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Spherical Ni/NiO nanoparticles decorated on nanoporous carbon (NNC) as an active electrode material for urea and water oxidation reactions†

 Parag P. Chavan,^{‡ab} Pratiksha D. Tanwade,^{‡a} Vijay S. Sapner^{id ac}
and Bhaskar R. Sathe^{id *ad}

Herein, we report a chemical method for scalable synthesis of spherical Ni/NiO nanoparticle-decorated nanoporous carbon (NNC) based electrocatalytic system using a simple and easy chemical method with ultra-high activity towards urea electrooxidation. Morphological analysis by scanning electron microscopy (SEM) and high-resolution transmission electron microscopy (HR-TEM) confirms the formation of Ni/NiO NPs on highly nanoporous carbon with an average size of ~50 nm. X-ray diffraction (XRD) confirms NNC with a face-centred cubic (FCC) crystal structure. Ni/NiO NPs intercalated with nanoporous carbon exhibited the best electrocatalytic performance towards urea oxidation with an ultra-low onset potential of ~0.33 V vs. SCE, and faster electrokinetic mechanism confirmed from Tafel slope (~45 mV dec⁻¹), EIS R_{ct} (~6.98 Ω), and long term durability for 7 h at 10 mA cm⁻² with high CO poisoning tolerance. This work affords noble metal-free electrocatalysts for novel appliances and remarkable potential for urea determination, hydrogen generation, real-time water remediation, and energy conversion.

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1. Introduction

One of the most vital tasks is finding an eco-friendly electrocatalyst and an efficient way to create H₂. Electrocatalytic water splitting is considered to be the most appropriate technology.¹ Although there are many advantages to OER, it is limited by the intrinsically slow kinetics of the process (which involves four electron-transfer processes and 1.23 V vs. RHE).^{2,3} This increases the cost as well as energy waste due to the use of noble metal-based electrocatalysts, which severely limits their use on a large scale. To overcome this obstacle, finding alternative strategies to fabricate high-performance OER electrocatalysts is desirable. In recent years, it has been shown that anodic OER can be replaced with electrocatalytic oxidation of small molecules, such as urea,⁴ benzyl alcohol,⁵ and hydrazine,⁶ which can

considerably reduce the overpotential required for H₂ production.

Denitrification in sewage treatment is currently a major concern since high nitrogen pollution contributes to odorous odours, aquatic bio-toxicity, and even dangerously eutrophic diseases. As a result, available urea decomposition is in great demand for wastewater treatment plants and the overall environment. However, conventional technologies, such as biological means, reverse osmosis, electrochemical, and photochemical methods, are still limited by the disadvantages of traditional management, and most importantly, they should easily convert urea to ammonia or nitrate instead of harmless N₂, resulting in terrible energy consumption and nitrogen recontamination. To this end, it is very urgent to propose a new method to directly and completely degrade urea to N₂ and CO₂.

Noble metals such as Pt,⁷ Ir,⁸ Ru,⁹ and others are expensive; hence, non-noble metal catalysts that are affordable are being studied for the oxidation of small organic molecules in alkaline media. Ni and Ni-based hybrid electrodes are widely employed as electrocatalysts in the oxidation and determination of organics such as methanol, ethanol, glucose, cyclohexanol, and aspirin. Aladeemy *et al.*¹⁰ successfully demonstrated the electrolysis of urea to produce H₂ using Ni electrodes in an alkaline medium. Several recent experiments and theoretical investigations have been conducted to uncover urea hydrolysis in the presence of urease (containing Ni active sites).¹¹ However, there has been little research into the electrochemical oxidation mechanism of urea on Ni-based systems in alkaline media.^{3,12}

^aDepartment of Chemistry, Dr Babasaheb Ambedkar Marathwada University, Aurangabad 431004, MS, India. E-mail: bhaskarsathe@gmail.com; Tel: +91-8275306471

^bDepartment of Chemistry, School of Science, Sandip University, Nashik, MS, India

^cDepartment of Chemistry, Shri Mathuradas Mohota Collage of Science, Nagpur-440024, MS, India

^dDepartment of Nanotechnology, Dr Babasaheb Ambedkar Marathwada University, Aurangabad 431004, MS, India

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‡ These authors contributed equally to this work.



500 cm^{-1} on a PerkinElmer Spectrum-I spectrometer. Field emission scanning electron microscopy and Raman spectroscopy were performed using a Raman spectrometer with a microscope (Seki Technotron Corp., Tokyo) with a 532 nm laser. UV-visible spectroscopy of the samples was done using a Jasco V-750 spectrometer. X-ray photoelectron spectroscopy (XPS) was analyzed by a Thermo Fischer Scientific ESCALAB Xi+ instrument. All electrochemical studies were performed using a CHI-660E potentiostat (CH-instrument, USA) and a three-electrode cell comprising a saturated calomel (SCE) reference electrode, a Pt wire counter electrode, and a working electrode.

3. Results and discussion

3.1 Scanning electron microscopy (SEM) analysis

Scanning electron microscopy (SEM) is used to characterize the morphology of NiO-NPs decorated on nanoporous carbon, *i.e.* (NNC) (Fig. 1(a) and (b)). The nanoporous carbon has a well-defined and interlinked 3D network, as well as a uniform distribution of Ni and NiO nanoparticles (yellow circle) ranging in size from 10 to 50 nm. This is consistent with the tiny size of the pores (green circle) depicted in Fig. 1(b), with pore sizes ranging from 50 to 70 nm. It is obvious from the images that NiO-NPs are uniformly dispersed over a mesh-like structure of carbon, *i.e.*, nanoporous carbon.

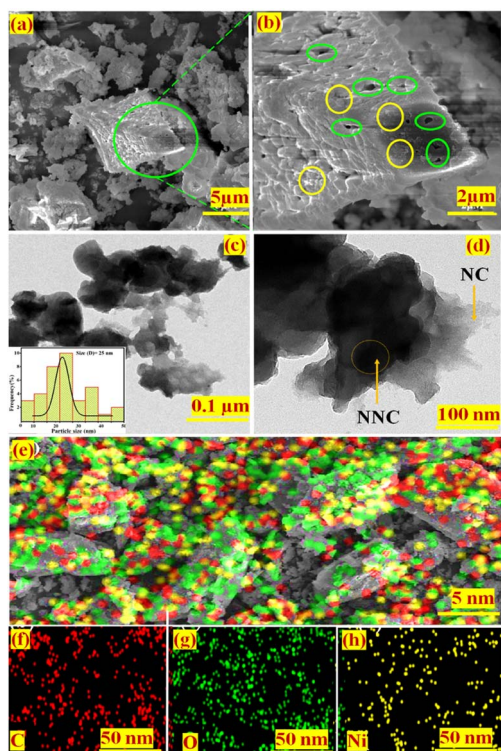


Fig. 1 (a and b) Different magnification field emission scanning electron microscopic (FE-SEM) images having porous carbon nanoparticles (green circle) along with the decorated NiO-NPs (yellow circle). (c and d) Different magnifications of HR-TEM images having porous carbon nanoparticles with successful doping of NiO-NPs, (e–h) are the EDAX images of NNC, C, O, and Ni, respectively.

3.2 Transmission electron microscopy (TEM) analysis

Furthermore, the morphology of the NNC material is scanned using transmission electron microscopy (TEM) and shown in Fig. 1(c) and (d). This demonstrates that the NNC catalyst has a highly porous carbon structure and well-distributed Ni/NiO spherical NPs, with an average particle size of ~ 25 nm, resulting in good functionalization of nano-porous carbon. Ni core-shell and surface are covered with O and are uniformly spread across the nanoporous carbon sheets.

3.3 Energy dispersive analysis of X-ray analysis

The elemental analysis of NNC is performed by SEM using energy dispersive spectral analysis of X-ray (EDAX), which evaluates the energy and intensity distribution of X-ray signals from the surface. Fig. 1(e) depicts the uniform distribution of Ni and NiO NPs over nanoporous carbon. While Fig. 1(f–h) depicts elemental analysis by EDAX containing C, O, and Ni, with corresponding elemental compositions of 48%, 35%, and 17% for NNC.

3.4 X-ray diffraction (XRD)

XRD pattern used to identify the crystal phases and crystallinity of NNC electrocatalysts. From Fig. 2(a) the planes of (111), (200) and (101) match with metallic Ni [JCPDS card no. 04-0850]. While intense planes of (002), and (111) with some weaker signals for (200) and (220) matching NiO formation JCPDS card no. 45-1027 forming a Face Centered Cubic Structure for NNC material. Accordingly, NC material exhibits peaks at the plane of (002) and (102) matching with [JCPDS card no. 00-041-1487]. Overall data is support to literature and conforming the formation of Ni/NiO NPs on NC support.^{16–18}

3.5 FTIR analysis

FTIR analysis validated the different functional groups of NC and changes after Ni/NiO decoration, *i.e.*, of NNC, as shown in Fig. 2(b). As a result, the overlaid FTIR spectra of NC and NNC demonstrated that C–H, C–C, C=O (carbonyl), and C=C have similar intense and well-defined distinctive peaks at 1030 cm^{-1} ,

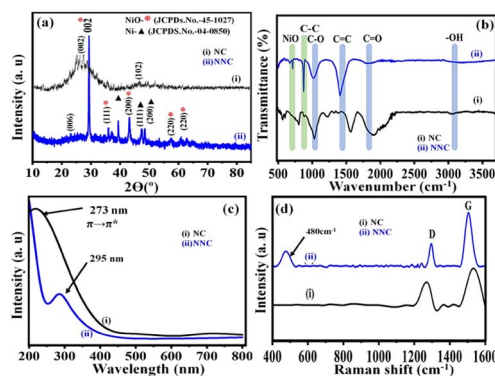


Fig. 2 Superimposed (a) X-Ray Diffraction (XRD), (b) FTIR spectra, (c) UV visible spectra and (d) Raman spectra of (i) NC and (ii) NNC, respectively.



1400 cm^{-1} , 1750 cm^{-1} , and 2200 cm^{-1} , respectively. The additional stretching frequencies 710 cm^{-1} and 875 cm^{-1} correspond to Ni–O–C and Ni–O, respectively, in NNC and are in good agreement with literature.^{15,19}

3.6 UV-visible spectral analysis

It indicates that the decoration of NiO-NPs onto a nano-porous carbon material is successfully achieved. One of the most important approaches for estimating the energy structures and optical properties of nanoparticles is UV-visible absorption spectroscopy. Accordingly, Fig. 2(c) depicts the absorbance by Ni/NiO nanoparticles as a function of wavelength at 302 nm. A faint band at 520 nm and 725 nm was attributed to the charge transfer due to the nanoporous carbon structures. These findings are supported by the physicochemical characterizations and similar systems from the literature.²⁰

3.7 Raman spectroscopy

In addition to this, the structural changes in the carbon framework were investigated by superimposed Raman spectra and shown in Fig. 2(d), monitoring sp^3 (defective) and sp^2 (graphitic) carbons before and after doping with Ni/NiO NPs. The peak at 480 cm^{-1} indicates the extent of Ni decoration on the surface of nonporous carbon. Accordingly, in the case of NC, signals appeared at 1305 cm^{-1} and 1530 cm^{-1} , corresponding to D and G bands, respectively, and their associated intensity ratios, *i.e.*, I_D/I_G are 0.64 and 1.03 for NC and NNC, respectively. The enhancement of the I_D/I_G ratio by more than double could be due to an increase in the disorders in NC after the decoration of Ni/NiO NPs and is also in good agreement with similar reports from the literature.²¹

Thermogravimetric (TG & DT) analysis is shown in (Fig. S1†). In brief, the TGA curve (i) shows two weight losses, *i.e.*, at 120 °C and 460 °C, whereas sharp DTA peaks are associated with weight losses at 450 °C and 560 °C (ii). The first weight loss observed at 100 °C could correspond to the removal of hygroscopic and volatile physically adsorbed species. The second weight loss is caused by the thermal decomposition of supported carbon from NC and the remaining weight of Ni/NiO particles.

3.8 X-ray photoelectron spectral analysis

The qualitative and elemental surface analysis of the NNC composite has been evaluated by using XPS and shown in Fig. 3. Accordingly, Fig. 3(a) shows the full survey having prominent signals corresponding to O 1s, C 1s, and Ni 2p, confirming the NNC formation. Moreover, Fig. 3(b) demonstrates the deconvoluted data for C 1s signal appeared at 284.8 eV, 286.8 eV, 287.5, and 288.2 eV, corresponding to C–C, C=C, C–O, C=O, and O–C=O bands respectively. Further, Fig. 3(c) represents the O 1s spectra having intense signals at 529, 530.2, 531.9, and 533.5 eV correspond to Ni–O, C–O–Ni, CO, and C–OH, confirming the oxygen environment presented on the nanoporous carbon surface.^{17,22} Finally, Fig. 3(d) indicates that Ni 2P peaks appearing at 856.07 and 873.9 eV are Ni 2P_{3/2} and peaks at 861.3 and 878.8 correspond to Ni 2P_{1/2}, confirming the Ni is in +2 and

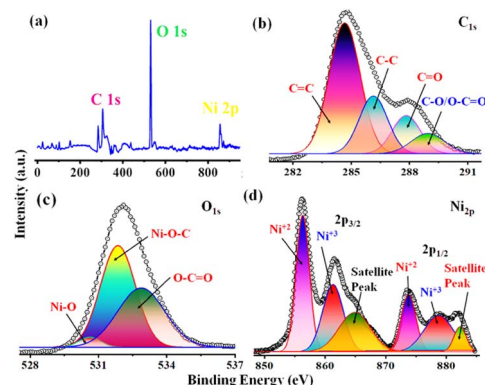


Fig. 3 X-ray photoelectron spectral data of NNC showed (a) full survey XPS of individual elements C, O, and Ni is (b–d), representing in C spectra of C=C, C–C, C–O, O–C=O, C=O, in O spectra C–O–Ni, C=O, O–Ni, C–OH and Ni spectra Ni is in +2 state respectively.

+3 oxidation states, respectively, and the additional informal peaks at 860.7 eV and 880.2 eV correspond to the satellite peaks arising from surface oxidation. The peak at 860.3 is broad due to surface adsorbed oxygen and also locally available oxygen of the nanoporous carbon surface and Ni atoms of nanoparticles and these findings are in good agreement with the literature.⁴ The surface area is one of the most important quantities for characterizing porous materials. The BET analysis is the standard method for defining surface areas from N_2 -adsorption isotherms and was originally derived for multilayer gas adsorption onto flat surfaces. NNC shows type-IV adsorption isotherm with a high BET (Brunauer–Emmett–Teller) specific surface area of 340 $\text{m}^2 \text{g}^{-1}$, and as shown in Fig. 4(a) inset with isotherm of multiple times BET at low pressure provide linear slope with $R^2 = 0.9863$. Fig. 4(b) shows that the pore diameter of NNC is ~ 70 nm. The presence of the H1 hysteresis loop suggests the existence of the mesoporous feature in the NNC due to the porous nature of rGO.

4. Electrochemical studies

Electrochemical features of NNC towards UOR in comparison with NiO-NPs benchmark catalyst were examined from cyclic voltammetry (CV), EIS, Tafel slope, and chronoamperometry ($i-t$) under different operational conditions. Accordingly, Fig. 5(a) displays LSVs at 50 mV s^{-1} for NNC, and the ultrahigh current

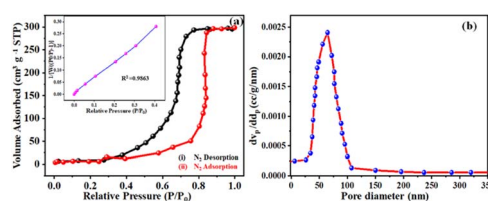


Fig. 4 Represents the (a) adsorption–desorption pattern of NNC, which provides a higher surface area $\sim 340 \text{ m}^2 \text{g}^{-1}$ inset with isotherm of multiple times BET at low pressure provides linear slope with $R^2 = 0.9863$ and (b) shows that pore diameter of NNC is ~ 70 nm.



density is of 10 mA cm^{-2} at an ultralow onset potential of 0.32 V vs. SCE than NiO-NPs (0.45 V vs. SCE). Moreover, for NNC, the oxidation peak for distinctive activation of Ni to Ni-OOH occurred with no back peak for CO_x adsorption in CV (Fig. S2†) and EIS, as shown in Fig. 5(b) for NNC having lower semicircle as compared with NiO-NPs and NC at 0.35 V vs. SCE. Low-frequency region semicircle represents solution resistance called Warburg impedance (W) caused by ion diffusion from the NNC-electrolyte interface. NNC exhibits the smallest R_{ct} value ($\sim 6.98 \Omega$) and provides active electrode surface interactions. As illustrated in Fig. 5(c), the Tafel slopes for NNC, NiO-NPs, and NC are 38 , 52 , and 78 mV dec^{-1} , respectively, indicating that NCC demonstrates superior UOR performance and kinetic activity over other systems.

In addition, NC nanocomposite stabilizes Ni NP's structural, functional as well as electrochemical stability even in harsh conditions, as displayed in Fig. 5(d) and inhibits structural damage from the continuous redox processes of electrolyte ions during chronoamperometric ($i-t$) stability test at 0.34 V of NNC in 0.5 M KOH with 0.3 M urea, as shown in Fig. S3.† NNC remains stable at 0.34 V for 10 mA cm^{-2} for 7 h . Interestingly, the current density increases without drop due to surface activation during electrochemical studies, indicating NNC has a strong ability to inhibit CO poisoning or surface deactivation during UOR. The enhancement in the electrocatalytic performance of NNC towards UOR is mainly attributed to the strong electronic coupling between NC and NiO-NPs and also could be due to the surface modification with NiO-NPs, which provides more exposure of active sites and increases specific surface area ESCA (Fig. S6†). Higher ESCA (42.89 cm^2), enhancement factor (3040), and roughness factor (142.89) for NNC demonstrate high electron transport ability over NC. This suggests that a low diffusion coefficient for NNC during UOR, owing to enhanced charge transfer rate and rapid kinetics. The synergetic interactions between Ni and NiO with NC help to increase the rate of charge transfer during electrocatalysis.

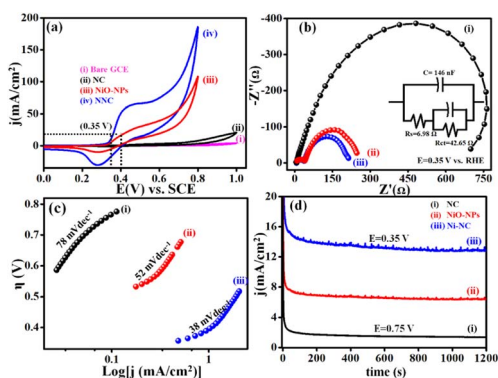


Fig. 5 Represents the (a) LSV (b) Nyquist plot with equivalent circuit fitted from EIS with the help of Z SIM software. (c) Tafel plot and (d) chronoamperometric $i-t$ stability of NNC, NiO-NPs, and NC at 10 mA cm^{-2} , respectively, in a solution of 0.3 M urea in 0.5 M KOH at 0.35 V vs. SCE towards UOR activity.

4.1 Effect of urea concentration

Interestingly, from Fig. 6(a), it is observed that the peak current of urea oxidation linearly increases while increasing the concentration of urea. This could be due to the diffusion of oxidized species on NNC increasing with increasing urea concentration. These observations can be co-related in (Fig. S4†), which shows a linear increment of anodic peak vs. concentration of urea. Hence, calibration data obtained from (Fig. S4b†) proved that the NNC revealed an excellent electrocatalytic activity toward UOR, and it can be used as a novel electrode active material for future applications in the sensitive and selective electrochemical UOR.

4.2 Effect of KOH concentration

When the concentration of KOH increases, the current density increases and this may be caused due to the availability of OH^- ions at the interface, resulting in the formation of an active intermediate Ni-OOH layer. It has also been discovered that hydroxide ions protect the NCC surface from CO_2 poisoning during urea oxidation. Accordingly, UOR must have a skin/layer on NNC. Fig. 6(b) shows a linear calibration curve, and these findings demonstrate that UOR at greater OH^- ion concentrations is more thermodynamically progressive and shows a linear deterioration plot in Fig. S5(a) and (b).† More intriguingly, at higher OH^- concentrations, cathodic current diminishes and this could be due to the surface covering of adsorbed OH^- ions, which limits the interaction of urea molecules with Ni sites.^{23,24}

4.3 Effect of scan rate

Fig. 6(c) demonstrates LSVs of the scan rate-dependent studies of NNC for the electrooxidation of urea in the range of 10 to 200 mV s^{-1} in 0.5 M KOH . Interestingly, it has been demonstrated that with scan rate, the current density increased linearly. Remarkably, the oxidation peaks increased with an

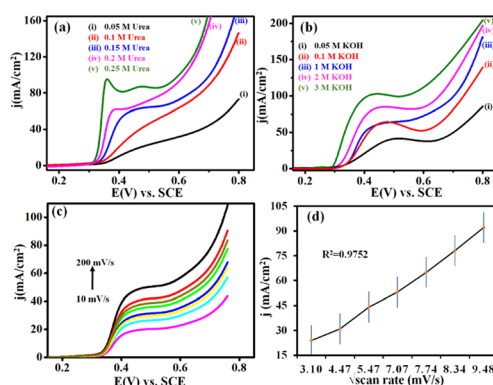


Fig. 6 (a) Superimposed concentration dependent LSV profile from 0.05 M to 0.25 M urea concentration in 0.5 M KOH . (b) Effect of KOH concentration in 0.1 M urea in 0.05 M , 0.1 M , 1 M , 2 M , and 3 M KOH at 50 mV s^{-1} . (c) Scan rate dependent LSV response ranging from 10 mV s^{-1} to 200 mV s^{-1} and (d) current density vs. scan rate linearity profile using NNC modified working electrode, SCE reference, and Pt foil counter electrodes, respectively.



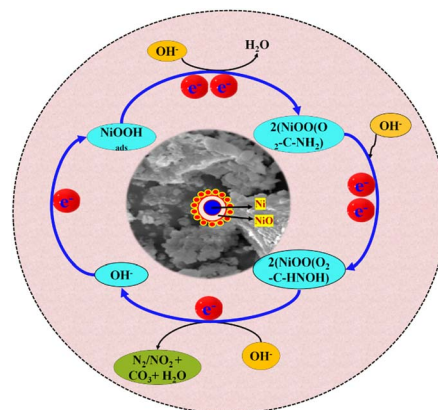
increase in the scan rate and demonstrated a linear relationship with the square root of the scan rate, as shown in Fig. 6(d), which indicates that the electrooxidation of urea is a diffusion-controlled process with first-order kinetics.

4.4 Difference between UOR and OER performance

Fig. 7(a) displays the LSVs scan for NNC in the presence of 0.3 M urea and absence of urea in 0.5 M KOH. Fig. 7(b) displays the Nyquist plot of EIS comparative for UOR and OER, showing a lower semicircle for charge transfer that gives higher electrochemical activity that is diffusion-controlled for UOR as compared with OER. Hence, NNC electrocatalytic system is best for UOR over OER activity because more favourable interactions lower the activation energy. NC provides a backbone to e^- delocalization due to the electronegative O-acceptor, which accepts the e^- from Ni/NiO to donate to urea (*via* Ni/NiOOH) in an electrolyte solution with supportive OH^- ions. Increasing delocalized electrons increases the d-band centre and O-binding strength, which triggers the electrocatalytic UOR kinetics.²⁵

4.5 Possible mechanistic pathway of UOR on NNC

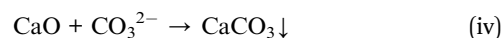
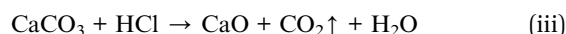
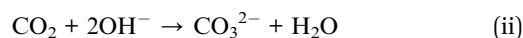
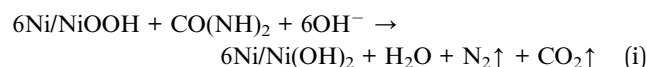
Electrocatalytic UOR at the Ni/NiO anode can be well understood by the electrocatalytic mechanism and is described as follows in Scheme 2. The overall reaction of the electrolysis process in an alkaline solution is the conversion of urea to CO_2 , H_2O , and N_2 catalysed by NNC. First, NNC interacts with OH^- ions and forms Ni/NiOOH as a stable active intermediate,^{4,21} and to find this interaction and formation of Ni/NiOOH, we also calculate the surface active sites, as discussed in Fig. S7(a)–(d).[†] It is confirmed that Ni/NiO with NC provides higher active sites than NiO-NPs and NC,²⁵ which have an important role in UOR at a specific applied potential. Herein, the urea molecules with electron-donating amino groups and electron-withdrawing carbonyl groups can be selectively adsorbed on NNC *via*. Ni/NiOOH–carbonyl group as an intermediate step,^{5,26–29} followed by the corresponding nucleophilic OH^- attack. As a result, cleavage of the C–N bond became more favourable, and UOR could proceed at a lower potential. In the KOH solution without urea, the electrode shows a pair of redox peaks. The redox peaks originate from redox reactions between Ni^{2+} and Ni^{3+} in an alkaline environment. Interestingly, the cathodic peak corresponds to the electrochemical reduction of Ni/NiOOH and not due to the reduction of the urea molecules, which has a faster reaction rate than the urea reduction reactions.²⁴



Scheme 2 Possible mechanistic pathways for UOR on NNC electrocatalyst.

4.6 After electrolysis product analysis test

To confirm CO_2 as the by-product, we performed the precipitation test using CaCO_3 after the *i-t* stability test.³⁰ CaCO_3 reacts with dilute HCl, evolving CO_2 molecules and forms CaO (*i.e.*, lime). CaO easily reacts with electrolyte solution and forms a white ppt of CaCO_3 . This indicates that the generation of CO_2 is one of the reactions and is well explained by the following eqn (i)–(iv)³¹



4.7 After electrolysis UV-spectra

After the UOR stability test on the NNC, we performed the UV-visible spectra analysis and compared it with standard urea solution. As usual, the urea solution shows 3 main peaks observed due to $\pi \rightarrow \pi^*$, $\eta \rightarrow \pi^*$, and intramolecular transitions for urea with KOH at ~ 250 , 350 , and 520 nm.³² After urea electrolysis, only two peaks at ~ 270 and 310 nm, corresponding to CO_3 and NO_x species, confirm the UOR oxidation (Fig. 8) and is in good agreement with the literature.³³

4.8 Electrocatalytic OER performance

LSVs scan for NNC, NiO-NPs, and NC in 1 M KOH at a scan rate of 50 mV s^{-1} is depicted in Fig. 9(a). EIS studies show lower onset potential and higher current density for NNC over NiO-NPs and NC counterparts. Tafel slope is represented in Fig. 9(b), with NNC having a lower Tafel slope (54 mV dec^{-1}) than NiO-NPs (69 mV dec^{-1}), which represents rapid kinetics towards OER. Nyquist plot of EIS showing a lower semicircle for

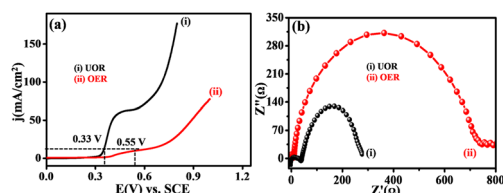


Fig. 7 (a) Superimposed LSV and (b) EIS response for UOR (i) and OER (ii) on NNC electrocatalyst in 0.3 M urea in 0.5 M KOH.

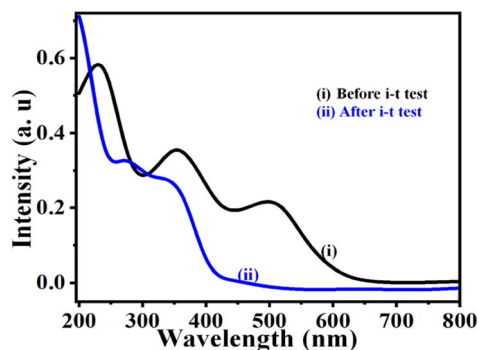


Fig. 8 UV-visible analysis of electrolyte solution of (i) 0.3 M urea in 0.5 M KOH and (ii) after urea electrolysis, i.e., $i-t$ at an applied potential of 0.55 V vs. SCE for 7 h (based on complete oxidation of urea molecules).

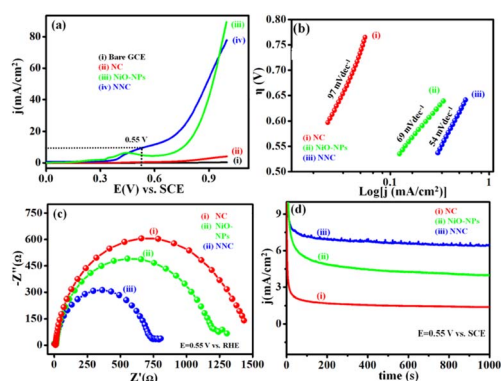


Fig. 9 (a) Superimposed LSVs for NNC, NiO-NPs, and NC recorded in 0.5 M KOH at a scan rate 50 mV s^{-1} , (b) Tafel slope represents rapid kinetics for NNC, (c) Nyquist plot for NNC, NiO-NPs, and NC at 0.55 V vs. SCE in 0.5 M KOH, (d) chronoamperometric $i-t$ plot for NNC, NiO-NPs, and NC at 0.55 V vs. SCE, respectively (potential chosen herein is based on potential required for 10 mA cm^{-2} current density on NCC electrocatalyst).

charge transfer gives higher electrochemical activity with diffusion-controlled electron transfer, as shown in Fig. 9(c). The comparative chronoamperometric $i-t$ stability for NNC, NiO-NPs, and NC at 0.55 V vs. SCE for 1000 s is shown in Fig. 9(d).

Moreover, comparative studies of the previously reported similar electrocatalytic systems for UOR are given in Tables S1 and S2 (ESI†). Hence, it is confirmed that the electrochemical enhancement in the activity is purely due to the decoration of NiO-NPs on the surface of NC. It can be stated that the NNC is an active system for UOR over OER.

5. Conclusion

In summary, herein, we represent a facile chemical synthesis of Ni/NiO-NPs decorated on a nanoporous carbon-based electrocatalytic system for urea and water oxidation reactions. The as-synthesized NNC was characterized *via* SEM, EDAX, HR-TEM, FTIR, UV-vis spectroscopy, BET analysis, TGA, and XPS spectroscopy. The analyses confirm that the NNC is spherical-

shaped ($\sim 25 \text{ nm}$), highly porous, with a specific surface area of $340 \text{ m}^2 \text{ g}^{-1}$ and pore size of $\sim 70 \text{ nm}$, and Ni in +2, +3 oxidation states in the FCC structure. These findings support the electrocatalytic performance of NNC. NC not only provides a conducting backbone but also shows remarkable stability and extremely high and long-term catalytic activity on NNC. After electrolysis, UV-vis spectra and precipitation test permit the determination of the main product of the reaction, namely carbonate with high selectivity after a 7 h electrolysis. Interestingly, NC donates e^- to Ni with O acceptor to form more delocalized electrons, increased d-band centre, and O-binding strength, triggering the electrocatalytic UOR kinetics, with a lower overpotential of 60 mV, and Tafel slope of 45 mV dec^{-1} required for achieving a current density of 10 mA cm^{-2} , which is good enough than that of other Ni-based systems and open a new path for further application of cheap Ni-based materials in energy conversion and storage devices.

Conflicts of interest

No conflicts of interest.

Acknowledgements

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