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# Annealing activated nickel–molybdenum oxide as an efficient electrocatalyst toward benzyl alcohol upgrading†

Shunda Hu,<sup>a</sup> Xiaoning Sun,<sup>b</sup> Zimeng Liu,<sup>b</sup> Lingfeng Gao,<sup>\*a</sup> Xiaoli Li,<sup>a</sup> Chunyu Yu,<sup>a</sup> Xiangkang Han,<sup>a</sup> Junfeng Xie<sup>ib</sup>\*<sup>b</sup> and Xu Sun<sup>\*a</sup>

Efficient coupling of economically favorable electro-oxidation reactions with the hydrogen evolution reaction (HER) has been considered as a promising way to realize synergistic production of hydrogen and value-added chemicals. In this work, a supported nickel molybdenum oxide catalyst was fabricated, which exhibits enhanced activity towards the benzyl alcohol oxidation reaction (BOR) benefited from the enriched active sites. Further investigations indicate that the nearly complete conversion from benzyl alcohol (BA) to the benzoic acid (BC) product can be achieved with simultaneously realized high selectivity and high faradaic efficiency (FE) for long-term operation. The efficient catalyst explored in this work could offer a new material platform for coupled production and value-added benzoic acid and hydrogen.

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## Introduction

With the concern of the global energy crisis as well as the depletion of fossil fuel reserves, there is an urgent need to explore clean and sustainable alternative energy sources. Hydrogen (H<sub>2</sub>), as a carbon-free and sustainable energy carrier, holds immense potential to alleviate the dependence on fossil fuels and related environmental concerns.<sup>1,2</sup> The electrocatalytic water splitting process that is composed of the hydrogen evolution reaction (HER) and oxygen evolution reaction (OER) is now a viable method for hydrogen production, with significant progress being made to improve the reaction efficiency.<sup>3–9</sup> However, the high overpotential and the sluggish kinetics of the OER process at the anode still impede the whole reaction efficiency for hydrogen production. Hence, optimizing the coupled anodic reaction is of great significance to promote the hydrogen production process.<sup>10–16</sup>

As discussed above, replacing the OER process with thermodynamically favorable electro-oxidation reactions of small molecular chemicals, such as methanol, ethanol, hydrazine and urea,<sup>17–28</sup> has been regarded as an advanced way of realizing efficient hydrogen production and even simultaneous valorization to produce value-added chemicals.<sup>29,30</sup> As a widely used chemical in the pharmaceutical, dye and fragrance industries, benzoic acid (BC) receives substantial demand, and can be produced from the oxidation of benzyl alcohol (BA).<sup>31</sup> However, the current synthesis of BC from BA requires harsh conditions such as the use of harmful reagents, high temperature and noble metal catalysts.<sup>32</sup> Therefore, developing a green and economic method for BC production is highly attractive and urgently desired.<sup>33,34</sup> With the above considerations, developing an advanced catalyst that could catalyze the BA-to-BC conversion will be of great significance to realize simultaneous production of hydrogen and value-added BC.<sup>35,36</sup>

Despite the optimization of the oxidation reaction, the design and selection of the appropriate electrocatalyst is also an important factor that would influence the whole reaction efficiency. Recently, transition metal-based compounds have emerged as the preferred choice due to their low cost, high activity, excellent stability, and abundant reserves.<sup>37–39</sup> In particular, the synergistic effect between different cationic sites in multi-metal oxides has been extensively studied, which could lead to optimized electronic structures and multi-site active centers.<sup>40–46</sup> Among the transition metal oxide catalysts, the Ni–Mo based oxides, have received substantial research interest owing to their abundant high-valence active sites with favorable electronic structure, which have

<sup>a</sup> Key Laboratory of Interfacial Reaction & Sensing Analysis in Universities of Shandong, School of Chemistry and Chemical Engineering Institution, University of Jinan, Jinan, Shandong, 250022, P. R. China. E-mail: chm\_gaolf@ujn.edu.cn, chm\_sunx@ujn.edu.cn

<sup>b</sup> College of Chemistry, Chemical Engineering and Materials Science, Key Laboratory of Molecular and Nano Probes (Ministry of Education), Collaborative Innovation Center of Functionalized Probes for Chemical Imaging in Universities of Shandong, Institute of Molecular and Nano Science, Shandong Normal University, Jinan, Shandong, 250014, P. R. China. E-mail: xiejf@sdsnu.edu.cn

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been applied as highly efficient electrocatalysts for the OER and the urea oxidation reaction (UOR).<sup>47</sup> For example, Solomon and co-workers fabricated nickel–molybdenum oxide microcrystals covered by  $\text{Co}_3\text{O}_4$  nanoparticles *via* atomic layer deposition, which exhibit high activity towards the OER.<sup>48</sup> Fang and co-workers proposed an electrochemical tuning strategy to fabricate  $\text{Ni}^{3+}$ -rich  $\text{NiMoO}_4 \cdot x\text{H}_2\text{O}$  nanorod arrays, and realized electrochemical tuning that effectively enriches the content of  $\text{Ni}^{3+}$ , finally resulting in the remarkable enhancement of the UOR activity.<sup>49</sup> However, the application of Ni–Mo oxides for organic molecule oxidation is still rare to date.<sup>50,51</sup> Developing the potential application of Ni–Mo based catalysts for the electrocatalytic oxidation of organic molecules is highly desired.<sup>52</sup> In this work, a Ni–Mo oxide catalyst supported on nickel foam ( $\text{NiMoO-Ar/NF}$ ) was successfully fabricated, which exhibits advanced catalytic activity for the benzyl alcohol oxidation reaction. The as-exploded  $\text{NiMoO-Ar/NF}$  catalyst exhibits a low potential of 1.39 V *vs.* RHE to reach a current density of  $10 \text{ mA cm}^{-2}$  in 1 M KOH solution with 15 mM BA. Gas chromatography results indicate that nearly complete conversion from BA to BC can be achieved after electrolysis at 1.5 V *vs.* RHE for 2 h, and the faradaic efficiency (FE) and yield of the product reach 93% and 95%, respectively. Meanwhile, excellent stability can be achieved even for 4-cycle chronoamperometric tests with a total operation time of 8 hours. This research developed a highly efficient electrocatalyst for coupled green synthesis of value-added benzoic acid and hydrogen.

## Results and discussion

The  $\text{NiMoO-Ar/NF}$  catalyst was synthesized *via* a two-step reaction (Fig. 1), and the experimental details are provided in the ESI.† Typically, the  $\text{NiMoO/NF}$  precursor was firstly fabricated *via* a hydrothermal reaction, followed by annealing in an Ar atmosphere, which finally leads to the formation of  $\text{NiMoO-Ar/NF}$ . To analyze the crystal structure of the product, the X-ray diffraction (XRD) pattern of the NF-supported  $\text{NiMoO}$  after calcination in Ar was provided in Fig. 2A, and the XRD pattern of the  $\text{NiMoO/NF}$  precursor without calcination was provided as well for comparison. As presented by the purple line, all the peaks can be indexed to the  $\text{NiMoO}_4 \cdot x\text{H}_2\text{O}$  phase as reported in the literature.<sup>48</sup> In contrast, a series of peaks appeared at  $14.2^\circ$ ,  $25.3^\circ$ ,  $28.8^\circ$  and  $32.5^\circ$  for  $\text{NiMoO-Ar/NF}$ , which match well with the standard card of  $\text{NiMoO}_4$  (JCPDS card no. 33-0948), proving

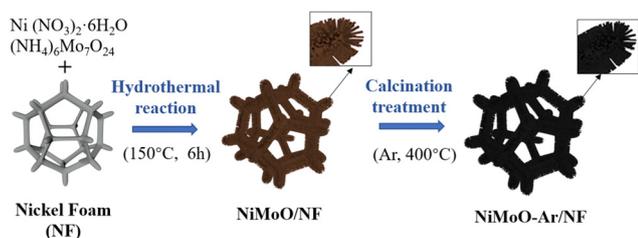


Fig. 1 Schematic illustration of the synthesis process of the  $\text{NiMoO-Ar/NF}$  catalyst.

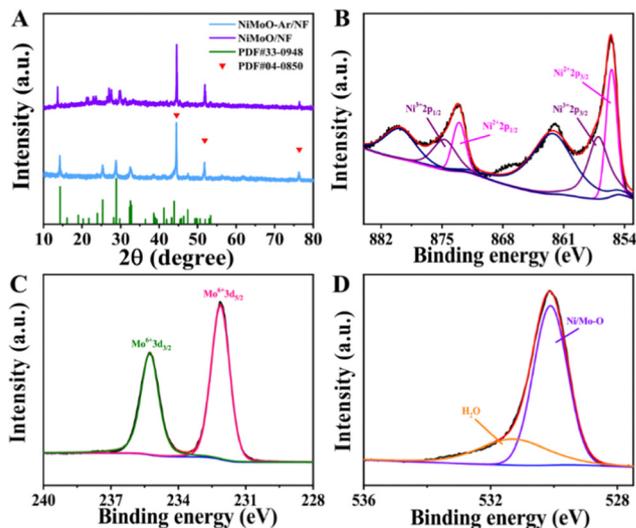


Fig. 2 (A) XRD patterns of  $\text{NiMoO-Ar/NF}$  and  $\text{NiMoO/NF}$ . (B) Ni 2p spectra. (C) Mo 3d spectra. (D) O 1s spectra.

the removal of crystal water in the  $\text{NiMoO}_4 \cdot x\text{H}_2\text{O}$  precursor. The peaks located at  $44.5^\circ$ ,  $51.8^\circ$  and  $76.3^\circ$  correspond to metallic nickel of the NF support (JCPDS card no. 04-0850). As reported in previous literature, the removing of crystal water is beneficial for the enhancement of the catalytic activity since crystal water may occupy the active sites and block the catalytic reaction.<sup>53,54</sup> Hence, enhanced catalytic activity can be expected for the  $\text{NiMoO}_4$ -based catalyst without crystal water.

To further analyze the chemical composition and valence states of the as-synthesized  $\text{NiMoO-Ar/NF}$  catalyst, X-ray photoelectron spectroscopy (XPS) was conducted. As shown in the survey spectrum, Fig. S1 (ESI†), the characteristic peaks corresponding to Ni, Mo and O elements can be observed, confirming the presence of these elements in the catalyst. As shown in the high-resolution Ni 2p spectrum in Fig. 2B, two intense peaks can be identified at 873.0 and 855.4 eV, which can be indexed to the  $2p_{1/2}$  and  $2p_{3/2}$  peaks of  $\text{Ni}^{2+}$  ions, respectively. Meanwhile, the peaks corresponding to  $\text{Ni}^{3+}$  emerge at 874.8 eV and 856.9 eV, confirming the coexistence of  $\text{Ni}^{2+}$  and  $\text{Ni}^{3+}$  in the  $\text{NiMoO-Ar/NF}$  catalyst.<sup>45,55</sup> The Mo 3d spectrum was provided as Fig. 2C, from which two peaks corresponding to  $\text{Mo}^{6+}$  species can be identified at 232.1 and 235.2 eV, respectively.<sup>55,56</sup> As reported in previous literature, the presence of Mo species with high oxidation valence can significantly influence the electronic structure of neighboring metals, which might bring about the variation of the catalytic performance.<sup>48,55</sup> Considering the above fact, the electro-oxidation performance can be expected for  $\text{NiMoO-Ar/NF}$ . In addition, the O 1s spectrum in Fig. 2D exhibits two peaks centered at 530.0 and 531.3 eV, assigned to the metal–oxygen bonds and the adsorbed water molecules, respectively.<sup>8,57</sup> Furthermore, considering the existence of  $[\text{MoO}_4]^{2-}$  oxyanion counterparts, which are vibrationally active, both the Raman and Fourier transform infrared (FTIR) spectra of  $\text{NiMoO-Ar/NF}$  and  $\text{NiMoO/NF}$  were provided for better structure analysis, as shown in Fig. S3A and B (ESI†).



As shown in the FTIR spectral in Fig. S3A (ESI<sup>†</sup>), it can be seen that an obvious peak appeared at  $3400.3\text{ cm}^{-1}$ , which should be attributed to the O–H stretching of water molecules.<sup>48</sup> Besides, the peak corresponding to the absorbed water molecules was observed as well at  $1621.8\text{ cm}^{-1}$ , which is well consistent with the previous literature.<sup>58,59</sup> This confirms the existence of water in the NiMoO/NF structure. Besides, a serious peak appeared at about  $449.8$  to  $964.7\text{ cm}^{-1}$ , which should be attributed to the symmetric and asymmetric stretching of Mo–O bonds.<sup>48,60</sup> As for the NiMoO-Ar/NF, similar peaks were observed as well at  $455.6\text{ cm}^{-1}$  to  $960.8\text{ cm}^{-1}$ . Meanwhile, the two peaks that were indexed to the water molecules were decreased in the NiMoO-Ar/NF catalyst, proving the removal of water molecules. From the Raman spectra in Fig. S3B (ESI<sup>†</sup>), as for the NiMoO/NF sample, an obvious peak appeared at about  $942.8\text{ cm}^{-1}$ , which should be attributed to the symmetric stretching of the Mo–O bands. Besides, two peaks that were indexed to the asymmetric stretching of O–Mo–O appeared as well at  $891$  and  $818.7\text{ cm}^{-1}$ , respectively. Furthermore, the band corresponding to the bending mode appeared as well at the low energy area of about  $300$ – $400\text{ cm}^{-1}$ . These values were well consistent with the previous literature, confirming the successful fabrication of NiMoO<sub>4</sub>·xH<sub>2</sub>O.<sup>58–60</sup> As for the NiMoO-Ar/NF obtained after annealing treatment, similar peaks were observed as well at  $952.8$ ,  $892$ ,  $817$  and  $342\text{ cm}^{-1}$ , confirming the well-kept Mo–O and O–Mo–O structure. Furthermore, a new peak corresponding to NiMoO<sub>4</sub> appeared at  $706\text{ cm}^{-1}$ , which is in agreement with the annealed NiMoO<sub>4</sub> sample as reported in the literature.<sup>61</sup> The peak at  $952.8\text{ cm}^{-1}$  corresponds to the symmetric stretching mode of the Mo–O bond, while the peaks at  $706$  and  $342\text{ cm}^{-1}$  correspond to the asymmetric stretching mode of the Ni–O–Mo bond and the bending mode of Mo–O, respectively.<sup>62</sup>

Scanning electron microscopy (SEM) and transmission electron microscopy (TEM) images are provided in Fig. 3. As shown in the SEM images in Fig. 3A, a thick oxide layer comprised from nanorods can be observed on the surface of the Ni foam. The magnified SEM image was provided in Fig. 3B, from which the square-shaped cross section of the nanorods can be observed. Meanwhile, the TEM images of the powdery sample detached from NiMoO-Ar/NF are shown in Fig. 3C and D, further proving the nanorod morphology with a width of about  $500\text{ nm}$ , which is consistent with the SEM result. To figure out the crystal structure, a high-resolution TEM (HRTEM) image is provided in Fig. 3E. As can be seen, uniform lattice fringes with an interplanar spacing of  $0.619\text{ nm}$  can be identified, which match well with the (110) plane of the NiMoO<sub>4</sub> phase. In addition, the elemental distribution based on the energy-dispersive X-ray spectroscopy (EDS) technique is shown in Fig. 3F. It can be observed that Ni (green), Mo (red), and O (blue) are homogeneously distributed across the nanorods. In addition, Fig. S4 (ESI<sup>†</sup>) illustrates the SEM of NiMoO/NF.

The electrochemical measurements were conducted by using a typical three-electrode system. The linear sweep voltammetry (LSV) curves of NiMoO-Ar/NF were firstly analyzed to unravel the different behavior between the OER and BOR. Typically, the

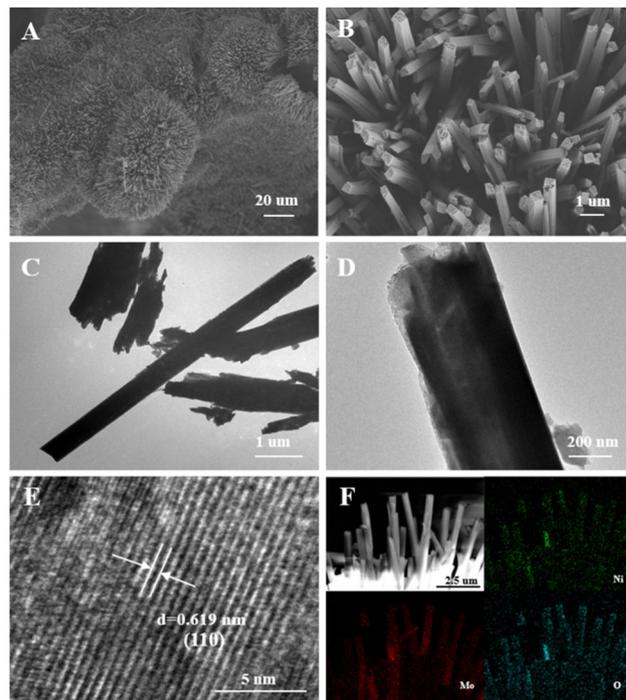


Fig. 3 (A) SEM images of NiMoO-Ar/NF. (B) Magnified SEM images. (C) and (D) TEM images of NiMoO-Ar/NF. (E) HRTEM images of NiMoO-Ar/NF. (F) Element mapping of Ni, Mo, and O in NiMoO-Ar/NF.

polarization curves for BOR and OER were obtained from  $1\text{ M KOH}$  electrolyte with and without  $15\text{ mM}$  benzyl alcohol. As can be seen from Fig. S6 (ESI<sup>†</sup>), the required potential for BOR to drive a  $10\text{ mA cm}^{-2}$  anodic current density is only  $1.39\text{ V vs. RHE}$ . In contrast, the required potential for the OER to reach the same current density is  $1.49\text{ V vs. RHE}$ .

Furthermore, compared with the OER process that only generates oxygen, the electro-oxidation of benzyl alcohol may achieve the production of the value-added benzoic acid, which is considered to be a more promising pathway for coupling with the HER half reaction to reach energy-saving production. To investigate the BOR performance of the catalysts, the LSV curve comparison of NiMoO-Ar/NF, NiMoO/NF and calcined NF for BOR was provided in Fig. 4A. It can be seen that as for the NiMoO-Ar/NF catalyst, the current density reaches  $10.72\text{ mA cm}^{-2}$  at  $1.4\text{ V vs. RHE}$ . Meanwhile, for NiMoO/NF and NF, the current density was only about  $0.266$  and  $1.452\text{ mA cm}^{-2}$ , respectively. Furthermore, the potentials at the fixed current density ( $20\text{ mA cm}^{-2}$ ) were compared as well. This illustrated that the NiMoO-Ar/NF catalyst only needed  $399\text{ mV}$  to reach  $20\text{ mA cm}^{-2}$ . The potentials for NiMoO/NF and NF to reach the same current density were  $563$  and  $747\text{ mV}$ , respectively, much larger than that of NiMoO-Ar/NF catalyst. Additionally, as shown in Fig. 4B, the Tafel plots also indicate the excellent activity of the NiMoO-Ar/NF catalyst. It can be seen that the Tafel slope of NiMoO-Ar/NF is as small as  $42\text{ mV dec}^{-1}$ , significantly lower than that of the NiMoO/NF catalyst ( $55\text{ mV dec}^{-1}$ ) and the NF-Ar catalyst ( $82\text{ mV dec}^{-1}$ ), further demonstrating the enhanced BOR performance of NiMoO-Ar/NF.



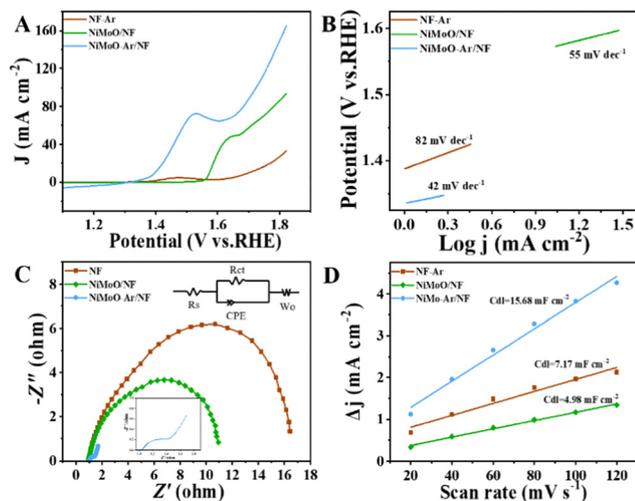


Fig. 4 (A) Polarization curve comparison of NF-Ar, NiMoO/NF, and NiMoO-Ar/NF in electrolyte containing 15 mM BA. (B) Tafel curves corresponding to figure A. (C) Comparison of the impedance of NF-Ar, NiMoO/NF and NiMoO-Ar/NF in electrolyte containing 15 mM BA. (D)  $C_{dl}$  curves of NF-Ar, NiMoO/NF and NiMoO-Ar/NF in an electrolyte containing 15 mM BA.

To further analyze the catalytic transfer kinetics of the NiMoO-Ar/NF catalyst, electrochemical impedance spectroscopy (EIS) analysis was performed at 1.5 V vs. RHE, with the corresponding results being shown in Fig. 4C. It reveals that the charge transfer resistance ( $R_{ct}$ ) of the NiMoO-Ar/NF catalyst is about 1.45  $\Omega$ , much smaller than that of NiMoO/NF (10.42  $\Omega$ ) and NF-Ar (16.98  $\Omega$ ), indicating the promoted charge transfer process during the BOR catalysis. According to the above analyses, it is concluded that the annealing treatment is beneficial for the enhancement of the catalytic activity. This might be owing to the following two reasons. Firstly, the annealing treatment could lead to the removal of crystal water within the  $\text{NiMoO}_4 \cdot x\text{H}_2\text{O}$  lattice, thus increasing the exposure of active sites, which finally results in enhanced catalytic activity. Secondly, the calcination treatment may induce the formation of a porous structure, thereby further increasing the surface area and enriching the catalytic activity. To prove the increment of the surface area, the nitrogen adsorption-desorption isotherms of NiMoO-Ar/NF and NiMoO/NF were measured. As revealed in Fig. S5 (ESI<sup>†</sup>), NiMoO-Ar/NF displays a larger BET specific surface area compared with that of NiMoO/NF, therefore confirming the enhancement of the surface area owing to the calcination. In addition, the electrochemical active surface area (ECSA) that is linearly proportional to the double-layer capacitance ( $C_{dl}$ ) is estimated as presented in Fig. 4D and Fig. S7 (ESI<sup>†</sup>). The results showed that the NiMoO-Ar/NF holds the largest  $C_{dl}$  value of 15.68  $\text{mF cm}^{-2}$ . Meanwhile, for the NiMoO/NF catalyst and NF-Ar catalyst, the  $C_{dl}$  values were just 4.98  $\text{mF cm}^{-2}$  and 7.17  $\text{mF cm}^{-2}$ , respectively. The much larger  $C_{dl}$  value of the NiMoO-Ar/NF catalyst confirmed its larger specific surface area, which is essential for increasing the exposure of active sites and thus improving the catalytic activity.

Based on the above analysis, it is concluded that the NiMoO-Ar/NF is an ideal catalyst for BOR. To further investigate the oxidation products of BOR, chronoamperometry measurement

was performed at 1.5 V vs. RHE with the gas chromatography (GC) results, as shown in Fig. S10 (ESI<sup>†</sup>). As can be seen from Fig. S10A (ESI<sup>†</sup>), before the reaction, an intensive peak that corresponds to BA was observed at about 10.2 min. After the reaction for half an hour, a new peak located at approximately 12.2 min emerges, corresponding to BC, as shown in Fig. S10B (ESI<sup>†</sup>). The formation of the BC product indicates the occurrence of the electro-oxidation of benzyl alcohol. Besides, another new peak that was attributed to benzaldehyde (BH) can be observed at 9.0 min, which was brought by the side reaction. It is noteworthy that the peak area of BC is larger than that of BH as presented in Table S2 (ESI<sup>†</sup>), suggesting that the side reaction is slight. Furthermore, as illustrated in Fig. S10C and D (ESI<sup>†</sup>), along with the increment of the reaction time to 1 h and 1.5 h, the peak for BC was further increased, indicating that more BA was converted to the product. Then, after the reaction for 2 h (Fig. S10E, ESI<sup>†</sup>), the peak for BA nearly disappeared, while the BC peak is dominant, indicating the high conversion efficiency from BA to BC. The concentration of BC was calculated according to the GC results (Fig. 5A). It can be seen that along with the increment of the reaction time, the concentration of BC increased, achieving a high value of 14.21 mM after reaction for 2 h. Hence, the high activity of NiMoO-Ar/NF catalyst towards the BOR can be confirmed.

To survey the influence of applied potential, chronoamperometry measurements at various potentials were conducted, and the corresponding GC results are provided in Fig. 5B and Fig. S11, Table S3 (ESI<sup>†</sup>). It can be seen that high selectivity (Sel.) can be achieved at various potentials, proving a wide potential window for high-performance BOR. In contrast, the conversion percentage (Con.), yield and faradaic efficiency (FE) display variation along with the change of potential, and the optimal performance is determined to be 1.5 V vs. RHE. In detail, the Con. of BA reaches 94.27%, and the Sel. to BC and FE are as high as 100% and 91.25%. When the applied potential

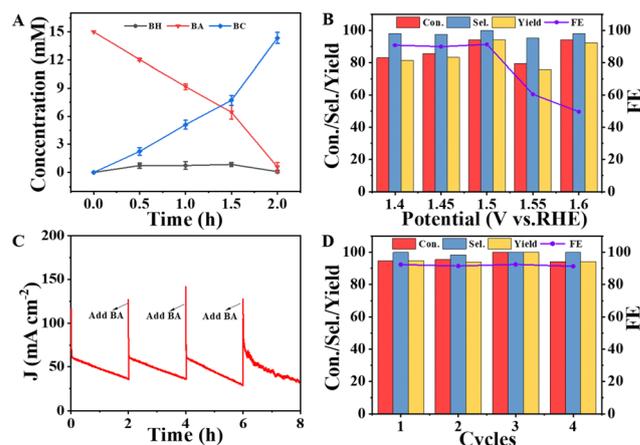


Fig. 5 (A) Evolution of the concentration of BA, BC and BH under various reaction times at 1.5 V vs. RHE. (B) Con., Sel., yield and FE at different potentials. (C) The chronoamperometry tests were repeated four times. (D) Con., Sel., yield and FE of BC in four electrochemical cycles.



further increases, the competitive OER process becomes obvious, and the BOR efficiency declines.

Operational stability is another important criterion to judge an advanced catalyst. We further performed cyclic voltammetry (CV) cycling tests and compared the polarization curves before and after 1000 CV cycles. The electrolyte for the CV stability test was 1 M KOH + 15 mM BA. As shown in Fig. S12 (ESI<sup>†</sup>), the anodic current density shows negligible degradation, confirming the high stability of the NiMoO-Ar/NF catalyst. In addition, a long-term chronoamperometry test was also performed, and the electrolyte was refreshed per 2 h to reduce the influence of the concentration of BA to the activity. As can be seen from Fig. 5C, the current density decreases along with the BOR catalysis, and the current density can be well recovered after refreshing the electrolyte, indicating that the decreasing current density originates from the efficient BOR process that rapidly consumes BA. The conversion percentage, selectivity, yield and FE were also measured after 4 cycles of 2 hour BOR operation. As shown in Fig. 5D, all the parameters exhibit negligible decay, and the FE still reaches ~93%, further confirming the superior operational stability of NiMoO-Ar/NF towards the BOR. The high activity and stability of NiMoO-Ar/NF for BOR catalysis make the catalyst a promising catalyst for the green production of value-added benzoic acid.

## Conclusions

In summary, a NiMoO-Ar/NF nanorod array catalyst was synthesized for an electrocatalytic benzyl alcohol oxidation reaction to produce benzoic acid. By means of annealing treatment, the crystal water within the lattice of the hydrated Ni-Mo oxide precursor can be removed, leading to highly exposed active sites and enhanced catalytic activity. Meanwhile, along with the calcination process, porous structure with increased surface area can be obtained, thereby further facilitating mass transport and enriching the active sites. Benefited from the above structural merits, the NiMoO-Ar/NF catalyst exhibited excellent electrocatalytic activity for BOR catalysis with a low required potential of 1.39 V *vs.* RHE to achieve a 10 mA cm<sup>-2</sup> current density. In addition, nearly complete conversion from BA to BC can be realized after 2-hour electrolysis at 1.5 V *vs.* RHE with high faradaic efficiency and yield of product. Superior operational stability can also be achieved even for 4-cycle chronoamperometric tests with only slight performance decay. This work presents a new electrocatalyst for green electrochemical synthesis of benzoic acid, which may shed light on the future design of advanced catalysts for the electro-oxidation of organic molecules.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 Y. Li, X. Wei, L. Chen and J. Shi, *Angew. Chem., Int. Ed.*, 2021, **60**, 19550–19571.
- 2 O. J. Guerra, J. Eichman, J. Kurtz and B.-M. Hodge, *Joule*, 2019, **3**, 2425–2443.
- 3 H. Sun, Z. Yan, F. Liu, W. Xu, F. Cheng and J. Chen, *Adv. Mater.*, 2020, **32**, 1806326.
- 4 Z. Chen, X. Duan, W. Wei, S. Wang and B.-J. Ni, *J. Mater. Chem. A*, 2019, **7**, 14971–15005.
- 5 X. Shang, B. Dong, Y.-M. Chai and C.-G. Liu, *Sci. Bull.*, 2018, **63**, 853–876.
- 6 J. Xie, X. Yang and Y. Xie, *Nanoscale*, 2020, **12**, 4283–4294.
- 7 J. Xie, J. Qi, F. Lei and Y. Xie, *Chem. Commun.*, 2020, **56**, 11910–11930.
- 8 J. Xie, X. Yang, Y. Wang, L. Kang, J. Li, Z. Wei, P. Hao, F. Lei, Q. Wang and B. Tang, *Chem. Commun.*, 2021, **57**, 11517–11520.
- 9 X. Han, H. Sheng, C. Yu, T. W. Walker, G. W. Huber, J. Qiu and S. Jin, *ACS Catal.*, 2020, **10**, 6741–6752.
- 10 M. Yu, E. Budiyo and H. Tüysüz, *Angew. Chem., Int. Ed.*, 2022, **61**, e202103824.
- 11 L. Li, P. Wang, Q. Shao and X. Huang, *Chem. Soc. Rev.*, 2020, **49**, 3072–3106.
- 12 Q. Hu, G. Li, Z. Han, Z. Wang, X. Huang, H. Yang, Q. Zhang, J. Liu and C. He, *J. Mater. Chem. A*, 2019, **7**, 14380–14390.
- 13 W. Sun, J. Li, W. Gao, L. Kang, F. Lei and J. Xie, *Chem. Commun.*, 2022, **58**, 2430–2442.
- 14 X. Yang, L. Kang, Z. Wei, S. Lou, F. Lei, P. Hao, J. Xie and B. Tang, *Chem. Eng. J.*, 2021, **422**, 130139.
- 15 M. Hao, J. Chen, J. Chen, K. Wang, J. Wang, F. Lei, P. Hao, X. Sun, J. Xie and B. Tang, *J. Colloid Interface Sci.*, 2023, **642**, 41–52.
- 16 J. Li, C. Dong, M. Guo, W. Gao, L. Kang, F. Lei, P. Hao, J. Xie and B. Tang, *Chem. Commun.*, 2022, **58**, 6845–6848.
- 17 Y. Wang, X. Yang, K. Wang, Z. Liu, X. Sun, J. Chen, S.-S. Liu, X. Sun, J. Xie and B. Tang, *Green Chem.*, 2023, **25**, 8216–8225.
- 18 Z. Liang, D. Jiang, X. Wang, M. Shakouri, T. Zhang, Z. Li, P. Tang, J. Llorca, L. Liu, Y. Yuan, M. Heggen, R. E. Dunin-Borkowski, J. R. Morante, A. Cabot and J. Arbiol, *Adv. Funct. Mater.*, 2021, **31**, 2106349.
- 19 R. Li, P. Kuang, L. Wang, H. Tang and J. Yu, *Chem. Eng. J.*, 2022, **431**, 134137.
- 20 G. Liu, W. Zhou, Y. Ji, B. Chen, G. Fu, Q. Yun, S. Chen, Y. Lin, P.-F. Yin, X. Cui, J. Liu, F. Meng, Q. Zhang, L. Song, L. Gu and H. Zhang, *J. Am. Chem. Soc.*, 2021, **143**, 11262–11270.
- 21 M. T. Bender, X. Yuan and K.-S. Choi, *Nat. Commun.*, 2020, **11**, 4594.
- 22 J. Xie, W. Liu, X. Zhang, Y. Guo, L. Gao, F. Lei, B. Tang and Y. Xie, *ACS Mater. Lett.*, 2019, **1**, 103–110.



- 23 J. Xie, L. Gao, S. Cao, W. Liu, F. Lei, P. Hao, X. Xia and B. Tang, *J. Mater. Chem. A*, 2019, **7**, 13577–13584.
- 24 J. Xie, H. Qu, F. Lei, X. Peng, W. Liu, L. Gao, P. Hao, G. Cui and B. Tang, *J. Mater. Chem. A*, 2018, **6**, 16121–16129.
- 25 L. Gao, J. Xie, S. Liu, S. Lou, Z. Wei, X. Zhu and B. Tang, *ACS Appl. Mater. Interfaces*, 2020, **12**, 24701–24709.
- 26 M. Li, W. Qi, J. Yu, L. Shen, X. Yang, S. Liu and M.-Q. Yang, *Chin. J. Struct. Chem.*, 2022, **41**, 15–24.
- 27 W. Qiao, X. Huang and L. Feng, *Chin. J. Struct. Chem.*, 2022, **41**, 16–34.
- 28 J. Li, J. Xie, X. Wang, Y. Dai, X. Xu, J. Liu, Z. Cai, X. Meng and J. Zou, *Chin. J. Struct. Chem.*, 2022, **41**, 59–67.
- 29 Y. Huang, X. Chong, C. Liu, Y. Liang and B. Zhang, *Angew. Chem., Int. Ed.*, 2018, **57**, 13163–13166.
- 30 M. T. Bender, Y. C. Lam, S. Hammes-Schiffer and K.-S. Choi, *J. Am. Chem. Soc.*, 2020, **142**, 21538–21547.
- 31 E. Tiburcio, R. Greco, M. Mon, J. Ballesteros-Soberanas, J. Ferrando-Soria, M. López-Haro, J. C. Hernández-Garrido, J. Oliver-Meseguer, C. Marini, M. Boronat, D. Armentano, A. Leyva-Pérez and E. Pardo, *J. Am. Chem. Soc.*, 2021, **143**, 2581–2592.
- 32 H. Huang, X. Song, C. Yu, Q. Wei, L. Ni, X. Han, H. Huang, Y. Han and J. Qiu, *Angew. Chem., Int. Ed.*, 2023, **62**, e202216321.
- 33 B. You, X. Liu, X. Liu and Y. Sun, *ACS Catal.*, 2017, **7**, 4564–4570.
- 34 D. Feng, Y. Dong, L. Zhang, X. Ge, W. Zhang, S. Dai and Z.-A. Qiao, *Angew. Chem., Int. Ed.*, 2020, **59**, 19503–19509.
- 35 J. Zheng, X. Chen, X. Zhong, S. Li, T. Liu, G. Zhuang, X. Li, S. Deng, D. Mei and J.-G. Wang, *Adv. Funct. Mater.*, 2017, **27**, 1704169.
- 36 H. Huang, C. Yu, X. Han, H. Huang, Q. Wei, W. Guo, Z. Wang and J. Qiu, *Energy Environ. Sci.*, 2020, **13**, 4990–4999.
- 37 Y. Xu, M. Liu, S. Wang, K. Ren, M. Wang, Z. Wang, X. Li, L. Wang and H. Wang, *Appl. Catal., B*, 2021, **298**, 120493.
- 38 J. Cao, H. Li, R. Zhu, L. Ma, K. Zhou, Q. Wei and F. Luo, *J. Alloys Compd.*, 2020, **844**, 155382.
- 39 Y. Wang, Y. Sun, F. Yan, C. Zhu, P. Gao, X. Zhang and Y. Chen, *J. Mater. Chem. A*, 2018, **6**, 8479–8487.
- 40 D. Li, F. Ruan, Y. Jin, Q. Ke, Y. Cao, H. Wang, T. Wang, Y. Song and P. Cui, *Catal. Sci. Technol.*, 2019, **9**, 418–424.
- 41 J. Zhang, S. Guo, B. Xiao, Z. Lin, L. Yan, D. Du and S. Shen, *Chem. Eng. J.*, 2021, **416**, 129127.
- 42 D. Yang, L. Yang, L. Zhong, X. Yu and L. Feng, *Electrochim. Acta*, 2019, **295**, 524–531.
- 43 X. Cui, M. Chen, R. Xiong, J. Sun, X. Liu and B. Geng, *J. Mater. Chem. A*, 2019, **7**, 16501–16507.
- 44 W. Sun, Y. Wang, S. Liu, F. Lei, J. Xie and B. Tang, *Chem. Commun.*, 2022, **58**, 11981–11984.
- 45 M. Guo, P. Li, A. Wang, J. Wang, J. Chen, F. Lei, P. Hao, X. Sun, J. Xie and B. Tang, *Chem. Commun.*, 2023, **59**, 5098–5101.
- 46 J. Li, M. Guo, X. Yang, J. Wang, K. Wang, A. Wang, F. Lei, P. Hao, J. Xie and B. Tang, *Prog. Nat. Sci.: Mater. Int.*, 2022, **32**, 705–714.
- 47 Z. Ma, H. Meng, M. Wang, B. Tang, J. Li and X. Wang, *ChemElectroChem*, 2018, **5**, 335–342.
- 48 G. Solomon, A. Landström, R. Mazzaro, M. Jugovac, P. Moras, E. Cattaruzza, V. Morandi, I. Concina and A. Vomiero, *Adv. Energy Mater.*, 2021, **11**, 2101324.
- 49 M. Fang, W.-B. Xu, Y. Shen, P. Cao, S. Han, W. Xu, D. Zhu, Y. Lu and W. Liu, *Appl. Catal., A*, 2021, **622**, 118220.
- 50 D. M. Morales, D. Jambrec, M. A. Kazakova, M. Braun, N. Sikdar, A. Koul, A. C. Brix, S. Seisel, C. Andronescu and W. Schuhmann, *ACS Catal.*, 2022, **12**, 982–992.
- 51 L. A. Wolzak, F. J. de Zwart, J.-P. H. Oudsen, S. A. Bartlett, B. de Bruin, J. N. H. Reek, M. Tromp and T. J. Korstanje, *ChemCatChem*, 2022, **14**, e202201033.
- 52 Y. Wang, B. Zhang, W. Pan, H. Ma and J. Zhang, *ChemSusChem*, 2017, **10**, 4170–4177.
- 53 P. Wang, D. Li, H. Chi, Y. Zhao, J. Wang, D. Li, S. Pang, P. Fu, J. Shi and C. Li, *Angew. Chem., Int. Ed.*, 2021, **60**, 6691–6698.
- 54 H. Wu, T. He, M. Dan, L. Du, N. Li and Z.-Q. Liu, *Chem. Eng. J.*, 2022, **435**, 134863.
- 55 C. Dong, M. Guo, W. Gao, P. Hao, F. Lei, J. Xie and B. Tang, *J. Colloid Interface Sci.*, 2022, **627**, 891–899.
- 56 S. Cao, J. Qi, F. Lei, Z. Wei, S. Lou, X. Yang, Y. Guo, P. Hao, J. Xie and B. Tang, *Chem. Eng. J.*, 2021, **413**, 127540.
- 57 J. Qi, J. Xie, Z. Wei, S. Lou, P. Hao, F. Lei and B. Tang, *Chem. Commun.*, 2020, **56**, 4575–4578.
- 58 A. P. de Moura, L. H. de Oliveira, I. L. V. Rosa, C. S. Xavier, P. N. Lisboa-Filho, M. S. Li, F. A. La Porta, E. Longo and J. A. Varela, *Sci. World J.*, 2015, **2015**, 315084.
- 59 K. Eda, Y. Kato, Y. Ohshiro, T. Sugitani and M. S. Whittingham, *J. Solid State Chem.*, 2010, **183**, 1334–1339.
- 60 H. Wan, J. Jiang, X. Ji, L. Miao, L. Zhang, K. Xu, H. Chen and Y. Ruan, *Mater. Lett.*, 2013, **108**, 164–167.
- 61 O. Popovych, I. Budzulyak, V. Vashchynskiy, M. Khemii, R. Ilnytskyi and L. Yablon, *Appl. Nanosci.*, 2023, **13**, 6803–6809.
- 62 P. R. Jothi, K. Shanthi, R. R. Salunkhe, M. Pramanik, V. Malgras, S. M. Alshehri and Y. Yamauchi, *Eur. J. Inorg. Chem.*, 2015, 3694–3699.

