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A Ru/RuO₂ heterostructure boosting electrochemistry-assisted selective benzoic acid hydrogenation†

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Electrocatalytic hydrogenation of benzoic acid (BA) to cyclohexanecarboxylic acid (CCA) at ambient temperature and pressure has been recognized as a promising alternative to thermal hydrogenation since water is required as the hydrogen source. So far, only a few Pt-based electrocatalysts have been developed in acidic electrolyte. To overcome the limitations of reactant solubility and catalyst corrosion, herein, carbon fiber-supported Ru electrocatalysts with abundant Ru/RuO₂ heterojunctions were fabricated *via* cyclic electrodeposition between −0.8 and 1.1 V vs. Ag/AgCl. In an alkaline environment, a Ru/RuO₂ catalyst achieves an excellent ECH reactivity in terms of high BA conversion (100%) and selectivity towards CCA (100%) within 180 min at a current density of 200/3 mA cm^{−2}, showing exceptional reusability and long-term stability. 1-Cyclohexenecarboxylic acid (CEA) was identified as the reaction intermediate, whose the selectivity is governed by the applied potential. Kinetic studies demonstrate that ECH of BA over Ru/RuO₂ follows a Langmuir–Hinshelwood (L–H) mechanism. *In situ* Raman spectroscopy and theoretical calculations reveal that the Ru/RuO₂ interface enhances the adsorption strength of CEA, thereby facilitating the production of fully hydrogenated CCA. This work provides a deep understanding of the ECH pathway of BA in alkaline media, and gives a new methodology to fabricate heterostructure electrocatalysts.

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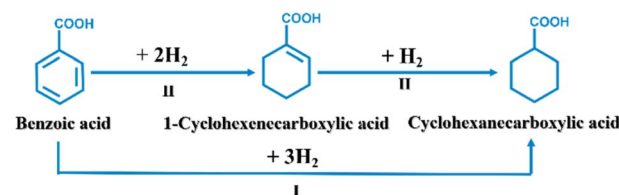
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Introduction

Cyclohexane carboxylic acid (CCA) is an important organic intermediate for therapeutic drugs, such as ansatrienin, praziquantel and caprolactam,^{1,2} and its derivatives are important intermediates for organic synthesis in fine chemical industries, such as *trans*-4-isopropylcyclohexyl acid and cyclohexylmethyl carbamate. For industrial scale synthesis of CCA, selective hydrogenation of benzoic acid (BA) to CCA has attracted growing research attention in the past few years^{3–12} (listed in Table S1†). At 80 °C and 1 MPa H₂, Qiu *et al.*¹³ realized a CCA selectivity of 100% at a conversion of 89.4% over Ru nanoparticles loaded on N-doped porous two-dimensional carbon nanosheets without hydrogenated intermediates (pathway I as illustrated in Scheme 1). For comparison, Ren *et al.*¹⁴ reported that the Ru/PPh₃@FDU catalyst achieved a CCA selectivity of 98% with 99% conversion at 60 °C and 1 MPa H₂, showing CEA as the reaction intermediate (as shown in Scheme 1). Guo *et al.*¹⁵

fabricated Pt/TiO₂ to achieve BA conversion (90%) with a CCA selectivity of 99% at 80 °C and 50 bar H₂ *via* pathway II (Scheme 1). Moreover, the RhPt/MCM-41 bimetallic catalyst¹⁶ offered a conversion of 63.5% and a selectivity of 97.8% at 40 °C and 1 MPa H₂, following reaction pathway II (Scheme 1). Accordingly, the reaction pathway from BA to CCA is highly dependent upon the feature of the catalyst. Hence, it is urgent to design high-efficiency catalysts with satisfactory conversion.

Electrocatalytic hydrogenation (ECH) of aromatic organics without external H₂ gas has aroused extensive research activities.^{17–25} For example, Peng *et al.* employed a ternary PtRhAu electrocatalyst to reach a high faradaic efficiency (58%) toward 2-methoxycyclohexanol from guaiacol.²⁶ Moreover, Ni–MoO₂ catalysts were designed to perform ECH of phenol to

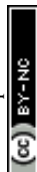


Scheme 1 Reaction pathways for thermal catalytic hydrogenation of BA.

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cyclohexanol and cyclohexanone at 60 °C.²⁷ Recently, several Pt-based electrocatalysts have been used for ECH of BA in an acidic environment.²⁸ Du *et al.*²⁹ developed a bimetallic PtRu alloy supported on carbon paper (CP), yielding a conversion of 98.6% with a CCA selectivity of 100%. The PtRu/CP alloy catalyst exhibits significantly higher conversion than that on Pt/CP (51.7%) or Ru/CP (5.8%). Moreover, a coral-shaped PtRu/Ni(OH)₂ electrocatalyst simultaneously achieved a 100% conversion of BA along with a 100% selectivity to CCA,³⁰ significantly outperforming its monometallic counterpart Pt/Ni(OH)₂ or Ru/Ni(OH)₂. Nevertheless, owing to low solubility of BA in acidic electrolyte as well as acid corrosion of the electrocatalyst, it's highly desirable to develop high-efficiency electrocatalysts for ECH of BA in alkaline media at ambient temperature and pressure.

In this work, self-supporting Ru-based electrocatalysts were *in situ* constructed on activated carbon cloths *via* a cyclic voltammetry electrodeposition technique. Through regulating the electrodeposition potential range, both the loading amount and electronic state can be facily tuned. An optimal electrocatalyst with abundant Ru/RuO₂ heterostructures exhibits unprecedented performance for ECH of BA in alkaline electrolyte. BA conversion and CCA selectivity are both greatly associated with the applied potential and KOH concentration. The ECH mechanism and the indispensable role of the Ru/RuO₂ interface were revealed by the kinetic study, *in situ* spectroelectrochemical technique and theoretical calculations.

Results and discussion

Cyclic electrodeposition enables the formation of a Ru/RuO₂ heterostructure

A series of Ru/CC electrode materials were synthesized *via* CV electrodeposition with various potential ranges as illustrated in Scheme 2. According to CV profiles during electrodeposition (Fig. S1†), there are four potentials at which an increase in deposition current can be observed in the cathodic scan: (i) the peak 'A' at 0.74 V *vs.* Ag/AgCl corresponding to the reduction of Ru³⁺ to metallic Ru ($\text{Ru}^{3+} + 3\text{e}^- \rightleftharpoons \text{Ru}$, $E = 0.74 \text{ V vs. Ag/AgCl}$);³¹ (ii) the peak 'B' observed at around 0.30–0.45 V *vs.* Ag/AgCl reflecting the potential range for underpotential-deposited hydrogen (UPD H) adsorption; (iii) the peak 'C' at –0.4 V *vs.* Ag/AgCl denoting the H absorption;³² (iv) the peak 'F'

attributable to H₂ evolution.³³ During the subsequent anodic scan, 'G' corresponds to H₂ oxidation, the peak 'D' appears due to H_{abs} desorption, the peak 'E' means H_{UPD} desorption, and 'H' indicates the oxidation of Ru³⁺ to Ru⁴⁺.³⁴

According to SEM and TEM images, it can be clearly seen that densely packed Ru particles uniformly cover the surface of CC when the electrodeposition potential lies in the range of –0.8 to 1.1 V *vs.* Ag/AgCl (Fig. 1a and S2a†). For comparison, the above-mentioned range is separated into relatively low potential (–0.8 to 0 V) and relatively high potential (0–1.1 V), respectively. Consequently, some large aggregates are observed on the deposited metal layer within either the low potential range (Fig. 1b and S2b†) or the high potential range (Fig. 1c and S2c†). As the potential moves negatively to –1.0 to 0.9 V and –1.3 to 0.6 V, the deposited metal species agglomerate significantly and disperse unevenly (Fig. S3 and S4†). As the potential changes

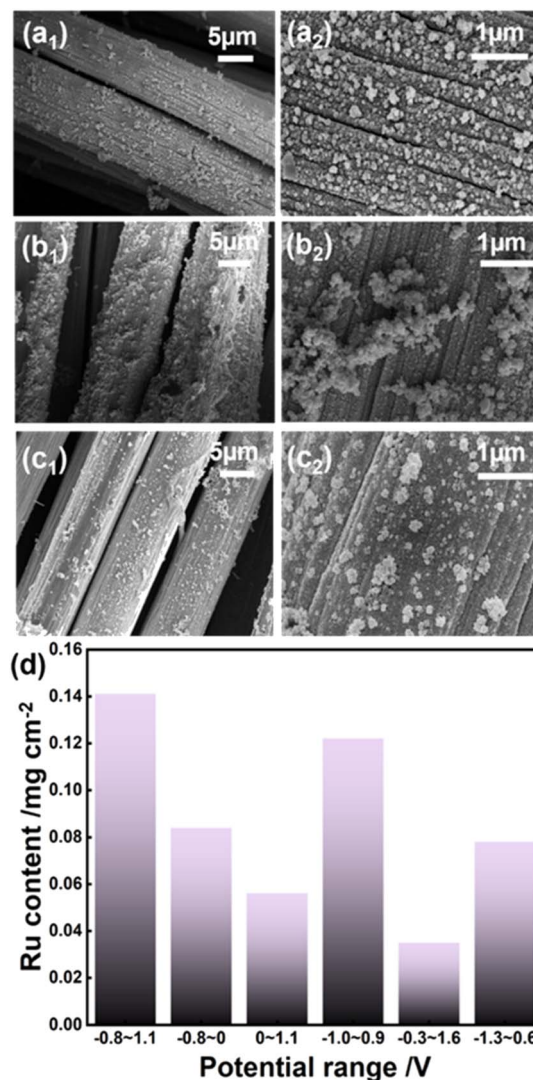
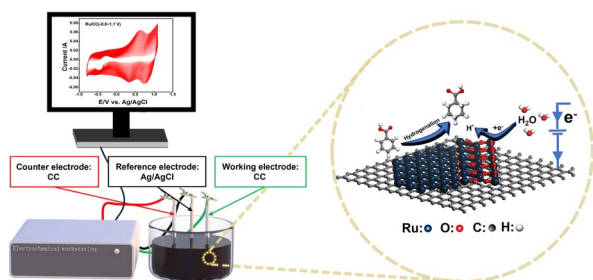


Fig. 1 SEM images of (a) Ru/CC (–0.8 to 1.1 V), (b) Ru/CC (–0.8 to 0 V), and (c) Ru/CC (0–1.1 V). (d) The mass loading of Ru for Ru/CC (–0.8 to 1.1 V), Ru/CC (–0.8 to 0 V), Ru/CC (0–1.1 V), Ru/CC (–1.0 to 0.9 V), Ru/CC (–0.3 to 1.6 V) and Ru/CC (–1.3 to 0.6 V).



Scheme 2 Schematic illustration for synthesis of Ru/CC (–0.8 to 1.1 V) *via* cyclic electrodeposition.



positively to -0.3 to 1.6 V, fewer deposits can be detected as compared to those obtained with more negative potentials.

The loading amounts of Ru prepared with different potential ranges were determined by ICP-OES. As shown in Fig. 1d, Ru/CC (-0.8 to 1.1 V) shows the maximum loading mass while Ru/CC (-0.3 to 1.6 V) exhibits the minimum content. Ru/CC (-0.8 to 1.1 V) contains 0.141 mg cm^{-2} of Ru species, which is significantly higher relative to 0.0812 mg cm^{-2} for Ru/CC (-0.8 to 0 V) and 0.0567 mg cm^{-2} for Ru/CC (0 – 1.1 V). Nevertheless, as the potential range moves negatively, the Ru content gradually decreases to 0.122 mg cm^{-2} for Ru/CC (-1.0 to 0.9 V) and 0.0787 mg cm^{-2} for Ru/CC (-1.3 to 0.6 V). Hence, an evenly distributed metal layer can form within a suitable potential range. The crystalline structures of different Ru electrode materials were analyzed by HRTEM. Interestingly, the Ru/RuO₂ heterointerface for Ru/CC (-0.8 to 1.1 V) was analyzed through interplanar spacing in HRTEM and the fast Fourier transform (FFT) diffraction pattern (shown in Fig. 2a). The interplanar spacing at the interface is different from that of Ru (101) or RuO₂ (210) at the edge. Two kinds of symmetries can be observed from FFT patterns (inset of Fig. 2a), verifying the coexistence of Ru (101) and RuO₂ (210). The tetragonal symmetry could be attributed to RuO₂,³⁵ and the lattice fringe ($d = 0.217$ nm) with an angle of 90° represents the (210) plane of RuO₂ (Fig. 2b). A lattice spacing of 0.208 nm with an angle of 60° corresponds well to the (101) plane of metallic Ru (Fig. 2c).³⁶

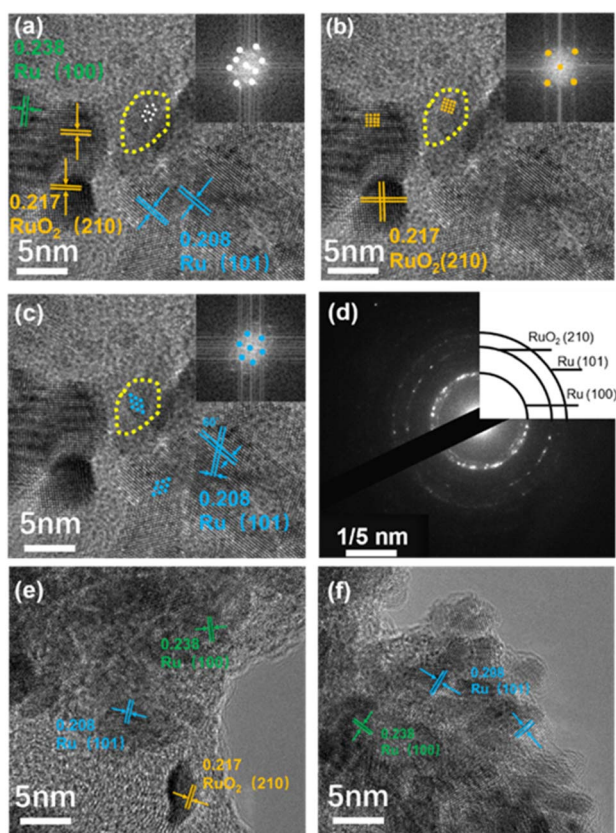


Fig. 2 (a–c) HRTEM images of Ru/CC (-0.8 to 1.1 V) and the corresponding FFT patterns. (d) SAED pattern of Ru/CC (-0.8 to 1.1 V). HRTEM images of (e) Ru/CC (0 – 1.1 V) and (f) Ru/CC (-0.8 to 0 V).

Notably, the lattice fringes of Ru and RuO₂ fuse together at the interface. For comparison, identical patterns can be observed around the interface, which are consistent with Ru and RuO₂ (Fig. 2b and c). So, it can be reasonably deduced that the Ru/RuO₂ interface has a distinctive structure with Ru (101)/RuO₂ (210) intercalation. The corresponding SAED pattern further confirms that the strongest orientations for Ru and RuO₂ are indexed to the (101) and (210) crystal planes, respectively (Fig. 2d). Moreover, Ru/CC (0 – 1.1 V) exhibits characteristic crystal planes of Ru (101), Ru (100) and RuO₂ (210), whereas Ru/CC (-0.8 to 0 V) displays typical lattice fringes for Ru (101) and Ru (100) (Fig. 2e and f). The corresponding SAED patterns verify the HRTEM analysis (Fig. S5†). The element mapping confirms the uniform distribution of Ru species (Fig. S6†).

The electronic states of the as-synthesized electrode materials were characterized by XPS (Fig. 3 and S7 and Table S2†). Except for Ru/CC (-0.8 to 0 V), all the Ru electrocatalysts are composed of metallic Ru⁰ atoms and high valent Ru⁴⁺ species, which are in perfect agreement with HRTEM analysis. The deconvolution spectra of Ru 3p for Ru/CC (-0.8 to 1.1 V) present two typical peaks at the binding energies (B.E.s) of 461.88 and 484.08 eV, respectively, which are ascribed to Ru⁰ 3p_{3/2} and Ru⁰ 3p_{1/2}.³⁷ The B.E.s located at 464.38 and 486.48 eV are attributable to Ru⁴⁺ 3p_{3/2} and Ru⁴⁺ 3p_{1/2} respectively, indicating the existence of RuO₂.³⁸ Accordingly, Ru/CC (-0.8 to 1.1 V) possesses 68.13% of Ru⁰ along with 31.87% of RuO₂. As the potential range shifts negatively, the relative proportion of Ru⁰ apparently increases (see Table S2†). Ru/CC (-0.8 to 0 V) is composed of metallic Ru⁰ atoms with the B.E.s at 461.58 and 483.88 eV, respectively. As a control, Ru/CC (0 – 1.1 V) exhibits two valence states involving Ru⁰ (64.26%) and Ru⁴⁺ (35.74%). Overall, the loading mass, dispersity and electronic state of Ru species are greatly associated with electrodeposition potential ranges.

Ru/RuO₂ promotes electrocatalytic hydrogenation of BA

First, the effects of applied potentials were investigated under the same passed charges (Fig. 4a). Within various potentials, CEA is the only intermediate product during ECH of BA. At 0 V vs. RHE, the conversion of BA is close to 17.6% , and the hydrogenated products contain 19.6% of CCA and 80.4% of CEA. As the potential drops to -0.1 V vs. RHE, either the conversion of BA or the selectivity of CCA shows a sudden increase. With the applied potential stepping from -0.1 to -0.4 V vs. RHE, both the conversion of BA and the selectivity of CCA exhibit gradual enhancement. At -0.4 V vs. RHE, a selectivity of 82.5% is realized toward CCA at a conversion of 75.7% . With the potential changing to -0.5 V vs. RHE, the conversion of BA slightly decreases to 69% . Next, five cycles of ECH at -0.2 V vs. RHE were performed. As a consequence, ECH of BA maintains a faradaic efficiency of around 47% (Fig. 4b), indicating that Ru/CC (-0.8 to 1.1 V) possesses high stability toward CCA production.

Notably, Ru-based electrode materials exhibit distinct ECH performance with different electrodeposition potential ranges (Fig. 5a). For Ru/CC (-0.8 to 1.1 V), the conversion reaches 95.8% after 120 min at $200/3$ mA cm^{-2} , giving a selectivity of 95.2% toward CCA. The catholyte after 30 min of ECH was analyzed by GC-MS (Fig. S8–S10†). It is demonstrated that the only reaction intermediate is CEA. Under the same electrolysis



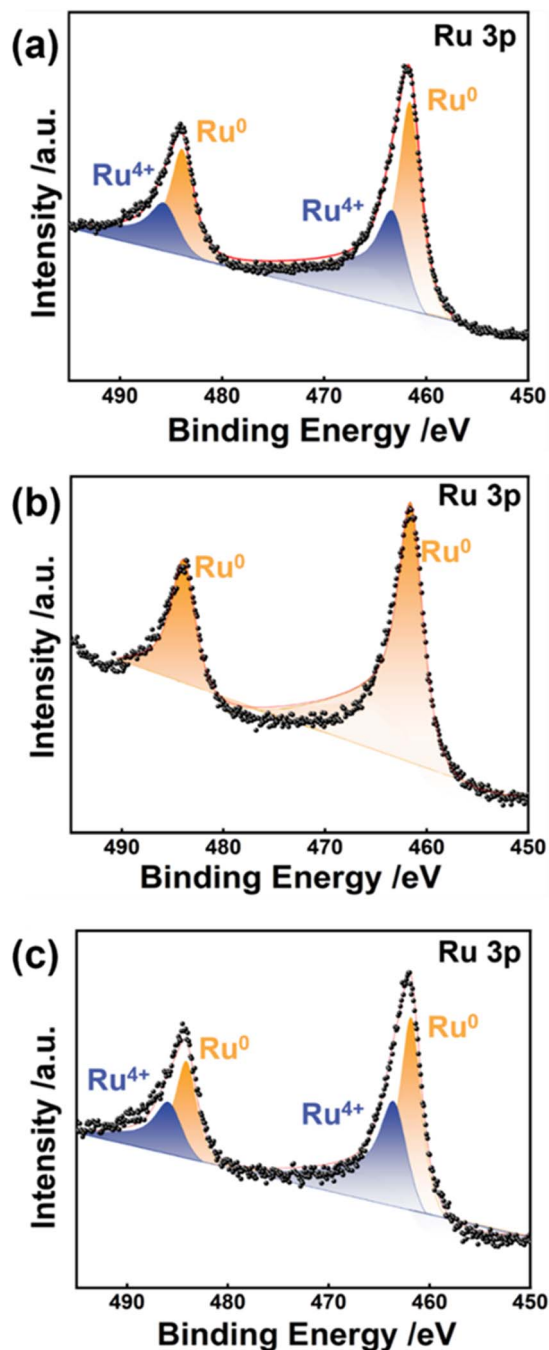


Fig. 3 Ru 3p XPS spectra of (a) Ru/CC (−0.8 to 1.1 V), (b) Ru/CC (−0.8 to 0 V), and (c) Ru/CC (0–1.1 V).

conditions, Ru/CC (−0.8 to 0 V) and Ru/CC (0–1.1 V) show a conversion of 22.5% and 33.2%, respectively, accompanied by a certain proportion of CEA. Additionally, Ru/CC (−1.0 to 0.9 V) shows a conversion of 61.6%, whereas Ru/CC (−0.3 to 1.6 V) and Ru/CC (−1.3 to 0.6 V) exhibit extremely low activity for ECH of BA. So, it can be assumed that different ECH performances may be attributable to various electrodeposition potential ranges, which affect the loading mass, dispersity and electronic state of Ru species.³⁹

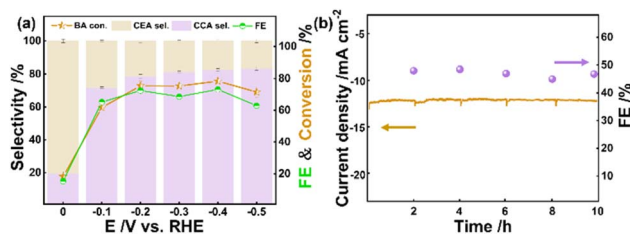


Fig. 4 (a) The catalytic performance of Ru/CC (−0.8 to 1.1 V) at different applied potentials in 0.1 M KOH with 110 C of charges passed. (b) The faradaic efficiency and current density during 10 h of ECH at −0.2 V vs. RHE; the electrolyte was refreshed every 2 h.

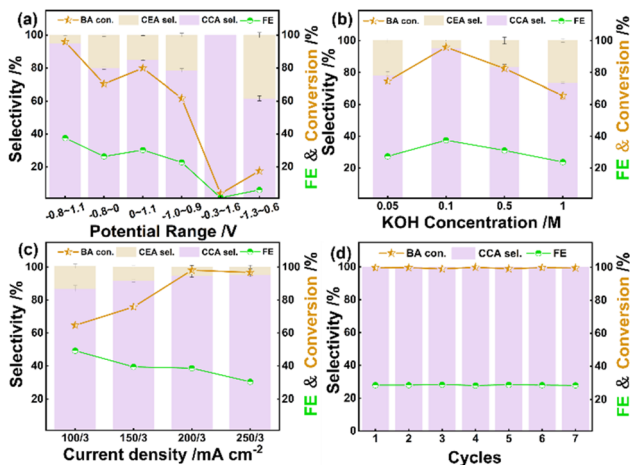


Fig. 5 (a) ECH of BA over the as-synthesized catalysts at a current density of 200/3 mA cm^{−2}; the reaction time is 120 min. (b) the conversion, FE and selectivity over Ru/CC (−0.8 to 1.1 V) at 200/3 mA cm^{−2} with different concentrations of KOH; the concentration of BA is 50 mM, and the reaction time is 120 min. (c) The conversion, FE and selectivity over Ru/CC (−0.8 to 1.1 V) at different current densities; the reaction time is 120 min. (d) Stability test over Ru/CC (−0.8 to 1.1 V) at 200/3 mA cm^{−2}; the reaction time is 180 min. The concentration of BA is 50 mM for all the tests.

Considering the concentration of electrolyte, it should be noted that ECH of BA achieves the optimal performance in 0.1 M KOH (Fig. 5b). The influence of current density and reaction time was further studied to optimize the electrolysis conditions (Fig. 5c and S11[†]). At 100/3 mA cm^{−2}, the conversion comes up to 100% within 600 min while the selectivity gradually increases from 62.0% to 100%. Apparently, the total reaction time significantly decreases to 180 min when the current density increases from 100/3 mA cm^{−2} to 200/3 mA cm^{−2}. At 250/3 mA cm^{−2}, it takes a longer time to reach a conversion of 100% relative to that at 200/3 mA cm^{−2}. This implies the fierce competition of the HER at large current density. The effect of reagent concentration was investigated at 200/3 mA cm^{−2} (Fig. S12[†]). It takes 120 min to accomplish 100% conversion for 40 mM BA. The total reaction time extends to 480 min for 80 mM BA.

Furthermore, we also employed 0.05 M H₂SO₄, 0.1 M NaOH, 0.1 M Na₂CO₃ and 0.1 M NaHCO₃ as the control electrolytes. Owing to the poor solubility of BA in H₂SO₄ solution, we used



relatively low concentration of BA (5 mM). As can be seen in Fig. S13 and Table S3,[†] the conversion of BA reaches 91.67% at 180 min in acidic media, giving 100% selectivity toward CCA. In NaOH electrolyte, the conversion reaches 93.3% with a CCA selectivity of 95.65%, which is a little lower than that in KOH electrolyte. Moreover, ECH of BA in Na₂CO₃ and NaHCO₃ achieves the conversions of 80.0% and 82.3%, respectively, accompanied by small fractions of CEA. Considering the influence of the aromatic ring structure of benzoic acid derivatives, ECH of *p*-methyl benzoic acid was studied at 200/3 mA cm⁻² in 0.1 M KOH. As a consequence, a conversion of 83.3% was achieved at 10 h with 100% selectivity toward 4-methyl cyclohexane carboxylic acid. Clearly, ECH of *p*-methyl benzoic acid is more time-consuming than that of BA. Future studies will focus on the investigation of the catalytic mechanism and the optimization of electrocatalysts for broad benzoic acid derivatives.

HRTEM and XPS analysis demonstrate that Ru/CC (-0.8 to 1.1 V) is composed of two phases (Ru and RuO₂) while Ru (-0.8 to 0 V) only contains metallic Ru⁰ species. To elucidate the role of the Ru/RuO₂ heterostructure, two catalysts involving Ru/CC (-0.8 to 1.1 V) with 100 CV cycles and Ru (-0.8 to 0 V) with 225 CV cycles were compared at the same current density. To ensure the same electrodeposition time as that of Ru/CC (-0.8 to 1.1 V), the number of CV cycles was increased to 225 for Ru/CC (-0.8 to 0 V). It is also worth noting that the selectivity toward CEA changes slightly on prolonging the reaction time (Fig. S14[†]), which indicates that full hydrogenation of BA proceeds slowly over Ru/CC (-0.8 to 0 V). For comparison, 100% selectivity toward CCA is rapidly realized over Ru/CC (-0.8 to 1.1 V) (Fig. S11c[†]). For ECH of CEA, Ru/CC (-0.8 to 1.1 V) also exhibits much higher activity as compared to Ru/CC (-0.8 to 0 V) (Fig. S15[†]). Accordingly, it can be reasonably deduced that Ru/CC (-0.8 to 1.1 V) embedded with Ru/RuO₂ heterointerfaces facilitates ECH of BA into CCA.

The stability test for Ru/CC (-0.8 to 1.1 V) was conducted at 200/3 mA cm⁻² *via* seven cycles of ECH, in which each cycle proceeds for 180 min. As can be seen in Fig. 5d, Ru/CC (-0.8 to 1.1 V) maintains outstanding performance without apparent loss of activity. The spent Ru/CC (-0.8 to 1.1 V) shows identical surface morphology, Ru loading mass (0.138 mg cm⁻²) and valence distribution (Ru⁰ accounts for 63.5%) (Fig. S16–S18[†]) to the as-synthesized sample. So, it is demonstrated that Ru/CC (-0.8 to 1.1 V) possesses outstanding durability during ECH of BA in alkaline electrolyte.

Mechanism study for ECH of BA

According to LSV profiles of Ru-based catalysts (Fig. S19, Table S4[†]), it can be found that Ru/CC (-0.8 to 1.1 V) shows a minimum overpotential (η_{10}) of ~110 mV with 50 mM BA added, which implies that electrochemical reduction of BA is kinetically favorable on Ru/CC (-0.8 to 1.1 V). What's more, addition of BA increases the overpotential of the Ru-based electrocatalyst, indicating that adsorption of BA on the electrode surface hinders hydrogen adsorption and evolution.^{40,41} As shown in Fig. 6a, Ru/CC (-0.8 to 1.1 V) exhibits the smallest Tafel slope of 176 mV dec⁻¹, which signifies the best performance for ECH of BA. Electrochemical impedance was

employed to demonstrate the electron transfer efficiency of the catalysts (Fig. 6b). Ru/CC (-0.8 to 1.1 V) shows the smallest R_{ct} (21 Ω cm⁻²) among the investigated samples, suggesting the most efficient electron transfer between the electrode and electrolyte.

Without BA added, the CV curve of Ru/CC (-0.8 to 1.1 V) exhibits a shoulder wave at around 0.1 V vs. RHE in the cathodic scan as well as a distinct peak at 0.2 V vs. RHE in the anodic scan, which can be attributed to H⁺ adsorption and desorption. On increasing the concentration of BA, both cathodic and anodic currents progressively decrease owing to competitive adsorption between H₂O and BA (Fig. 6c).⁴² To unveil the reaction mechanism, reaction rates were assessed at different initial concentrations of BA. As a result, a negative reaction order of -1.02 was obtained for ECH of BA in 0.1 M KOH over Ru/CC (-0.8 to 1.1 V), which is indicative of a Langmuir–Hinshelwood (L–H) mechanism (Fig. 6d).^{43–45} Combined with the CV results, it is indicated that ECH of BA consumes active H⁺ on the catalyst surface.

To further explore the reaction mechanism, *in situ* Raman spectroscopy was performed with different applied potentials. As shown in Fig. 7a, Raman spectra collected for negatively charged BA on the catalyst surface exhibit three distinguishable peaks:^{46,47} (i) aromatic ring vibration $\nu(\text{C-C})_{\text{ring}}$ at 1600 cm⁻¹, (ii) carboxylate group vibration $\nu(\text{COO}^-)_{\text{sym}}$ at 1350 and 1390 cm⁻¹, (iii) ring breathing at 1005 cm⁻¹. On applying positive potentials (0.1–0.2 V), the above-mentioned typical peaks are intensified on the catalyst surface, which implies that a positive potential enhances the adsorption strength of a negatively charged reactant. It is worth noting that the intensities of these typical bands are greatly weakened at 0 V vs. RHE. Combined with ECH performance (see Fig. 4a), the decreased signal could be explained by the hydrogenation of benzoate. Moreover, these

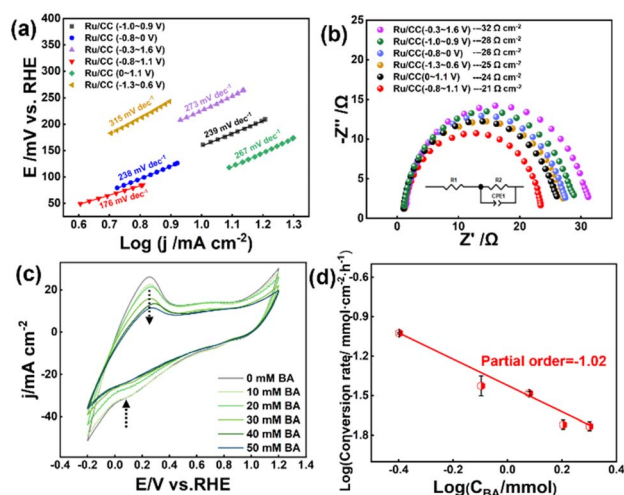


Fig. 6 (a) Tafel slopes and (b) Nyquist plots of Ru/CC (-0.8 to 1.1 V), Ru/CC (-0.8 to 0 V), Ru/CC (0–1.1 V), Ru/CC (-1.0 to 0.9 V), Ru/CC (-1.3 to 0.6 V) and Ru/CC (-0.3 to 1.6 V). (c) CV curves of Ru/CC (-0.8 to 1.1 V) with different concentrations of BA. (d) Kinetic study for ECH of BA over Ru/CC (-0.8 to 1.1 V); the conversion of BA is controlled less than 15%.



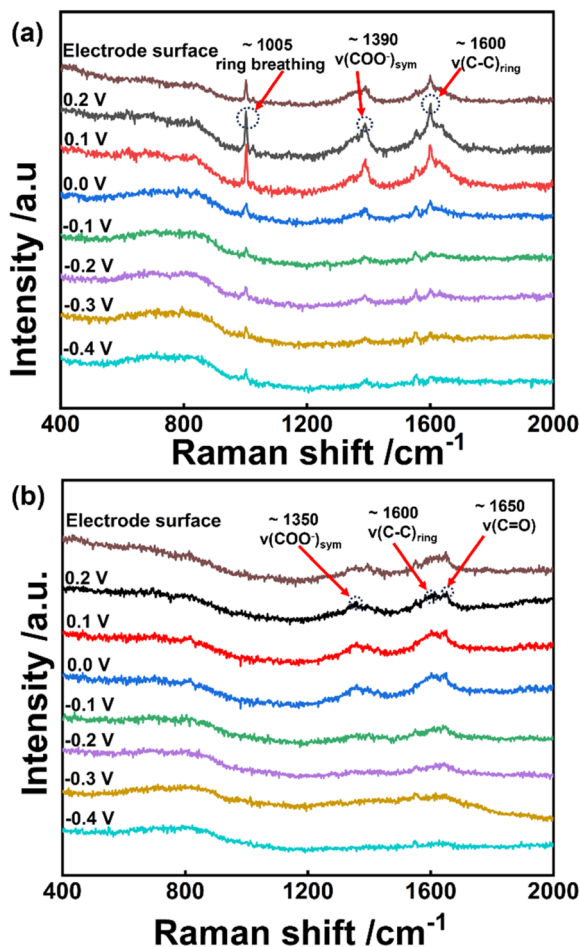


Fig. 7 *In situ* Raman spectra collected on the Ru/CC (−0.8 to 1.1 V) surface for (a) benzoate and (b) cyclohexene carboxylate in 0.1 M KOH at different applied potentials.

peaks almost remain unchanged within −0.1 to −0.4 V. To further explore the effect of applied potentials, *in situ* Raman spectroscopy was performed on non-catalytic carbon cloth (Fig. S20†). The typical peaks basically remain unchanged in the potential range of 0.2 to −0.4 V. This implies that the peak intensity will not decrease significantly if no reaction occurs. For comparison, the characteristic bands for cyclohexene carboxylate show a sudden decrease at −0.1 V, and gradually descend as the potential negatively shifts from −0.1 to −0.4 V (Fig. 7b). So, it is indicated that the hydrogenation of cyclohexene carboxylate requires a more negative potential than that of benzoate.

To further reveal the role of the Ru/RuO₂ heterostructure during ECH, the adsorption energies of potassium benzoate and potassium cyclohexene carboxylate were calculated on the Ru (101)/RuO₂ (210) model (Fig. 8a, b, S21 and S22†). As shown in Fig. 8a, potassium benzoate adsorbing on the Ru (101) surface of the heterostructure model (*Ru/RuO₂) exhibits a superior adsorption energy of −3.57 eV, followed by the figures for interface and Ru/RuO₂* sites (−2.24 eV in comparison with −2.41 eV). Fig. 8b demonstrates that the adsorption energy of potassium cyclohexene carboxylate on the interface

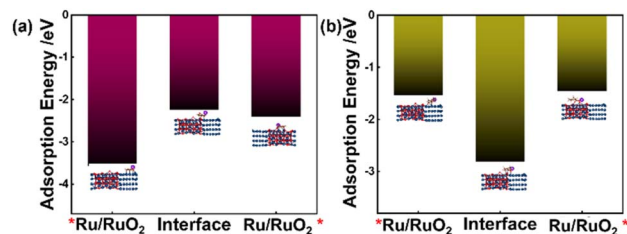


Fig. 8 Adsorption energy of (a) potassium benzoate and (b) potassium cyclohexene carboxylate on different sites of the Ru (101)/RuO₂ (210) model.

(−2.81 eV) is much larger than that on *Ru/RuO₂ (−1.53 eV) or Ru/RuO₂* (−1.45 eV). It is noteworthy that the adsorption energy of potassium cyclohexene carboxylate on the interface is only −0.57 eV (Fig. S23†), which is much smaller than that of potassium cyclohexene carboxylate (−2.81 eV). Accordingly, the Ru site facilitates the adsorption of potassium benzoate while the interface promotes the adsorption of potassium cyclohexene carboxylate. DFT calculations reveal that the critical role of the Ru/RuO₂ heterointerface is to promote the adsorption strength of the intermediate potassium cyclohexene carboxylate, which is in accordance with ECH results (Fig. S15†).

Conclusion

A facile CV electrodeposition was adopted to fabricate self-supporting Ru-based electrocatalysts on activated carbon cloths through regulating the potential ranges. Both the loading amount and electronic state can be facilely tuned. The optimal electrocatalyst with abundant Ru/RuO₂ heterostructures exhibits unprecedented performance for ECH of BA in alkaline electrolyte. In an alkaline environment, the Ru/RuO₂ catalyst achieves high BA conversion (100%) and selectivity towards CCA (100%) within 180 min at 200/3 mA cm^{−2}, showing excellent reusability and long-term stability. 1-Cyclohexenecarboxylic acid (CEA) was identified as the reaction intermediate, whose the selectivity is governed by the applied potential. Kinetic studies demonstrate that ECH of BA over Ru/RuO₂ follows a Langmuir–Hinshelwood (L–H) mechanism. *In situ* Raman spectroscopy and theoretical calculations unravel that the Ru/RuO₂ interface enhances the adsorption strength of cyclohexene carboxylate, thereby facilitating the production of fully hydrogenated cyclohexene carboxylate. This work provides a deep understanding of the ECH pathway of BA in alkaline media and gives a new methodology to fabricate heterostructure electrocatalysts.

Author contributions

Zifan Cao carried out the synthesis and characterization, performed electrochemical experiments and wrote the paper; Chenhui Wang undertook DFT calculations for the article; Yifan Sun and Menghui Liu assisted with electrocatalytic experiments and participated in discussion; Wei Li and Jinli Zhang acquired funding; Yan Fu conceived the project and supervised the work.



All authors have approved the published version of the manuscript.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Notes and references

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