






Cite this: *Sustainable Energy Fuels*,  
2025, 9, 1552

# Improving the electrocatalytic activity of Pd nanoparticles through electronic coupling interaction with a Ni<sub>2</sub>P–MoS<sub>2</sub> hybrid support for ethanol electro-oxidation in an alkaline medium†

Thabo Matthews,  \*abc Makhaokane Paulina Chabalala,  bc  
Siyabonga Patrick Mbokazi,  bc Memory Zikhali,  bc Tarekegn Heliso Dolla,  d  
Anatolijs Šarakovskis,  e Guntars Vaivars,  efg Tunde Lewis Yusuf,  h Rhiyaad Mohamed  a  
and Nobanathi Wendy Maxakato  \*bc

To improve the performance of direct ethanol fuel cells (DEFCs), which are hindered by traditional catalysts, having matters pertaining to stability, activity, and selectivity in reaction environments, various electrocatalysts such as Pd/Ni<sub>2</sub>P, Pd/MoS<sub>2</sub>, and Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> were synthesized using the microwave-assisted NaBH<sub>4</sub>–ethylene glycol reduction method. The research findings suggest that the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> catalyst we developed had the highest activity (1579 mA mg<sub>Pd</sub><sup>−1</sup>), approximately 21 times greater than that of commercial Pd/C. The stability of the electrocatalysts were examined using chronoamperometry (CA) and cyclic voltammetry (CV) measurements, which indicated that the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst had good stability towards the ethanol oxidation reaction (EOR) in alkaline electrolyte. Electrochemical impedance spectroscopy (EIS) analysis showed that the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst had lower charge transfer resistance, indicating better electrochemical kinetics. According to XRD, HR-TEM, XPS, and electrochemical analysis, the enhanced electrocatalytic activity, long-term stability of the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst were attributable to the interface synergism as well as electronic and strain effects between the Pd, Ni<sub>2</sub>P, and MoS<sub>2</sub> interactions. This resulted in a downshift in the d-band center of the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst, weakening intermediate adsorption and the adsorbate metal interaction.

Received 4th September 2024  
Accepted 6th January 2025

DOI: 10.1039/d4se01223b

rsc.li/sustainable-energy

## 1 Introduction

The widespread use of CO<sub>2</sub> emitting fossil fuels has made meeting the world's energy demand a significant challenge.

<sup>a</sup>HySA/Catalysis Centre of Competence, Catalysis Institute, Department of Chemical Engineering, University of Cape Town, Cape Town 7701, South Africa. E-mail: matthews.thabo@uct.ac.za; matthewsThabo@gmail.com

<sup>b</sup>Department of Chemical Sciences, University of Johannesburg, Doornfontein, 2028, South Africa. E-mail: nmaxakato@uj.ac.za

<sup>c</sup>Center for Nanomaterials Research, University of Johannesburg, South Africa

<sup>d</sup>Institute for Catalysis and Energy Solutions (ICES), College of Science, Engineering and Technology, University of South Africa (UNISA), Private Bag X6, Florida, 1710, South Africa

<sup>e</sup>Institute of Solid State Physics, University of Latvia, Kengaraga Iela 8, LV-1063 Riga, Latvia

<sup>f</sup>Faculty of Chemistry, University of Latvia, Jelgavas Iela 1, LV-1004 Riga, Latvia

<sup>g</sup>Institute of Chemical Physics, University of Latvia, Jelgavas Iela 1, LV-1004 Riga, Latvia

<sup>h</sup>Department of Chemistry, Faculty of Natural and Agricultural Sciences, University of Pretoria, Private Bag X20, Hatfield 0028, Pretoria, South Africa

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4se01223b>

This has put a strain on the planet's natural resources. Therefore, developing renewable energy resources globally, is essential to ensure sustainable growth, especially in developing countries. Direct alcohol fuel cells are promising energy conversion and storage technologies, suitable for portable and mobile devices. Both monohydric and polyhydric alcohols have appreciable high energy densities, high boiling points, and low volatilities, which makes them easy to store. Direct ethanol fuel cell (DEFCs) have several advantages.<sup>1,2</sup> It has an existing infrastructure for production, high specific energy, and abundance. However, ethanol usage is limited due to the slow kinetics of the ethanol oxidation reaction (EOR) and the requirement to cleave the C–C bond, which has presented significant challenges.<sup>1–5</sup>

Electrocatalyst design is essential in developing DEFC electrocatalysts with improved performance. It is possible to modify electrocatalysts by changing their shape, composition, support materials, electronic state, structural effects, and preparatory methods. Various synthesis methods have been used to control the compositions and structures of bimetallic electrocatalysts, aiming to enhance the electrocatalyst activity, CO selectivity, and durability of Pd-based electrocatalysts. For example, in



their study, Shu *et al.*<sup>6</sup> demonstrated that the PdCuCo electrocatalyst, showed a significantly higher peak current density for the electro-oxidation of fuel under alkaline conditions. The performance was ten times greater than that of the Pd/C catalyst. It is well known that the use of appropriate metals and supports can adjust the electronic structure and affect the intrinsic activity of electrocatalysts. Doping and alloying protocols have been shown to impact the electronic structure of Pd and improve the performance of the electro-oxidation reaction. Junfeng Liu *et al.*<sup>7</sup> evaluated these catalyst design strategies, reporting that a P-doped PdMo bimetallic exhibits superior performance in the electro-oxidation of various alcohols due to the altered Pd electronic structures.

For enhanced Pd electrocatalytic activity, Wang *et al.*<sup>8</sup> synthesized a Pd–Ni–P ternary catalyst (by introducing Ni and P into Pd) that showed high EOR electrocatalytic performance. In another study, Ye *et al.*<sup>9</sup> prepared stable 3D hierarchical Pd@CoP nanosheets to promote the oxidation or removal of CO and CH<sub>3</sub>CO species. Similarly, Chen *et al.*<sup>10</sup> improved the EOR performance in alkaline media by introducing P into a CuPdNi alloy, forming a CuPdNiP catalyst. From these studies, we concluded that P promotes bifunctional effects, shortens the interatomic distance, and promotes metal support interactions (MSIs). Improving the activity and stability of Pd, with lower Pd loading, promotes the prospects for commercialization.

Both experimental and theoretical evaluations have shown that Ni<sub>2</sub>P exhibits higher electrocatalytic activity and stability for various reactions, such as the hydrogen evolution reaction (HER) and formic acid oxidation reaction (FAOR). These improved electrocatalytic or photochemical catalytic properties are due to the electronic charge transfer from the interaction of Ni and P and active metal site exposure effects. Chang *et al.*<sup>11</sup> synthesized Pt–Ni<sub>2</sub>P/C electrocatalysts, wherein the 30 wt% loading of Ni<sub>2</sub>P exhibited higher performances relative to Pt/C, Ni-promoted Pt/C, and P-promoted Pt/C catalysts, as assessed by electrochemical evaluation. On the other hand, metal dichalcogenides, such as MoS<sub>2</sub>, WS<sub>2</sub>, *etc.*, have attracted attention for catalytic applications due to their plausible electrical and electrochemical properties. In particular, MoS<sub>2</sub> is interesting because it is easy to synthesize and cost-effective and has longer active edges. All these features make MoS<sub>2</sub> an efficient component for preparing a hybrid electrocatalyst. For example, Tang *et al.*<sup>12</sup> synthesized Pt-decorated porous coral reef-like MoS<sub>2</sub>/nitrogen-doped biocarbon, which showed superior CO tolerance and was an effective methanol oxidation reaction (MOR) catalyst. In a different study, Meng Li *et al.*<sup>13</sup> reported the use of ultrafine Pt/MoP-NC for the MOR, and attributed the improved performance to a strong synergistic interface effect between platinum (Pt) and molybdenum phosphide (MoP) which was strengthened by the electronic donation properties of MoP and in turn enhanced both the catalytic activity and stability. The same phenomenon was put forward by Yubin K. *et al.*<sup>14</sup> where they alluded to Pt doping and interfacial engineering with MoSe<sub>2</sub> which accelerated MOR kinetics.

In light of the aforementioned efforts and drawing inspiration from previous findings as the foundation, we report for the first time the promotion of Pd by Ni<sub>2</sub>P–MoS<sub>2</sub> for the ethanol

oxidation reaction. We show that the Ni<sub>2</sub>P–MoS<sub>2</sub> promotion effect on Pd was significant. In this study, we detail the synthesis of the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst using an ethylene glycol/NaBH<sub>4</sub> microwave-assisted reduction method and its characterization as well as electrocatalytic properties. Through the interface of MoS<sub>2</sub> with Ni<sub>2</sub>P and Pd nanoparticles, we achieved a synergy between Pd and Ni<sub>2</sub>P–MoS<sub>2</sub>. The hybrid Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst demonstrated superior electrocatalytic activity with a mass activity of 1579 mA mg<sub>Pd</sub><sup>−1</sup> and an onset potential of 0.48 (V vs. RHE) while also exhibiting remarkable durability after 500 cycles, retaining 88.5% of the initial current.

## 2 Experimental

### 2.1 Chemicals

The following chemicals were all purchased from Sigma-Aldrich. Molybdenum(IV) sulphide (MoS<sub>2</sub>) powder (<2 μm, 99%), nickel chloride hexahydrate, NiCl<sub>2</sub>·6H<sub>2</sub>O, red phosphorus, ethanol, 99.9%, palladium(II) chloride PdCl<sub>2</sub> (98.0%), Pd/C commercial 10 wt%, dimethyl ether (DME), ethylene glycol (EG), 99.5.0%, potassium hydroxide (KOH) pellets, ferrocene C<sub>10</sub>H<sub>10</sub>Fe, and hydrogen peroxide H<sub>2</sub>O<sub>2</sub>.

**2.1.1 Synthesis of nickel phosphide, Ni<sub>2</sub>P.** Pristine Ni<sub>2</sub>P was synthesized using a modified solvothermal approach.<sup>12</sup> In brief, 1.0 mmol of NiCl<sub>2</sub>·6H<sub>2</sub>O was stirred with 4.0 mmol of red phosphorus in ethylene glycol for 30 minutes. This was followed by ultrasonic irradiation at ambient temperatures for 30 minutes. The solution was carefully poured into a stainless-steel Teflon-lined autoclave (80 mL) and reacted at 180 °C for 12 h in a furnace. The product was cooled and washed multiple times with distilled H<sub>2</sub>O/CH<sub>3</sub>CH<sub>2</sub>OH until a clear filtrate was observed. A dry sample was obtained by oven drying at 80 °C for 8 h.

**2.1.2 Synthesis of the Ni<sub>2</sub>P–MoS<sub>2</sub> hybrid support.** A 1 : 1 ratio of Ni<sub>2</sub>P–MoS<sub>2</sub> was used in the experiment. First, 800 mg of Ni<sub>2</sub>P and 800 mg of MoS<sub>2</sub> were mixed using a pestle and mortar for 15 minutes. The resulting mixture was then dissolved in 50 mL of a solution containing equal parts of ethanol and distilled water. This solution was placed in a glass beaker and stirred continuously for 1 hour while bubbling N<sub>2</sub>/Ar gas to prevent oxide formation. The solution was then transferred to a stainless-steel autoclave and heated to 150 °C for 3 hours. The contents were then centrifuged and dried at 80 °C for 6 hours. Finally, the resulting powder was named Ni<sub>2</sub>P–MoS<sub>2</sub>.

**2.1.3 Synthesis of the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> nanocomposite.** To prepare a 5 wt% Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> nanocomposite, 83.33 mg of PdCl<sub>2</sub> was dissolved in 40 mL of double distilled water. The Pd solution was added dropwise to the hybrid support Ni<sub>2</sub>P–MoS<sub>2</sub> while stirring for 3 h. Afterwards, a mixture of ethylene glycol/NaBH<sub>4</sub> (40 mL: 1 g) was added dropwise to the above mixture, and the pH was increased to between 10 and 12 using a 3 M NaOH solution. The 5 wt% Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> solution was microwave irradiated for 4 minutes with 1-minute sonication intervals. After irradiation, the black suspension was washed multiple times with a solution of (1 : 1) ethanol and double distilled water. The black solid was dried in a vacuum oven at



60 °C for 24 h and labeled as Pd/Ni<sub>2</sub>P–MoS<sub>2</sub>. The other electrocatalysts were synthesized using the same method with appropriate chemicals, specifically Pd/Ni<sub>2</sub>P without the addition of MoS<sub>2</sub> and Pd/MoS<sub>2</sub> without the addition of Ni<sub>2</sub>P.

## 2.2 Physicochemical instrumentation

The samples were analysed for their crystalline nature using an X'Pert PRO PANalytical diffractometer at room temperature, with CuK $\alpha$  radiation ( $\lambda = 1.5406 \text{ \AA}$ ), covering the  $2\theta$  range of 10–90°. Other tests performed included FTIR spectroscopy using a PerkinElmer FTIR spectrophotometer and RAMAN spectroscopy using a WITec alpha300R confocal Raman spectrometer with a laser wavelength of 532 nm and 600 grooves per mm, using a 50X objective and a 100  $\mu\text{m}$  fiber. Images were recorded using scanning transmission electron microscopy (STEM) with a JEOL JEM-ARM200F at 200 kV. Pd concentrations were measured using inductively coupled plasma-optical emission spectrometry (ICP-OES), while X-ray photoemission spectroscopy (XPS) measurements were performed in an ultra-high vacuum (UHV) chamber equipped with a SPECS PHOIBOS 150 hemispherical electron energy analyser, with an Al K $\alpha$  excitation line ( $h\nu = 1486.71 \text{ eV}$ ).

## 2.3 Electrochemical characterization

A Dropsens ustat 4000 and a standard three-electrode setup were used for the electrochemical measurements. The glassy carbon working electrode (WE) had an area of 0.07065 cm<sup>2</sup>, while the counter was made of a Pt sheet and Ag/AgCl (3 M KCl) was used as the reference electrode. To prepare the electrocatalyst ink, 10 mg of the electrocatalyst was dispersed in 50/50 (v/v) ultrapure H<sub>2</sub>O/CH<sub>3</sub>CH(OH)CH<sub>3</sub> solution containing 30  $\mu\text{L}$  of DME. The catalyst ink was then ultrasonicated at room temperature for 3 hours. 5  $\mu\text{L}$  of the electrocatalyst ink was drop-coated on the active area of the GCE and dried at 80 °C in an oven. A freshly prepared N<sub>2</sub> gas-saturated mixture of 0.5 M CH<sub>3</sub>CH<sub>2</sub>OH and 1 M KOH was used as the electrolyte. The CV measurements were performed between a potential window of 0.0 to 1.1 V (V vs. RHE) at 50 mV s<sup>-1</sup> (scan direction from left to right) at room temperature, and the third scans were recorded. Electrochemical impedance spectra (EIS) at 0.59 V (vs. RHE) in the frequency range of 10 Hz to 100 000 kHz were also recorded. Chronoamperometry measurements were conducted in N<sub>2</sub>-saturated 1 M KOH + 0.5 M ethanol at 0.59 V (vs. RHE) using a Gamry interface 1010E 27143. To convert all potentials to the RHE scale, the Nernst equation  $E_{\text{RHE}} = E_{\text{Ag/AgCl}} + 0.059 \times \text{pH} + 0.197$  was used. A 10 wt% commercial Pd/C electrocatalyst was used as the benchmark for all electrochemical tests under similar reaction conditions.

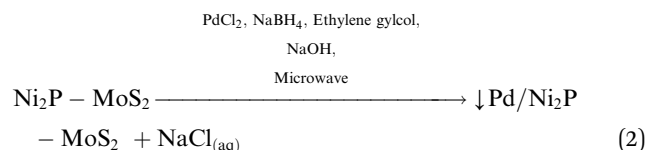
# 3 Results and discussion

## 3.1 Synthesis chemistry of Pd/Ni<sub>2</sub>P–MoS<sub>2</sub>

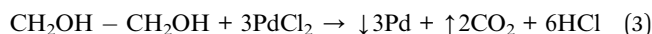
Initially, Ni<sub>2</sub>P and MoS<sub>2</sub> were mixed in water and ethanol.



For the deposition of Pd nanoparticels (NPs) onto the Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst (assuming the anchoring of Pd NPs onto the catalyst surface), we postulate that on anchoring the Pd NPs they should mostly be in the zero state *via* the reduction of PdCl<sub>2</sub>. This reduction process is carried out in a mixture of NaBH<sub>4</sub>/EG and is facilitated by microwave irradiation. We propose that Na<sup>+</sup> from NaBH<sub>4</sub>/NaOH reacts with Cl from PdCl<sub>2</sub> to form NaCl, which is then washed off. The representative equation is shown below:



During our study, we assumed that the precursor reduction to a metal results in the complete oxidation of EG into CO<sub>2</sub> and H<sub>2</sub>O, indicating the maximum reducing power of the alcohol.



## 3.2 Physicochemical characterization

The prepared electrocatalysts' arrangement and composition were determined using XRD, FTIR, and Raman spectroscopy. Fig. 1(a) presents the XRD patterns for the electrocatalytic materials. The presence of sharp and intense diffraction peaks in all samples attests to their high crystallinity. Fig. 1(a) illustrates MoS<sub>2</sub> characteristic diffraction peaks at 14.1°, 29.0°, 33.68°, 44.0°, 59.56°, and 60.2°, indexed to (002), (004), (100), (006), (110), and (008) planes, respectively, of the 2H-phase of MoS<sub>2</sub> (JCPDS 37-1492).<sup>15</sup> Additionally, there is an extra peak at around  $2\theta \approx 28.8^\circ$  corresponding to the (004) peak of 1T-phase MoS<sub>2</sub>, suggesting the presence of a minor amount of 1T-phase molybdenum disulfide (MoS<sub>2</sub>) alongside the predominant 2H-phase. Molybdenum disulfide has different phases, with the natural 2H-phase being semiconducting and thermodynamically favored. This phase consists of two S–Mo–S layers formed from edge-sharing MoS<sub>6</sub> trigonal prisms, whereas the metallic 1T phase features a single S–Mo–S layer of edge-sharing MoS<sub>6</sub> octahedra. Recent studies show that adding metallic atoms to MoS<sub>2</sub> nanosheets can induce the transformation from the 2H phase into the 1T phase, leading to lower charge transfer resistance and improved catalytic performance.<sup>16–18</sup>

The (002) peak predominance indicates a preferential orientation along the (001) direction in the prepared materials. For Ni<sub>2</sub>P, the diffraction peaks at 40.6°, 44.7°, 47.3°, 54.2°, and 54.7° are indexed to (111), (021), (210), (300), and (002) planes, respectively, and relate to the hexagonal phase of Ni<sub>2</sub>P (JCPDS Card No. 65-1989).<sup>19,20</sup> The Pd diffraction lines are invisible on Pd/MoS<sub>2</sub>, Pd/Ni<sub>2</sub>P, and Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> samples due to overlap, low loading, smaller sizes, and good dispersion of Pd nanoparticles; as seen from the HR-TEM. This is seen by the broadening of the peak base where Pd appears. Thus, the XRD pattern of Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> confirms the successful formation of the hybrid catalyst.



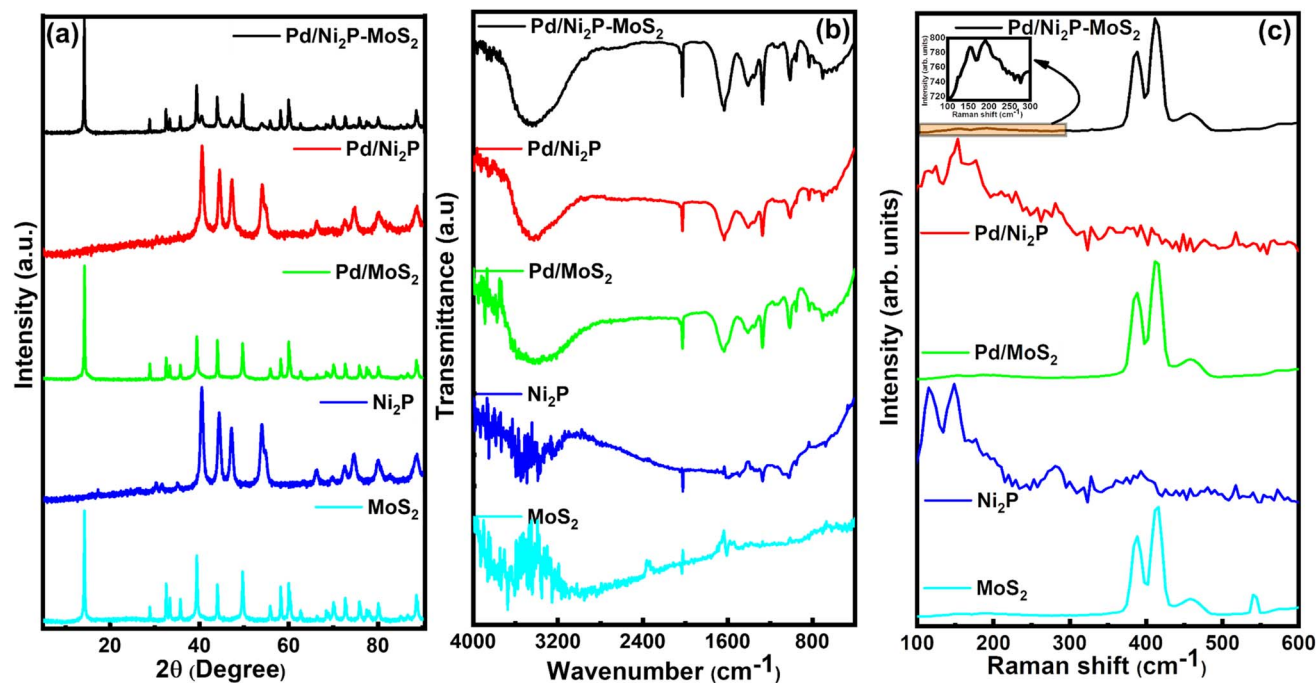


Fig. 1 (a) XRD patterns, (b) FTIR spectra, and (c) Raman spectra of the synthesised electrocatalysts.

The surface chemical states of the samples were analyzed using Fourier transform infrared (FTIR) spectroscopy. The FTIR spectra of  $\text{Ni}_2\text{P}$ ,  $\text{MoS}_2$ ,  $\text{Pd/Ni}_2\text{P}$ ,  $\text{Pd/MoS}_2$ , and  $\text{Pd/Ni}_2\text{P-MoS}_2$  are shown in Fig. 1(b). The peak at  $3419\text{ cm}^{-1}$  is a strong signal from the adsorbed  $-\text{OH}$  groups. The peaks from  $1300\text{ cm}^{-1}$  to  $1700\text{ cm}^{-1}$  indicate the presence of water and the O-H group bending vibration.<sup>21–24</sup>  $\text{MoS}_2$  exhibits characteristic Mo-S and S-S vibrations at  $600$  and  $905\text{ cm}^{-1}$ . The two bands at  $1042\text{ cm}^{-1}$  and  $918\text{ cm}^{-1}$  are attributed to species with phosphorus.<sup>21,25</sup>

Raman spectra shown in Fig. 1(c) were used to assess the structural properties of the electrocatalysts. The distinct peaks at  $381$  and  $410\text{ cm}^{-1}$  in  $\text{MoS}_2$ ,  $\text{Pd/MoS}_2$ , and  $\text{Pd/Ni}_2\text{P-MoS}_2$  samples are assigned to the  $\text{E}_{2g}^1$  and  $\text{A}_{1g}$  of Mo-S phonon mode belonging to the 2H crystalline phase of  $\text{MoS}_2$ .<sup>26,27</sup> Previous research studies have shown similar findings to the ones presented here.<sup>28</sup> This further proves that the composite structure of  $\text{MoS}_2$  and  $\text{Ni}_2\text{P}$  has successfully formed. The deposition of Pd atoms onto  $\text{Ni}_2\text{P}$ ,  $\text{MoS}_2$ , and  $\text{Ni}_2\text{P-MoS}_2$  leads to a shift of approximately  $2.0\text{ cm}^{-1}$  in the  $\text{A}_{1g}$  peak. This indicates that Pd interacts with support materials and can considerably modify the electronic structure, as confirmed from XPS analysis in Fig. 3 and 4. Similarly, the  $\text{E}_{2g}^1$  and  $\text{A}_{1g}$  peaks of  $\text{Ni}_2\text{P-MoS}_2$  exhibit a slight red shift compared to  $\text{MoS}_2$  due to interfacial stress resulting from the interaction between  $\text{Ni}_2\text{P}$  and  $\text{MoS}_2$ . This shift suggests that the formation of the  $\text{MoS}_2$ - $\text{Ni}_2\text{P}$  heterojunction induces changes in the Raman spectra of  $\text{MoS}_2$ , confirming the successful preparation of  $\text{MoS}_2$ - $\text{Ni}_2\text{P}$ . Furthermore, the peak separations ( $\text{A}_{1g} - \text{E}_{2g}^1$ ) in the bare supports are  $29.7$ ,  $25.4$ , and  $23.2\text{ cm}^{-1}$ , with Pd-loaded supports showing a slight decrease in the peak separation attributable to introducing S vacancies during chemical synthesis; this peak

separation further proves successful Pd anchoring and support hybridization and that  $\text{MoS}_2$  is a good candidate for Pd anchoring.<sup>28–30</sup>

The morphology of the electrocatalysts was studied using HR-TEM, as seen in Fig. 2.  $\text{Ni}_2\text{P}$  and  $\text{MoS}_2$  have a sheet-like morphology, as shown in Fig. 2(a) and (b), respectively, with SAED pattern insets. Fig. 2(a) shows lattice fringes, confirming that the desired  $\text{Ni}_2\text{P}$  phase was successfully formed. The distinct lattice fringes with a  $d$ -spacing of  $0.20\text{ nm}$  were observed for the  $\text{Ni}_2\text{P}$  crystal, corresponding to the (201) lattice plane of the hexagonal  $\text{Ni}_2\text{P}$  phase.<sup>31</sup> The SAED pattern of  $\text{Ni}_2\text{P}$  has three diffraction rings representing the (111), (300), and (321) planes of  $\text{Ni}_2\text{P}$ , from the inner to the outer ring. Fig. 2(c) and (d) show Pd NPs anchored onto the surface of  $\text{Ni}_2\text{P}$  and  $\text{MoS}_2$ , respectively, with  $2$  and  $3.4\text{ nm}$  sizes. The different sizes of the Pd NPs indicate that the supports interact with the NPs differently. The HR-TEM image in Fig. 2(e) confirms the microstructure of the  $\text{Ni}_2\text{P-MoS}_2$  nanosheet hybrids with Pd NPs on the surface, demonstrating the feasibility of achieving the hybrid electrocatalyst using an  $\text{NABH}_4/\text{EG}$  microwave-assisted reduction method. The Pd NPs' particle size on the  $\text{Ni}_2\text{P-MoS}_2$  is  $3.4\text{ nm}$ , as that for  $\text{Pd/MoS}_2$ , which proves that  $\text{MoS}_2$  plays a major role in the anchoring of Pd NPs, as explained based on the shift and separation from Raman spectra (Fig. 1(c)). The synthesized materials are highly crystalline, as seen from the SAED patterns shown in Fig. 2 (a and b insets) and (b–d) (iii), as well as the XRD patterns.

XPS was used to examine the surface chemical composition of  $\text{Pd/MoS}_2$ - $\text{Ni}_2\text{P}$ ,  $\text{Pd/MoS}_2$ ,  $\text{Pd/Ni}_2\text{P}$ ,  $\text{Ni}_2\text{P}$ , and  $\text{MoS}_2$  hybrid nano-materials. In comparison, the survey spectra in Fig. 3(a)



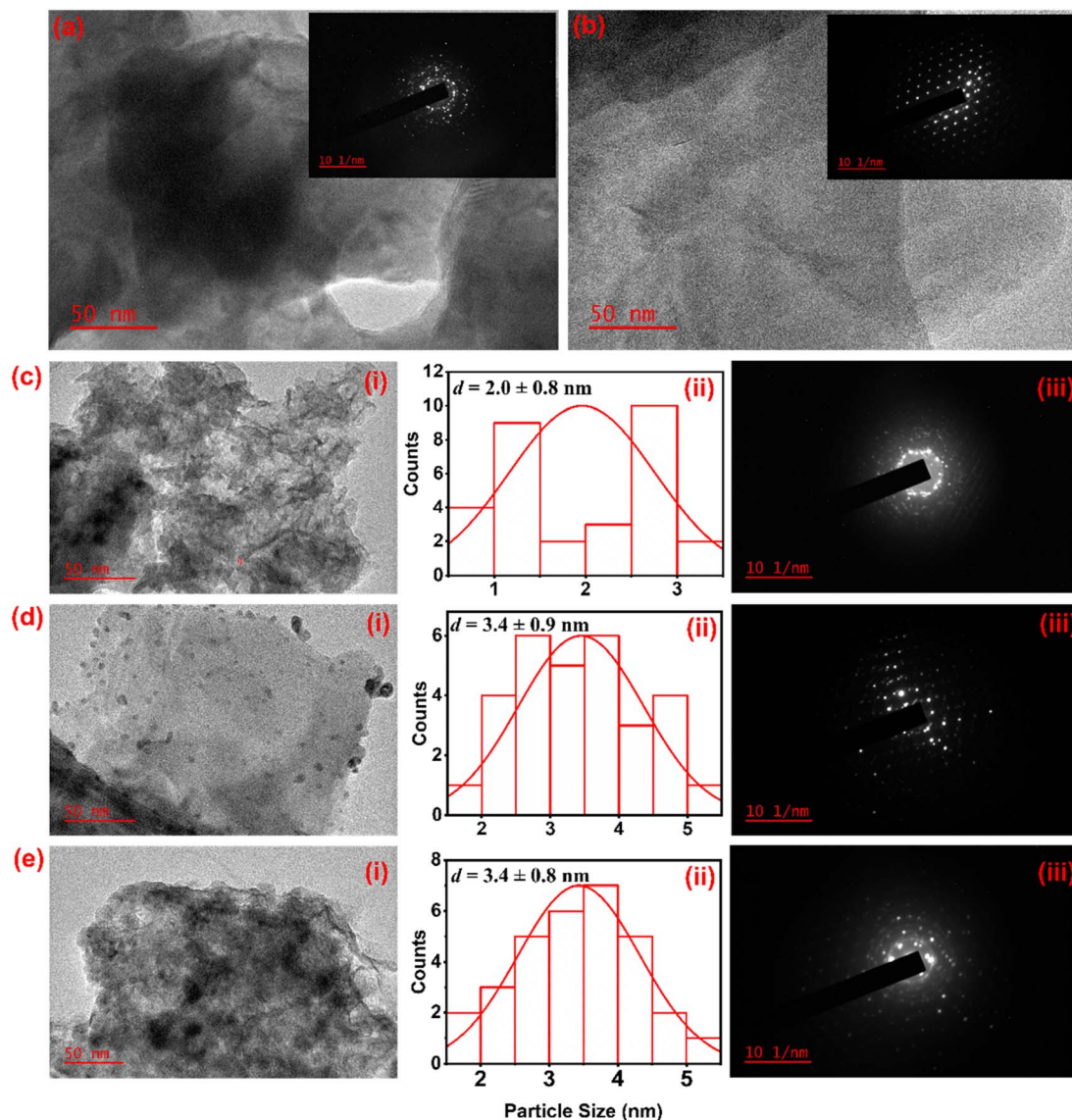


Fig. 2 HR-TEM images of (a)  $\text{Ni}_2\text{P}$ , (b)  $\text{MoS}_2$ , (c) (i)  $\text{Pd}/\text{Ni}_2\text{P}$ , (d) (i)  $\text{Pd}/\text{MoS}_2$ , and (e) (i)  $\text{Pd}/\text{Ni}_2\text{P}-\text{MoS}_2$ , (ii) particle size distributions, and (iii) SAED patterns.

demonstrate the presence of C, O, P, Pd, S, Mo, and Ni for the main composite and subsequent elements for the other nanomaterials. Furthermore, the presence of C and O elements can be seen in the survey spectra and Fig. S4 in the ESI,<sup>†</sup> which could be a result of absorbed gaseous molecules. The deconvoluted spectra of S 2p exhibit twin peaks of S  $2p_{1/2}$  and S  $2p_{3/2}$  at the corresponding BEs of 163.8 and 162.6 eV, which is attributed to  $\text{S}^{2-}$  in  $\text{MoS}_2$  of  $\text{Pd}/\text{MoS}_2-\text{Ni}_2\text{P}$ ,  $\text{Pd}/\text{MoS}_2$ , and  $\text{MoS}_2$  nanomaterials, Fig. 3(b).  $\text{Pd}/\text{MoS}_2-\text{Ni}_2\text{P}$  and  $\text{Pd}/\text{MoS}_2$  have slightly different binding energy (BE) values than pure  $\text{MoS}_2$  (S  $2p_{3/2}$  = 162.4 eV), indicating charge transfer from Pd to  $\text{MoS}_2$ .<sup>32</sup> The distinctive peak of phosphate species is centered at 133.67 eV for P 2p, as shown in Fig. 3(c) for the nanocomposites with the nickel phosphide constituent.<sup>33</sup> The P 2p is shifted to a higher BE relative to  $\text{Ni}_2\text{P}$ ; we postulate that this represents the interaction of Pd with  $\text{Ni}_2\text{P}$  to form species such as Ni-P-Pd or Pd-P.

Fig. 4(a) shows high-resolution spectra of the Mo 3d for  $\text{Pd}/\text{MoS}_2-\text{Ni}_2\text{P}$ ,  $\text{Pd}/\text{MoS}_2$ , and  $\text{MoS}_2$  electrocatalysts. The doublets of the  $\text{Mo}^{4+}$  bands at 229.5 eV and 232.7 eV in the Mo 3d region are ascribed to Mo  $3d_{5/2}$  and Mo  $3d_{3/2}$  orbitals, respectively, and correspond well with the literature.<sup>34</sup> When  $\text{Pd}/\text{MoS}_2-\text{Ni}_2\text{P}$  is formed, the major composite species undergoes changes in binding energy (BE). The core-level Mo peaks in  $\text{Pd}/\text{MoS}_2-\text{Ni}_2\text{P}$  indicate a shift to lower BEs relative to pristine  $\text{MoS}_2$  and  $\text{Pd}/\text{MoS}_2$ . This change is attributable to a decrease in the Fermi level (EF) caused by p-type doping, as previously seen in another research study. This demonstrates successful Mo substitution with Pd dopants, and the covalent bonds within the lattice are reported to be responsible for the stabilization of Pd atoms.<sup>35</sup>

The high-resolution spectrum's Pd 3d regions display four peaks, as seen in Fig. 4(b). Pd  $3d_{5/2}$  peaks at 335.8 eV and 336.8 eV, respectively, correspond to  $\text{Pd}^{0/2+}$ .<sup>36</sup> It is interesting to



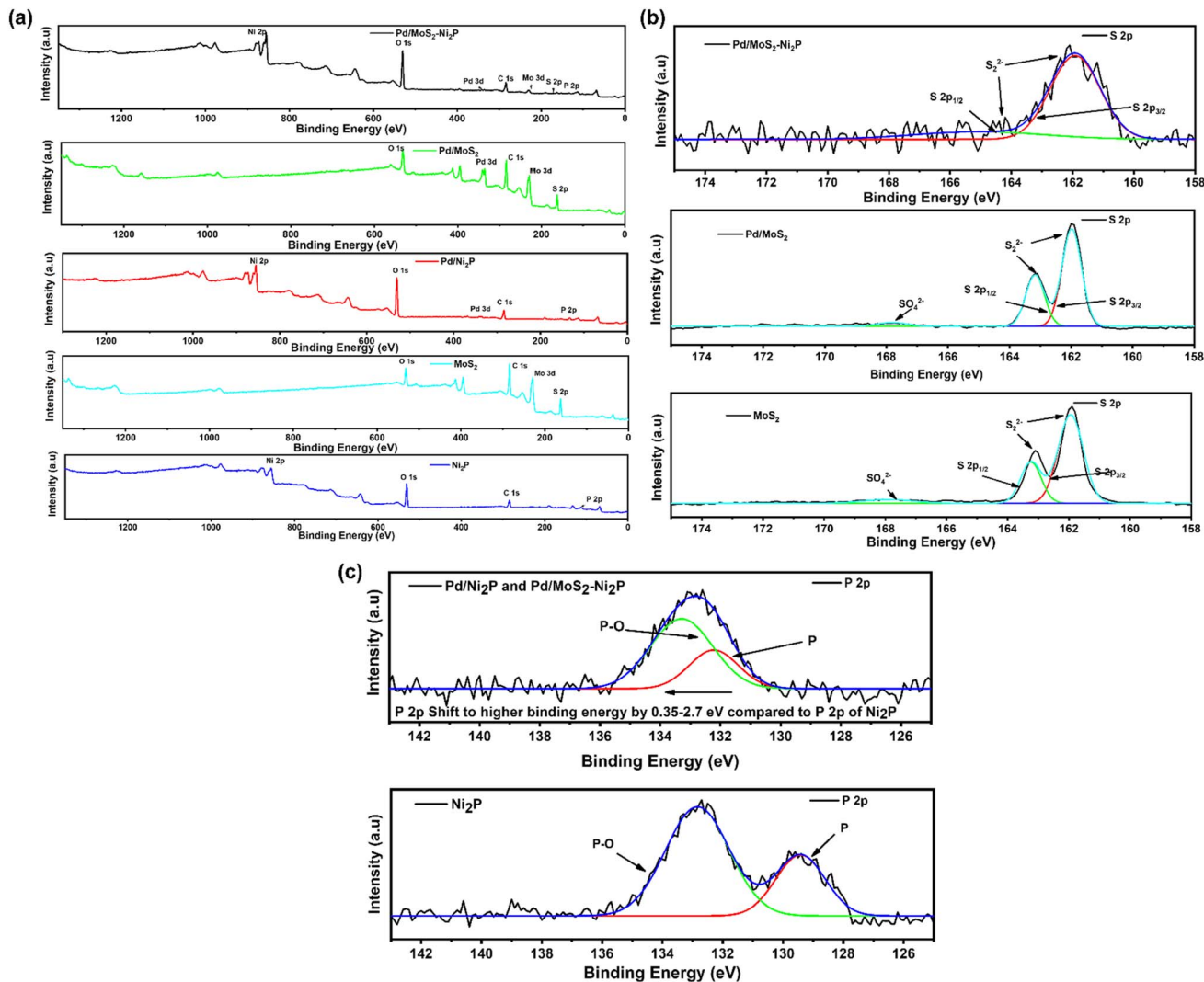


Fig. 3 XPS spectra. (a) Survey scans, (b) S 2p and (c) P 2p.

note that the Pd BEs in Pd/Ni<sub>2</sub>P and Pd/MoS<sub>2</sub> occur at the same BE, and there is a notable shift in Pd/MoS<sub>2</sub>-Ni<sub>2</sub>P, indicating great electronic coupling interaction. Thus, there is a shift in BEs for Pd<sup>0</sup> of Pd/MoS<sub>2</sub>-Ni<sub>2</sub>P compared to metallic Pd (3d<sub>5/2</sub> = 335.1 eV) on Pd/Ni<sub>2</sub>P and Pd/MoS<sub>2</sub>, which implies charge transfer from Pd to double support MoS<sub>2</sub>-Ni<sub>2</sub>P.<sup>37</sup> Therefore, substantial Pd and MoS<sub>2</sub> contacts occur for Pd/MoS<sub>2</sub>-Ni<sub>2</sub>P, implying that Pd catalysts for ethanol electro-oxidation have enhanced catalytic characteristics. The shift in Pd binding energy for Pd/MoS<sub>2</sub>-Ni<sub>2</sub>P indicated that the insertion of Pd particles resulted in a substantial electronic alteration with Ni<sub>2</sub>P due to the strain effect.<sup>38</sup>

The Ni 2p deconvoluted spectra, Fig. 4(c), contain peaks at 855.15 (Ni 2p<sub>3/2</sub>) and 872.83 eV (Ni 2p<sub>1/2</sub>) for Ni<sup>2+</sup> presence. Satellite peaks were seen at 860.57 and 878.57 eV. The change in the electronic environment of the nickel atoms during composite creation causes the shift in BE for Ni 2p for Pd/MoS<sub>2</sub>-Ni<sub>2</sub>P, indicating successful hybridization of the nanocomposite, which had a great effect as seen in the shifting of 3d Pd BE in Fig. 4(b).

### 3.3 Electrochemical characterization

Cyclic voltammetry was utilized to evaluate the performance of the synthesized electrocatalysts in an alkaline medium. The commercial Pd/C, Pd/Ni<sub>2</sub>P-MoS<sub>2</sub>, Pd/Ni<sub>2</sub>P, and Pd/MoS<sub>2</sub> electrocatalysts were assessed in 1 M KOH, Fig. 5(a). The PdO reduction peak in the potential range of 0.6–0.8 (V vs. RHE) for all catalysts indicates efficient alkaline electrocatalysis, with Pd/Ni<sub>2</sub>P-MoS<sub>2</sub> having the highest catalytic activity for electrochemical reactions in an alkaline medium, as observed in the negative scan. The current density of the electrocatalysts in the negative scan is arranged in ascending order as follows: Pd/MoS<sub>2</sub> < Pd/C < Pd/Ni<sub>2</sub>P < Pd/Ni<sub>2</sub>P-MoS<sub>2</sub>.

The electrochemically active surface area (ECSA) (eqn (4)) for the electrocatalysts was determined using the PdO reduction peak based on the literature as depicted in Fig. 5(a)

$$\text{ECSA} = \frac{Q}{SI} \quad (4)$$



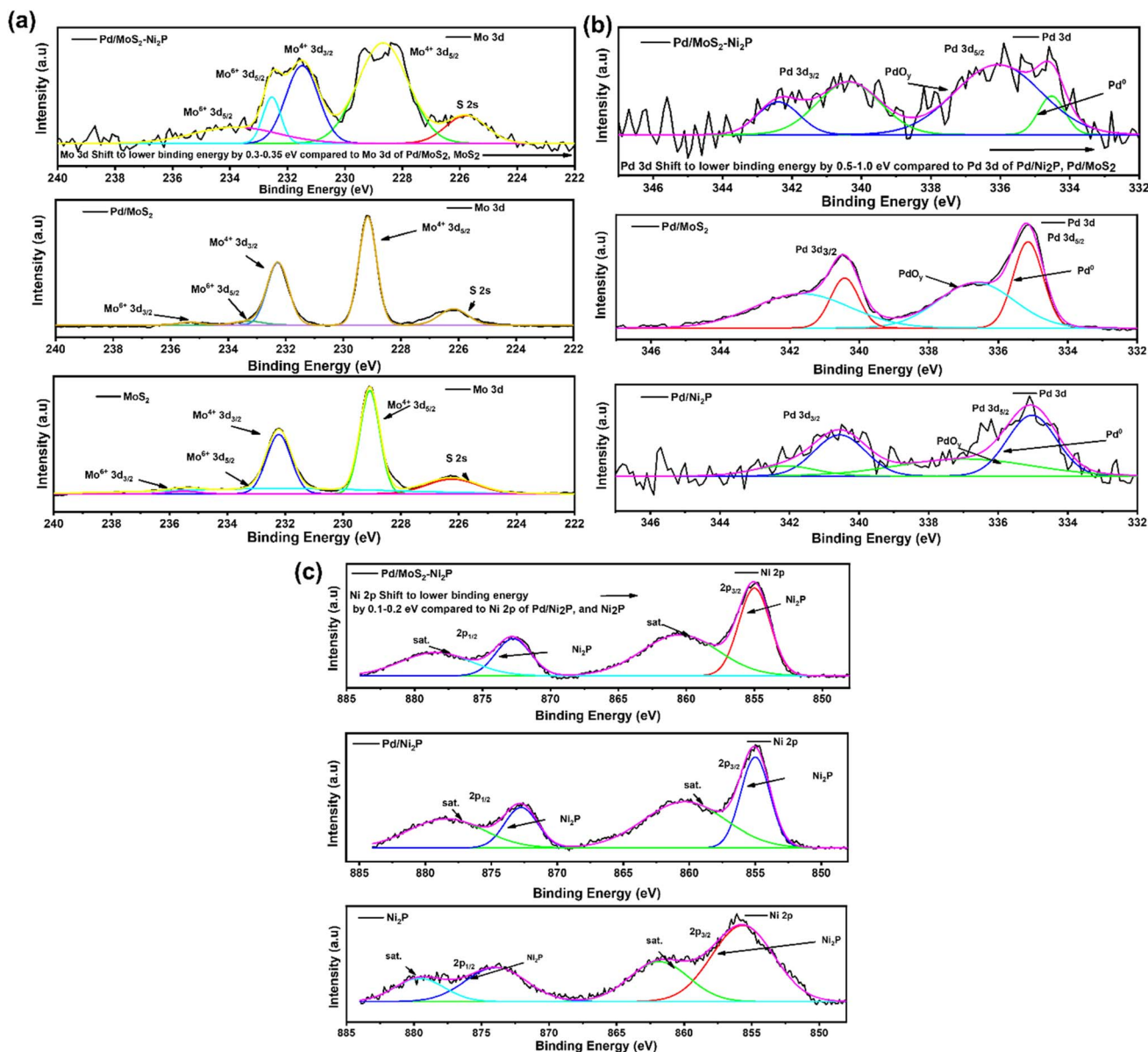


Fig. 4 XPS spectra. (a) Survey, (b) Pd 3d, (c) O 1S, and (d) C 1S for 1% Pd at 10%, 20%, and 30%  $V_2O_5-C$ , 3% Pd at 10%, 20% and 30%  $V_2O_5-C$ , and 5% Pd/at 10%, 20% and 30%  $V_2O_5-C$ .

where  $Q$  is integrated coulombic charge (reduction of PdO (in mC)),  $S$  is a proportionality constant ( $0.424 \text{ mC cm}^{-2}$ ), and  $I$  is the loading in  $\mu\text{g}$ . The ECSA of Pd/Ni<sub>2</sub>P/MoS<sub>2</sub> is calculated to be  $113.38 \text{ m}^2 \text{ g}^{-1}$ , which is 1.44, 2.99 and 16.70 times larger than that of Pd/Ni<sub>2</sub>P ( $78.30 \text{ m}^2 \text{ g}^{-1}$ ), Pd/C ( $37.88 \text{ m}^2 \text{ g}^{-1}$ ) and Pd/MoS<sub>2</sub> ( $6.79 \text{ m}^2 \text{ g}^{-1}$ ) under identical conditions, which indicates that incorporating Pd nanoparticles with MoS<sub>2</sub> and Ni<sub>2</sub>P inorganic support materials discloses more catalytically active sites for electrochemical reactions under alkaline conditions.

The CV cycles of the multi-metallic electrocatalysts Pd/Ni<sub>2</sub>P-MoS<sub>2</sub>, Pd/Ni<sub>2</sub>P, and Pd/MoS<sub>2</sub> and commercial Pd/C were studied in 0.5 M ethanol and 1 M KOH; their cyclic voltammograms are displayed in Fig. 5(b). All the corresponding catalysts show distinct anodic scans, as expected. The observed forward anodic

scan is ascribed to the electro-oxidation of fresh  $\text{CH}_3\text{CH}_2\text{OH}_{\text{ads}}$  on electrocatalyst, whilst the reverse scan is attributed to the removal of intermediates species such as  $\text{CO}_{\text{ad}}$ ,  $\text{COH}_{\text{ad}}$ ,  $\text{HCO}_{\text{ad}}$ , and  $\text{HCOO}_{\text{ad}}$  that accumulate on the electrocatalyst surface resulting from the EOR. The ternary metal-based electrocatalyst Pd/Ni<sub>2</sub>P-MoS<sub>2</sub> recorded the highest current density of  $1579 \text{ mA mg}_{\text{Pd}}^{-1}$  with the lowest onset potential, meaning better kinetics and low activation energy, which appears to be 5, 22, and 23 times more than that of Pd/Ni<sub>2</sub>P ( $314.8 \text{ mA mg}_{\text{Pd}}^{-1}$ ), Pd/C ( $71.9 \text{ mA mg}_{\text{Pd}}^{-1}$ ) and Pd/MoS<sub>2</sub> ( $69.2 \text{ mA mg}_{\text{Pd}}^{-1}$ ) respectively. The current densities are plausible relative to others reported in the literature, as shown in Table S2.† The onset potential ( $E_{\text{onset}}$ ) of the catalysts, as shown in Table 1, followed the descending order as follows: Pd/MoS<sub>2</sub> ( $0.64 \text{ V vs. RHE}$ ) > Pd/C ( $0.54 \text{ V vs.}$



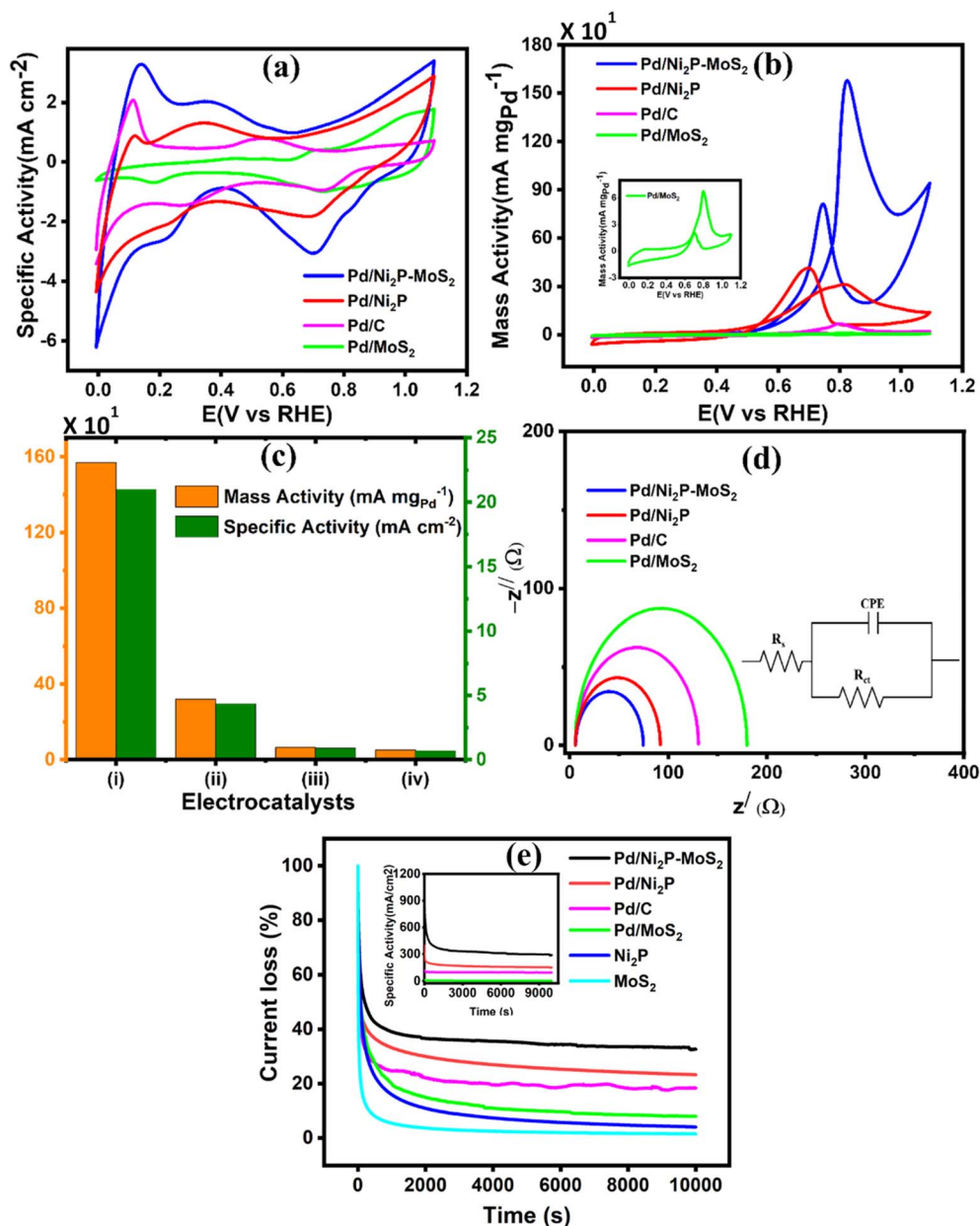


Fig. 5 (a) Cyclic voltammograms in 1 M KOH, (b) ethanol oxidation at a scan rate of  $50 \text{ mV s}^{-1}$ , (c) the maximum mass activity and specific activity for (i) Pd/Ni<sub>2</sub>P–MoS<sub>2</sub>, (ii) Pd/Ni<sub>2</sub>P, (iii) Pd/C and (iv) Pd/MoS<sub>2</sub>, (d) Nyquist plots at 0.55 V in 1 M KOH + 0.5 M CH<sub>3</sub>CH<sub>2</sub>OH, and (e) chronoamperometric curves measured at 0.59 V vs. RHE).

RHE) > Pd/Ni<sub>2</sub>P (0.52 V vs. RHE) > Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> (0.48 V vs. RHE).

Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> recorded the most negative  $E_{\text{onset}}$  compared to the other electrocatalysts; this observation can be due to high reaction kinetics and low overpotential, hence its high catalytic activity. Furthermore, the good active characteristics of Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> may be due to the increased number of active sites as a result of combining nickel phosphide and molybdenum disulfide. The two inorganic support materials in Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> have demonstrated a remarkable electrocatalytic improvement for the ethanol electrooxidation reaction, incorporating their well-reported electrocatalytic characteristics, such as high conductivity and mass transport of Ni<sub>2</sub>P, and the unique

properties of MoS<sub>2</sub>, such as large surface area and high electrochemical accessibility, and, most importantly, their bifunctional effect promotes electron transfer between the electrode and the reactants, leading to enhanced electrocatalytic performance.

As shown in Fig. 5(b), Pd/C appears to outperform Pd/MoS<sub>2</sub>; this observation might be due to the poor conductivity of MoS<sub>2</sub>. Fig. 5(c) shows mass activity and specific activity of (i) Pd/Ni<sub>2</sub>P–MoS<sub>2</sub>, (ii) Pd/Ni<sub>2</sub>P, (iii) Pd/C, and (iv) Pd/MoS<sub>2</sub>. Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> recorded the largest mass activity ( $1579 \text{ mA mg}_{\text{Pd}}^{-1}$ ) and specific activity ( $216.1 \text{ mA cm}^{-2}$ ) when compared to other reference electrocatalysts; this indicates its improved efficiency for the EOR under alkaline conditions. The remarkable activity of



Table 1 Electrochemical characterization results of the as-prepared catalysts in 1 M KOH + 0.5 M ethanol<sup>a</sup>

Catalysts	ICP-OES	$E_{\text{onset}}$ (V vs. RHE)	Forward scan		ECDSA <sup>b</sup> (m <sup>2</sup> g <sup>-1</sup> )	EIS <sup>c</sup>		Chronoamperometry	Cycle studies
			$i_f$	$I_b$		$R_{\text{ct}}$ ( $\Omega$ )	$R_s$ ( $\Omega$ )	Retained current <sup>d</sup> (%)	$\left(\frac{I}{I_0}\right)^e$
Pd/Ni <sub>2</sub> P-MoS <sub>2</sub>	5.24	0.480	1579	812	113.4	68.36	5.68	33.19	0.885
Pd/Ni <sub>2</sub> P	4.98	0.520	314.8	413.2	78.30	86.62	5.79	23.97	0.868
Pd/C	10.00	0.540	71.9	25.7	37.88	125.13	5.98	17.37	0.674
Pd/MoS <sub>2</sub>	5.77	0.640	69.2	22.2	6.79	174.01	6.04	8.47	0.654

<sup>a</sup>  $i_f$  and  $I_b$  (mA mg<sup>-1</sup><sub>Pd</sub>). <sup>b</sup> Electrochemical surface area (based on the PdO peak). <sup>c</sup> Electrochemical Impedance parameters:  $R_{\text{ct}}$  charge transfer resistance values from EIS fitting (Nyquist) and  $R_s$  solution resistance values from EIS fitting (Nyquist). <sup>d</sup> @ 1000 s from CA. <sup>e</sup>  $\left(\frac{I}{I_0}\right)$  electrocatalyst deactivation rate, from cyclic studies.

Pd/Ni<sub>2</sub>P-MoS<sub>2</sub> is attributed to interface synergism and electronic and strain effects among Pd, Ni<sub>2</sub>P, and MoS<sub>2</sub>. Also, the ultimate electron coupling within the nanocomposite boosted the Pd NPs' electrocatalytic activity. The Pd/Ni<sub>2</sub>P-MoS<sub>2</sub> electrocatalyst exhibits metal support interactions that enhances its durability, stability, electrocatalytic activity, and selectivity. The electronic effect at the interface promotes electron transfer towards the direct C1-12e-ethanol electro-oxidation route for DEFCs. The proposed Pd/Ni<sub>2</sub>P-MoS<sub>2</sub> with interface synergism and overall nano-structural cooperation gives maximum electrocatalytic power and operates for more than 10 000 s, surpassing the performance of other electrocatalysts.

Electrochemical impedance spectroscopy was used to further investigate the remarkable reaction kinetics of the

electrocatalysts by determining their charge transfer ( $R_{\text{ct}}$ ) and solution resistance ( $R_s$ ) in relation to their Nyquist arch diameter. The obtained Nyquist plots, together with the corresponding circuit, are shown in Fig. 5(d). The arch diameter of the Nyquist plots was used to measure electrocatalyst reaction kinetics by determining their  $R_{\text{ct}}$  at the electrode-electrolyte interface. The arch diameter presented decreased as follows: Pd/MoS<sub>2</sub> > Pd/C > Pd/Ni<sub>2</sub>P > Pd/Ni<sub>2</sub>P-MoS<sub>2</sub>. This was due to the extent of electronic coupling interaction with Ni<sub>2</sub>P, MoS<sub>2</sub>, and Ni<sub>2</sub>P-MoS<sub>2</sub>. Thus, Pd/Ni<sub>2</sub>P-MoS<sub>2</sub> has the smallest  $R_{\text{ct}}$  and  $R_s$ , as indicated in Table 1; this further proves the composite catalyst's great kinetics reactions due to increased reaction active sites, leading to exceptional catalyst performance seen in Fig. 5(b). The excellent catalytic reaction kinetics can further be ascribed

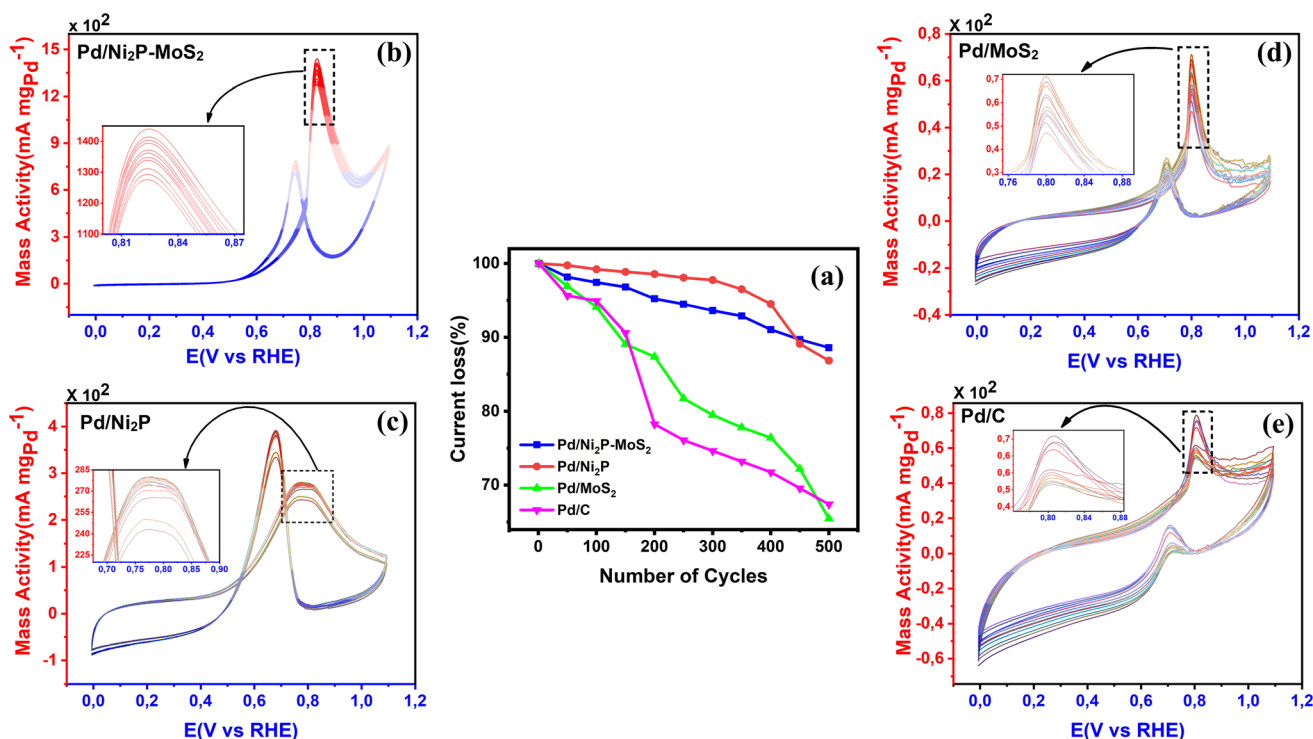


Fig. 6 (a) Line profile of CV cycles and (b–e) the durability tests performed by CV continuous cyclization in 0.5 M ethanol and 1 M KOH solution for 500 cycles of Pd/Ni<sub>2</sub>P-MoS<sub>2</sub>, Pd/Ni<sub>2</sub>P, Pd/MoS<sub>2</sub>, and Pd/C.



to the synergistic effect of the Pd–Ni–Mo alloy, that is, there is easy movement of electrons within the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> and the electrode–electrolyte interface. The Pd/MoS<sub>2</sub> electrocatalyst recorded the highest  $R_{ct}$  and  $R_s$  values with a large Nyquist arch as demonstrated; this shows sluggish reaction kinetics and high overpotential, and the least catalytic activity was obtained in CV. This can be ascribed to the layered structure of MoS<sub>2</sub>, which hinders electron transfer between the catalyst surface and reactants.

The vulnerability of the catalysts to CO poisoning is one of the few drawbacks limiting fuel cell commercialization. Thus, potentiostatic chronoamperometry was used to evaluate electrocatalyst stability at a constant potential of 0.59 (V vs. RHE) for 10 000 s in 0.5 M CH<sub>3</sub>CH<sub>2</sub>OH and 1 M KOH electrolyte solution as shown in Fig. 5(e). The inset in Fig. 5(e) shows the CA curve as a function of current density and time, but for a better comparison, the CA curve was plotted as a function of % current loss and time. A high initial current density was observed due to a transient potential jump that induced a significant oxidation current. However, this current density gradually decreased because of the diffusion of reactants and the accumulation of poisoning species, such as adsorbed carbon monoxide (CO<sub>ads</sub>), on the electrode surface, which deactivate the active sites of the electrocatalyst.<sup>39,40</sup> The electrocatalysts initially experienced a sharp decline in current density within the first 1000 seconds before stabilizing at a constant current level from 2500 s. The constant current density region represents the current density each electrocatalyst retained after CO poisoning. The retained current density increased in the order: Pd/MoS<sub>2</sub> > Pd/C > Pd/Ni<sub>2</sub>P > Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> as shown in Table 1. Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> retained the highest current density and is more stable than commercial Pd/C. The strong metal–phosphorus interaction can explain this enhanced anti-poisoning ability, which prevents leaching and corrosion.<sup>41</sup> The modulation of the electronic structure may also influence this effect<sup>14</sup> from the Ni<sub>2</sub>P–MoS<sub>2</sub> hybrid substrate. This modulation promotes the electro-oxidation and removal of adsorbed CO-like species on the Pd surface. These findings further suggest that using Ni<sub>2</sub>P and MoS<sub>2</sub> inorganic materials as hybrid support materials improves the electrocatalysts' stability for the EOR in alkaline media. According to a recent argument by Hofstead-Duffy *et al.*,<sup>42</sup> assessing electrocatalysts using  $I_f/I_b$  is still debatable. They proposed that  $I_f$  and  $I_b$  share the same chemical origin, which means that the empirical criterion of  $I_f/I_b$  is inadequate

for assessing the resistance of electrocatalysts to poisoning. As we concur with their findings, we complemented the above findings in this study by calculating the poisoning rate,  $\sigma$  (%s<sup>-1</sup>) (eqn (5)). From CA, the linear decay current density portion was used to calculate the long-term poisoning rate of adsorbed species:

$$\sigma (\%s^{-1}) = \frac{100}{I_0} \left( \frac{dI}{dt} \right)_{t > 2500s} \quad (5)$$

where  $\left( \frac{dI}{dt} \right)_{t > 2500s}$  = linear portion of CA,  $t$  = time in seconds and  $I_0$  = extrapolated current density % at the point  $\left( \frac{dI}{dt} \right)_{t > 2500s}$ . Jiang *et al.*<sup>43</sup> proposed the procedure for the MOR on PtRu electrocatalysts. The same procedure was used by Guo *et al.*<sup>32</sup> for the MOR and Chen *et al.*<sup>10</sup> for glucose oxidation. From these studies, we have postulated that the slope should be calculated from  $t = t_s$  where  $t_s$  is the time when the CA curve starts to stabilize. The  $\sigma$  (%s<sup>-1</sup>) values followed the order: Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> < Pd/Ni<sub>2</sub>P < Pd/C < Pd/MoS<sub>2</sub>, as shown in Table 1. Among the synthesized electrocatalysts and commercial Pd/C, Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> exhibited the highest tolerance to poisonous intermediates and the lowest poisoning rate during the EOR. The  $\sigma$  (%s<sup>-1</sup>) values give a clear view of the contribution of the hybrid support relative to the reduction in poisoning rates for the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalyst. The low poisoning rate for Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> is due to synergistic effects, electronic coupling between Pd and Ni<sub>2</sub>P/MoS<sub>2</sub>, and MSI.<sup>44,45</sup> These findings also suggest that Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> has high chemical stability despite the initial drop in current density. Thus, the nanocomposite utilizes the synergy between Ni<sub>2</sub>P and MoS<sub>2</sub> to slow the decay and promote active site regeneration by removing poison. The Pd/MoS<sub>2</sub> electrocatalyst has the highest  $\sigma$  (%s<sup>-1</sup>) value, indicating that it is the most unstable and susceptible to intermediate poisoning due to a lack of Pd NP protection. This is evident from the significant migration and coalescence/Ostwald ripening shown in Fig. 7(a); this causes a reduction of the electrochemical active centers, which is also apparent in Fig. 6(a).

CV was further used to assess the durability of the synthesized electrocatalysts. Fig. 6(a) shows that Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> had the highest cycle durability relative to other electrocatalysts and commercial Pd/C. Fig. 6(b–e) show the actual CV cycles with the respective insets for forward oxidation peaks. The insets clearly

### After stability and durability testing

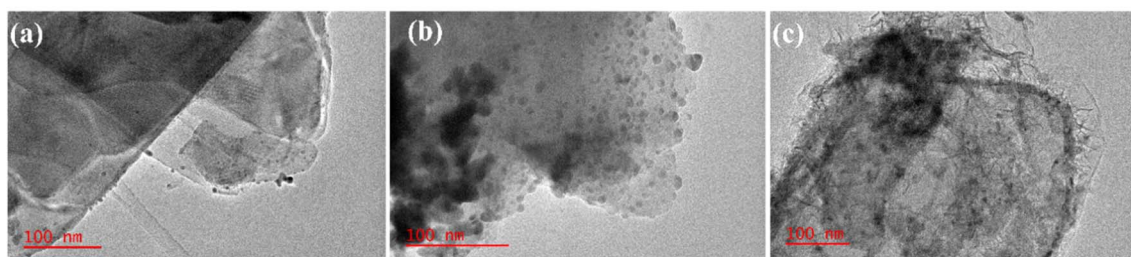


Fig. 7 HR-TEM images after durability and stability testing for (a) Pd/Ni<sub>2</sub>P, (b) Pd/MoS<sub>2</sub>, and (c) Pd/Ni<sub>2</sub>P–MoS<sub>2</sub>.



show the current decay profile as pseudo-contours. From the inset of Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> (Fig. 6(b)) we see a gradual drop of the CV oxidation peak with an increasing cycle number, which we postulated MSI and better dispersion on the hybrid support system as compared to randomized and non-uniform for Fig. 6(c–e) based on the CV cycle profiles. From the cycle studies, we complemented the findings by calculating the electrocatalyst deactivation rate, where  $I$  (current density) was normalized by  $I_0$  (initial current)<sup>46</sup> and the  $\left(\frac{I}{I_0}\right)$  ratios are shown in Table 1. For the Pd supported on Ni<sub>2</sub>P–MoS<sub>2</sub> containing the electrocatalyst had a  $\left(\frac{I}{I_0}\right)$  of 0.885 > Pd/C (0.674) and all the electrocatalysts indicating excellent cycling durability, which is summarized in Table S1.† On the other hand, Pd-supported electrocatalysts on mono-supports had a low  $\left(\frac{I}{I_0}\right)$  rate, which proves that the synergy and collaboration effect in a hybrid support system is fundamental in ensuring durability. The  $\left(\frac{I}{I_0}\right)$  ratios match CA curve findings, confirming the significance of hybrid support formation.

To further confirm the structural and morphological preservation of the electrocatalysts. Fig. 7(a and b) show HR-TEM images after the 10 000 s CA stability test and 500 CV cycles, respectively. The Pd NP distribution is not greatly affected, with minimal morphological changes. Of note is the small increase in the size of Pd NPs in Pd/MoS<sub>2</sub> assumed to be caused by the coalescence of the Pd NPs as they retained the spherical-like morphology. The major change seen was a reduction in the inter Pd NP distance. Thus, there was very little Pd NP agglomeration after the CA and cycle test, showing the electrocatalyst robustness, thus confirming the existence of MSI with the electrocatalyst matrix.

### 3.4 Complementary stability test (1 year cycle)

From the above-discussed results, we chose the most active electrocatalyst, Pd/Ni<sub>2</sub>P–MoS<sub>2</sub>, for further electrochemical evaluation to corroborate stability. The Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> stability was investigated using a modified GCE termed “GCE\_1” that was kept for a period of one year in an open environment. The Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> modified GCE was tested using cyclic voltammetry and chronoamperometry every 3 months. As seen from Fig. 8, the current density of GCE\_1 fluctuates between the nominal value in Fig. 5(c); we postulate that active site regeneration is possible. The presented bars in Fig. 8 are an average of current density obtained after every 3 months. The GCE\_1 was kept in an open environment, and the results clearly indicated that external weather factors minimally affected the electrocatalytic activity. This result means that the electrocatalytically active sites can be maintained over a period of one year, as per this study. As for the CA current retained, there was not much loss of current, which supports the results of cycle studies in Fig. 6(b). As shown in Fig. 8, the %CA current loss was from ~45% → ~38.9%. This plausible minimal current loss can be explained by MSI as proved by BE shifts in XPS,

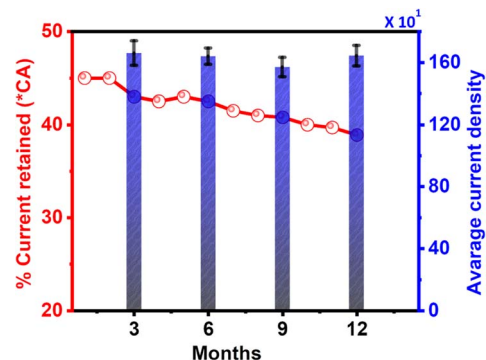


Fig. 8 Average current density for every three months of storage and the % retained current from chronoamperometry (\*CA) for the Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalysts in 1 M KOH + 0.5 M ethanol.

confirming the interaction of materials in the catalyst matrix and probable active site regeneration during the CA run.

## 4 Conclusion

A structural design has been developed to enhance the efficiency and lifespan of direct ethanol fuel cells (DEFCs) by engineering the synergistic interface of Pd/Ni<sub>2</sub>P–MoS<sub>2</sub>. Initial findings indicate that Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> exhibits greater electrocatalytic activity and improved steady-state performance for the ethanol oxidation reaction (EOR). The incorporation of the hybrid Ni<sub>2</sub>P–MoS<sub>2</sub> support significantly boosts the electrocatalytic activity of Pd for ethanol electro-oxidation through interface synergism and electronic coupling interaction. The Pd/Ni<sub>2</sub>P–MoS<sub>2</sub> electrocatalysts demonstrate the highest electrochemical catalytic activity for the EOR, with a mass activity of 1579 mA mg<sub>Pd</sub><sup>-1</sup>, a more negative onset potential of 0.48 V vs. RHE, and excellent durability, retaining 89% of the initial current after 500 cycles. This study illustrates that the Ni<sub>2</sub>P–MoS<sub>2</sub> hybrid support enhances electrocatalytic performance in comparison to the Pd/C electrocatalyst for the EOR in an alkaline medium. Nevertheless, it is essential to further assess other binary or ternary Pd-based electrocatalysts supported on Ni<sub>2</sub>P–MoS<sub>2</sub>@carbon to increase the reaction rate and achieve high selectivity towards CO<sub>2</sub> formation while breaking the C–C bond.

## Data availability

All data and materials used in this article are available upon reasonable request.

## 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.

## Acknowledgements

This research was funded by the University Research Council, International Postgraduate Scholarships (2021–2023), the



Centre for Nanomaterials Science Research, University of Johannesburg, South Africa, University of Cape Town 2025–2026 URC Postdoctoral Fellowship, Nurturing Emerging Scholars Programme, National Research Foundation of South Africa: Grant Numbers 118148, and 138083 and the Centre for Nanomaterials Science Research, University of Johannesburg, South Africa. We acknowledge the University of Johannesburg (UJ) FRC for financial support and Faculty of Science, Department of Chemical Sciences at the University of Johannesburg, South Africa. Funding was also provided by the Department of Science and Innovation (DSI), through the Hydrogen South Africa (HySA) Catalysis Centre of Competence Programme.

## References

- 1 L. An, T. S. Zhao and Y. S. Li, Carbon-neutral sustainable energy technology: Direct ethanol fuel cells, *Renewable Sustainable Energy Rev.*, 2015, **50**, 1462–1468, DOI: [10.1016/j.rser.2015.05.074](#).
- 2 J. Guo, R. Chen, F.-C. Zhu, S.-G. Sun and H. M. Villullas, New understandings of ethanol oxidation reaction mechanism on Pd/C and Pd<sub>2</sub>Ru/C catalysts in alkaline direct ethanol fuel cells, *Appl. Catal., B*, 2018, **224**, 602–611, DOI: [10.1016/j.apcatb.2017.10.037](#).
- 3 B. Cermenek, J. Ranninger and V. Hacker, *Alkaline direct ethanol fuel cell*, Elsevier, 2019, vol. 1904, DOI: [10.1016/B978-0-12-811458-2.00015-8](#).
- 4 Y. Zheng, X. Wan, X. Cheng, K. Cheng, Z. Liu and Z. Dai, Advanced catalytic materials for ethanol oxidation in direct ethanol fuel cells, *Catalysts*, 2020, **10**(2), 166, DOI: [10.3390/catal10020166](#).
- 5 M. Z. F. Kamarudin, S. K. Kamarudin, M. S. Masdar and W. R. W. Daud, Review: Direct ethanol fuel cells, *Int. J. Hydrogen Energy*, 2013, **38**(22), 9438–9453, DOI: [10.1016/j.ijhydene.2012.07.059](#).
- 6 Y. Shu, *et al.*, Hollow Echinus-like PdCuCo Alloy for Superior Efficient Catalysis of Ethanol, *ACS Appl. Mater. Interfaces*, 2018, **10**(5), 4743–4749, DOI: [10.1021/acsami.7b17731](#).
- 7 J. Liu, *et al.*, Phosphorus doped PdMo bimetallic as a superior bifunctional fuel cell electrocatalyst, *Chem. Eng. J.*, 2024, **486**, 150258, DOI: [10.1016/j.cej.2024.150258](#).
- 8 Y. Wang, F.-F. Shi, Y.-Y. Yang and W.-B. Cai, Carbon supported Pd–Ni–P nanoalloy as an efficient catalyst for ethanol electro-oxidation in alkaline media, *J. Power Sources*, 2013, **243**, 369–373, DOI: [10.1016/j.jpowsour.2013.06.021](#).
- 9 S.-H. Ye, J.-X. Feng and G.-R. Li, Pd Nanoparticle/CoP Nanosheet Hybrids: Highly Electroactive and Durable Catalysts for Ethanol Electrooxidation, *ACS Catal.*, 2016, **6**(11), 7962–7969, DOI: [10.1021/acscatal.6b02263](#).
- 10 D. Chen, *et al.*, Enhancing Ethanol Oxidation Reaction Activity of P-Doped CuPdNi Nanocatalyst by Optimizing Surface-Atom Distribution, *ACS Appl. Energy Mater.*, 2019, **2**(8), 5525–5533, DOI: [10.1021/acsaem.9b00715](#).
- 11 J. Chang, L. Feng, C. Liu, W. Xing and X. Hu, Ni<sub>2</sub>P enhances the activity and durability of the Pt anode catalyst in direct methanol fuel cells, *Energy Environ. Sci.*, 2014, **7**(5), 1628–1632, DOI: [10.1039/c4ee00100a](#).
- 12 B. Tang, *et al.*, Porous coral reefs-like MoS<sub>2</sub>/nitrogen-doped bio-carbon as an excellent Pt support/co-catalyst with promising catalytic activity and CO-tolerance for methanol oxidation reaction, *Electrochim. Acta*, 2017, **246**, 517–527, DOI: [10.1016/j.electacta.2017.06.052](#).
- 13 M. Li, F. Yang, J. Chang, A. Schechter and L. Feng, MoP-NC Nanosphere Supported Pt Nanoparticles for Efficient Methanol Electrolysis, *Acta Phys.-Chim. Sin.*, 2023, 2301005, DOI: [10.3866/PKU.WHXB202301005](#).
- 14 Y. Kuang, W. Qiao, S. Wang, F. Yang and L. Feng, Doping and Interfacial Engineering of MoSe<sub>2</sub> Nanosheets by NH<sub>3</sub> Plasma Promoted Pt for Methanol Electrolysis, *ACS Mater. Lett.*, 2024, **6**(5), 1722–1731, DOI: [10.1021/acsmaterialslett.4c00244](#).
- 15 M. Liu, *et al.*, Interfacial electronic structure engineering on molybdenum sulfide for robust dual-pH hydrogen evolution, *Nat. Commun.*, 2021, **12**(1), 5260.
- 16 D. Tsikritzis, N. Tsud, T. Skála and L. Sygellou, Unravelling the phase transition of 2H-MoS<sub>2</sub> to 1T-MoS<sub>2</sub> induced by the chemical interaction of Pd with molybdenum disulfide–graphene hybrids, *Appl. Surf. Sci.*, 2022, **599**, 153896, DOI: [10.1016/j.apsusc.2022.153896](#).
- 17 Y. Yan, X. Ge, Z. Liu, J.-Y. Wang, J.-M. Lee and X. Wang, Facile synthesis of low crystalline MoS<sub>2</sub> nanosheet-coated CNTs for enhanced hydrogen evolution reaction, *Nanoscale*, 2013, **5**(17), 7768, DOI: [10.1039/c3nr02994h](#).
- 18 Y. Li, H. Wang, L. Xie, Y. Liang, G. Hong and H. Dai, MoS<sub>2</sub> Nanoparticles Grown on Graphene: An Advanced Catalyst for the Hydrogen Evolution Reaction, *J. Am. Chem. Soc.*, 2011, **133**(19), 7296–7299, DOI: [10.1021/ja201269b](#).
- 19 E. Ramírez-Mondragón, O. E. Contreras, U. J. Tamayo-Pérez and M. T. Oropeza-Guzmán, Synthesis and characterization of Ni<sub>2</sub>P and MoS<sub>2</sub> on MWCNT as an innovative catalytic material for hydrogen generation, *Appl. Surf. Sci.*, 2020, **503**, 144163.
- 20 H. Sun, *et al.*, Porous multishelled Ni<sub>2</sub>P hollow microspheres as an active electrocatalyst for hydrogen and oxygen evolution, *Chem. Mater.*, 2017, **29**(19), 8539–8547.
- 21 M. Asim, *et al.*, Synergetic effect of Au nanoparticles and transition metal phosphides for enhanced hydrogen evolution from ammonia-borane, *J. Colloid Interface Sci.*, 2023, **638**, 14–25, DOI: [10.1016/j.jcis.2023.01.122](#).
- 22 Y. Xiao, *et al.*, Ball-milled Ni<sub>2</sub>P/g-C<sub>3</sub>N<sub>4</sub> for improved photocatalytic hydrogen production, *Int. J. Hydrogen Energy*, 2023, **48**(41), 15460–15472, DOI: [10.1016/j.ijhydene.2023.01.086](#).
- 23 S. Carencu, C. Boissière, L. Nicole, C. Sanchez, P. Le Floch and N. Mézailles, Controlled Design of Size-Tunable Monodisperse Nickel Nanoparticles, *Chem. Mater.*, 2010, **22**(4), 1340–1349, DOI: [10.1021/cm902007g](#).
- 24 W. Wang, *et al.*, Earth-abundant Ni<sub>2</sub>P/g-C<sub>3</sub>N<sub>4</sub> lamellar nanohybrids for enhanced photocatalytic hydrogen evolution and bacterial inactivation under visible light irradiation, *Appl. Catal., B*, 2017, **217**, 570–580, DOI: [10.1016/j.apcatb.2017.06.027](#).



- 25 Y. Xu, R. Wang, Z. Liu, L. Gao, T. Jiao and Z. Liu, Ni<sub>2</sub>P/MoS<sub>2</sub> interfacial structures loading on N-doped carbon matrix for highly efficient hydrogen evolution, *Green Energy Environ.*, 2022, 7(4), 829–839, DOI: [10.1016/j.gee.2020.12.008](https://doi.org/10.1016/j.gee.2020.12.008).
- 26 D. Vikraman, *et al.*, Theoretical evaluation and experimental investigation of layered 2H/1T-phase MoS<sub>2</sub> and its reduced graphene-oxide hybrids for hydrogen evolution reactions, *J. Alloys Compd.*, 2021, **868**, 159272.
- 27 X. Yu, Y. Feng, Y. Jeon, B. Guan, X. W. Lou and U. Paik, Formation of Ni–Co–MoS<sub>2</sub> nanoboxes with enhanced electrocatalytic activity for hydrogen evolution, *Adv. Mater.*, 2016, **28**(40), 9006–9011.
- 28 F. Sultana, *et al.*, An insight to catalytic synergic effect of Pd–MoS<sub>2</sub> nanorods for highly efficient hydrogen evolution reaction, *Arabian Journal of Chemistry*, 2022, **15**(5), 103735, DOI: [10.1016/j.arabj.2022.103735](https://doi.org/10.1016/j.arabj.2022.103735).
- 29 D. M. Sim, *et al.*, Controlled Doping of Vacancy-Containing Few-Layer MoS<sub>2</sub> via Highly Stable Thiol-Based Molecular Chemisorption, *ACS Nano*, 2015, **9**(12), 12115–12123, DOI: [10.1021/acs.nano.5b05173](https://doi.org/10.1021/acs.nano.5b05173).
- 30 H. Lv, *et al.*, Pd decorated MoS<sub>2</sub> nanoflowers as photothermal catalyst for enhanced NIR-induced 4-nitrophenol reduction, *J. Environ. Chem. Eng.*, 2023, **11**(5), 110375, DOI: [10.1016/j.jece.2023.110375](https://doi.org/10.1016/j.jece.2023.110375).
- 31 F.-C. Pan, H. He, Z.-X. Yang, Q. Zheng, D. Lin and Y. Huo, Rationally designed Ni<sub>2</sub>P/WS<sub>2</sub>/Co<sub>9</sub>S<sub>8</sub>@C multi-interfacial electrocatalyst for efficient overall water splitting, *Chem. Eng. J.*, 2022, **446**, 136961, DOI: [10.1016/j.cej.2022.136961](https://doi.org/10.1016/j.cej.2022.136961).
- 32 Y. Guo, *et al.*, Preparation of MoS<sub>2</sub> nanosheets to support Pd species for selective steerable hydrogenation of acetylene, *J. Mater. Sci.*, 2021, **56**, 2129–2137.
- 33 R. Bernasconi, *et al.*, Electrocatalytic layers for hydrogen evolution reaction based on nickel phosphides: cost-effective fabrication and XPS characterization, *J. Mater. Sci.*, 2022, **57**(20), 9370–9388.
- 34 M. Acerce, D. Voiry and M. Chhowalla, Metallic 1T phase MoS<sub>2</sub> nanosheets as supercapacitor electrode materials, *Nat. Nanotechnol.*, 2015, **10**(4), 313–318.
- 35 L. Z. Hao, *et al.*, Self-powered broadband, high-detectivity and ultrafast photodetectors based on Pd–MoS<sub>2</sub>/Si heterojunctions, *Phys. Chem. Chem. Phys.*, 2016, **18**(2), 1131–1139.
- 36 A. Drelinkiewicz, M. Hasik and M. Kloc, Pd/polyaniline as the catalysts for 2-ethylanthraquinone hydrogenation. The effect of palladium dispersion, *Catal. Lett.*, 2000, **64**, 41–47.
- 37 H. Li, K. Yu, X. Lei, B. Guo, H. Fu and Z. Zhu, Hydrothermal synthesis of novel MoS<sub>2</sub>/BiVO<sub>4</sub> hetero-nanoflowers with enhanced photocatalytic activity and a mechanism investigation, *J. Phys. Chem. C*, 2015, **119**(39), 22681–22689.
- 38 R. Kottayintavida and N. K. Gopalan, Nickel phosphate modified carbon supported Pd catalyst for enhanced alcohol electro oxidation, *Int. J. Hydrogen Energy*, 2020, **45**(19), 11116–11126.
- 39 Y. Zhou, Q. Wang, X. Tian, J. Chang and L. Feng, Electron-enriched Pt induced by CoSe<sub>2</sub> promoted bifunctional catalytic ability for low carbon alcohol fuel electro-reforming of hydrogen evolution, *J. Energy Chem.*, 2022, **75**, 46–54, DOI: [10.1016/j.jechem.2022.08.009](https://doi.org/10.1016/j.jechem.2022.08.009).
- 40 W. Qiao, L. Yu, J. Chang, F. Yang and L. Feng, Efficient bifunctional catalysis of coupled MoSe<sub>2</sub> nanosheet/Pt nanoparticles for methanol-assisted water splitting, *Chin. J. Catal.*, 2023, **51**, 113–123, DOI: [10.1016/S1872-2067\(23\)64469-9](https://doi.org/10.1016/S1872-2067(23)64469-9).
- 41 Y. Shi and B. Zhang, Recent advances in transition metal phosphide nanomaterials: Synthesis and applications in hydrogen evolution reaction, *Chem. Soc. Rev.*, 2016, **45**, 1529–1541, DOI: [10.1039/c5cs00434a](https://doi.org/10.1039/c5cs00434a).
- 42 A. M. Hofstead-Duffy, D.-J. Chen, S.-G. Sun and Y. J. Tong, Origin of the current peak of negative scan in the cyclic voltammetry of methanol electro-oxidation on Pt-based electrocatalysts: a revisit to the current ratio criterion, *J. Mater. Chem.*, 2012, **22**(11), 5205, DOI: [10.1039/c2jm15426a](https://doi.org/10.1039/c2jm15426a).
- 43 J. Jiang and A. Kucernak, Electrooxidation of small organic molecules on mesoporous precious metal catalysts. II: CO and methanol on platinum–ruthenium alloy, *J. Electroanal. Chem.*, 2003, **543**, 187–199, DOI: [10.1016/S0022-0728\(03\)00046-9](https://doi.org/10.1016/S0022-0728(03)00046-9).
- 44 H. An, *et al.*, Synthesis and performance of palladium-based catalysts for methanol and ethanol oxidation in alkaline fuel cells, *Electrochim. Acta*, 2013, **102**, 79–87, DOI: [10.1016/j.electacta.2013.03.142](https://doi.org/10.1016/j.electacta.2013.03.142).
- 45 J. Goel and S. Basu, Effect of support materials on the performance of direct ethanol fuel cell anode catalyst, *Int. J. Hydrogen Energy*, 2014, **39**(28), 15956–15966, DOI: [10.1016/j.ijhydene.2014.01.203](https://doi.org/10.1016/j.ijhydene.2014.01.203).
- 46 J. Han, *et al.*, Role of Au–TiO<sub>2</sub> interfacial sites in enhancing the electrocatalytic glycerol oxidation performance, *Electrochem. Commun.*, 2018, **96**, 16–21, DOI: [10.1016/j.elecom.2018.09.004](https://doi.org/10.1016/j.elecom.2018.09.004).

