₄ SAC in the HP RR, especially over the ORR, we next applied this catalyst to develop a microsensor for *in vivo* selective sensing of H₂O₂ in a living rat brain. To do this, a carbon fiber electrode (CFE) with a diameter of 7 μm was prepared and used for fabrication of the microsensor.^{68,69} The microsensor exhibits high electrocatalytic performance towards H₂O₂ reduction (Fig. S22 and S23†). To evaluate the priority of the HP RR over the ORR, potential-controlled amperometry was performed with the microsensor toward H₂O₂ and O₂. As shown in Fig. 5a, there is negligible current change upon the addition of 50 μM O₂, while an obvious current decrease was observed for 5 μM H₂O₂, suggesting that H₂O₂ reduction occurs at a potential more positive than that of O₂ reduction. In addition, the microsensor responds well to H₂O₂ with a good linearity between current response and H₂O₂ concentration (I (nA) = $-5.454 - 0.155 C$ (μM), $R^2 = 0.992$) (Fig. 5b and S24†). Considering that there are other redox-active species coexisting in the cerebral system, interference from these neurochemicals was studied. Compared with H₂O₂, the species studied here give rise to negligible current responses (Fig. 5c and S25†), indicating high selectivity of the microsensor for the detection of H₂O₂. Moreover, the microsensor shows a high stability toward H₂O₂ detection over 5 h (Fig. 5d), which is beneficial for *in vivo* monitoring H₂O₂ in living animals.

To study the possibility of using the microsensor for *in vivo* sensing of H₂O₂ in a rat brain, the microsensor was implanted into the cortex of a guinea pig and the current response to the locally microinfused H₂O₂ was recorded (Fig. 6a). As illustrated in Fig. 6b, local infusion of 100 μM H₂O₂ solution into the rat cortex results in a fast amperometric response that can recover quickly after stopping the infusion, while no current response was observed with infusion of pure aCSF. These results show that the sensing platform with an SAC as the catalyst can be used for selective monitoring of H₂O₂ fluctuation *in vivo*.

Having shown the validity of our microsensor for *in vivo* sensing, we then performed quantitative analysis of H₂O₂-related biochemical processes induced by mercaptosuccinate (MCS) and glutathione monoethyl ester (GSOEt). As one kind of glutathione peroxidase inhibitor, MCS can increase the levels of H₂O₂ in the brain, while GSOEt can suppress H₂O₂ production with the generation of antioxidant glutathione.^{70,71} However, these biochemical processes still necessitate a quantitative study at a living animal level. To *in vivo* study the dynamics of H₂O₂ fluctuation in these biochemical processes induced by the two drugs in a living rat brain, we locally perfused MCS and GSOEt into the rat cortex and real-time tracked the level of H₂O₂ with the SAC-based microsensor developed here. As typically shown in Fig. 6c and d, the local microinjection of MCS into the cortex led to an increase in the current response of the microsensor ($n = 3$), corresponding to the increase of the H₂O₂ level in the brain. Quantitatively, microinjection of 10 mM MCS (4 μL) gives rise to an increase of *ca.* 75.0 ± 13.5 μM H₂O₂. In the other experiments, GSOEt was locally infused 30 min before MCS infusion and the current change induced by MCS was

recorded. We found that pre-infusion of GSOEt to some extent suppressed the increase of current for H₂O₂, indicating that the H₂O₂ production induced by MCS was inhibited. The pre-infusion of 10 mM GSOEt (15 μL) results in the decrease of H₂O₂ production induced by MCS to only *ca.* 33.7 ± 11.9 μM (Fig. 6c and d). These quantitative description of H₂O₂ dynamics in a living rat brain provides a direct basis for the study of biochemical processes associated with H₂O₂ chemistry.

Conclusions

In summary, we have discovered a single-atom Cu catalyst (Cu₁/C₃N₄) that gives priority to the electrochemical HP RR over the ORR and successfully established an O₂-interference-free electrochemical sensing platform for selective detection of H₂O₂ *in vivo*. The isolated Cu–N₂ site and surrounding C atom from the C₃N₄ support are responsible for the preferable HP RR over the ORR through synergistic adsorption of the intermediates during the reaction, enabling highly selective H₂O₂ detection. In addition, the microsensor with the Cu₁/C₃N₄ SAC responds quickly to H₂O₂ fluctuation *in vivo* and is successfully used for the quantitative study of H₂O₂-related biochemical processes. This study not only provides a new solution to the long-standing challenge in the electrochemical HP RR prior to the ORR, but also provides a novel sensing platform for understanding neurochemical processes by real-time monitoring of molecular events involved in physiological and pathological processes.

Data availability

All experimental and computational data is available in the ESI.†

Author contributions

L. M., J. M. and W. M. came up with the idea. X. G. conducted the sample preparation, physical characterization and electrochemical measurements. C. H. helped with the theoretical calculation and discussion. W. J. helped to conduct the animal experiment and discussion. W. C. helped with the analysis of XAS measurements. Z. C. and W. W. contributed to the synthesis experiment. P. Y. contributed to the discussion of the electrocatalysis part. X. G. and W. M. wrote the initial draft of this manuscript, and L. M. revised and finalized the manuscript. All the authors participated in the discussion of the results and commented on the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

All *in vivo* experimental procedures complied with the guidelines of the Animal Advisory Committee at the State Key Laboratory of Cognitive Neuroscience and Learning, and were approved by the Institutional Animal Care and Use Committee



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Notes and references

- B. Morgan, K. V. Laer, T. N. E. Owusu, D. Ezeriņa, D. Pastor-Flores, P. S. Amponsah, A. Tursch and T. P. Dick, *Nat. Chem. Biol.*, 2016, **12**, 437–443.
- G. S. Shadel and T. L. Horvath, *Cell*, 2015, **163**, 560–569.
- M. C. Sobotta, W. Liou, S. Stöcker, D. Talwar, M. Oehler, T. Ruppert, A. N. D. Scharf and T. P. Dick, *Nat. Chem. Biol.*, 2015, **11**, 64–70.
- D. Peralta, A. K. Bronowska, B. Morgan, É. Dóka, K. V. Laer, P. Nagy, F. Gräter and T. P. Dick, *Nat. Chem. Biol.*, 2015, **11**, 156–163.
- J. Noh, B. Kwon, E. Han, M. Park, W. Yang, W. Cho, W. Yoo, G. Khang and D. Lee, *Nat. Commun.*, 2015, **6**, 6907–6915.
- S. E. Weinberg and N. S. Chandel, *Nat. Chem. Biol.*, 2015, **11**, 9–15.
- B. C. Dickinson and C. J. Chang, *Nat. Chem. Biol.*, 2011, **7**, 504–511.
- A. J. Shuhendler, K. Pu, L. Cui, J. P. Uetrecht and J. Rao, *Nat. Biotechnol.*, 2014, **32**, 373–380.
- Y. Cheng, J. Dai, C. Sun, R. Liu, T. Zhai, X. Lou and F. Xia, *Angew. Chem., Int. Ed.*, 2018, **57**, 3123–3127.
- R. Wang, C. Yan, H. Zhang, Z. Guo and W.-H. Zhu, *Chem. Sci.*, 2020, **11**, 3371–3377.
- E. W. Miller, O. Tulyathan, E. Y. Isacoff and C. J. Chang, *Nat. Chem. Biol.*, 2007, **3**, 263–267.
- S. G. Rhee, *Science*, 2006, **312**, 1882–1883.
- A. K. Nath, A. Ghatak, A. Dey and S. G. Dey, *Chem. Sci.*, 2021, **12**, 1924–1929.
- Z. Wang, X. Ai, Z. Zhang, Y. Wang, X. Wu, R. Haindl, E. K. L. Yeow, W. Drexler, M. Gao and B. Xing, *Chem. Sci.*, 2020, **11**, 803–811.
- D. Pramanik, C. Ghosh and S. G. Dey, *J. Am. Chem. Soc.*, 2011, **133**, 15545–15552.
- S. Wu, R. Snajdrova, J. C. Moore, K. Baldenius and U. T. Bornscheuer, *Angew. Chem., Int. Ed.*, 2021, **60**, 88–119.
- C. Martin, M. Trajkovic and M. W. Fraaije, *Angew. Chem., Int. Ed.*, 2020, **59**, 4869–4872.
- S. Ye, N. Hananya, O. Green, H. Chen, A. Q. Zhao, J. Shen, D. Shabat and D. Yang, *Angew. Chem., Int. Ed.*, 2020, **59**, 14326–14330.
- W. Li, M. Khan, L. Lin, Q. Zhang, S. Feng, Z. Wu and J. Lin, *Angew. Chem., Int. Ed.*, 2020, **59**, 9282–9287.
- X.-W. Zhang, A. Oleinick, H. Jiang, Q.-L. Liao, Q.-F. Qiu, I. Svir, Y.-L. Liu, C. Amatore and W.-H. Huang, *Angew. Chem., Int. Ed.*, 2019, **58**, 7753–7756.
- H. Zhu, A. Sigdel, S. Zhang, D. Su, Z. Xi, Q. Li and S. Sun, *Angew. Chem., Int. Ed.*, 2014, **53**, 12508–12512.
- X. Xu, S. Jiang, Z. Hu and S. Liu, *ACS Nano*, 2010, **4**, 4292–4298.
- Y. Qu, Z. Li, W. Chen, Y. Lin, T. Yuan, Z. Yang, C. Zhao, J. Wang, C. Zhao, X. Wang, F. Zhou, Z. Zhuang, Y. Wu and Y. Li, *Nat. Catal.*, 2018, **1**, 781–786.
- C. H. Choi, W. S. Choi, O. Kasian, A. K. Mechler, M. T. Sougrati, S. Brüller, K. Strickland, Q. Jia, S. Mukerjee, K. J. J. Mayrhofer and F. Jaouen, *Angew. Chem., Int. Ed.*, 2017, **56**, 8809–8812.
- R. Li, X. Liu, W. Qiu and M. Zhang, *Anal. Chem.*, 2016, **88**, 7769–7776.
- Y. Lin, K. Liu, P. Yu, L. Xiang, X. Li and L. Mao, *Anal. Chem.*, 2007, **79**, 9577–9583.
- Z. Li, S. Ji, Y. Liu, X. Cao, S. Tian, Y. Chen, Z. Niu and Y. Li, *Chem. Rev.*, 2020, **120**, 623–682.
- L. Jiao, W. Xu, Y. Wu, H. Yan, W. Gu, D. Du, Y. Lin and C. Zhu, *Chem. Soc. Rev.*, 2021, **50**, 750–765.
- L. Li, X. Chang, X. Lin, Z.-J. Zhao and J. Gong, *Chem. Soc. Rev.*, 2020, **49**, 8156–8178.
- W. Ma, J. Mao, X. Yang, C. Pan, W. Chen, M. Wang, P. Yu, L. Mao and Y. Li, *Chem. Commun.*, 2019, **55**, 159–162.
- J. Zhang, C. Zheng, M. Zhang, Y. Qiu, Q. Xu, W.-C. Cheong, W. Chen, L. Zheng, L. Gu, Z. Hu, D. Wang and Y. Li, *Nano Res.*, 2020, **13**, 3082–3087.
- X. Liu, H. Pang, X. Liu, Q. Li, N. Zhang, L. Mao, M. Qiu, B. Hu, H. Yang and X. Wang, *The Innovation*, 2021, **2**, 100076.
- R. Lang, X. Du, Y. Huang, X. Jiang, Q. Zhang, Y. Guo, K. Liu, B. Qiao, A. Wang and T. Zhang, *Chem. Rev.*, 2020, **120**, 11986–12043.
- B. Qiao, A. Wang, X. Yang, L. F. Allard, Z. Jiang, Y. Cui, J. Liu, J. Li and T. Zhang, *Nat. Chem.*, 2011, **3**, 634–641.
- G. Kyriakou, M. B. Boucher, A. D. Jewell, E. A. Lewis, T. J. Lawton, A. E. Baber, H. L. Tierney, M. Flytzani-Stephanopoulos and E. C. H. Sykes, *Science*, 2012, **335**, 1209–1212.
- Y. Xiong, J. Dong, Z.-Q. Huang, P. Xin, W. Chen, Y. Wang, Z. Li, Z. Jin, W. Xing, Z. Zhuang, J. Ye, X. Wei, R. Cao, L. Gu, S. Sun, L. Zhuang, X. Chen, H. Yang, C. Chen, Q. Peng, C.-R. Chang, D. Wang and Y. Li, *Nat. Nanotechnol.*, 2020, **15**, 390–397.
- W. Zhu, L. Zhang, S. Liu, A. Li, X. Yuan, C. Hu, G. Zhang, W. Deng, K. Zang, J. Luo, Y. Zhu, M. Gu, Z.-J. Zhao and J. Gong, *Angew. Chem., Int. Ed.*, 2020, **59**, 12664–12668.
- D. Yang, H. Yu, T. He, S. Zuo, X. Liu, H. Yang, B. Ni, H. Li, L. Gu, D. Wang and X. Wang, *Nat. Commun.*, 2019, **10**, 3844.
- C. Fang and W. An, *Nano Res.*, 2021, **14**, 4211–4219.
- L. Jiao, R. Zhang, G. Wan, W. Yang, X. Wan, H. Zhou, J. Shui, S.-H. Yu and H.-L. Jiang, *Nat. Commun.*, 2020, **11**, 2831.
- H. Shang, Z. Jiang, D. Zhou, J. Pei, Y. Wang, J. Dong, X. Zheng, J. Zhang and W. Chen, *Chem. Sci.*, 2020, **11**, 5994–5999.
- W. Ma, F. Wu, P. Yu and L. Mao, *Chem. Sci.*, 2021, **12**, 7908–7917.



- 43 W. Wu, Y. Liu, D. Liu, W. Chen, Z. Song, X. Wang, Y. Zheng, N. Lu, C. Wang, J. Mao and Y. Li, *Nano Res.*, 2021, **14**, 998–1003.
- 44 W. Fu, Y. Wang, W. Tian, H. Zhang, J. Li, S. Wang and Y. Wang, *Angew. Chem., Int. Ed.*, 2020, **59**, 23791–23799.
- 45 J. Mao, C.-T. He, J. Pei, W. Chen, D. He, Y. He, Z. Zhuang, C. Chen, Q. Peng, D. Wang and Y. Li, *Nat. Commun.*, 2018, **9**, 4958–4965.
- 46 C. Zhou, J. Y. Zhao, P. F. Liu, J. Chen, S. Dai, H. G. Yang, P. Hu and H. Wang, *Chem. Sci.*, 2021, **12**, 10634–10642.
- 47 H. Fei, J. Dong, Y. Feng, C. S. Allen, C. Wan, B. Voloskiy, M. Li, Z. Zhao, Y. Wang, H. Sun, P. An, W. Chen, Z. Guo, C. Lee, D. Chen, I. Shakir, M. Liu, T. Hu, Y. Li, A. I. Kirkland, X. Duan and Y. Huang, *Nat. Catal.*, 2018, **1**, 63–72.
- 48 P. Li, M. Wang, X. Duan, L. Zheng, X. Cheng, Y. Zhang, Y. Kuang, Y. Li, Q. Ma, Z. Feng, W. Liu and X. Sun, *Nat. Commun.*, 2019, **10**, 1711–1721.
- 49 K. Kamiya, *Chem. Sci.*, 2020, **11**, 8339–8349.
- 50 J. Gu, C.-S. Hsu, L. Bai, H. M. Chen and X. Hu, *Science*, 2019, **364**, 1091–1094.
- 51 H. B. Yang, S.-F. Hung, S. Liu, K. Yuan, S. Miao, L. Zhang, X. Huang, H.-Y. Wang, W. Cai, R. Chen, J. Gao, X. Yang, W. Chen, Y. Huang, H. M. Chen, C. M. Li, T. Zhang and B. Liu, *Nat. Energy*, 2018, **3**, 140–147.
- 52 J. Zhang, W. Cai, F. X. Hu, H. Yang and B. Liu, *Chem. Sci.*, 2021, **12**, 6800–6819.
- 53 Z. Wang, C. Wang, Y. Hu, S. Yang, J. Yang, W. Chen, H. Zhou, F. Zhou, L. Wang, J. Du, Y. Li and Y. Wu, *Nano Res.*, 2021, **14**, 2790–2796.
- 54 Z. Li, R. Wu, L. Zhao, P. Li, X. Wei, J. Wang, J. S. Chen and T. Zhang, *Nano Res.*, 2021, **14**, 3795–3809.
- 55 M. Zhou, Y. Jiang, G. Wang, W. Wu, W. Chen, P. Yu, Y. Lin, J. Mao and L. Mao, *Nat. Commun.*, 2020, **11**, 3188–3196.
- 56 W. Wu, L. Huang, E. Wang and S. Dong, *Chem. Sci.*, 2020, **11**, 9741–9756.
- 57 Z. Guo, Y. Xie, J. Xiao, Z.-J. Zhao, Y. Wang, Z. Xu, Y. Zhang, L. Yin, H. Cao and J. Gong, *J. Am. Chem. Soc.*, 2019, **141**, 12005–12010.
- 58 F. Luo, A. Roy, L. Silvioli, D. A. Cullen, A. Zitolo, M. T. Sougrati, I. C. Oguz, T. Mineva, D. Teschner, S. Wagner, J. Wen, F. Dionigi, U. I. Kramm, J. Rossmeisl, F. Jaouen and P. Strasser, *Nat. Mater.*, 2020, **19**, 1215–1223.
- 59 L. Jiao, W. Yang, G. Wan, R. Zhang, X. Zheng, H. Zhou, S.-H. Yu and H.-L. Jiang, *Angew. Chem., Int. Ed.*, 2020, **59**, 20589–20595.
- 60 X. Li, H. Rong, J. Zhang, D. Wang and Y. Li, *Nano Res.*, 2020, **13**, 1842–1855.
- 61 H. Jing, Z. Zhao, C. Zhang, W. Liu, D. Wu, C. Zhu, C. Hao, J. Zhang and Y. Shi, *Nano Res.*, 2021, **14**, 4025–4032.
- 62 F. Wu, C. Pan, C.-T. He, Y. Han, W. Ma, H. Wei, W. Ji, W. Chen, J. Mao, P. Yu, D. Wang, L. Mao and Y. Li, *J. Am. Chem. Soc.*, 2020, **142**, 16861–16867.
- 63 Y. Wang, G. Jia, X. Cui, X. Zhao, Q. Zhang, L. Gu, L. Zheng, L.-H. Li, Q. Wu, D. J. Singh, D. Matsumura, T. Tsuji, Y.-T. Cui, J. Zhao and W. Zheng, *Chem*, 2021, **7**, 1–14.
- 64 X. Chen, L. Zhao, K. Wu, H. Yang, Q. Zhou, Y. Xu, Y. Zheng, Y. Shen, S. Liu and Y. Zhang, *Chem. Sci.*, 2021, **12**, 8865–8871.
- 65 A. Savateev, N. V. Tarakina, V. Strauss, T. Hussain, K. Brummelhuis, J. M. S. Vadiello, Y. Markushyna, S. Mazzanti, A. P. Tyutyunnik, R. Walczak, M. Oschatz, D. M. Guldi, A. Karton and M. Antonietti, *Angew. Chem., Int. Ed.*, 2020, **59**, 15061–15068.
- 66 A. Bakandritsos, R. G. Kadam, P. Kumar, G. Zoppellaro, M. Medved', J. Tuček, T. Montini, O. Tomanec, P. Andryšková, B. Drahoš, R. S. Varma, M. Otyepka, M. B. Gawande, P. Fornasiero and R. Zboril, *Adv. Mater.*, 2019, **31**, 1900323.
- 67 S. Tian, Z. Wang, W. Gong, W. Chen, Q. Feng, Q. Xu, C. Chen, C. Chen, Q. Peng, L. Gu, H. Zhao, P. Hu, D. Wang and Y. Li, *J. Am. Chem. Soc.*, 2018, **140**, 11161–11164.
- 68 H. Hou, Y. Jin, H. Wei, W. Ji, Y. Xue, J. Hu, M. Zhang, Y. Jiang and L. Mao, *Angew. Chem., Int. Ed.*, 2020, **59**, 18996–19000.
- 69 J. Jin, W. Ji, L. Li, G. Zhao, W. Wu, H. Wei, F. Ma, Y. Jiang and L. Mao, *J. Am. Chem. Soc.*, 2020, **142**, 19012–19016.
- 70 M. V. Avshalumov and M. E. Rice, *Proc. Natl. Acad. Sci. U. S. A.*, 2003, **100**, 11729–11734.
- 71 V. P. Wellner, M. E. Anderson, R. N. Puri, G. L. Jensen and A. Meister, *Proc. Natl. Acad. Sci. U. S. A.*, 1984, **81**, 4732–4735.

