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Nickel–cobalt oxalate as an efficient non-precious electrocatalyst for an improved alkaline oxygen evolution reaction†

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The quest for developing next-generation non-precious electrocatalysts has risen in recent times. Herein, we have designed and developed a low cost electrocatalyst by a ligand-assisted synthetic strategy in an aqueous medium. An oxalate ligand-assisted non-oxide electrocatalyst was developed by a simple wet-chemical technique for alkaline water oxidation application. The synthetic parameters for the preparation of nickel–cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$) were optimized, such as the metal precursor (Ni/Co) ratio, oxalic acid amount, reaction temperature, and time. Microstructural analysis revealed a mesoporous block-like architecture for nickel–cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$). The required overpotential of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ for the alkaline oxygen evolution reaction (OER) was found to be 330 mV for achieving 10 mA $\text{cm}_{\text{geo}}^{-2}$, which is superior to that of NiC_2O_4 , CoC_2O_4 , NiCo_2O_4 and the state-of-the-art RuO_2 . The splendid performance of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ was further verified by its low charge transfer resistance, impressive stability performance, and 87% faradaic efficiency in alkaline medium (pH = 14). The improved electrochemical activity was further attributed to double layer capacitance (C_{dl}), which indefinitely divulged the inferiority of NiCo_2O_4 compared to $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ for the alkaline oxygen evolution reaction (OER). The obtained proton reaction order (p_{RHE}) was about 0.80, thus indicating the proton decoupled electron transfer (PDET) mechanism for OER in alkaline medium. Post-catalytic investigation revealed the formation of a flake-like porous nanostructure, indicating distinct transformation in morphology during the alkaline OER process. Further, XPS analysis demonstrated complete oxidation of Ni^{2+} and Co^{2+} centres into Ni^{3+} and Co^{3+} , respectively under high oxidation potential, thereby indicating active site formation throughout the microstructural network. Additionally, from BET-normalised LSV investigation, the intrinsic activity of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ was also found to be higher than that of NiCo_2O_4 . Finally, $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ delivered a TOF value of around $3.28 \times 10^{-3} \text{ s}^{-1}$, which is 5.56 fold that of NiCo_2O_4 for the alkaline OER process. This report highlights the unique benefit of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ over NiCo_2O_4 for the alkaline OER. The structure–catalytic property relationship was further elucidated using density functional theory (DFT) study. To the best of our knowledge, nickel–cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$) was introduced for the first time as a non-precious non-oxide electrocatalyst for alkaline OER application.

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1. Introduction

Water splitting is becoming the most promising approach for hydrogen fuel production and energy storage.^{1–5} The

requirement of a large overpotential for four electron trajectories all through the oxygen evolution reaction (OER) restricts the application in bulk mode.^{6,7} The ideal category of catalyst for OER is still RuO_2 in alkaline and IrO_2 in acidic solution.⁸ However, the cost of scalable design and availability in nature limit the use of such precious elements in day to day industrial applications. Non-precious transition metal based electrocatalysts particularly oxide and hydroxide have attracted great interest particularly owing to their simple synthetic strategy, extraordinary OER activity, and cheap market price.^{9,10} The improved alkaline OER performance can be attributed to the corrosion resistant physiochemical properties of the working electrode leading to substantial stability in alkaline medium.¹¹ However, non-precious non-oxide transition metal based

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electrocatalysts are prone to quick *in situ* electrochemical active surface transformation, thereby rendering rapid kinetics, a low overpotential barrier and high stability. The performance of this catalytic system originates from the presence of microstructural defects, porosity and pore architecture, conductivity and oxidation (chemical) states of active elements.^{12–15} For example, non-oxide electrocatalysts based on cobalt (Co) are well-known for their metallic nature, which encourages their wide practical application.^{16–18} There are quite a few literature studies on non-precious non-oxide electrocatalysts for alkaline OER application, such as Co_4N , $(\text{Co}_{0.7}\text{Fe}_{0.3})_2\text{B}$, $(\text{Co}_x\text{Fe}_{1-x})_2\text{P}$ and $\text{Co}_{0.85}\text{Se}/\text{Co}_9\text{Se}_8$.^{19–22} Thus, non-precious non-oxide electrocatalysts have attracted great attention for the alkaline OER.

A ligand assisted soft-chemical strategy has been employed for the design of electrocatalysts in recent years. Feng *et al.* demonstrated an ionic liquid assisted microwave irradiation technique for the preparation of nickel–cobalt fluoride for water electrolysis.²³ Next, hierarchical hydrous cobalt phosphate micro-flower was employed as an efficient electrocatalyst for the alkaline OER.²⁴ Recently, our group has discussed the superior OER electrocatalytic activity of $\text{NiCo}_2\text{O}_x\text{S}_{4-x}$ over NiCo_2O_4 .²⁵ On the other hand, transition metal based oxalate is renowned as a carbon sink, suggesting it as a more sustainable way to design energy storage materials. Numerous bio- and artificial-synthesis routes have already been reported for the synthesis of oxalate anion ($\text{C}_2\text{O}_4^{2-}$) from CO_2 .²⁶ Therefore, utilization of oxalate-based materials is an impactful way from the clean and green synthesis viewpoint. To our knowledge, there are only few reports on oxalate based materials for OER applications. For instance, cobalt oxalate has shown promising activity as an electrocatalyst for alkaline OER applications.^{27,28} Liu *et al.* reported the most convenient and widely used solvothermal synthetic procedure for the preparation of $\text{CoC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$ with a tuneable rod and 3-D polyhedron like morphology. Post-catalytic transformation into CoOOH species resulted in active sites for the alkaline OER with 436 and 492 mV overpotential (for 10 $\text{mA cm}_{\text{geo}}^{-2}$) for the micro-rod and 3-D polyhedron respectively.²⁹ However, the high overpotential and sluggish kinetics of the catalyst are identified as major limitations for practical implementation in large scale production. To date, no such reports were found for nickel–cobalt oxalate, which might be an excellent electrocatalyst for alkaline water oxidation. In addition, fabrication of non-precious non-oxide transition metal based electrocatalysts that meet the requirement of 10 $\text{mA cm}_{\text{geo}}^{-2}$ current density, thereby reaching a benchmark of 12% solar-to-hydrogen efficiency, is a challenging task.

Herein, we have demonstrated an oxalic acid assisted non-precious non-oxide transition metal (nickel and cobalt) based electrocatalyst for the alkaline OER. Nickel–cobalt oxalate was primarily synthesized by a wet chemical technique, followed by annealing to form nickel–cobalt oxide. The preparation technique was optimised in terms of Ni/Co ratio, oxalic acid amount, temperature and processing time. The overpotential requirement for the block-like $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ was found to be 330 mV for reaching the benchmark of 10 $\text{mA cm}_{\text{geo}}^{-2}$. The improved performance of the catalyst was further verified by low charge transfer resistance, long term stability and 87% faradic

efficiency. Interestingly, the control studies and the change in double layer capacitance (C_{dl}) because of electrochemical preconditions clearly suggest that the alkaline OER catalytic performance of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ is superior to that of its high temperature oxide counterpart *i.e.* NiCo_2O_4 . The structure–catalytic property relationship was further elucidated by density functional theory (DFT) study which was performed to get insight into the improved electrochemical performance of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ compared to the CoC_2O_4 , NiC_2O_4 and NiCo_2O_4 , and atomic interpretation was also proposed from computed results. In addition, electrocatalytic measurements in electrolyte solutions with different pH values were performed to understand the mechanistic details of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ for the alkaline OER. This investigation exclusively unlocks a strategy to fabricate oxalate ligand assisted non-precious nickel–cobalt based bimetallic electrocatalysts for water oxidation application in alkaline medium.

2. Results and discussion

2.1 Structural characterization

The synthesis of nickel oxalate (NiC_2O_4), cobalt oxalate (CoC_2O_4) and nickel–cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$) samples was performed by a one-pot method using metal nitrate salts and oxalic acid in aqueous media. For comparative study, nickel cobalt oxide (NiCo_2O_4) was prepared by calcination of nickel–cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$) at 350 °C for 2 h (see the details in the Experimental and Characterization section, ESI, Table S1†). The elemental phase was examined by the X-ray diffraction (XRD) method. As shown in Fig. 1a, the peak position resembles with the appearance of cobalt oxalate hydrate ($\text{CoC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$; ICDD no. #25-0250) and nickel oxalate hydrate ($\text{NiC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$; ICDD no. #01-0299). The XRD pattern of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ is quite identical to that of phase-pure cobalt oxalate hydrate, thereby indicating the existence of substituted nickel at some portion of cobalt sites throughout the mixed-metal oxalate crystal framework.³⁰ Furthermore, the XRD pattern of a physical mixture comprising CoC_2O_4 and NiC_2O_4 was examined, revealing doublet peaks at 35.1° and 35.59° that can be labelled as CoC_2O_4 and NiC_2O_4 respectively. However, $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ shows a single broad peak around 34.9°, manifesting the (022) plane of CoC_2O_4 . The resemblance of XRD patterns between $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ and CoC_2O_4 further supports the existence of substituted nickel at some portion of cobalt sites throughout the mixed-metal oxalate crystal framework. In addition, the (022) peak of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ shifts to a lower 2θ value w.r.t CoC_2O_4 , revealing [022] oriented unit cell expansion due to the presence of the nickel component within the composite network (see Fig. S1, ESI†).³¹

Thermal stability of the as-prepared sample was corroborated by DTA-TG analysis. Fig. 1b shows an endothermic peak around 205 °C, indicating the removal of lattice water, resulting in 19% weight loss. The exothermic peak around 294 °C represents the decomposition of oxalate ligand with a 57% weight loss compared to the as-prepared mixed oxalate sample.³² No further weight loss is observed in the 300 to 600 °C temperature range. Thus, for the preparation of the NiCo_2O_4



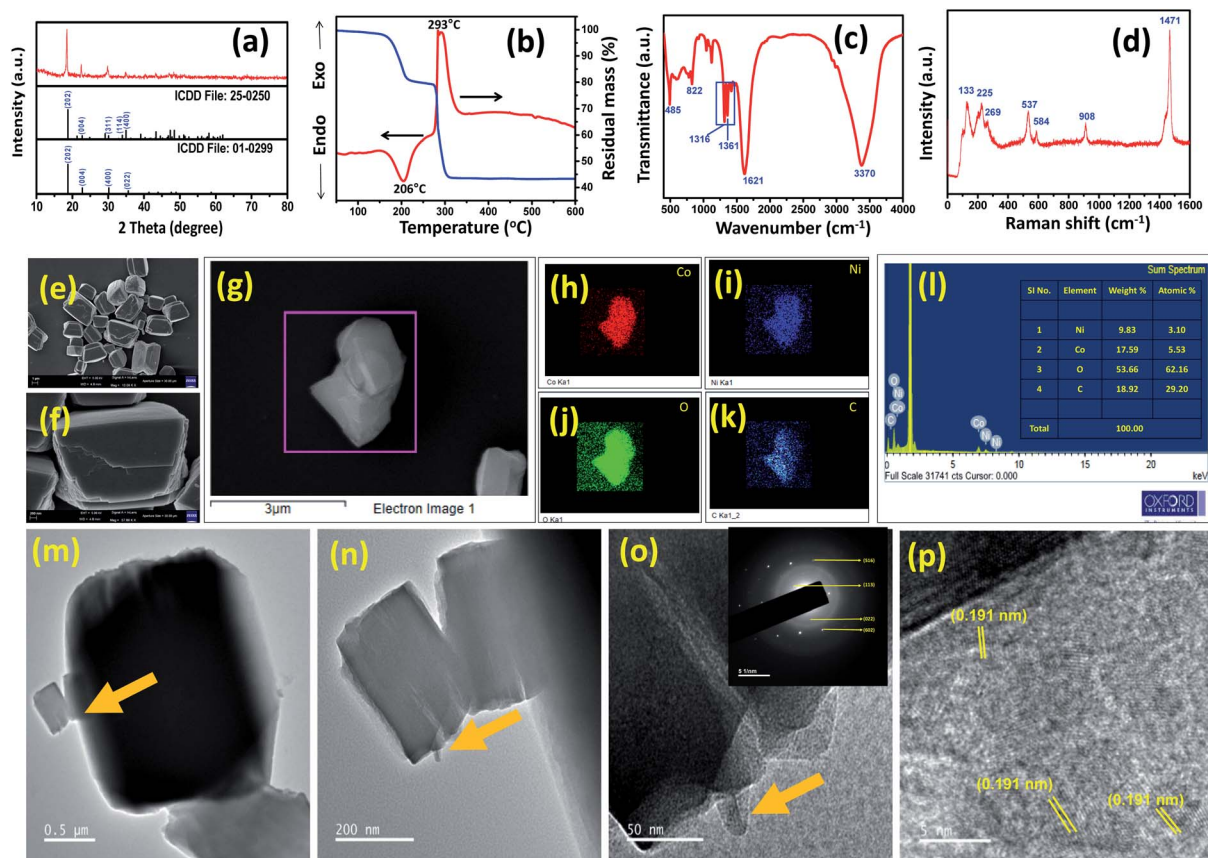


Fig. 1 (a) PXRD with the standard patterns of $\text{NiC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$ (ICDD no. 01-0299) and $\text{CoC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$ (ICDD no. 25-0250), (b) DTA-TG, (c) FTIR and (d) Raman pattern of the as-prepared nickel-cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$) sample. Morphological study including (e and f) FESEM images, (g–k) elemental mapping, (l) EDS study, (m–o) TEM images (the inset showing SAED) and (p) HRTEM image of the as-prepared nickel-cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$) sample.

sample, the as-prepared $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ was calcined at 350°C . Fourier-transform infrared (FTIR) spectroscopy study was carried out for further structural characterization of the sample (Fig. 1c). The FTIR peak positioned around 485 cm^{-1} is assigned to metal–oxygen vibration mode. The C–O vibration mode is obtained around 1316 (symmetric) and 1361 cm^{-1} (asymmetric) along with a strong vibrational band at 1621 cm^{-1} , which is a signature characteristic peak for carbonyl stretching vibrational mode, confirming the presence of the oxalate moiety in the as-prepared sample.²⁹ The broad vibrational band around 3370 cm^{-1} is attributed to water molecules, while 822 cm^{-1} is to asymmetric O–C–O vibrational mode. Raman measurements were also performed to obtain more structural information (Fig. 1d). The intense peak around 912 cm^{-1} is attributed to C–C stretching mode, which is IR-silent. The presence of C–C–O bending mode is concluded from the observed 532 cm^{-1} peak. In addition, appearance of peaks around 225 and 266 cm^{-1} is ascribed to M–O stretching and O–M–O ring-bending modes respectively, where M stands for the metal centre (cobalt and nickel).^{16,33} The XPS technique was employed for the analysis of chemical state and probable elemental composition of the as-prepared sample (see Fig. S2, ESI†). The C 1s spectrum exhibits peaks around 286.48 , 287.69 and 290.3 eV particularly

for C–O, C=O and C–F respectively (see Fig. S2a, ESI†). The appearance of 290.3 eV binding energy could originate from Nafion residue which is used as a binder.^{34,35} As shown in Fig. S2b (ESI†), peaks at 782.93 and 798.87 eV can be designated as Co $2p_{3/2}$ and Co $2p_{1/2}$, respectively with a clear separation of 15.94 eV , confirming the presence of Co^{2+} .³⁶ The satellite signals around 786.79 and 803.86 eV further support the same (see Fig. S2b, ESI†). The presence of Ni^{2+} can be suspected from the binding energy peaks around 857.71 and 875.35 eV assigned to Ni $2p_{3/2}$ and Ni $2p_{1/2}$ respectively, along with satellite peaks at 862.61 and 881.14 eV (see Fig. S2c, ESI†).³⁷ The narrow scan of the O 1s peak is deconvoluted into two peaks at 533.62 and 534.65 eV representing C–O and lattice water molecules, respectively (see Fig. S2d, ESI†).³⁸ Inductively coupled plasma-atomic emission spectroscopy (ICP-AES) was also employed to get quantitative information regarding the elemental composition of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$. The Ni/Co ratio is found to be $1 : 2.018$ (see Table S2, ESI†).

2.2 Morphological characterization

Microstructural analysis was performed by FESEM and TEM investigation. The displayed FESEM image presents block shaped nanostructures of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (Fig. 1e and f). Close



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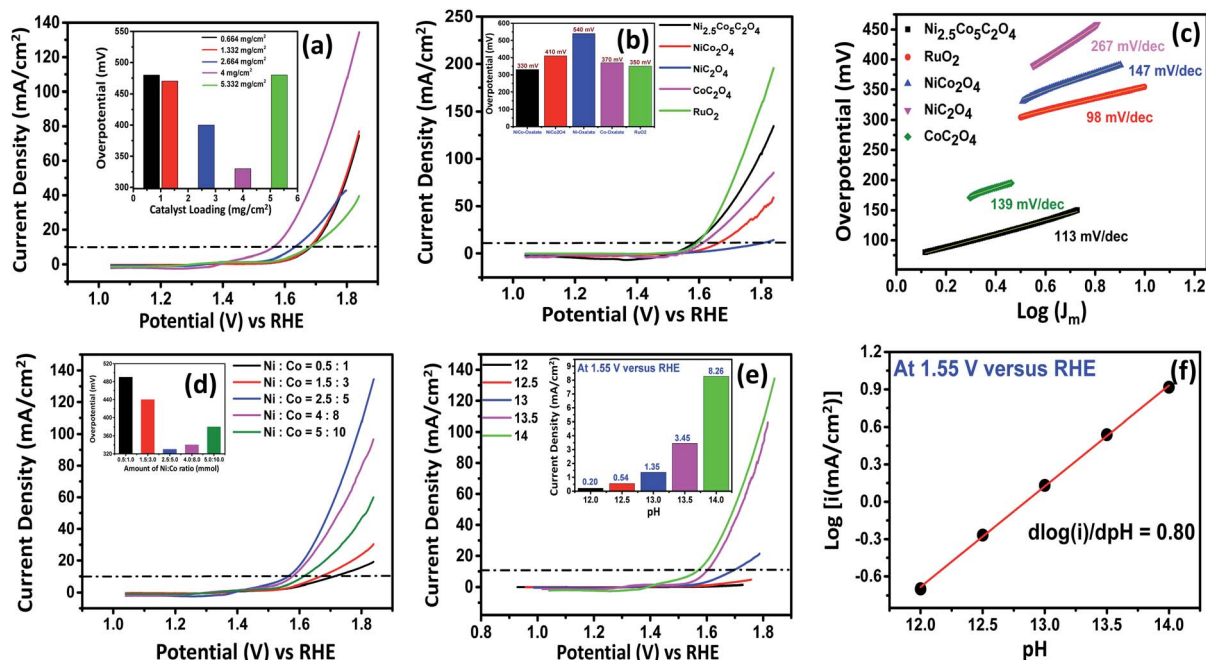


Fig. 2 (a) iR -corrected LSV scan for different loading amounts for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$, (b) iR -corrected LSV scan for different electrocatalysts [the inset showing the overpotential value to reach the benchmark of $10 \text{ mA cm}_{\text{geo}}^{-2}$], (c) Tafel slope analysis of different electrocatalysts, (d) iR -corrected LSV scan for different Ni/Co ratios, (e) LSV scan and (f) linear fit of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ with the variation of pH.

vs. RHE for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$, whereas NiCo_2O_4 delivered 1.64 V vs. RHE for the same. It unequivocally implies the higher geometrical OER activity of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ than NiCo_2O_4 . To gather further support, double layer capacitance (C_{dl}) was calculated that was indefinitely assumed as a non-destructive parameter for the electrochemical active surface area (ECSA) calculations.⁴³ The C_{dl} value for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ is calculated to be 1.31 mF cm^{-2} , whereas NiCo_2O_4 shows 2.84 mF cm^{-2} just before precondition (Fig. 3a and d). However, the C_{dl} value is significantly increased up to 10.61 mF cm^{-2} after electrochemical precondition for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$, while the corresponding value for NiCo_2O_4 remains nearly the same at 2.93 mF cm^{-2} (Fig. 3b and e). Thus development of a higher number of

accessible active sites is observed during electrochemical precondition for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ compared to the NiCo_2O_4 sample (Fig. 3c and f).⁴⁴ This also suggests more ECSA and thus more surface coverage (θ) that provides a higher value of current (j) according to the following expression:

$$j = [\text{OH}^-] \times \theta \times e^{-\frac{\Delta G}{RT}}$$

where the current (j) value is directly proportional to surface coverage (θ) involving $^*\text{OOH}$ or $^*\text{OH}$ sites, $[\text{OH}^-]$ concentration and exponential factor which actually depends on the adsorbed surface of intermediate species for the alkaline OER.⁴⁰ The change in C_{dl} is also estimated for CoC_2O_4 and NiC_2O_4 samples that further supports the better electrochemical performance of CoC_2O_4 compared to NiC_2O_4 for the alkaline OER (see Fig. S14 and S15, ESI†).

To further study the electrochemical properties, electrochemical impedance spectroscopy (EIS) was employed to evaluate the charge transfer resistance (R_{ct}) and transport kinetics. As expected, $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ delivers a lower R_{ct} value of about 6.34 ohm in comparison with NiCo_2O_4 (26.14 ohm), thereby indicating the occurrence of rapid charge transport phenomena for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (Fig. 4a).⁴⁵ Likewise, NiC_2O_4 shows a higher R_{ct} value than CoC_2O_4 , justifying the higher conducting behaviour of CoC_2O_4 in alkaline medium (see Fig. S16, ESI†). To study the intrinsic activity of the drop-cast catalyst, BET normalized electrochemical performance was already reported as the most convenient analytical technique by Shao-Horn and co-workers.⁴⁶ As shown in Fig. 4b, the nitrogen adsorption-desorption isotherm displays a type-IV isotherm and H-3 hysteresis, reflecting a mesoporous network and slit-like pore geometry

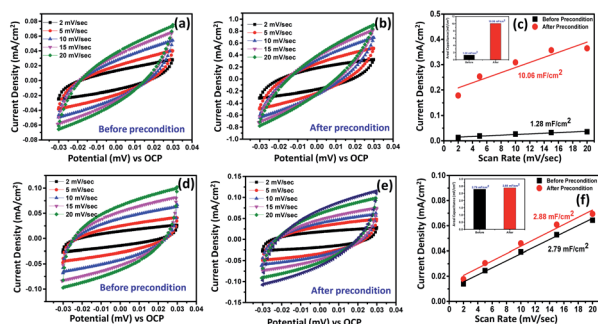


Fig. 3 CV curves recorded at different scan rates to estimate the double layer capacitance (C_{dl}) for (a and b) $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ and (d and e) NiCo_2O_4 . Linear fit for the estimation of double layer capacitance (C_{dl}) from the DLC current vs. scan rate plot for (c) $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ and (f) NiCo_2O_4 .



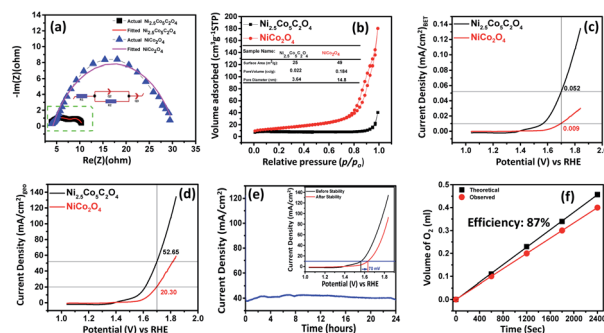


Fig. 4 (a) Nyquist plots of different electrocatalysts, (b) N_2 adsorption-desorption isotherm of $Ni_{2.5}Co_5C_2O_4$ and $NiCo_2O_4$ samples, catalytic performance in terms of (c) BET surface area normalised LSV plot and (d) geometrical area normalised LSV plot for $Ni_{2.5}Co_5C_2O_4$ and $NiCo_2O_4$ samples, (e) stability study of the best catalyst $Ni_{2.5}Co_5C_2O_4$ [the inset showing the overpotential value to reach the benchmark of $10 \text{ mA cm}_{\text{geo}}^{-2}$], and (f) analysis of faradaic efficiency for the $Ni_{2.5}Co_5C_2O_4$.

respectively.⁴⁷ The BET surface area is found to be 25 and $49 \text{ m}^2 \text{ g}^{-1}$ for $Ni_{2.5}Co_5C_2O_4$ and $NiCo_2O_4$ respectively, which could be attributed to thermal decomposition of oxalate ligand during calcination treatment at high-temperature. Fig. 4c presents the comparative survey of intrinsic alkaline OER performance for both the samples. From the BET normalized LSV plot, $Ni_{2.5}Co_5C_2O_4$ shows a current density of about $0.052 \text{ mA cm}_{\text{BET}}^{-2}$, which is 5.7 fold greater than that of $NiCo_2O_4$ at 1.7 V vs. RHE. In addition, the geometrical performance of $Ni_{2.5}Co_5C_2O_4$ is also superior to that of $NiCo_2O_4$ for the OER in alkaline medium (Fig. 4d). This result indicates the superiority of $Ni_{2.5}Co_5C_2O_4$ over its widely used high temperature calcined product of $NiCo_2O_4$ from both intrinsic and geometrical aspects for the OER under alkaline conditions.⁴⁸ To evaluate the applicability for long-term usage, the stability test of the $Ni_{2.5}Co_5C_2O_4$ catalyst was performed for 24 hours by the chronoamperometry technique (Fig. 4e). Alkaline OER performance is relatively stable up to the initial 12 hours of the experiment; thereafter, the performance slightly decreased with time. After the stability test, the $Ni_{2.5}Co_5C_2O_4$ sample achieves $10 \text{ mA cm}_{\text{geo}}^{-2}$ with around 350 mV overpotential, which is roughly 20 mV more than that of the as-prepared electrocatalyst (inset of Fig. 4e). Interestingly, the Ni/Co ratio is found to be 1 : 2.38, indicating mechanical inertness of the electrocatalyst even after the long-term 24 h stability test at pH = 14.

Finally, faradaic efficiency was studied and the obtained value for $Ni_{2.5}Co_5C_2O_4$ is 87% (Fig. 4f). It indeed denotes sufficiently high activity of the electrocatalyst for alkaline water oxidation reaction. Turnover frequency (TOF) is one of the essential parameters to examine the intrinsic performance of the electrocatalyst. The TOF is expressed as follows:

$$\text{TOF} = j \frac{SA_{\text{geo}}}{4Fn}$$

where F , n and j denote the Faraday constant (96485 C mol^{-1}), moles of the catalytically active species and geometrical area (SA_{geo}) normalized current density (mA cm^{-2})_{geo}.⁴⁹ The amount

of active elemental species was calculated from ICP-AES study (see Tables S2 and S3, ESI†). The TOF value is found to be $3.28 \times 10^{-3} \text{ s}^{-1}$ for $Ni_{2.5}Co_5C_2O_4$, whereas the value for $NiCo_2O_4$ is $5.89 \times 10^{-4} \text{ s}^{-1}$ at 1.63 V vs. RHE, indicating 5.56 fold increased intrinsic superiority of $Ni_{2.5}Co_5C_2O_4$ over $NiCo_2O_4$, which is also previously supported by BET normalised LSV investigation. Furthermore, the electrochemical activity of $Ni_{2.5}Co_5C_2O_4$ is almost found to be comparable with that of earlier reported non-precious electrocatalysts and standard RuO_2 for the alkaline OER (see Table S10, ESI†).^{19–24,29,50–53}

2.4 Post-catalytic analysis

To understand the post-catalytic transformation, the carbon paper supported catalyst was subjected to composition and microstructural analysis. As evident from Fig. S17 (ESI†), no characteristic XRD pattern of the post-catalytic sample is observed except the sharp conspicuous diffraction signal which is assigned to the peaks from carbon paper. However, XPS measurements clearly show the structural features and elemental composition of the post-catalytic sample (see Fig. S18, ESI†). The C 1s pattern shows the appearance of two new signals around 291.77 and 294.76 eV, indicating the existence of a K^+ adsorbed microstructural framework for the post-catalytic sample (see Fig. S18a, ESI†).³⁷ From the Co 2p narrow scan, the peaks around 780.25 and 796.15 eV could corroborate the existence of Co^{3+} in the post-catalytic network (see Fig. S18b, ESI†).³⁶ Further, the presence of satellite peaks around 785.33 and 802.35 eV confirms the same.⁵⁴ Likewise, distinct XPS signals around 855.95 and 874.1 eV along with satellite peaks around 861.54 and 880.01 eV confirm the presence of Ni^{3+} (see Fig. S18c, ESI†).³⁷ However, the O 1s spectrum shows peaks around 530.01, 531.08 and 534.69 eV particularly for lattice oxygen, hydroxyl groups (adsorbed) and lattice water molecules respectively (see Fig. S18d, ESI†). Thus, composition analysis clearly shows that almost all Ni^{2+} and Co^{2+} oxidizes into Ni^{3+} and Co^{3+} respectively under high oxidation potential gradient during the OER process. Furthermore, inductively coupled plasma-atomic emission spectroscopy (ICP-AES) study reveals a Ni/Co ratio of about 1 : 2.15 for the post-catalyst sample, thereby confirming the absence of any such metal ion leaching during the OER for $Ni_{2.5}Co_5C_2O_4$ that further confirms the mechanical stability of the working electrode in alkaline medium. The morphology of the post-OER sample was investigated by FESEM and TEM. The FESEM image shows flake-like assembly, which confirms structural transformation from a block to flake-like nanostructure under alkaline OER conditions (see Fig. S19a–d, ESI†). In addition, TEM investigation displays microstructural aggregation throughout the mesoporous network of the post-catalytic sample with lattice fringes of 0.220 nm representing the (006) plane of $CoOOH$ (ICDD no. 014-0673).⁴⁸ EDS and elemental mapping further reveal the uniform spreading of Ni, Co, C and O for the post-catalytic sample (see Fig. S19e–j, ESI†). For further validation, Raman measurements were performed to examine the *in situ* transformation of the catalyst during electrochemical precondition treatment. The Raman signals appear at 186, 470, 510 and



666 cm^{-1} for A_g , E_g , F_{2g} and A_{1g} respectively, confirming the formation of oxyhydroxide species from nickel cobalt oxalate (see Fig. S20, ESI†).^{20,29}

2.5 Computational study

To get better insight into the structure–catalytic property relationship of the designed materials, we have performed density functional theory (DFT) study based on first-principles calculations using the Vienna *Ab Initio* Simulation Package (VASP).⁵⁵ As revealed from PXRD analysis, the highest intensity planes namely (311) for NiCo_2O_4 and (202) for NiC_2O_4 , CoC_2O_4 and $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ are considered to be the active planes for the electrochemical OER and all the DFT studies were performed on NiCo_2O_4 (311), NiC_2O_4 (202), CoC_2O_4 (202) and $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) surfaces. The NiCo_2O_4 (311) surface contains surface exposed Ni^{2+} and Co^{3+} ions which are bi and tri-coordinated respectively while both NiC_2O_4 (202) and CoC_2O_4 (202) surfaces consist of tri-coordinated surface exposed Ni^{2+} and Co^{2+} respectively (see Fig. S21a–c, ESI†). On the contrary, the $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) surface contains both surface exposed Ni^{2+} and Co^{2+} (see Fig. S21d, ESI†). However, all of the oxalates have bulk tetra and penta-coordinated Ni^{2+} and Co^{2+} ions (see Fig. S21, ESI†). The M–OH bond strength for the metal-based electrocatalysts is considered to be the key descriptor for OER activity.^{56–60} The weaker the M–OH bond strength, the higher will be the activity. The more stable OH adsorption site for the NiCo_2O_4 (311) surface is Co^{3+} with a binding energy (B. E.) of -0.65 eV while those for NiC_2O_4 and CoC_2O_4 (202) surfaces are -1.15 and -0.46 eV respectively (see Fig. 5a–c, S22 and Table S11, ESI†). On the other hand, tri-coordinated surface Ni^{2+} ion is

the most stable site for OH adsorption with a B.E. of -0.23 eV (see Fig. S23 and Table S11, ESI†). Hence, the M–OH bond strength follows the order $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) < CoC_2O_4 (202) < NiCo_2O_4 (311) < NiC_2O_4 (202). In addition, Bader charge analysis^{61,62} exhibits a charge transfer of $0.28|e|$ from adsorbed OH (OH^*) to the $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) catalyst surface, indicating relatively weaker charge transfer interaction compared to CoC_2O_4 ($0.4|e|$), NiCo_2O_4 ($0.65|e|$) and NiC_2O_4 ($0.84|e|$) surfaces. The charge density difference (CDD) plot further confirms the charge transfer from OH^* to different surfaces (Fig. S24†). Therefore, with relatively weaker M–OH bonding interaction, the $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) catalyst surface is superior to the other surfaces and consequently the order of OER activity is found to be NiC_2O_4 (202) < NiCo_2O_4 (311) < CoC_2O_4 (202) < $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202), which very nicely corroborates the experimental findings (*vide infra*). The bonding interaction between adsorbed OH and interacting Ni in $\text{OH}^*-\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) can be further confirmed from the density of states (DOS) plot (asterisk indicates the contribution from O-2p) as shown in Fig. 5d. The generation of the O-2p state can be clearly visualized near -5.95 eV (as marked with a black box) in the DOS plot for $\text{OH}^*-\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) (Fig. 5e). Besides, projected density of states (PDOS) analysis further confirms the bonding interaction between O-2p_z and Ni-d_{xy} orbitals which indicates that hybridization mainly occurs between these orbitals (Fig. 5f).

From the above discussion, it can be concluded that the adsorption of OH is weaker on the $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) catalyst surface and hence exhibiting better OER activity and this conclusion can be further justified by a d-band model which correlates the d-band centre (DBC) with the adsorption energies of the intermediates (here OH).^{63–65} The positions of DBCs are

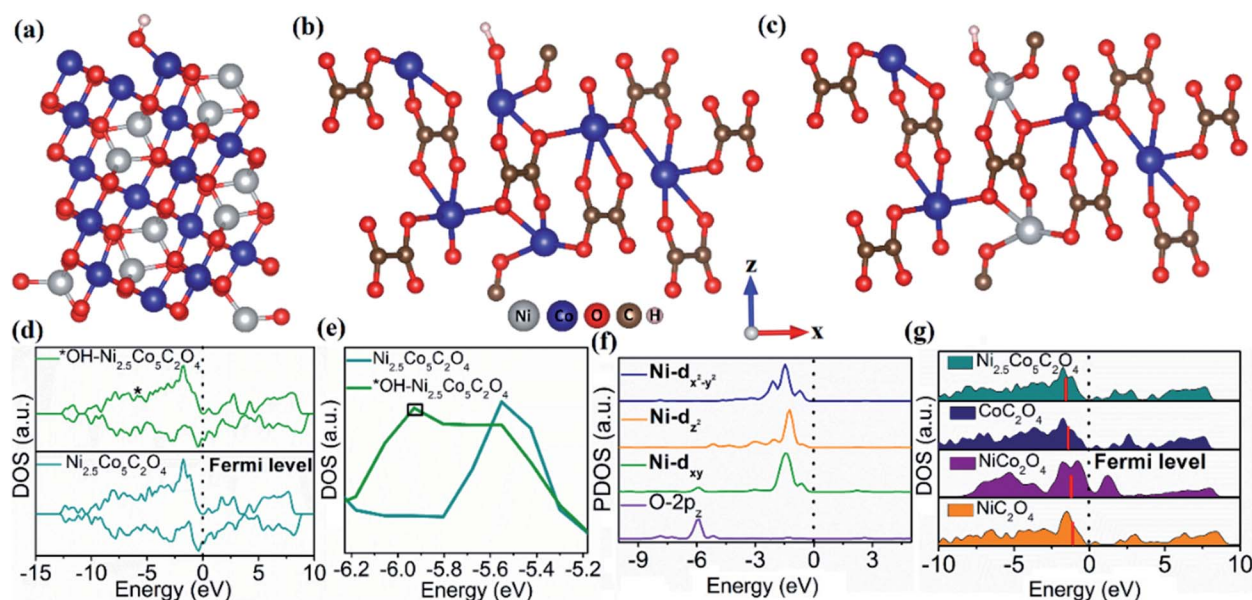


Fig. 5 Optimized structure of OH^* adsorbed (a) NiCo_2O_4 (311), (b) CoC_2O_4 (202), and (c) $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) surfaces, (d) DOS plots for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) and $\text{OH}^*-\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) surfaces (asterisk indicates the contribution from O-2p), (e) magnified DOS (the O-2p state is situated at around -5.95 eV as marked by a black box in the $\text{OH}^*-\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202) DOS), (f) projected density of states (PDOS) for O-2p_z, Ni-d_{xy}, Ni-d_z² and Ni-d_{x²-y²} of $\text{OH}^*-\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202), and (g) DOS plots for different catalyst surfaces. The thick red bar indicates the position of the d-band centre (E_d) on catalyst surfaces.



−1.50, −1.37, −1.26 and −1.07 eV respectively for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ (202), CoC_2O_4 (202), NiCo_2O_4 (311) and NiC_2O_4 (202) surfaces (Fig. 5g) which vividly shows that the DBC gradually shifts towards the Fermi level from $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ to the NiC_2O_4 (202) surface. The closer the DBC to the Fermi level, the stronger will be the adsorption of OH and hence deterioration of OER activity. This finding overwhelmingly supports our conclusion that the $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ catalyst is the best surface for the OER with weaker OH adsorption ability while NiC_2O_4 is the least active with stronger OH binding ability. The present study highlights the benefit of nickel–cobalt oxalate over nickel–cobalt oxide for the alkaline OER process. We also believe that the use of oxalate ligand could be a promising strategy for the fabrication of new-generation electrocatalysts for electrochemical water oxidation in alkaline medium.

3. Conclusions

In summary, nickel–cobalt oxalate ($\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$) based block-like nanostructures were prepared by a wet-chemical technique in an aqueous medium and further treated by calcination at 350 °C for the synthesis of nickel cobalt oxide (NiCo_2O_4). The preparation method was optimized in terms of Ni/Co ratio, oxalic acid amount, reaction temperature and time. The working electrode fabrication was performed on carbon paper as the substrate surface maintaining an optimum loading of the electrocatalyst. Next, $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ achieves 10 $\text{mA cm}_{\text{geo}}^{-2}$ with 330 mV overpotential, whereas NiCo_2O_4 requires 410 mV for the same. The comparative study suggests the best electrochemical catalytic efficiency for $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$, and that can be attributed to lower charge transfer resistance, more changes in double layer capacitance during precondition in alkaline medium, and high surface coverage. DFT study reveals that comparatively weaker OH adsorption as well as favourable d-band centre (DBC) position makes the $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ catalyst the best surface for the OER compared to individual CoC_2O_4 and NiC_2O_4 as well as well studied NiCo_2O_4 . This report thoroughly compares the activity of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ and NiCo_2O_4 for the alkaline OER process and highlights the benefit of $\text{Ni}_{2.5}\text{Co}_5\text{C}_2\text{O}_4$ over NiCo_2O_4 . The prudent choice of the oxalate based ligand-assisted synthetic approach could be used for other transition metals to execute bulk-scale alkaline water oxidation for a clean ecosystem and sustainable living.

Conflicts of interest

There are no conflicts to declare.

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