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In situ fabricated nickel vanadate/N-doped reduced graphene oxide hybrid as an advanced electrocatalyst in alkaline hydrogen evolution reaction†

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In the current work, we reported the *in situ* fabrication of a nickel vanadate ($\text{Ni}_3\text{V}_2\text{O}_8$)-anchored N-doped reduced graphene oxide (NRGO) hybrid by a simple one-step reflux method. Subsequently, the electrocatalytic performance of the $\text{Ni}_3\text{V}_2\text{O}_8$ /NRGO hybrid was investigated in an alkaline medium (1.0 M KOH) for the hydrogen evolution reaction (HER). It was noted that the optimization of the NRGO content (~5.6 wt%) highly influenced the homogeneous distribution of quasi-spherical $\text{Ni}_3\text{V}_2\text{O}_8$ nanoparticles over NRGO sheets and enhanced the water reduction ability. This hybrid material exhibited a sufficiently high electrochemical active surface area (517.5 cm^2) and remarkably low charge transfer resistance (~1.6 Ω). Furthermore, the very low overpotential (~43 mV) and the high exchange current density (~1.24 mA cm^{-2}) of $\text{Ni}_3\text{V}_2\text{O}_8$ /NRGO (5.6 wt%) demonstrated its promising HER performance. Additionally, superior long-term and accelerated stability compared to that of the benchmark Pt/C in a strong basic medium clearly signified that $\text{Ni}_3\text{V}_2\text{O}_8$ /NRGO (5.6 wt%) can act as an efficient, cost-effective, and durable electrocatalyst for water electrolyzers.

Hydrogen is one of the most important alternative fuels compared to many conventional energy resources.¹ In this regard, the electrolysis of water in an alkaline medium remains the most cost-effective, sustainable and industrially accepted method for hydrogen production.² However, the high overpotential of the hydrogen evolution reaction (HER) affects the efficiency of hydrogen generation.³ Therefore, it is mandatory to lower the overpotential of HER in the presence of suitable electrocatalyst(s). As a result, platinum (Pt) has been reported to be the most active HER electrocatalyst of recent times.⁴

However, the widespread use of Pt for industrialization and commercialization is limited due to its scarcity, very high cost and inferior durability in strong alkaline solutions.⁵ According to the available literature, mixed transition metal oxides compared to the corresponding monometallic oxides show better electrochemical activity due to the presence of a large number of active sites.^{6,7} In this regard, transition metal vanadates, especially $\text{Ni}_3\text{V}_2\text{O}_8$, $\text{Co}_3\text{V}_2\text{O}_8$, and CoVO_x , exhibit enhanced electrochemical activity owing to the multiple valence oxidation states of vanadium.^{8–11} In view of this, $\text{Ni}_3\text{V}_2\text{O}_8$ nanowires grown on a Ti-foil,¹² $\text{Ni}_3\text{V}_2\text{O}_8$ /carbon cloth,⁸ urchin-shaped $\text{Ni}_3(\text{VO}_4)_2$ hollow nanospheres,¹³ nickel vanadate nanoparticles,¹⁴ and $\text{Ni}_3(\text{VO}_4)_2$ /rGO nanocomposites¹⁵ have been successfully investigated for their applications in lithium-ion batteries, sodium-ion batteries, and supercapacitors. However, there exists very limited literature related to the application of $\text{Ni}_3\text{V}_2\text{O}_8$ in electrocatalytic energy conversions. Shang *et al.*¹⁶ observed a moderate overpotential of ~113 mV at 10 mA cm^{-2} current density for HER in a 1.0 M KOH solution in the nickel foam-supported combination of two mixed transition metal oxides ($\text{Ni}_3(\text{VO}_4)_2$ @ NiCo_2O_4 /NF). In addition, sea-urchin-like $\text{Ni}_3(\text{VO}_4)_2$ nanostructures showed promising electrocatalytic HER activity with a very small overpotential (η_{10}) of about 90 mV at 10 mA cm^{-2} and a small Tafel slope (50 mV dec^{-1}) in 0.5 M H_2SO_4 .¹⁷ However, the low charge transfer resistance and stability of electrocatalysts (non-precious) in a strong alkaline medium are few other desirable parameters to be taken into consideration for their superior electrocatalytic performance.^{4,18}

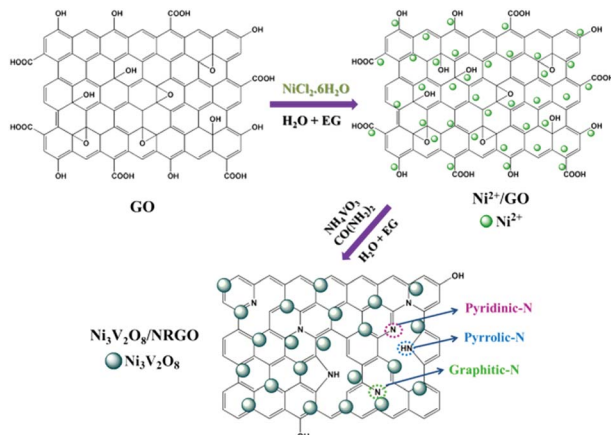
Motivated by these facts, we focused our work on the fabrication of a nickel vanadate ($\text{Ni}_3\text{V}_2\text{O}_8$) nanoparticle-decorated nitrogen-doped RGO (NRGO) hybrid through a simple one-step reflux method (ESI†). Scheme 1 clearly demonstrates the *in situ* fabrication of the $\text{Ni}_3\text{V}_2\text{O}_8$ /NRGO hybrids. The choice of NRGO in the present work was mainly guided by its high electrical conductivity and the presence of larger defects. Therefore, it is anticipated that the structural stability of the electrode material can be enhanced in the presence of NRGO as the conducting backbone in a strong alkaline medium. Notably, to

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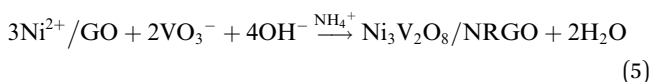
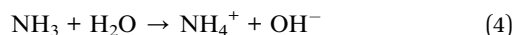




Scheme 1 Probable formation pathway of $\text{Ni}_3\text{V}_2\text{O}_8/\text{NRGO}$ hybrids.

the best of our knowledge, no reports are available on evaluating the electrocatalytic HER performance of an $\text{Ni}_3\text{V}_2\text{O}_8/\text{NRGO}$ hybrid in an alkaline solution. In this context, our findings showed that the $\text{Ni}_3\text{V}_2\text{O}_8/\text{NRGO}$ (~5.6 wt%) hybrid demonstrated superior HER activity in an alkaline medium.

The formation pathway of the $\text{Ni}_3\text{V}_2\text{O}_8/\text{NRGO}$ hybrids through a simple one-step reflux method, as displayed in Scheme 1, consists of the adsorption of Ni^{2+} ions on the surface of graphene oxide (GO). This is most likely due to the electrostatic attraction between the positively charged metal ions (Ni^{2+}) and the negatively charged oxygen functionalities ($-\text{COOH}$, $-\text{OH}$) of GO (Fig. S2†). Subsequently, the solutions of NH_4VO_3 (2 mmol) and urea (12 mmol) prepared in a mixed solvent (water + EG) were added to the earlier dispersion of Ni^{2+}/GO . As a result, NH_3 (source of nitrogen) was formed during the hydrolysis and the transesterification of urea in the presence of water and ethylene glycol, respectively.¹⁹ In addition, Ni^{2+} and VO_3^- ions originating from $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ and NH_4VO_3 combined together to form $\text{Ni}_3\text{V}_2\text{O}_8$ in the basic medium. Furthermore, ammonia generated during the hydrolysis and the transesterification of urea successfully reduced GO to RGO, followed by simultaneous N doping to form the $\text{Ni}_3\text{V}_2\text{O}_8/\text{NRGO}$ hybrid. The plausible reactions involved in the fabrication of the $\text{Ni}_3\text{V}_2\text{O}_8/\text{NRGO}$ hybrid can be summarized as follows:^{20,21}



Thermogravimetric analysis (TGA) in air revealed that the NV/NRGO1, NV/NRGO2, and NV/NRGO3 hybrids contained ~3.3, 5.6, and 6.6 wt% NRGO, respectively (ESI†). Furthermore,

the results of the CHN analyses of the hybrids were also in agreement with the TGA results (Table S1†). The X-ray diffraction (XRD) patterns of the NV, NRGO, and NV/NRGO hybrids were obtained and the corresponding findings are displayed in Fig. 1a. The XRD pattern of NRGO showed the appearance of peaks at $\sim 23.8^\circ$ (002) and $\sim 43.0^\circ$ (100) similar to that reported for reduced graphene oxide.²² It is also evident from Fig. 1a that the XRD patterns of NV as well as NV/NRGO hybrids show the formation of the pure orthorhombic phase (JCPDS no. 74-1484) of $\text{Ni}_3\text{V}_2\text{O}_8$ without any impurity peaks corresponding to NiO or V_2O_5 . Interestingly, the intensity of the NRGO peaks almost disappeared for NV/NRGO hybrids, indicating the possible formation of $\text{Ni}_3\text{V}_2\text{O}_8$ on highly exfoliated NRGO sheets. This could be further ascertained by the morphological investigations of the NV/NRGO hybrids through electron microscopy. The Raman spectra (Fig. 1b) show that the peaks appearing at 760 cm^{-1} (low intensity) and 823 cm^{-1} (high intensity) for NV correspond to the asymmetric and symmetric stretching vibration modes of the V–O bonds.^{15,17} However, the intensity of the peak at 823 cm^{-1} is considerably reduced for all the hybrids, while the peak corresponding to $\sim 760\text{ cm}^{-1}$ almost disappears for the NV/NRGO3 hybrids compared to that for NV. The Raman spectra of the NV/NRGO hybrids also exhibit the presence of D ($\sim 1596\text{ cm}^{-1}$) and G ($\sim 1359\text{ cm}^{-1}$) bands appearing at nearly identical positions. This finding suggests that the graphitic structure remains unaltered in the hybrids.²³ Furthermore, the I_D/I_G ratios corresponding to the NV/NRGO hybrids were calculated and followed the order NV/NRGO2 (0.86) > NV/NRGO1 (0.84) > NV/NRGO3 (0.76). The relatively higher value of the I_D/I_G ratio in NV/NRGO2 could be ascribed to the presence of higher defects/disorders.²⁴

The X-ray photoelectron spectroscopy (XPS) survey scans of NV and NV/NRGO2 are displayed in Fig. 2a. Furthermore, the high-resolution Ni 2p and V 2p spectra of NV were compared with that of NV/NRGO2 (Fig. S4†). It was noted that the Ni 2p and V 2p peaks present in NV/NRGO2 shifted to higher binding energies with respect to that for NV due to the strong electronic interaction between $\text{Ni}_3\text{V}_2\text{O}_8$ and NRGO present in NV/NRGO2. Such interaction between C/N (NRGO) and Ni/V ($\text{Ni}_3\text{V}_2\text{O}_8$) that leads to an increase in the respective binding energies can be accounted on the basis of the reduction in the electron density around Ni/V in NV/NRGO2 as a result of their electronegativity (χ) differences ($\chi_{\text{Ni}} = 1.91$, $\chi_{\text{V}} = 1.63$, $\chi_{\text{C}} = 2.55$, $\chi_{\text{N}} = 3.04$).²⁵ The survey spectrum (Fig. 2a) of NV/NRGO2 clearly shows the

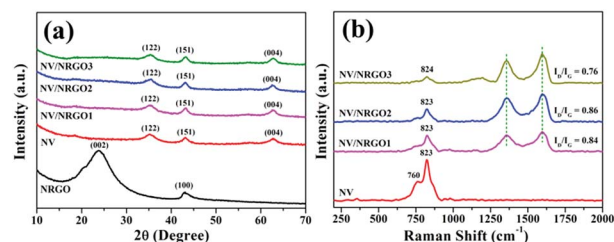


Fig. 1 XRD patterns (a) and Raman spectra (b) of NRGO, NV, NV/NRGO1, NV/NRGO2, and NV/NRGO3.



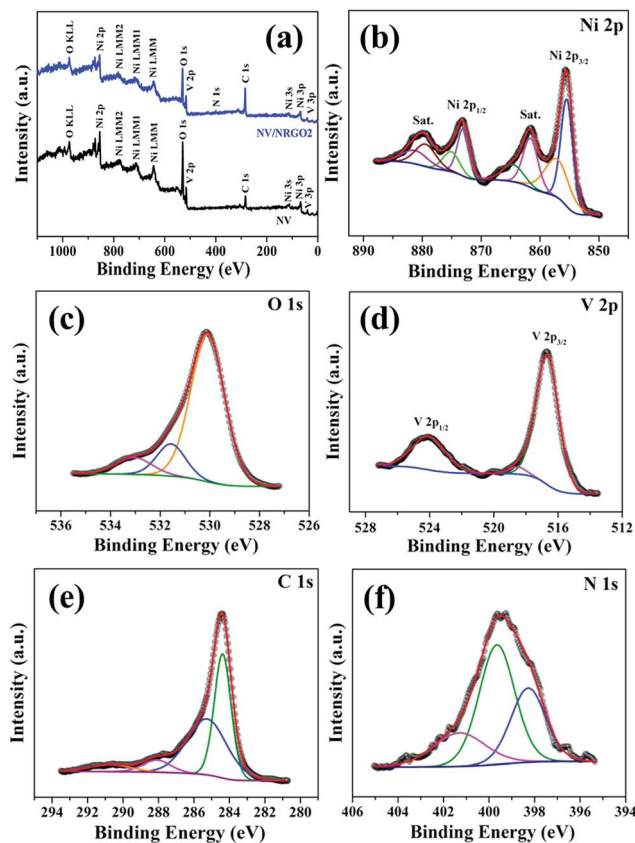


Fig. 2 (a) XPS survey scans of NV and NV/NRGO2 and deconvoluted high-resolution XPS spectra of (b) Ni 2p, (c) O 1s, (d) V 2p, (e) C 1s, and (f) N 1s of NV/NRGO2.

presence of Ni, V, O, C, and N. Fig. 2b–f show the high-resolution core levels and the corresponding deconvoluted spectra of Ni 2p, O 1s, V 2p, C 1s, and N 1s, respectively. The peaks at 873.0 eV (Ni 2p_{1/2}) and 855.4 eV (Ni 2p_{3/2}) in Ni 2p spectra correspond to the presence of Ni²⁺ in NV/NRGO2.^{16,17} This is also supplemented by the presence of satellite peaks at 881.2 and 861.6 eV.¹⁶ In addition, the peaks at 875.1 eV (Ni 2p_{1/2}) and 857.1 eV (Ni 2p_{3/2}) can be assigned to the existence of Ni³⁺.²⁶ In the case of O 1s, the peaks at ~530.1, 531.6, and 533.0 eV correspond to the metal–oxygen bond, oxygen vacancies, and surface –OH groups.¹⁵ Furthermore, the peaks at ~524.0, 518.5, and 516.7 eV in the V 2p spectrum can be attributed to the +5 oxidation state of V.^{17,27} The intense peak at 284.4 eV in the C 1s spectrum corresponds to sp² hybridized graphitic C originating from NRGO.⁴ The other additional peaks at ~290.7, 288.1, and 285.3 eV can be assigned to C=O, N–sp³C, and N–sp²C, respectively.¹⁵ The peaks in the spectrum of N 1s can be indexed to graphitic-N (401.3 eV), pyrrolic-N (399.6 eV), and pyridinic-N (398.3 eV).⁴ These findings clearly demonstrate the nitrogen doping (~1.6 at%) of RGO and the formation of highly pure Ni₃V₂O₈/NRGO hybrids. Also, the presence of pyrrolic-N is responsible for the high electrical conductivity of NRGO, while pyridinic-N facilitates electrocatalytic activity.²⁸ Therefore, the optimised NRGO content in NV/NRGO2 is most likely to enhance the electrocatalytic HER performance of Ni₃V₂O₈.

Field emission scanning electron microscopy (FESEM) and transmission electron microscopy (TEM) were employed to reveal the formation of the aggregated sub-micrometre hierarchical spherical particles of NV (Fig. 3a and e) and the distribution of Ni₃V₂O₈ particles on NRGO sheets for the NV/NRGO hybrids (Fig. 3b–h). It was clearly seen that the addition of NRGO resulted in a narrow size and better distribution of the Ni₃V₂O₈ particles on the NRGO sheets compared to that for pure NV. Furthermore, a lower degree of exfoliation of NRGO in the NV/NRGO1 hybrid was observed (Fig. 3b and f). It was noted that the agglomerated Ni₃V₂O₈ particles covered entire NRGO sheets and bare sheets of NRGO were hardly visible. When the amount of GO was increased during the preparation of the NV/NRGO2 hybrid, relatively better exfoliation of NRGO and uniform distribution of Ni₃V₂O₈ particles were observed over the NRGO sheets, as displayed in Fig. 3c and g. The FESEM/TEM images of the NV/NRGO3 hybrid in Fig. 3d and h indicate the irregular distribution of NV particles on NRGO sheets due to their self-agglomeration with the visible bare surface of NRGO, indicating its incomplete coverage. Considering all these observations, it is anticipated that the optimization of the GO/

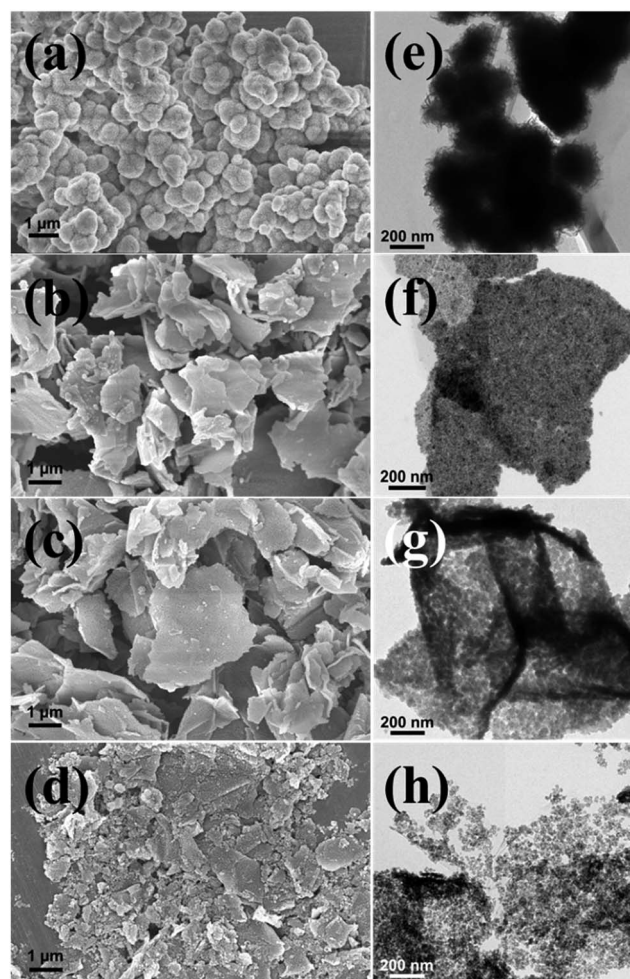


Fig. 3 FESEM images (a–d) and corresponding TEM images (e–h) of NV, NV/NRGO1, NV/NRGO2, and NV/NRGO3.



$\text{Ni}_3\text{V}_2\text{O}_8$ ratio in the fabrication of the NV/NRGO hybrids is the key factor to achieve uniform distribution of the NV particles on the NRGO sheets. Furthermore, the high-magnification FESEM and TEM images (Fig. S5a and b†) of the NV/NRGO2 hybrid also reaffirmed our earlier contention on the homogeneous distribution of the $\text{Ni}_3\text{V}_2\text{O}_8$ nanoparticles on the NRGO sheets. The corresponding HRTEM image of the NV/NRGO2 hybrid is also displayed in Fig. S5c.† It clearly indicates the growth of the (122) plane ($d = 0.249$ nm), and the presence of concentric rings in the selected area diffraction pattern (SAED) (in the inset of Fig. S5c†) demonstrates its polycrystalline nature. The elemental composition analysis of the marked region of NV/NRGO2 in Fig. S5d† was carried out through STEM-elemental mapping and the results are displayed in Fig. S5(e-h),† which clearly demonstrate the presence of Ni, V, O, and N in NV/NRGO2. It may be noted that the mapping of C was not taken into account due to its presence in the carbon-coated copper grid. The energy dispersive X-ray (EDX) analysis of NV/NRGO2 in Fig. S6† also confirms that the Ni-to-V atomic ratio is $\sim 3 : 2$, indicating the formation of $\text{Ni}_3\text{V}_2\text{O}_8$, as observed from the XRD analysis (Fig. 1a). In addition, the Ni-to-V molar ratio (~ 1.59) obtained from the ICP-MS analysis (Table S2†) confirmed our observations obtained from EDX.

The electrocatalytic HER polarization curves of Pt/C (20 wt%) and the NV, NRGO, NV/RGO and NV/NRGO hybrids (scan rate: 5 mV s^{-1}) in 1.0 M KOH are displayed in Fig. 4a. The overpotentials (η) required to achieve 10 mA cm^{-2} cathodic current density were found to be $\sim 16, 97, 194, 236, 94, 43,$ and 81 mV for Pt/C, NV, NRGO, NV/RGO, NV/NRGO1, NV/NRGO2, and NV/NRGO3, respectively. Our investigations on the HER activity of NV ($\eta_{10} = 97 \text{ mV}$) and NRGO sheets ($\eta_{10} = 194 \text{ mV}$) established the synergistic effect in NV/NRGO2 ($\eta_{10} = 43 \text{ mV}$). Furthermore, control experiments were also performed on NiO and V_2O_5 to evaluate their performance in HER (Fig. S7†). It was observed that NiO ($\eta_{10} = 200 \text{ mV}$) and V_2O_5 ($\eta_{10} = 163 \text{ mV}$) individually exhibit lower activities compared to NV ($\eta_{10} = 97 \text{ mV}$) in HER. Therefore, the synergistic contribution of NiO and V_2O_5 in addition to that of NV and NRGO as stated earlier can account for the improved hydrogen evolution ability owing to the synergistic effect in NV/NRGO2. It was noted that NV/NRGO1 showed a slightly better performance compared to NV in the low overpotential region although its performance deteriorated at higher overpotentials (Table S3†). This phenomenon could be attributed to the low exfoliation of NRGO as well as the aggregation and surface blockage of the $\text{Ni}_3\text{V}_2\text{O}_8$ particles in NV/NRGO1. Alternatively, the possibility of a less number of active sites coming in contact with the electrolyte as a result of the faster rate of hydrogen evolution at the high overpotential regions could not be overruled. In contrast, NV/NRGO2 loaded with 5.6 wt% of NRGO achieved the lowest value of η_{10} (43 mV), demonstrating its superior electrocatalytic HER activity. Such superior HER activity of NV/NRGO2 could be assigned to the uniform distribution of quasi-spherical $\text{Ni}_3\text{V}_2\text{O}_8$ nanoparticles on NRGO sheets, thereby facilitating a better contact between active sites and the electrolyte (OH^- ions).²⁹ However, the NV/NRGO3 (6.6 wt%) hybrid exhibited lower electrocatalytic activity due to the restacking of NRGO sheets, self-

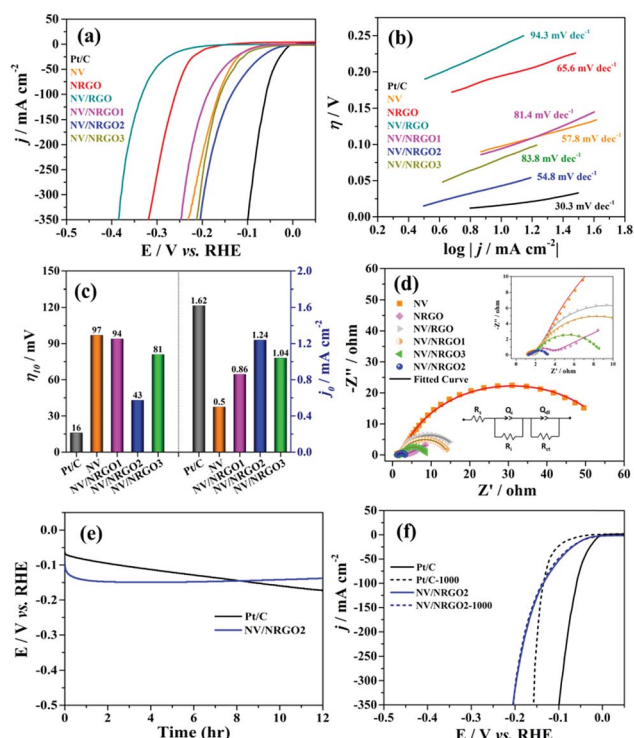
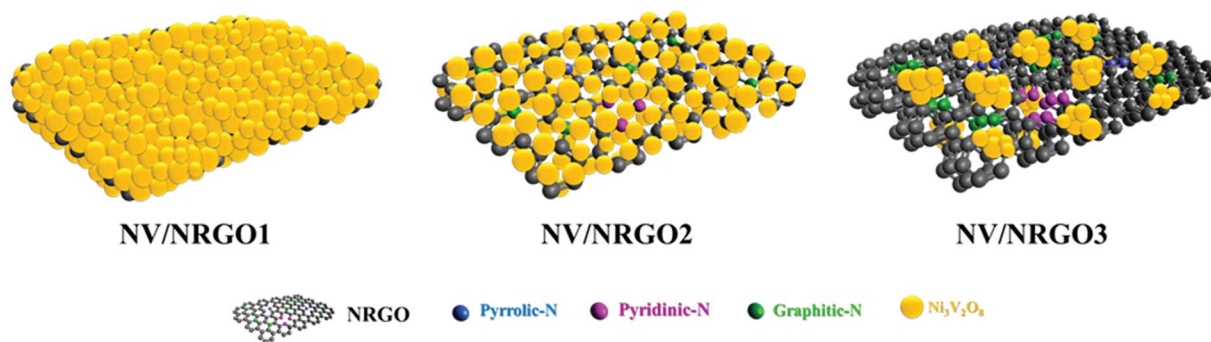


Fig. 4 (a) LSV curves at 5 mV s^{-1} . (b) Tafel plots derived from LSV curves of Pt/C, NV, NRGO, NV/RGO, NV/NRGO1, NV/NRGO2, and NV/NRGO3. (c) Overpotential (η_{10}) and exchange current density (j_0) of Pt/C, NV, NV/NRGO1, NV/NRGO2, and NV/NRGO3. (d) Nyquist plots at $\eta = 244 \text{ mV}$ (equivalent circuit and the magnified plot are shown in the inset) of NV, NRGO, NV/RGO, NV/NRGO1, NV/NRGO2, and NV/NRGO3. (e) CP curves for 12 h without iR-correction and (f) initial LSV curves (solid lines) and LSV curves after 1000 cycles (dotted lines) of Pt/C and NV/NRGO2. Unless mentioned, all the measurements were carried out with iR-compensation at room temperature in 1.0 M KOH.

agglomeration, and the irregular distribution of catalytic active sites.²⁹ Scheme 2 shows the distribution of the active sites in NV/NRGO1, NV/NRGO2, and NV/NRGO3 considering these aspects and the TEM studies discussed earlier. Furthermore, the HER performance of NV/RGO was also compared with that of the NV/NRGO2 hybrid to ascertain the advantages of NRGO if any. Our findings clearly demonstrated the relative inferior performance of NV/RGO ($\eta_{10} = 236 \text{ mV}$) due to the poor exfoliation/aggregation of RGO sheets and NV particles, as evident from the FESEM image (Fig. S8b†). In contrast, NV/NRGO2 displayed a superior HER performance due to the homogeneous distribution of NV particles in the NRGO matrix, as indicated by the TEM image shown in Fig. S5b.† Alternatively, the fact that the presence of enhanced defect sites in NRGO can contribute to the improvement in its HER activity also cannot be ruled out. According to our findings, NV/NRGO2 also outperformed other HER electrocatalysts such as $\text{Ni}_3(\text{-VO}_4)_2@\text{NiCo}_2\text{O}_4/\text{NF}$ (113 mV),¹⁶ $\text{Ni}_3\text{N}@\text{VN-NF}$ (56 mV),³⁰ N-Co-C/RGO (130 mV),³¹ Sr_2RuO_4 (61 mV),³² $\beta\text{-Ni}(\text{OH})_2/\text{Pt}$ (108 mV),³³ and nest-like NiCoP (62 mV)³⁴ (Table S4†).

A Tafel slope is useful to gain insights into the mechanistic pathway through the rate-determining step (RDS). According to





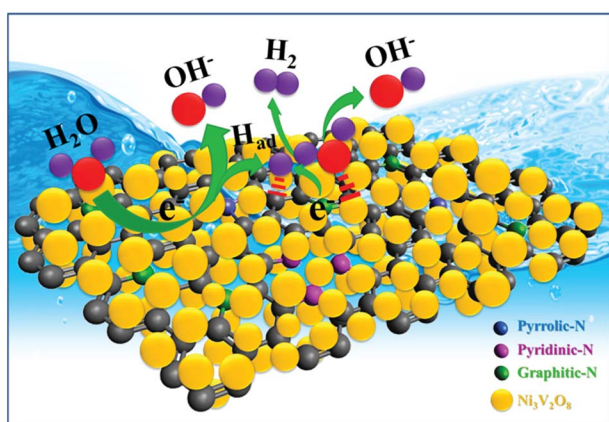
Scheme 2 Distribution of active sites in NV/NRGO hybrids.

the available literature, HER in an alkaline medium proceeds through the Volmer–Heyrovsky or Volmer–Tafel pathways.³³ In the Volmer step, the dissociation of a surface-adsorbed water molecule results in the formation of surface-adsorbed hydrogen atoms ($S\text{-H}_{\text{ad}}$) and the generation of OH^- ($\text{H}_2\text{O} + \text{S} + \text{e}^- \rightarrow \text{S}\text{-H}_{\text{ad}} + \text{OH}^-$). Subsequently, $\text{S}\text{-H}_{\text{ad}}$ either combines with another H_2O molecule in the Heyrovsky step ($\text{H}_2\text{O} + \text{S}\text{-H}_{\text{ad}} + \text{e}^- \rightarrow \text{H}_2 + \text{S} + \text{OH}^-$) or recombines with another $\text{S}\text{-H}_{\text{ad}}$ to release molecular hydrogen in the Tafel step ($2\text{S}\text{-H}_{\text{ad}} \rightarrow 2\text{S} + \text{H}_2$).^{35,36} According to the Butler–Volmer kinetics, the rate-determining step can be identified depending on the Tafel slope as the Volmer (120 mV dec^{-1}), Heyrovsky (40 mV dec^{-1}), or Tafel (30 mV dec^{-1}) reaction.^{33,35} We observed the Tafel slope (Fig. 4b) values of ~ 30.3 , 57.8 , 65.6 , 94.3 , 81.4 , 54.8 , and 83.8 mV dec^{-1} corresponding to Pt/C, NV, NRGO, NV/RGO, NV/NRGO1, NV/NRGO2, and NV/NRGO3. These findings suggested Tafel and Heyrovsky reactions as the rate determining steps, confirming the overall Volmer–Tafel and Volmer–Heyrovsky pathways in Pt/C and NV/NRGO2 (Scheme 3), respectively. It was anticipated that the relatively lowest Tafel slope of NV/NRGO2 (54.8 mV dec^{-1}) among those of NV and all its NRGO hybrids, would indicate faster reaction kinetics in HER. It was also noted that the Tafel slope for NV/NRGO2 (54.8 mV dec^{-1}) was lower in comparison to the results of the electrocatalysts studied in the contemporary

works reported in the literature such as NiO/Ni-CNT (84.6 mV dec^{-1}),³⁷ VOOH hollow nanospheres (104 mV dec^{-1}),³⁸ $\text{Fe}_{17.5\%}\text{-Ni}_3\text{S}_2/\text{NF}$ (95 mV dec^{-1}),³⁹ $\text{P}_{8.6}\text{-Co}_3\text{O}_4/\text{NF}$ (86 mV dec^{-1}),⁴⁰ Cu NDs/ Ni_3S_2 NTs-CFs (76.2 mV dec^{-1}),⁴¹ and $\text{Ni}_3\text{N}@$ CQDs (108 mV dec^{-1})⁴² for HER in an alkaline medium.

Moreover, exchange current density (j_0) is an important parameter to assess the intrinsic electrocatalytic activity of a catalyst. In this regard, the Tafel plot was extrapolated⁴³ to $\eta = 0$ and the j_0 values of NV (0.05 mA cm^{-2}), NV/NRGO1 (0.86 mA cm^{-2}), NV/NRGO2 (1.24 mA cm^{-2}) and NV/NRGO3 (1.04 mA cm^{-2}) were evaluated, as shown in Fig. 4c. The excellent HER activity of NV/NRGO2 is reflected from its j_0 value, which is closer to that of Pt/C (1.62 mA cm^{-2}). Furthermore, the intrinsic activity of an electrocatalyst can be correlated to the turnover frequency (TOF). Accordingly, the TOF values at $\eta = 100 \text{ mV}$ follow the order NV/NRGO2 (0.021 s^{-1}) > NV (0.013 s^{-1}) > NV/NRGO1 (0.008 s^{-1}) > NV/NRGO3 (0.004 s^{-1}). The highest TOF value of NV/NRGO2 further confirmed our earlier contention on its superior intrinsic HER activity. The electrochemical active surface area (ECSA) is an important parameter for the surface redox reactions at the electrode.⁴⁴ Accordingly, the obtained ECSA values (ESI†) follow the order (Fig. S11b†) NV (92.5 cm^2) < NV/NRGO1 (297.5 cm^2) < NV/NRGO3 (455.0 cm^2) < NV/NRGO2 (517.5 cm^2). These findings clearly suggest that NV/NRGO2 consists of a large number of active sites owing to its highest ECSA compared to all other synthesized electrocatalysts. The Nyquist plots ($\eta = 244 \text{ mV}$) of NV, NRGO, NV/RGO, NV/NRGO1, NV/NRGO2, and NV/NRGO3 in Fig. 4d demonstrate the corresponding R_{ct} values of ~ 57.1 , 2.2 , 15.4 , 12.6 , 1.6 , and 6.9Ω . The high electronic conductivity of the NRGO sheets accounted for the significantly decreased R_{ct} value for the NV/NRGO hybrids in comparison to that for NV. Furthermore, the remarkably low R_{ct} value (1.6Ω) of NV/NRGO2 also supported its highly enhanced HER activity.

The long-term stability and the durability of an electrocatalyst are considered to be the most important criteria for its practical applications.^{45,46} In view of this, Pt/C and NV/NRGO2 were investigated by the chronopotentiometry (CP) technique (without iR-compensation) at 10 mA cm^{-2} for their long-term durability (12 h); potential (E) versus time (t) is displayed in Fig. 4e. Our findings demonstrated the appearance of three regions in the E vs. t plot of NV/NRGO2: 92 mV ($t = 0$) to 149 mV



Scheme 3 Volmer–Heyrovsky pathway of hydrogen evolution in an alkaline medium for NV/NRGO2.



($t = 2.3$ h), 149 mV ($t = 2.3$ h) to 149 mV ($t = 5.6$ h) and 149 mV ($t = 5.6$ h) to 137 mV ($t = 12$ h). This clearly showed that the increase in E was more pronounced in the beginning up to 2.3 h, followed by an almost constant E up to 5.6 h. Thereafter, E , in contrast to that of the Pt/C electrode, decreased from 149 to 137 mV, corresponding to the increase in time from 5.6 to 12 h. A similar behaviour was also noticed for NiCo₂S₄/NF⁴⁷ and CoS|P/CNT.⁴⁸ In contrast to the result for NV/NRGO₂, the poor long-term durability of Pt/C in 1.0 M KOH was evident from the continuous increase in the overpotential from 67 mV to 172 mV. Subsequently, accelerated stability of the electrocatalysts was achieved by subjecting them to 1000 CV cycles at a scan rate of 100 mV s⁻¹. The corresponding polarization curves (iR-compensated) of Pt/C and NV/NRGO₂ (Fig. 4f) exhibited an almost unchanged η_{10} value for NV/NRGO₂ in contrast to the increase in η_{10} by 59 mV for Pt/C. Therefore, NV/NRGO₂ demonstrated much better long-term and accelerated stability in comparison to Pt/C in alkaline HER. Further investigations were carried out to study the performance of NV/NRGO₂ as a water-splitting electrocatalyst for a real alkaline (1.0 M KOH) electrolyzer in a two-electrode system (NV/NRGO₂||NV/NRGO₂) (Fig. S12†). It was noted that a current density value of 10 mA cm⁻² was achieved in the cell at 1.77 V, which was found to be lower than that reported in previously published works on Ni(OH)₂/NF||Ni(OH)₂/NF (1.82 V)⁴⁹ and commercial water electrolyzers (1.8 to 2.0 V).⁴⁹

The structural stability of NV/NRGO₂ after the durability tests was studied by XRD, FESEM and Raman spectroscopy; the corresponding findings are displayed in Fig. S13.† The XRD patterns of NV/NRGO₂ before and after HER are compared in Fig. S13(i).† The appearance of prominent peaks at 35.1° (122), 43.0° (151), and 62.5° (004) is noted in the XRD pattern of NV/NRGO₂ after electrocatalysis (Fig. S13(ib)†), corresponding to the orthorhombic phase of Ni₃V₂O₈ (JCPDS no. 74-1484). In addition, the appearance of the (002) peak at 23.1° can be ascribed to the presence of graphitic carbon in NRGO. Alternatively, the possibility of a peak appearing due to carbon black used in the preparation of the working electrode also cannot be ruled out. An additional low intense peak also appeared at 11.3°, which could be assigned to the (003) plane of NiOOH (JCPDS no. 06-0075) due to the continuous electrolysis of NV/NRGO₂ in the alkaline medium.¹⁶ Furthermore, the Raman spectrum of the sample (after HER) remains more or less unaltered except for the slightly reduced I_D/I_G ratio (0.82) with respect to that for pristine NV/NRGO₂. This could be attributed to the decrease in the density of the defects in NV/NRGO₂ after the HER durability test. The XPS analysis of NV/NRGO₂ was also performed after the stability test and the corresponding findings are displayed in Fig. S14.† The XPS survey scan of the post-HER sample demonstrated the presence of Ni, V, O, C, and N similar to that for the fresh catalyst. The high-resolution Ni 2p spectrum of the sample after the stability test shows negligible shifts in Ni 2p_{1/2} (872.9 eV) and Ni 2p_{3/2} (855.0 eV) peaks compared to the spectrum of the sample before HER. In contrast, two additional peaks appear at 873.5 and 856.2 eV in the deconvoluted spectrum of Ni 2p. This can be ascribed to the formation of NiOOH owing to the surface oxidation of the catalyst by adsorbed OH⁻

ions as a result of prolonged use in a strong alkaline medium.¹⁶ Similarly, the new peaks observed in the V 2p region at 521.8 and 515.2 eV can be assigned to VOOH.³⁸ In addition, the O 1s spectrum shows peaks at 531.1, 530.2, and 528.6 eV due to H₂O, O–H, and the metal–oxygen bond, respectively.³⁸ In addition, the comparison of the FESEM images of NV/NRGO₂ before (Fig. 3c) and after (Fig. S13(ii)†) HER shows some changes in the morphology due to the presence of carbon black and the polymeric binder (PVDF) used in electrode preparation. All these findings further strengthen our earlier inference: NV/NRGO₂ is a suitable electrocatalyst for practical applications.

The superior HER activity of NV/NRGO₂ to those of other electrocatalysts may be ascribed to the synergistic effect of Ni₃V₂O₈ particles with NRGO sheets, uniform particle distribution, N doping, and the optimised NRGO content. Alternatively, the remarkable increase in the charge transportation of NRGO due to high electrical conductivity also cannot be ruled out. Furthermore, the higher ECSA value of NV/NRGO₂ (517.5 cm²) most likely originates from the better exposure of the electrolyte to the active sites. Additionally, NRGO also imparted structural stability in its long-term application, as evidenced by post-HER characterizations. Therefore, NV/NRGO₂ can act as an effective and stable electrocatalyst in an alkaline medium for the hydrogen evolution reaction.

Our findings clearly showed that NV/NRGO₂ exhibited superior durability in contrast to other electrochemical properties compared to commercial Pt/C. Most likely, Pt/C facilitates the exposure of relatively more active sites to enable rapid electron and mass transfer.⁵⁰ Similar HER activity could be achieved for NV/NRGO₂ by generating more active sites *via* tuning its morphology or by the modification of the electronic structure by doping.³⁵ According to the available literature, an effective alkaline HER electrocatalyst should have the ability of efficient water dissociation at the surface and proper binding with hydrogen intermediates.^{35,51} This could be facilitated by adding another active component on the surface of NV/NRGO₂ to achieve similar/better HER activity to/than that of Pt/C in an alkaline medium.^{52–54}

In summary, an Ni₃V₂O₈/NRGO hybrid with 5.6 wt% NRGO content was successfully fabricated through refluxing and employed as an electrocatalyst for HER in an alkaline medium (1.0 M KOH). The superior performance of the Ni₃V₂O₈/NRGO (5.6 wt%) hybrid manifested a very low overpotential (~43 mV), small Tafel slope (~54.8 mV dec⁻¹), and quite high exchange current density (~1.24 mA cm⁻²) for hydrogen evolution in 1.0 M KOH. Furthermore, the value of the Tafel slope suggested that HER proceeded through the Volmer–Heyrovsky mechanism. In addition, this hybrid electrocatalyst also demonstrated good long-term and accelerated stability in a strong alkaline solution. Therefore, Ni₃V₂O₈/NRGO (5.6 wt%) can be used as a low-cost, highly efficient, and stable alternative to the commercial Pt/C electrocatalyst for the industrial production of hydrogen through alkaline water electrolysis.

Conflicts of interest

There are no conflicts of interest to declare.



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