



Cite this: *Green Chem.*, 2023, **25**, 2846

Ionic liquid-based electrolysis-deposition for modulating Pb crystal facets to boost CO₂ electroreduction†

Chongyang Jiang,^{a,b} Shaojuan Zeng,^{*a} Jiaqi Feng,^c Guilin Li,^a Zongxu Wang,^a Kuilin Peng,^a Lu Bai^{ID}^a and Xiangping Zhang^{ID}^{*a,b}

The crystal facet of electrodes is one of the main factors affecting the activity of CO₂ electroreduction. Herein, a new ionic liquid (IL)-based electrolysis-deposition (ED) method was developed for modulating the Pb crystal facet to boost CO₂ electroreduction. ED-Pb with an electrodeposition time of 900 s showed a high formate partial current density of 110.8 mA cm⁻² with over 80% formate faradaic efficiency at -2.4 V (vs. Ag/Ag⁺), which is much higher than values using a Pb planar electrode with 19.8 mA cm⁻² partial current density and 70% faradaic efficiency. Experimental results and density functional theory calculations revealed that ED-Pb catalysts have the dominant Pb(111) crystal facet due to the different adsorption energies of the IL on the crystal facet, and the improved performance is attributed to the low Gibbs free energy of OCHO* intermediates on the Pb(111) crystal facet. This work provides a new strategy for regulating the structures of electrocatalysts for high-activity CO₂ electroreduction.

Received 5th December 2022,

Accepted 1st March 2023

DOI: 10.1039/d2gc04622a

rsc.li/greenchem

Introduction

Electrochemical CO₂ reduction reaction (CO₂RR) to value-added products using renewable energy has been considered as an economically attractive approach for reducing the amount of CO₂ in the atmosphere.^{1–4} Among the possible CO₂ electroreduction products, formate as an important chemical intermediate and hydrogen carrier, is one of the most economical and viable products of the CO₂RR.^{5–9} Over the past decades, vigorous efforts have been made to develop electroreduction systems for high-efficiency CO₂RR to formate. To date, some active metal electrocatalysts, such as Bi, Sn, In, and Pb, have been used for CO₂ electroreduction to formate with significant progress.^{10–12} Among these metal electrocatalysts, Pb-based electrocatalysts are stable, hardly react with water and air, and are cheaper than those of other metals. At the same time, Pb-based catalysts have been widely studied for CO₂ electroreduction due to their high selectivity for formate products

and intrinsic inertness of the hydrogen evolution reaction (HER).^{13–15} However, most of the reported Pb-based catalysts still suffer from a low formation rate that leads to unsatisfactory CO₂ electroreduction performance.^{16–18}

In recent works, some means have been proposed to stabilize the reaction intermediates and improve the activity of CO₂ electroreduction to formate by Pb-based catalysts.^{19–22} In terms of catalyst design, the selectivity of Pb-based catalysts is dependent on surface morphology, crystal surface structure, and structural composition.^{23–25} Electrodeposition can lead to special morphology with large electrochemically active surface areas that are hypothesized to be catalytically active sites.²⁶ Guay *et al.*²⁷ designed honeycomb porous Pb films, which can significantly increase the formate current density (j_{formate} , 7.5 mA cm⁻²) and faradaic efficiency (FE_{formate}, 97%). The enhanced performance of the porous Pb films was attributed to the dendritic secondary structure that could help to stabilize the intermediates in the boundary layer. In addition, ionic liquids (ILs), as new solvents, can be directly used as reaction solvents to regulate the crystal facets of catalysts.^{28,29} For example, Zhang *et al.*²⁹ used 1-hexyl-3-methylimidazolium tetrafluoroborate ([Hmim][BF₄]) as the solvent to prepare the flowerlike In₂S₃ catalyst with a dominant (440) crystal facet, exhibiting a formate formation rate of 478 μmol cm⁻² h⁻¹ with 86% FE_{formate}. Moreover, ILs not only modified the catalyst crystal facet but also improved the current density and product selectivity as CO₂RR electrolytes.^{30–33} Although the CO₂ electroreduction to formate system has been extensively studied, it is

^aBeijing Key Laboratory of Ionic Liquids Clean Process, State Key Laboratory of Multiphase Complex Systems, Institute of Process Engineering, Chinese Academy of Sciences, Beijing 100190, China. E-mail: xpzhang@ipe.ac.cn, sjzeng@ipe.ac.cn;

Fax: +8610 6255 8174

^bSchool of Future Technology, University of Chinese Academy of Sciences, Beijing 100049, China

^cKey Laboratory of Colloid and Interface and Thermodynamics, Institute of Chemistry, Chinese Academy of Sciences, Beijing 100190, China

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2gc04622a>

still extremely challenging to achieve satisfactory formate formation rates using Pb-based materials.

Herein, a novel IL-based electrolysis-deposition method for modulating Pb crystal facets (ED-Pb) was proposed, which can significantly enhance the activity of CO₂ electroreduction to formate. This electrolysis-deposition process was observed by an *operando* optical microscope (OM), and ED-Pb catalysts with different crystal facet ratios were obtained by altering the electrodeposition conditions. Furthermore, combined with experimental characterization techniques and density functional theory (DFT) calculations, the mechanism of high CO₂ electrochemical performance to formate by ED-Pb electrodes was systematically investigated, which will provide a new and efficient method for modifying the catalyst crystal facet to boost CO₂RR activity.

Experimental section

Fabrication of ED-Pb electrodes

ED-Pb electrodes were prepared by the IL-based electrolysis-deposition method at room temperature. The cell is an ordinary H-type electrolytic cell separated by Nafion 117 membrane. Before electrolysis-deposition experiments, Pb planar electrodes ($\geq 99.9\%$, area: $1 \times 1 \text{ cm}^2$, thickness: 0.5 mm) were polished with #600 and #2000 sandpaper until the surface was smooth and then washed with water and acetone in sequence. The pre-treated Pb planar electrodes, Ag/Ag⁺, and platinum gauze were used as the working electrode, reference electrode, and counter electrode, respectively. Unless stated otherwise, all the potentials in this study were *versus* the Ag/Ag⁺ reference electrode. The working and reference electrodes were placed in the cathode chamber with 0.5 M 1-butyl-3-methylimidazolium trifluoroacetate ([Bmim][TFA], Fig. S1†)/acetonitrile (AcN) solution, while the counter electrode was placed in the anode chamber with a 0.1 M H₂SO₄ solution as the electrolyte.

A two-step electrolysis-deposition process with switching cathode/anode of the H-type cell was used to prepare highly active ED-Pb electrodes. First, the electrolysis process was carried out at -0.1 A for 900 s to form Pb ions by polarizing the Pb planar electrode. Second, the deposition process was carried out at a constant current (such as 0.1 A for 900 s) and the Pb ions were reduced to form the active ED-Pb electrode. The deposition morphology and crystal facet of ED-Pb electrodes could be tuned by applying different deposition times and currents (450 s-0.2 A; 900 s-0.1 A; 1800 s-0.05 A; and 3600 s-0.025 A; total charge of 90 C) during the deposition process, and the corresponding electrodes were annotated as ED-Pb-450, ED-Pb-900, ED-Pb-1800, and ED-Pb-3600, respectively. After the electrolysis-deposition process, the obtained black-colored ED-Pb electrodes were further washed with AcN and dried under an N₂ atmosphere.

Physical characterization

An *operando* optical microscope (OM) study was carried out using a SangNond light microscope (SN0745-SN108060SDU).

The morphologies of the ED-Pb electrodes were detected by scanning electron microscopy (SEM, Hitachi SU8020) and transmission electron microscopy (TEM, JEOL JEM-2100 system). The crystal structures of the ED-Pb electrodes were investigated by X-ray diffraction (XRD, Rigaku Smartlab diffractometer) using a Cu K α ($\lambda = 1.5418 \text{ \AA}$) radiation source with a scanning rate of $15^\circ \text{ min}^{-1}$.

Evaluation of CO₂ electroreduction performance

All the CO₂ electrochemical experiments were performed in a H-type cell using an electrochemical workstation (CHI660E, Shanghai Chenhua Instrument Co, Ltd). The ED-Pb electrodes were used as the working electrode, and Ag/Ag⁺ and platinum wire were used as the reference and counter electrodes, respectively. The working and reference electrodes were placed in the cathode chamber with 1-butyl-3-methylimidazolium hexafluorophosphate ([Bmim][PF₆])/AcN-H₂O solution mixture, while the counter electrode was placed in the anode chamber with a 0.1 M H₂SO₄ solution as the electrolyte. The electrochemical performances were characterized by linear sweep voltammetry (LSV), cyclic voltammetry (CV), and electrochemical impedance spectroscopy (EIS). LSV was carried out in a potential range of -1.5 to -2.4 V at a scan rate of 20 mV s^{-1} in N₂ or CO₂-saturated IL electrolyte. The electrochemically active surface area (ECSA) was determined by the double-layer capacitances calculated from a plot of discharging current density *vs.* scan rate, while the electric double-layer capacitance was determined by CV measurement at various scan rates (10 to 100 mV s^{-1}) from potentials of -0.7 to -0.8 V . EIS was performed within a frequency range of 0.01 to 0.1 MHz . CO₂ electroreduction was carried out at various constant potentials (-2.0 to -2.4 V) for 1 h. During the reduction process, CO₂ was continuously bubbled into the reactor with a gas flow rate of 30 mL min^{-1} .

Results and discussion

A schematic diagram of the ED-Pb electrode preparation process is shown in Fig. S2.† Compared with the catalyst preparation methods reported in previous literature (solvent-thermal reaction, chemical vapor deposition and chemical etching),^{29,34,35} the preparation of ED-Pb catalysts in IL electrolytes by electrolysis-deposition requires mild conditions, short time, and a simple preparation process. The obtained catalyst can be directly used for catalytic reactions, avoiding the addition of binders.^{36,37} Conventional electrodeposition was carried out in a mixed solution containing metal salts with other solutes.^{27,38,39} We obtained Pb²⁺ metal ions by electrolysis, and then electrodeposition was performed to produce ED-Pb electrodes to achieve efficient utilization of metal ions in this work. At the same time, the electrolyte undergoes no change and can be recycled after the whole electrolysis-deposition process.

To monitor the dynamic Pb electrolysis-deposition reaction in the [Bmim][TFA] electrolyte, an *in situ* OM investigation was

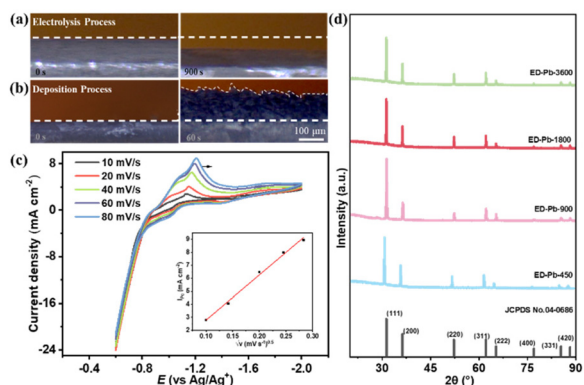


Fig. 1 *In situ* OM images of the Pb planar electrode in 0.5 M [Bmim][TFA]/AcN electrolyte (a) at -0.1 A for 900 s in the electrolysis process and (b) at 0.1 A for 900 s in the electrodeposition process. (c) Cyclic voltammograms of the Pb planar electrode at -0.6 to -2.0 V after the Pb electrolysis process (the inset illustration: relationship between \sqrt{v} and j_{pc}). (d) XRD patterns of the Pb standard PDF card (No. 04-0686) and ED-Pb electrodes with different deposition times.

conducted. A set of typical visualization results are shown in Fig. 1a and b (see ESI II† for the video of the electrolysis-deposition process). The Pb planar electrode was quite uniform before electrolysis. The surface of the Pb planar electrode drops gradually relative to its starting position at a current of -0.1 A, indicating that the Pb planar electrode is polarized to form Pb ions. Subsequently, at a current of 0.1 A, a black material was deposited and grew quickly to cover the Pb planar electrode surface, which could be due to the reduction of Pb ions in the solution to Pb. The same electrolysis-deposition process was carried out in an aqueous solution and IL electrolytes, and the potential curve over time is shown in Fig. S3.† In aqueous solutions, the oxidation potential was approximately 2 V (Fig. S3b†) and the experimental results showed that a large number of bubbles are generated on the Pb electrode surface, which is more prone to the oxygen evolution reaction. The IL electrolytes included 1-butylpyridinium chloride ([BPy][Cl]), tetrabutylphosphine chloride ([P₄₄₄₄][Cl]), 1-butyl-3-methylimidazolium chloride ([Bmim][Cl]), 1-butyl-3-methylimidazolium acetate ([Bmim][Ac]), and 1-butyl-3-methylimidazolium bistrifluoromethanesulfonate ([Bmim][NTf₂]) (Fig. S4†). As can be seen from Fig. S3,† the potential curves of the three ILs ([BPy][Cl], [P₄₄₄₄][Cl], and [Bmim][Cl]) containing the Cl⁻ anions with different cations were similar to [Bmim][TFA], and the black material was found on the Pb planar electrode in all three IL electrolytes. It can be seen that the cations of ILs have little effect on the electrolysis-deposition process. At a current density of 0.1 A in the [Bmim][Ac] electrolyte, the applied potential exceeds the range of the electrochemical station (CHI 660E, potential range of ± 10 V) and the electrolysis process is terminated (Fig. S3f†). When the IL contains halogens ([Bmim][NTf₂] and [Bmim][Cl]), a similar curve appears, and a black material appears on the Pb planar electrode surface, mainly because the halogen ion radius is small and the reactivity is high. Then the substrate can be

quickly corroded in the early stage of polarization, intensifying the dissolution of Pb in the form of ions.^{40,41} These results indicated that the electrolysis-deposition process depends on the composition of the electrolyte, and the IL electrolyte containing halogens can promote the oxidation of Pb to Pb ions.

To study the valence state of the Pb ions in the IL electrolyte, cyclic voltammograms (CV) after the Pb electrolysis process at different sweep rates, including 10 , 20 , 40 , 60 and 80 mV s⁻¹, were obtained as shown in Fig. 1c. The reduction potential of Pb was about -1.2 V, which is close to the standard electrode potential of Pb²⁺ (-0.1251 V vs. RHE).⁴² Subsequently, an equal molar amount of Pb(CH₃COO)₂ was added to the 0.5 M [Bmim][AC]/AcN electrolyte to observe the deposition CV curve with a reduction potential of -1.23 V (Fig. S3h†). From the electrolysis-deposition process, it could be inferred that Pb is firstly oxidized to Pb²⁺, and then Pb²⁺ is reduced to form metallic Pb. The Pb electrodeposition reaction is controlled by multiple facets, including kinetics, crystallographic thermodynamics, and ion diffusion. Thus, based on the CV results, it can be seen that as the scan rate increased from 10 to 80 mV s⁻¹, the reduction potential peak moved from -1.11 to -1.21 V and the reduction current density peak (j_{pc}) gradually increased from 3.49 to 8.94 mA cm⁻². Therefore, the Pb electrodeposition process is irreversible. The linear relationship between the square root of the scan rate (\sqrt{v}) and j_{pc} is shown in the inset of Fig. 1c, which indicates that the Pb electrodeposition process is controlled by diffusion. These results show that the electrodeposition process is complex and the parameters of the electrolysis-deposition process directly affect the structure of ED-Pb.^{43,44}

To understand the structures of electrodes, the powder X-ray diffraction (XRD) technique was utilized to reveal the crystal facet composition of the Pb electrode. As shown in Fig. S5,† the peak positions of the Pb planar electrode before and after electrolysis are 31.3° , 36.3° , 52.2° , 62.1° , 65.2° , 76.9° , 85.4° and 88.2° , corresponding to the Pb crystal facets of (111), (200), (220), (311), (222), (400), (331) and (420) (JCPDS card No. 04-0686), respectively. The relative peak intensity ratio of the Pb planar electrode at Pb(111) : Pb(200) ($I_{111} : I_{200}$) was also calculated. The $I_{111} : I_{200}$ of the Pb planar electrode before electrolysis was 2.0 , while the $I_{111} : I_{200}$ of the Pb planar electrode after electrolysis was 1.67 , indicating that metallic Pb is polarized to form Pb²⁺ ions during the electrolysis process. The diffraction peaks of all the ED-Pb electrodes were the same as those of Pb (JCPDS card No. 04-0686) (Fig. 1d), and the $I_{111} : I_{200}$ of the ED-Pb electrodes were calculated as 2.17 , 3.20 , 2.40 , and 2.38 from ED-Pb-450 to ED-Pb-3600. The clear lattice fringes were observed on the TEM images of the ED-Pb-900 electrode, and the fringe spacing of 0.285 nm was consistent with the interplanar spacing of the Pb(111) crystal facet (Fig. S6†). Therefore, ED-Pb-900 electrodes have a dominant crystal facet of Pb(111) compared to that of Pb(200). The experimental results indicated that ED-Pb electrodes with different relative peak intensity ratios can be adjusted by varying the applied deposition parameters. In order to understand the ED-Pb electrode with the dominant Pb(111) crystal

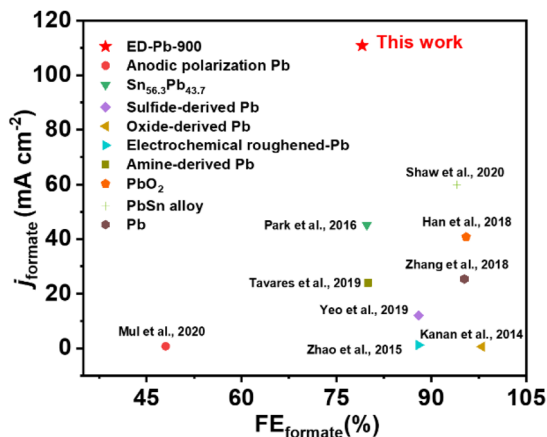


Fig. 3 Comparison of FE_{formate} and j_{formate} of ED-Pb-900 with those of other Pb-based catalysts as reported in the literature.

Whether the CO_2 electroreduction performance of an electrode can maintain long-term stability is of great importance for practical applications. Therefore, the stability of the ED-Pb-900 electrode was measured by five sequential CO_2 electroreduction experiments under corresponding optimal potentials in the $[\text{Bmim}][\text{PF}_6]/\text{AcN}/\text{H}_2\text{O}$ electrolyte (Fig. S13[†]). The results indicated that there is no obvious decay in either FE_{formate} or j_{formate} during the stability test. The XRD and SEM results after the electroreduction experiments indicated that the crystal structure and the morphology of ED-Pb-900 remained unchanged (Fig. S14[†]). These experimental results further confirmed the excellent stability of the ED-Pb-900 electrode.

To quantitatively analyze and understand the catalytic performance of Pb-based electrodes, the electrochemically active surface area (ECSA) of the ED-Pb electrodes and the Pb planar electrode was calculated (Fig. S15–S19[†]). The ECSA of ED-Pb-900 ($16.1 \mu\text{F cm}^{-2}$) was 4.73-fold higher than that of the Pb planar electrode ($3.4 \mu\text{F cm}^{-2}$) (Fig. 4a), suggesting that the electrolysis-deposition process can significantly increase the number of exposed active sites, which is beneficial to improve the current density of the CO_2RR .⁵¹ Electrochemical impedance spectroscopy (EIS) experiments were performed to investigate the effect of ED-Pb electrodes on the kinetics of the CO_2RR (Fig. 4b). It was found that the R_{ct} of ED-Pb-900 ($0.49 \Omega \text{ cm}^{-2}$) is 78% lower than that of the Pb planar electrode (2.3Ω

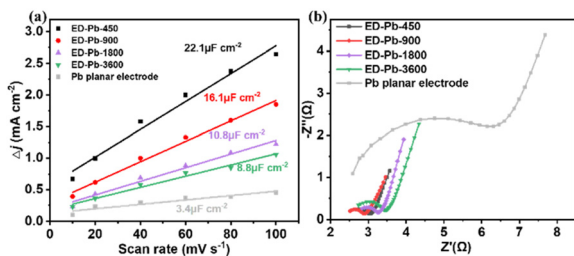


Fig. 4 (a) Charging current density differences plotted against scan rate and (b) the Nyquist diagrams of ED-Pb electrodes and the Pb planar electrode.

cm^{-2}), which is the smallest value among the ED-Pb electrodes. The lower R_{ct} value of ED-Pb-900 indicated a faster charge transfer rate between the electrode and electrolyte, implying that the adsorbed CO_2 molecules easily obtain electrons to form intermediates. Therefore, tuning the structure of ED-Pb electrodes can efficiently enhance the ECSA and decrease the interfacial charge transfer of the CO_2RR .

The reaction pathway for CO_2 electroreduction to formate mainly contains two steps: step 1 is the transfer of a proton–electron pair to CO_2 to form the $^*\text{OCHO}$ intermediate; step 2 is the formation of formic acid/formate product from the intermediate $^*\text{OCHO}$ with a second proton–electron pair.⁵² To understand the mechanism underpinning the outstanding performance of ED-Pb-900, DFT calculations were performed on the reaction energetics over the Pb(111) and Pb(200) crystal facets, which include competitive $^*\text{OCHO}$, $^*\text{COOH}$ and $^*\text{H}$ paths. Comparing the projected density of states (PDOS) of a Pb atom on Pb(111) with that on Pb(200) before $^*\text{OCHO}$ adsorption, the position of the d -band center for Pb(111) is closer to the Fermi level than that of Pb(200), indicating that Pb(111) is more favorable to the binding and activation of CO_2 (Fig. 5a). The optimized configurations of $^*\text{OCHO}$ and $^*\text{COOH}$ intermediates on the Pb(111) and Pb(200) crystal facets are displayed in Fig. S20[†]. The $^*\text{OCHO}$ intermediates were adsorbed on both crystal facets *via* two oxygen atoms. The bond lengths and angles of the optimized configuration of intermediates are summarized in Table S2[†]. As shown in Fig. 5b, Pb(111) exhibited the most negative adsorption energy on the $^*\text{OCHO}$ intermediate, which indicated that Pb(111) is favored to stabilize $^*\text{OCHO}$, which is beneficial to decrease the onset potential and enhance selectivity of formate. In addition, the Pb(111) crystal facet (0.299 eV) showed a lower free energy barrier for formate formation than the Pb(200) crystal facet (0.398 eV), which meant that the Pb(111) crystal facet is more energy-favorable for producing formate (Fig. 5c and d). Furthermore, the free energy diagram of competitive $^*\text{H}$ and $^*\text{COOH}$ inter-

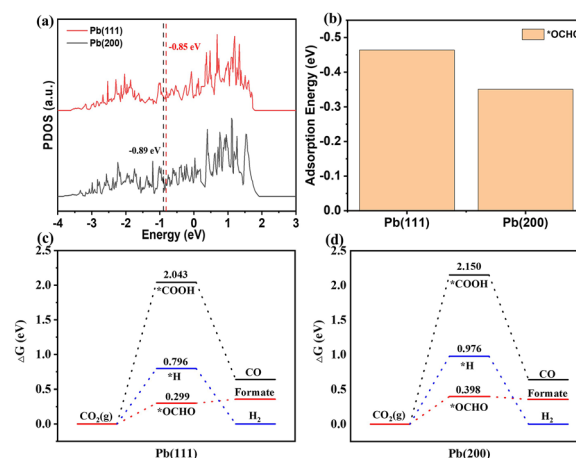


Fig. 5 (a) Projected density of states of Pb(111) and Pb(200). (b) The adsorption energies of $^*\text{OCHO}$ on Pb(111) and Pb(200). Calculated Gibbs free energy diagrams of $^*\text{OCHO}$, $^*\text{COOH}$, and $^*\text{H}$ on (c) Pb(111) and (d) Pb(200).

mediates is also provided as a descriptor as shown in Fig. 5c and d. The relatively high free energy barriers of *H and *COOH intermediates on the two Pb crystal facets indicated that they are not conducive to the occurrence of the hydrogen evolution reaction and CO₂ electroreduction to CO. To further explore the interaction between the Pb(111) crystal facet and these intermediates, we calculated the DOS of the *OCHO and *COOH intermediates adsorbed on Pb(111). The PDOS of the O atom in the adsorption of *OCHO and *COOH by Pb(111) were analyzed by decomposing the electron density and the wave function into the atomic orbital contributions. As shown in Fig. S21,† large overlaps between the O of *OCHO and Pb atoms of the Pb(111) surface indicated that Pb(111) has a strong interaction effect with the O atom in the *OCHO intermediates, which contributes to the absorption and stabilization of the *OCHO intermediates on the Pb(111) crystal facet. In summary, ED-Pb-900 with more Pb(111) crystal facets than Pb(200) crystal facets shows higher CO₂RR activity.

Conclusions

In this work, the IL-based electrolysis-deposition preparation of Pb electrodes for effectively promoting CO₂ electroreduction to formate was proposed and proved by experiments and simulation calculations. The electrolysis-deposition process depends on the composition of the electrolyte. The halogen-containing IL electrolytes could promote the oxidation of Pb to Pb²⁺ and the ED-Pb electrodes with different relative peak intensity ratios were prepared by altering the deposition conditions. Therefore, due to the low surface energy of Pb(111) and the low adsorption energy of [Bmim][TFA] on Pb(111), ED-Pb electrodes have a dominant Pb(111) crystal facet. The performance of the CO₂RR to formate was gradually improved with increasing $I_{111}:I_{200}$ of the ED-Pb electrodes. At -2.4 V, compared with the Pb planar electrode (70% FE, 19.8 mA cm⁻²), ED-Pb-900 had significantly improved CO₂RR performance (80.0% FE, 110.8 mA cm⁻²). In particular, an excellent formate formation rate of up to 2067.2 μmol h⁻¹ cm⁻² at -2.4 V was obtained for ED-Pb-900, which is 5.6 times higher than that of the Pb planar electrode. Based on the relationship between the structures and electrochemical activity, the excellent performance of ED-Pb electrodes was attributed to the preferential exposure of the Pb(111) crystal facet, the larger electrochemically active surface area, and the lower interfacial charge-transfer resistance. The DFT calculation results proved that the free energy of *OCHO on Pb(111) (0.299 eV) is smaller than that on Pb(200) (0.398 eV), resulting in a high intrinsic activity for CO₂ electroreduction to formate. This electrolysis-deposition strategy provides a new and efficient method of modifying catalysts for the CO₂RR.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

This work was supported by the National Key R&D Program of China (2020YFA0710200), the National Natural Science Foundation of China (21890764, 21838010 and 22122814), the Youth Innovation Promotion Association of the Chinese Academy of Sciences (2018064), the Instrument Developing Project of the Chinese Academy of Sciences (YJKYYQ20200062), the 2021 Industrial Technology Basic Public Service Platform Project—Public Service Platform for Emission Peak and Carbon Neutral of Key Raw Material Industries (2021-H029-1-1), and the State Key Laboratory of Multiphase Complex Systems, Institute of Process Engineering, Chinese Academy of Sciences (No. MPCS-2022-A-03).

References

- X. Zhang, G. Ren, C. Zhang, J. Xue, Q. Zhao, R. Li, Y. Wang and C. Fan, *Green Energy Environ.*, 2021, **6**, 693–702.
- F. Chang, G. Zhan, Z. Wu, Y. Duan, S. Shi, S. Zeng, X. Zhang and S. Zhang, *ACS Sustainable Chem. Eng.*, 2021, **9**, 9045–9052.
- B. Shao, Y. Zhang, Z. Sun, J. Li, Z. Gao, Z. Xie, J. Hu and H. Liu, *Green Chem. Eng.*, 2022, **3**, 189–198.
- A. Paulillo, M. Pucciarelli, F. Grimaldi and P. Lettieri, *Green Chem.*, 2021, **23**, 6639–6651.
- C. Jiang, J. Feng, S. Zeng and X. Zhang, *Chin. Sci. Bull.*, 2020, **66**, 716–727.
- J. Feng, S. Zeng, J. Feng, H. Dong and X. Zhang, *Chin. J. Chem.*, 2018, **36**, 961–970.
- J. Feng, H. Gao, L. Zheng, Z. Chen, S. Zeng, C. Jiang, H. Dong, L. Liu, S. Zhang and X. Zhang, *Nat. Commun.*, 2020, **11**, 4341.
- H. Liu, B. Miao, H. Chuai, X. Chen, S. Zhang and X. Ma, *Green Chem. Eng.*, 2022, **3**, 138–145.
- D.-G. Yu and L.-N. He, *Green Chem.*, 2021, **23**, 3499.
- P. Ding, H. Zhao, T. Li, Y. Luo, G. Fan, G. Chen, S. Gao, X. Shi, S. Lu and X. Sun, *J. Phys. Chem. A*, 2020, **8**, 21947–21960.
- C. Peng, X. Wu, G. Zeng and Q. Zhu, *Chem. – Asian J.*, 2021, **16**, 1539–1544.
- Y. Tian, D. Li, C. Li, J. Liu, J. Wu, G. Liu and Y. Feng, *Chem. Eng. J.*, 2021, **414**, 128671.
- Q. G. Zhu, J. Ma, X. C. Kang, X. Sun, H. Liu, J. Hu, Z. Liu and B. Han, *Angew. Chem., Int. Ed.*, 2016, **55**, 9012–9016.
- J. D. Watkins and A. B. Bocarsly, *ChemSusChem*, 2014, **7**, 284–290.
- Z. Yang, F. E. Oropeza and K. H. L. Zhang, *APL Mater.*, 2020, **8**, 060901.
- Z. He, J. Shen, Z. Ni, J. Tang, S. Song, J. Chen and L. Zhao, *Catal. Commun.*, 2015, **72**, 38–42.
- W. Yu, L. Wen, J. Gao, S. Chen, Z. He, D. Wang, Y. Shen and S. Song, *Chem. Commun.*, 2021, **57**, 7418–7421.

- 18 H. Wang, Y. Chen, X. Hou, C. Ma and T. Tan, *Green Chem.*, 2016, **18**, 3250–3256.
- 19 C. Chen, Y. Li, S. Yu, S. Louisia, J. Jin, M. Li, M. B. Ross and P. Yang, *Joule*, 2020, **4**, 1688–1699.
- 20 F. Yang, H. Yu, X. Mao, Q. Meng, S. Chen, Q. Deng, Z. Zeng, J. Wang and S. Deng, *Chem. Eng. J.*, 2021, **425**, 131661.
- 21 P. Yue, Q. Fu, J. Li, L. Zhang, L. Xing, Z. Kang, Q. Liao and X. Zhu, *Chem. Eng. J.*, 2021, **405**, 126975.
- 22 H. Wu, J. Song, C. Xie, Y. Hu and B. Han, *Green Chem.*, 2018, **20**, 1765–1769.
- 23 R. M. Arán-Ais, F. Scholten, S. Kunze, R. Rizo and B. R. Cuenya, *Nat. Energy*, 2020, **5**, 317–325.
- 24 J. Feng, L. Zheng, C. Jiang, Z. Chen, L. Liu, S. Zeng, L. Bai, S. Zhang and X. Zhang, *Green Chem.*, 2021, **23**, 5461–5466.
- 25 P. P. Guo, Z. H. He, S. Y. Yang, W. Wang, K. Wang, C. C. Li, Y. Y. Wei, Z. T. Liu and B. Han, *Green Chem.*, 2022, **24**, 1527–1533.
- 26 P. De Luna, R. Quintero-Bermudez, C. T. Dinh, M. B. Ross, O. S. Bushuyev, P. Todorović, T. Regier, S. O. Kelley, P. Yang and E. H. Sargent, *Nat. Catal.*, 2018, **1**, 103–110.
- 27 M. Fan, S. Garbarino, G. A. Botton, A. C. Tavares and D. Guay, *J. Mater. Chem. A*, 2017, **5**, 20747–20756.
- 28 Q. Zhu, D. Yang, H. Liu, X. Sun, C. Chen, J. Bi, J. Liu, H. Wu and B. Han, *Angew. Chem., Int. Ed.*, 2020, **59**, 8896–8901.
- 29 J. Q. Feng, H. S. Gao, J. P. Feng, L. Liu, S. Zeng, H. Dong, Y. Bai, L. Liu and X. P. Zhang, *ChemCatChem*, 2019, **12**, 926–931.
- 30 J. Feng, S. Zeng, H. Liu, J. Feng and H. Gao, *ChemSusChem*, 2018, **11**, 3191–3197.
- 31 S. K. Shukia, S. G. Khokarale, T. Q. Bui and J. P. T. Mikkola, *Front. Mater.*, 2019, **6**, 42.
- 32 Y. Yang, H. Gao, J. Feng, S. Zeng, L. Liu, L. Liu, B. Ren, T. Li, S. Zhang and X. Zhang, *ChemSusChem*, 2020, **13**, 4900–4905.
- 33 X. Sun, Q. Zhu, X. Kang, H. Liu, Q. Qian, J. Ma, Z. Zhang, G. Yang and B. Han, *Green Chem.*, 2017, **19**, 2086–2091.
- 34 Q. Gong, P. Ding, M. Xu, X. Zhu, M. Wang, J. Deng, Q. Ma, N. Han, Y. Zhu, J. Lu, Z. Feng, Y. Li, W. Zhou and Y. Li, *Nat. Commun.*, 2019, **10**, 2807.
- 35 R. Feng, Q. Zhu, M. Chu, S. Jia, J. Zhai, H. Wu, P. Wu and B. Han, *Green Chem.*, 2020, **22**, 7560–7565.
- 36 J. Liu, P. Li, J. Bi, Q. Zhu and B. Han, *Chemistry*, 2022, **28**, e202200242.
- 37 Q. Zhu, D. Yang, H. Liu, X. Sun, C. Chen, J. Bi, J. Liu, H. Wu and B. Han, *Angew. Chem., Int. Ed.*, 2020, **59**, 8896–8901.
- 38 B. Ren, G. Wen, R. Gao, D. Luo, Z. Zhang, W. Qiu, Q. Ma, X. Wang, Y. Cui, L. Ricardez-Sandoval, A. Yu and Z. Chen, *Nat. Commun.*, 2022, **13**, 2486.
- 39 Z. Li, B. Sun, D. Xiao, Z. Wang, Y. Liu, Z. Zheng, P. Wang, Y. Dai, H. Cheng and B. Huang, *Angew. Chem., Int. Ed.*, 2023, **62**, e202217569.
- 40 P. Ramachandran, K. V. Venkateswaran and V. Nandakumar, *Bull. Electrochem.*, 1996, **12**, 346–348.
- 41 X. C. Zhong, R. X. Wang, Q. S. Liu, L. X. Jiang, X. J. Lv, Y. Q. Lai and J. Li, *Chin. J. of Nonferrous Met.*, 2018, **28**, 792–801.
- 42 D. Wu, D. J. Solanki, J. L. Ramirez, W. Yang, A. Joi, Y. Dordi, N. Dole and S. R. Brankovic, *Langmuir*, 2018, **34**, 11384–11394.
- 43 Z. Cai, J. Wang, Z. Lu, R. Zhan, Y. Ou, L. Wang, M. Dahbi, J. Alami, J. Lu, K. Amine and Y. Sun, *Angew. Chem., Int. Ed.*, 2022, **61**, e202116560.
- 44 Y. Zhao, X. Tan, W. Yang, C. Jia, X. Chen, W. Ren, S. C. Smith and C. Zhao, *Angew. Chem., Int. Ed.*, 2020, **59**, 21493–21498.
- 45 H. Zhong, Y. Qiu, X. Li, L. Pan and H. Zhang, *J. Energy Chem.*, 2021, **55**, 236–243.
- 46 J. E. Pander, J. W. J. Lum and B. S. Yeo, *J. Phys. Chem. A*, 2019, **7**, 4093–4101.
- 47 Z. Zeng, P. Barai, S. Y. Lee, J. Yang, X. Zhang, W. Zheng, Y. S. Liu, K. C. Bustillo, P. Ercius, J. Guo, Y. Cui, V. Srinivasan and H. Zheng, *Nano Energy*, 2020, **72**, 104721.
- 48 P. Grosfils and J. F. Lutsko, *Crystals*, 2020, **11**, 4.
- 49 G. Wang, T. Liu, X. Fu, Z. Wu, M. Liu and X. Xiong, *Chem. Eng. J.*, 2021, **414**, 128698.
- 50 Y. Zhang, Z. Huang, K. Wu, F. Yu, M. Zhu, G. Wang, G. Xu, M. Wu, H. K. Liu, S. X. Dou and C. Wu, *Chem. Eng. J.*, 2022, **430**, 133042.
- 51 D. Li, J. Wu, T. Liu, J. Liu, Z. Yan, L. Zhen and Y. Feng, *Chem. Eng. J.*, 2019, **375**, 122024.
- 52 Y.-H. Wang, W.-J. Jiang, W. Yao, Z.-L. Liu, Z. Liu, Y. Yang and L.-Z. Gao, *Rare Met.*, 2021, **40**, 2327–2353.