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Supercapacitor performance of porous nickel cobaltite nanosheets

Xin Chen¹, Rui Xie², Hui Li³, F. Jaber⁴, F. Musharavati⁵, E. Zalnezhad⁶✉, S. Bae⁷✉, K. S. Hui⁸ & K. N. Hui⁹

In this work, nickel cobaltite (NiCo₂O₄) nanosheets with a porous structure were fabricated on nickel foam as a working electrode for supercapacitor applications. The nanosheets were fabricated by electrochemical deposition of nickel–cobalt hydroxide on the nickel foam substrate at ambient temperature in a three-electrode cell followed by annealing at 300 °C to transform the coating into a porous NiCo₂O₄ nanosheet. Field emission scanning electron microscopy and transmission electron microscopy revealed a three-dimensional mesoporous structure, which facilitates ion transport and electronic conduction for fast redox reactions. For one cycle, the NiCo₂O₄ electrodeposited nickel foam has a high specific capacitance (1734.9 F g⁻¹) at a current density (CD) of 2 A g⁻¹. The electrode capacitance decreased by only approximately 12.7% after 3500 cycles at a CD of 30 A g⁻¹. Moreover, a solid-state asymmetric supercapacitor (ASC) was built utilising the NiCo₂O₄ nanosheets, carbon nanotubes, and a polyvinyl alcohol-potassium hydroxide gel as the anode, cathode, and solid-state electrolyte, respectively. The ASC displayed great electrochemical properties with a 42.25 W h kg⁻¹ energy density at a power density of 298.79 W kg⁻¹.

Due to rapidly increasing logical pollution, fossil fuel depletion, and the fast growth of the worldwide economy, it is vital to develop clean, sustainable, and efficient energy resources, along with new technologies for energy conversion and storage^{1–5}. In recent years, ultracapacitors, or electrochemical supercapacitors (ESs), have generated substantial interest, owing mostly to their long lifecycles, high power density, and ability to bridge the power/energy gap in conventional batteries/fuel cells and dielectric capacitors because of their high energy storage capacity and high power output^{6,7}.

Lately, many researches have concentrated on binary metal oxides. Their outstanding specific capacity and great electrical conductivity are better than single component oxides due to their attainable oxidation states for multiple redox reaction⁸. Furthermore, binary metal oxides have many other benefits, such as relative abundance, low cost, and environmental friendliness⁹. In recent years, researches have shown that binary metal oxides, such as nickel cobaltite (NiCo₂O₄), ZnCo₂O₄, and Zn₂SnO₄ are favourable materials that show improved electrochemical performance. Moreover, they are scalable replacements owing to their ample surface active sites, high electrical conductivity, strong permeability, and attainable oxidation states. Thus, many works have been performed to synthesize dissimilar bimetallic oxide nanomaterials for supercapacitor applications with great rate capabilities. The binary metal oxide NiCo₂O₄ was recently studied for use as an electrode owing to its electrochemical activity, stability, and higher electronic conductivity compared to single metal oxides^{10–13}. NiCo₂O₄ has a structure similar to Co₃O₄ (spinel structure), and both the Co and Ni ion have a mixed oxidation state^{14,15}. The electrical conductivity of NiCo₂O₄ is more than twice that of the Co or Ni oxide alone because the replacement of Co with Ni brings additional electrons into the 3d orbital, which subtly changes the density of electrons in the crystal structure. The NiCo₂O₄ working potential window is often very slim, near 0–0.5 V, in comparison with alkaline

¹Department of Mechanical Convergence Engineering, Hanyang University, 222 Wangsimni-ro, Seongdong-gu, Seoul 04763, South Korea. ²Department of Electronic Information, Shandong Xiandai University, Jinan 250000, People's Republic of China. ³Department of Chemical Engineering, Hanyang University, 222 Wangsimni-ro, Seongdong-gu, Seoul 04763, South Korea. ⁴Department of Biomedical Engineering, Ajman University, 2758, Ajman, UAE. ⁵Department of Mechanical and Industrial Engineering, College of Engineering, Qatar University, 2713, Doha, Qatar. ⁶Department of Chemical and Biomedical Engineering, University of Texas At San Antonio, San Antonio, TX, USA. ⁷Department of Architectural Engineering, Hanyang University, Seoul 04763, Korea. ⁸School of Engineering, University of East Anglia, Norwich NR4 7TJ, UK. ⁹Joint Key Laboratory of the Ministry of Education, Institute of Applied Physics and Materials Engineering, University of Macau, Avenida da Universidade, Taipa, Macau SAR 999078, China. ✉email: erfanzalnezhad@utsa.edu; sbae@hanyang.ac.kr

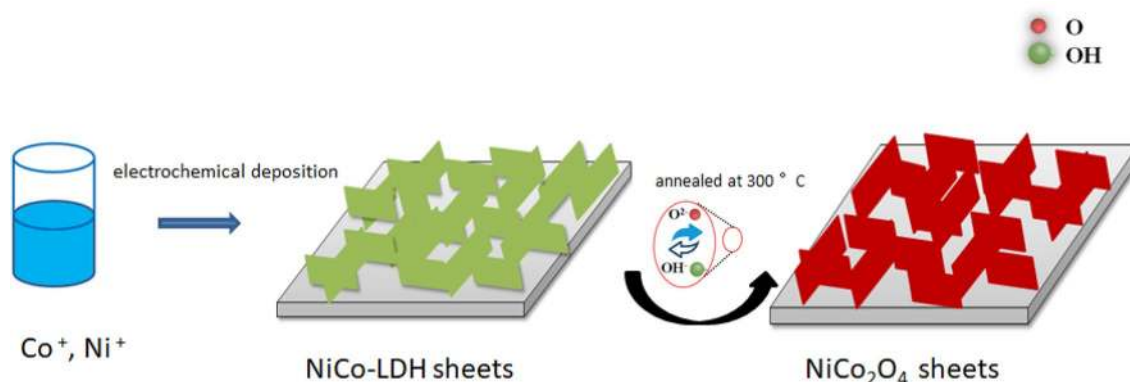


Figure 1. Schematic illustration of porous NiCo₂O₄ sheets synthesis.

solutions (Ag/AgCl)^{16–19}. It has been proved that the introduction of graphene can effectively enhance the total capacitance and stability, primarily because graphene can withstand the basic structures of polyaniline and evade mechanical deformation in the redox process²⁰. Therefore, ternary composites with carbon nanomaterials, using transition metal oxides and polymers, have been studied and showed an enhancement in the electrochemical performance that open a new fabrication pathway for next generation high-performance electrochemical electrodes²¹. Ternary CuCo₂S₄ is well known as an electrode material for electrochemical capacitors because of its synergistic effects, high conductivity, and low cost. The supercapacitive performance of CuCo₂S₄ electrodes for electrochemical capacitors was investigated by Xu et al. In their study, thin layers of CuCo₂S₄ were deposited on conductive substrates, and the results showed better properties than single crystal CuCo₂S₄²².

Electrodeposition has been utilised for different applications, such as microelectronics and energy conversion. Because of the need for enhanced performance and device miniaturization, nuanced control of the growth process is needed; electrochemical deposition is a technique that can fulfil those requirements²³.

Chen et al. successfully deposited Ni–Co–S nanosheet arrays on carbon fibres through one-step electro-deposition of the ternary sulfides to provide an effective and facile approach for large scale applications, which was a significant advantage compared to other multistep synthesis techniques. They found that the Ni–Co–S–4 interconnected nanosheet exhibits the best electrochemical performance as a supercapacitor electrode²⁴.

One effective method of enlarging the potential window of the NiCo₂O₄ electrode to attain a high energy density is the utilisation of asymmetric devices^{16,25–27}. For example, Yedluri et al. fabricated chain-like NiCo₂O₄/NiCo₂O₄ nanofile arrays using a facile hydrothermal and thermal decomposition approach and reported a specific capacitance of 2312 F g^{−1} at a current density (CD) of 2 mA cm^{−2}²⁸. They also studied the electrochemical performance of NiCo₂O₄@NiCo₂O₄ composite nanoplates and NiCo₂O₄ nanoplates decorated with NiMoO₄ honeycombs for high performance supercapacitor applications^{29,30}. Herein, we developed a solid asymmetric supercapacitor with a pre-synthesised NiCo₂O₄ mesoporous electrode without binders, where a nickel foam (with high conductivity) was selected as a current collector, and the NiCo₂O₄ provided a large surface area along with a unique mesoporous nanostructure. Our sample delivered a specific capacitance of 1734.9 F g^{−1} at a 2 A g^{−1} CD with a retention rate of 87.3% at a 30 A g^{−1} CD for 3500 cycles. Furthermore, the application of a NiCo₂O₄-based binder-free electrode to a solid asymmetric supercapacitor resulted in a 42.25 W h kg^{−1} energy density at a power density of 298.79 W kg^{−1} (Fig. 1).

Results and discussion

Figure 2 depicts SEM images of NiCo-LDH and NiCo₂O₄. The images show that a smooth, uniform array of NiCo₂O₄ nanosheets is grown on the nickel foam surface, and the nanosheets are interlaced to form a mesoporous structure. Figure 2 (a) shows that the NiCo-LDH possesses an interconnected nanosheet microstructure. Figure 2 (b, c) shows SEM images of NiCo₂O₄. These nanosheets, which are several hundred nanometers in size, have a porous structure that is intercrossed, which contains electroactive surface sites and plentiful vacancies^{31–33}.

The products were further examined by X-ray diffraction (XRD) analysis. Figure 3 depicts the XRD pattern of the NiCo₂O₄ nanosheets deposited on the Ni foam. The peaks (from the (111), (200), and (220) planes, respectively) at 44.7°, 52.1°, and 76.5°, denoted by asterisks, are created from the nickel foam. The peaks at 18.9°, 36.6°, 59.1°, and 64.9° can be clearly observed and are well indexed to the (111), (311), (511), and (440) planes, respectively, belonging to NiCo₂O₄ (JCPDS Card No. 20-0781)^{34,35}.

Transmission electron microscopy (TEM) measurements were performed to further investigate the structure of the synthesised NiCo₂O₄ nanosheets, as shown in Fig. 4. Figure 4(a, b) shows a NiCo₂O₄ nanosheet with a folding, silk-like morphology and transparent features, indicating its interconnected nature. Due to the significant difference between the lateral size and thickness, bending and crumpling are clearly observed. The spacing between adjacent fringes is ~0.29 nm, which is close to the theoretical interplane spacing of spinel NiCo₂O₄ (311) planes. Thus, the interconnected nanosheets are composed of 1–3 layers of NiCo₂O₄ atomic sheets. The selected area electron diffraction (SAED) pattern (Fig. 4 (c)) presents distinct diffraction rings, indicating polycrystalline characteristics. Furthermore, several interparticle mesopores, with sizes ranging from 1 to 3 nm, in these interconnected nanosheets can be evidently seen (Fig. 4 (a,b)). It is believed that the mesoporous structures in nanosheets are imperative in facilitating the electrolytes' mass transport within the electrodes for double-layer

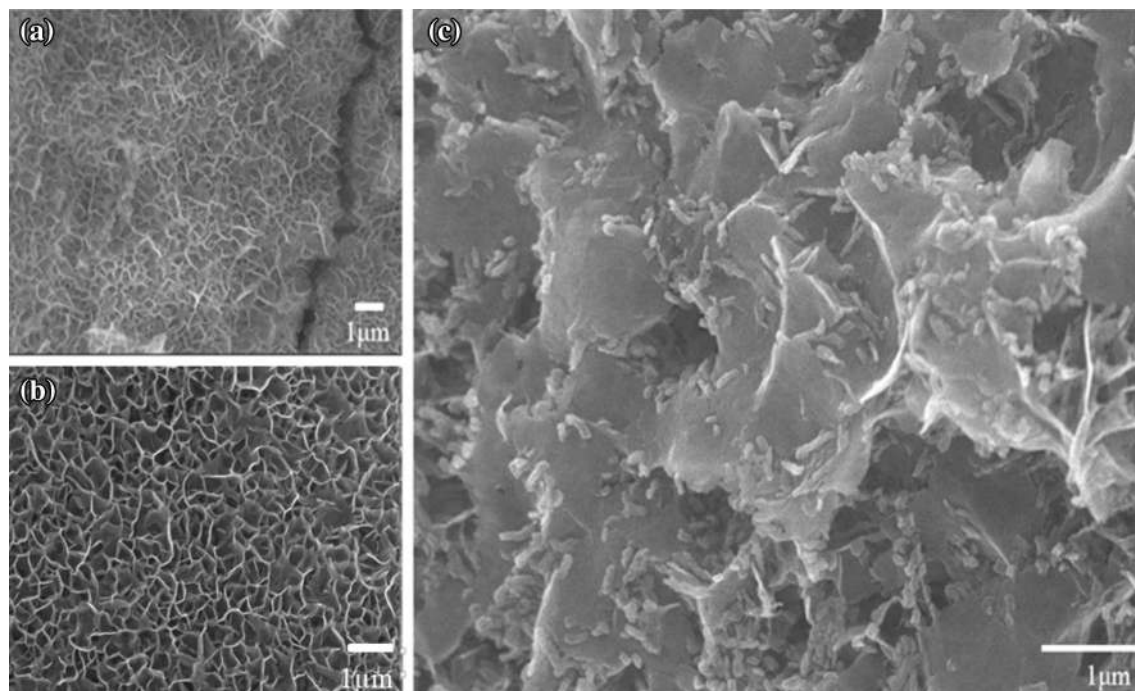


Figure 2. FESEM images of (a) Ni foam covered by the bimetallic (Ni, Co) hydroxide precursor, and (b, c) the derived NiCo₂O₄ interconnected sheets.

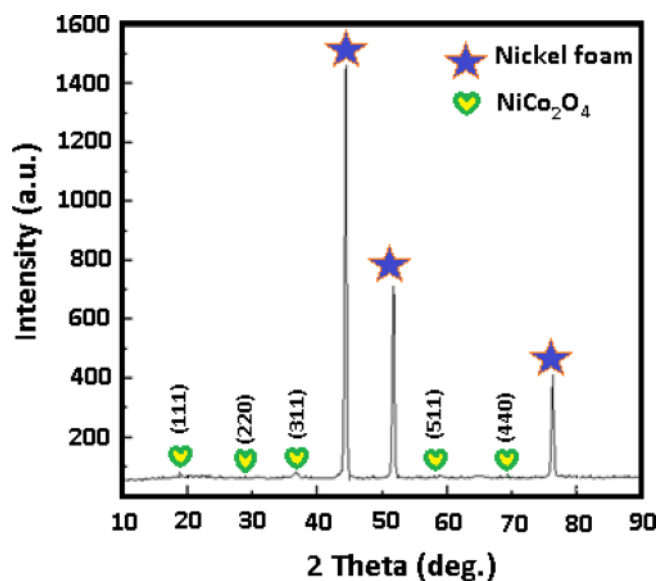


Figure 3. XRD pattern of NiCo₂O₄.

charging/discharging and quick redox reactions⁸. The porous structure also significantly increases the contact area of electrolyte/electrode, and consequently enhances the electrochemical performance^{9–15}. To show the advantages of this architecture, the interconnected mesoporous NiCo₂O₄ nanosheets with hybrid structure fabricated on Ni foam was directly applied as an electrode for a supercapacitor. Figure 4(d) show the energy dispersive X-ray spectroscopy (EDS) analysis of the NiCo₂O₄ sample; Co, Ni, and O are detected. This result confirms the chemical composition of the NiCo₂O₄ structure.

X-ray photoelectron spectroscopy (XPS) tests, along with the corresponding fitting results are shown in Fig. 5. Gaussian fitting method was used to best fit the Ni 2p and Co 2p (with two spin-orbit doublets for each), characteristic of Ni²⁺ and Ni³⁺ and Co²⁺ and Co³⁺, respectively and two shake-up satellite (indicated as “Sat.”) for both Ni 2p and Co 2. These data show that the surface of the as-prepared NiCo₂O₄ contains Co²⁺, Co³⁺, Ni²⁺, and Ni³⁺, where the atomic ratio of Co to Ni elements is *ca.* 2.2:1, which is close to that in the precursor electrolyte.

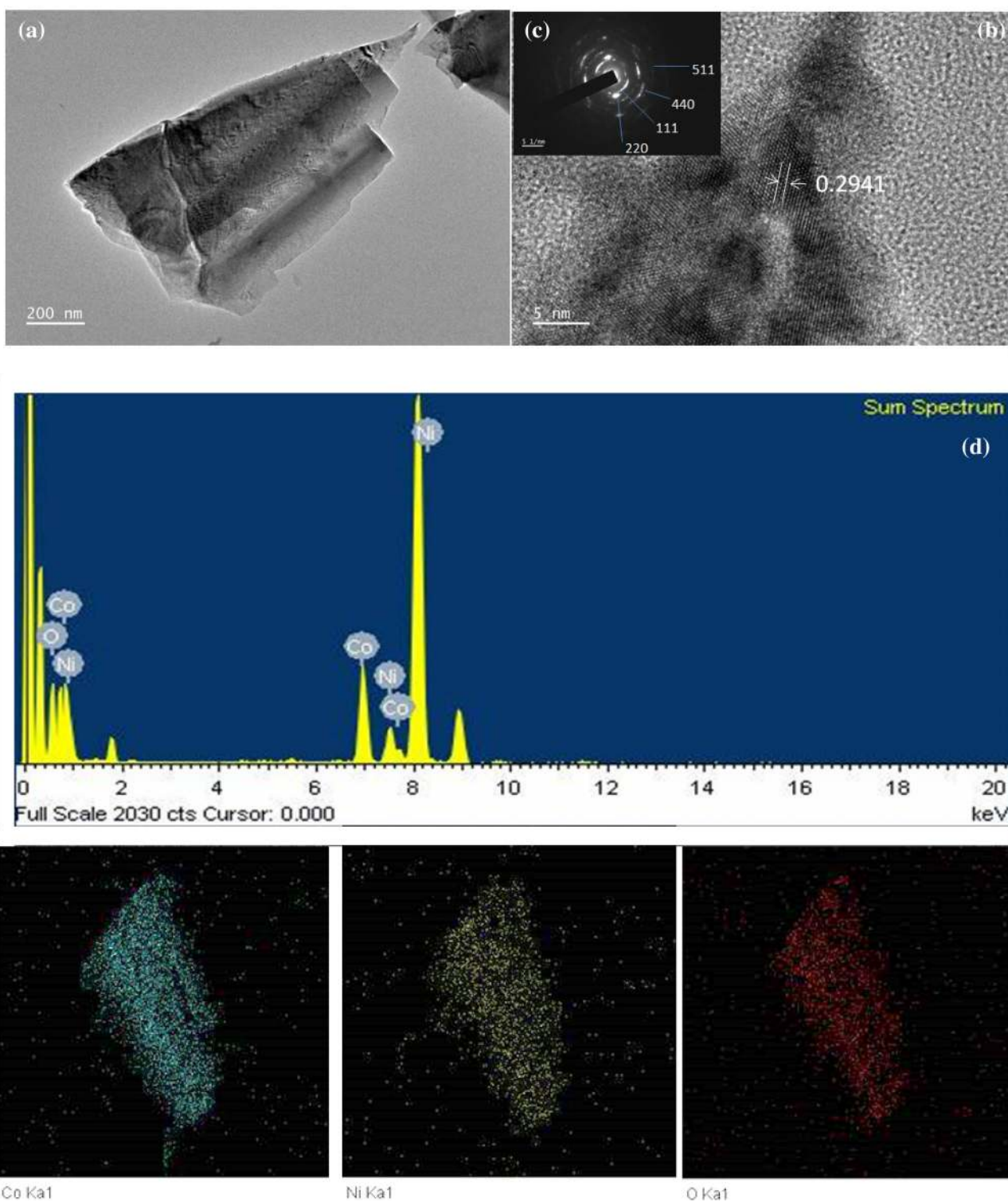
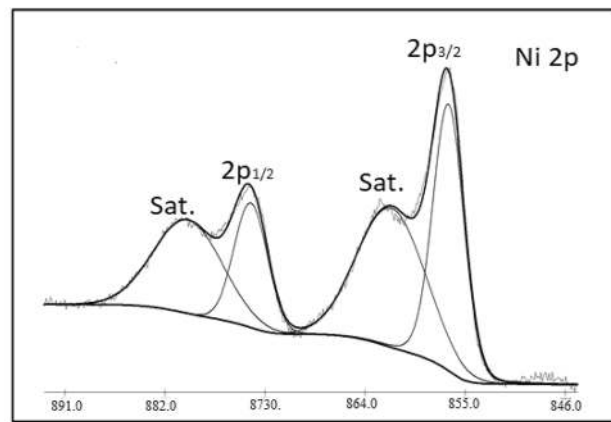
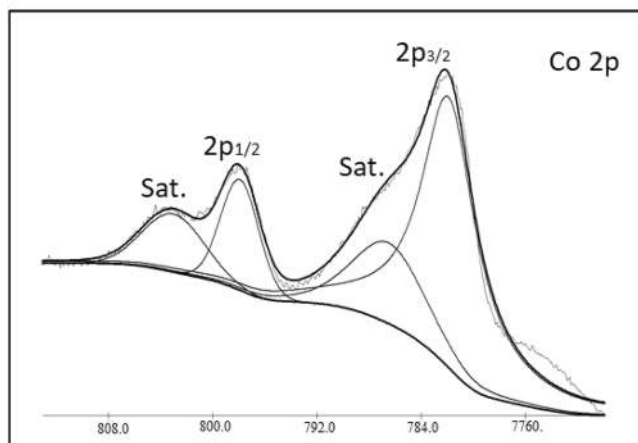


Figure 4. (a) TEM image and (b) HRTEM image of an individual NiCo_2O_4 nanosheet, along with (c) the SAED pattern (inset), and (d) the corresponding EDS mapping results.

Nitrogen adsorption/desorption measurements were utilised to examine the porosity and BET surface area and of samples. All the N_2 adsorption/desorption isotherms in Fig. 6 exhibit a typical IV isotherm with a hysteresis loop in the P/P_0 range of 0.25–1.0, suggesting the materials have a mesoporous structure. These curves were based on the IUPAC classification of type IV isotherms with loop hysteresis. The specific surface area calculated for the nanocomposite created in the current density of -6.0 mA/cm^2 were $40.2 \text{ m}^2/\text{g}$. The resulting structure had several advantages in electrochemical supercapacitors. Interconnected NiCo_2O_4 sheets grown directly on



Binding Energy [eV]



Binding Energy [eV]

Figure 5. High-resolution XPS spectra of Ni 2p and Co 2p.

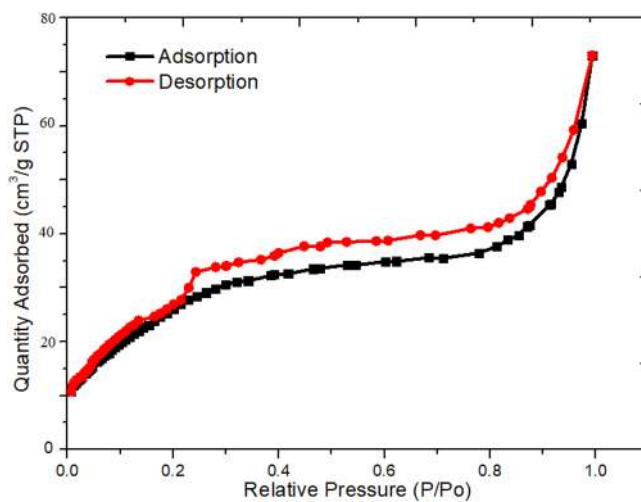


Figure 6. Nitrogen adsorption and desorption isotherms for the NiCo₂O₄ sheets.

the nickel foam using electrochemical deposition method provided an integrated and orderly electrode that facilitated the transport of ion and electrons, thereby reducing the electrode resistance.

Figure 7(a) shows the cyclic voltammetry (CV) curves of NiCo₂O₄ with different numbers of electrodeposition cycles at 5 mV s⁻¹. The NiCo₂O₄_1 electrode demonstrates higher peak currents and larger integrated areas compared with the NiCo₂O₄_2, NiCo₂O₄_4, and NiCo₂O₄_6 electrodes. Further, the properties of NiCo₂O₄_1 shown in Fig. 7(b) indicate better electrochemical capacity than those of Co₃O₄ and NiO. Figure 7(c) shows the CV curves of the NiCo₂O₄ electrode at scan rates from 5 to 100 mV s⁻¹ with a 0–0.6 V potential window (vs. HgCl reference electrode). Even at a relatively high scan rate, a couple of redox peaks are evident, a battery characteristic signifying quick kinetics¹⁶. The current response enhanced with increasing sweep rate without noticeable alteration in the trend of the CV curves. Furthermore, these peaks are attributed primarily to the faradaic redox reactions involving M–O–OH/M–O (where M signifies Co or Ni)³⁶. Figure 7(d, e) presents the galvanostatic charge–discharge (GCD) measurements of Co₃O₄, NiO, and NiCo₂O₄_1 with different numbers of electrodeposition cycles at 2 A g⁻¹ from 0 to 0.55 V. Among the samples, NiCo₂O₄_1 had the best charge–discharge properties. The GCD curves at various CDs are demonstrated in Fig. 7(f). These CD curves are nonlinear, indicating typical battery-type capacitive behaviour^{37,38}. These results confirm that the NiCo₂O₄_1 electrode had a much higher specific capacitance than the NiCo₂O₄ electrodes with more deposition cycles. The remarkable electrochemical performance of the NiCo₂O₄_1 electrode could be attributed to the excellent adhesion to the nickel foam substrate, with a large surface area and electrical connection of the active material to the current collector to ensure effective accessibility of the electrolyte ions and electrons.

Equation 1 defines a numerical calculation of specific capacitance during current density characterization³⁹:

$$C = \frac{I\Delta t}{m\Delta V} \quad (1)$$

where m , I , ΔV , Δt , and C are the active materials mass, discharge current, drop in potential, total discharge time, and specific capacitance, respectively.

Figure 8(a) shows the specific capacitance of NiCo₂O₄ with different numbers of electrodeposition cycles at different current densities. NiCo₂O₄_1 shows a superior specific capacitance. Figure 8(b) shows the specific capacitance of NiCo₂O₄_1, NiO, and Co₃O₄, and the NiCo₂O₄_1 nanosheet electrode shows excellent capacitance values of 1734.9, 1590.5, 1514.7, 1391.3, 1302.4, and 1201.8 F g⁻¹ at CDs of 2, 6, 10, 20, 30, and 50 A g⁻¹, respectively. This shows that when the charge–discharge rate increases from 2 to 50 A g⁻¹, around 87.3% of the capacitance is retained.

Electrochemical impedance spectroscopy (EIS) tests were carried out to examine the ion transport properties of the synthesised materials in the 0.01 Hz–100 kHz frequency range. Figure 9 shows the EIS results for the NiCo₂O₄_n ($n = 1, 2, 4, \text{ and } 6$), NiO, and Co₃O₄ materials. The Nyquist plots of the electrodes contain straight and semicircular curves in the low and high frequency regions, respectively^{40–43}. The intercept of Z_0 (the real axis) with the semicircle, in the high-frequency region, is identical to R_s (the internal resistance), which comprises of the ohmic resistance of the active materials, the resistance of the electrolyte, and the contact resistance at the interface of the active material/nickel foam. The internal resistance values of the NiCo₂O₄_1, NiO, and Co₃O₄ electrodes were 0.35, 0.22, and 1.58 Ω , respectively. The semicircle represents C_{dl} (the double-layer capacitance), which is related to the surface properties of the electrode, and the semicircle diameter indicates R_{ct} (the charge transfer resistance), which is related to the corresponding faradaic reactions at the interface of the electrode–electrolyte. On the contrary, the slope of the curves in the low frequency region signifies the Warburg resistance, which is related to the diffusion of the electrolyte in the electrodes. It is known that the electrochemical performance of supercapacitors can be effectively enhanced by reducing this resistance. Based on the results of GCD, an extraordinary specific capacitance of 1734.5 F g⁻¹ was obtained at a current density of 2 A g⁻¹. The CV curves of NiCo₂O₄ with different numbers of electrodeposition cycles at 5 mV s⁻¹ are given in Fig. 9(a). The NiCo₂O₄_1 electrode demonstrates higher peak currents and larger integrated areas compared with the NiCo₂O₄_2, NiCo₂O₄_4, and NiCo₂O₄_6 electrodes. The internal resistance of the NiCo₂O₄_1 electrode was only 0.35 Ω . These results indicate that NiCo₂O₄_1 electrode has good electrochemical performance.

The cycle life performance of NiCo₂O₄ at a CD of 30 A g⁻¹ for 3500 cycles is presented in Fig. 10. The NiCo₂O₄ deposited on the nickel foam electrode shows steady cycling stability. From the calculation for the discharge curves, an approximate decrease of only 12.7% in the specific capacitance value after 3500 cycles is obtained. At low current densities, some side reactions occurred during the electrochemical redox reaction, leading to incomplete discharge. As the current density increased, the charge and discharge time decreased. The electrochemical process was mainly affected by the electric double layer, so the Coulomb efficiency increased as well⁹. Given the superior electrochemical behaviour of the NiCo₂O₄ electrode, it is the best choice for use in electrochemical supercapacitors characterised by both excellent rate capability and long cycle life.

Furthermore, to study the practical performance of the synthesised electrode in this work, asymmetric supercapacitors (ASCs) were assembled, in which carbon nanotubes (CNTs) and NiCo₂O₄ were the cathode and anode, respectively, with a polyvinyl alcohol-KOH gel polymer electrolyte. Figure 11(a) displays the CV curves of the anode and cathode at 5–100 mV s⁻¹ scan rates. Figure 12(b) indicates that the NiCo₂O₄ and CNT electrodes operate in voltage ranges of 0.0–0.6 V and –1.0 to 0.0 V, respectively. Consequently, the NiCo₂O₄/CNT ASC can operate in a 1.6 V voltage range¹². As the scan rate increases from 5 to 100 mV s⁻¹, the shape of the CV curve does not change, indicating that the device has good and rapid charge–discharge properties. The charge–discharge behaviour of the device is depicted in Fig. 11(b). These charge–discharge curves are nonlinear, indicating battery-type capacitive behaviour. Equations 2 and 3 were used to further investigate the excellent rate capability and high capacitance to evaluate the performance indicators of specific power density (P) and specific energy density (E) from the discharge curves^{13,44–47}.

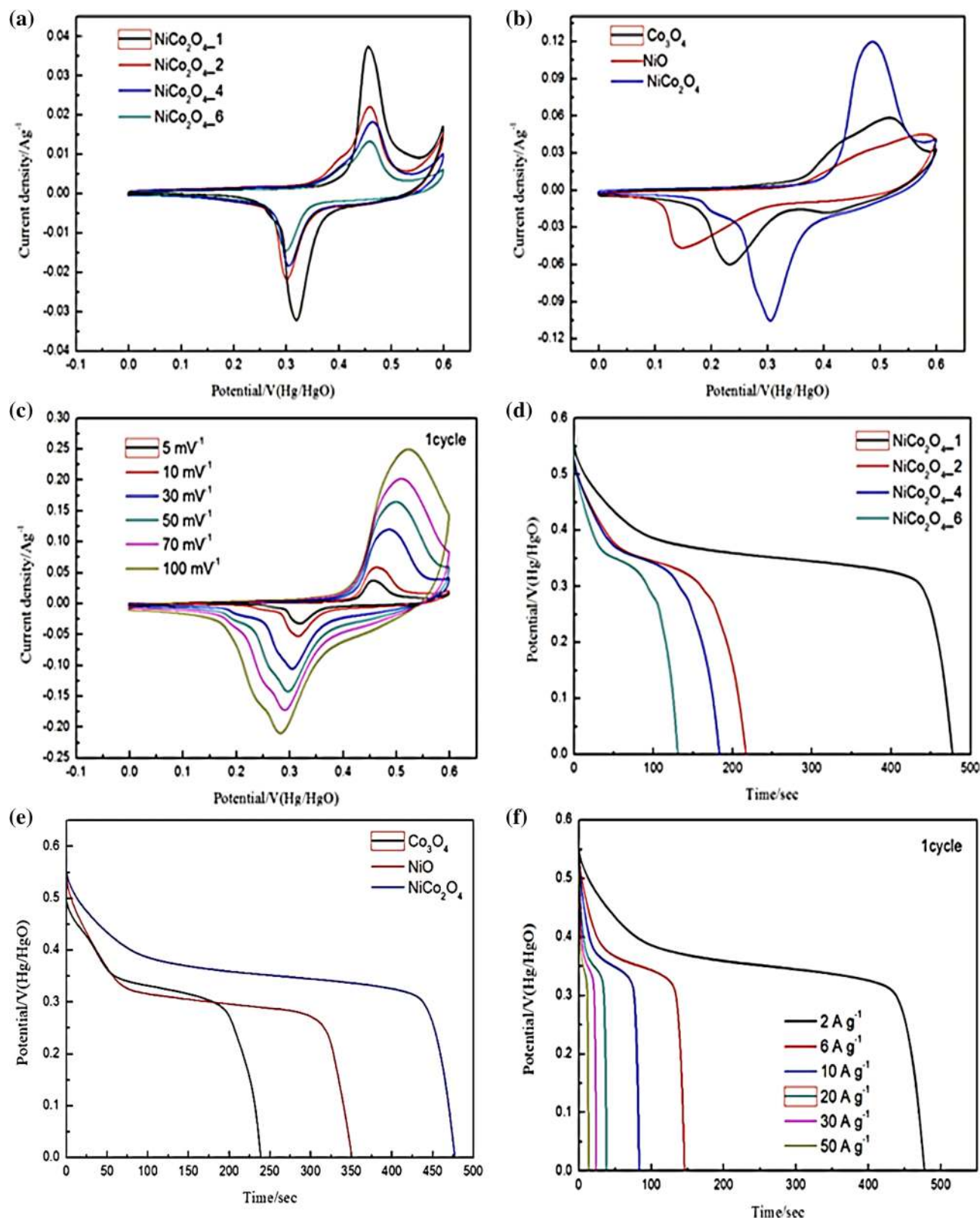


Figure 7. CV curves of (a) NiCo₂O₄ with different numbers of electrodeposition cycles at 5 mV s⁻¹; (b) NiCo₂O₄, Co₃O₄, and NiO as the electrode material at 30 mV s⁻¹; and (c) NiCo₂O₄ at different scan rates. GCD curves of (d) NiCo₂O₄ with different numbers of electrodeposition cycles at 2 A g⁻¹; (e) NiCo₂O₄, Co₃O₄, and NiO as the electrode material at 2 A g⁻¹; and (f) NiCo₂O₄ at different CDs.

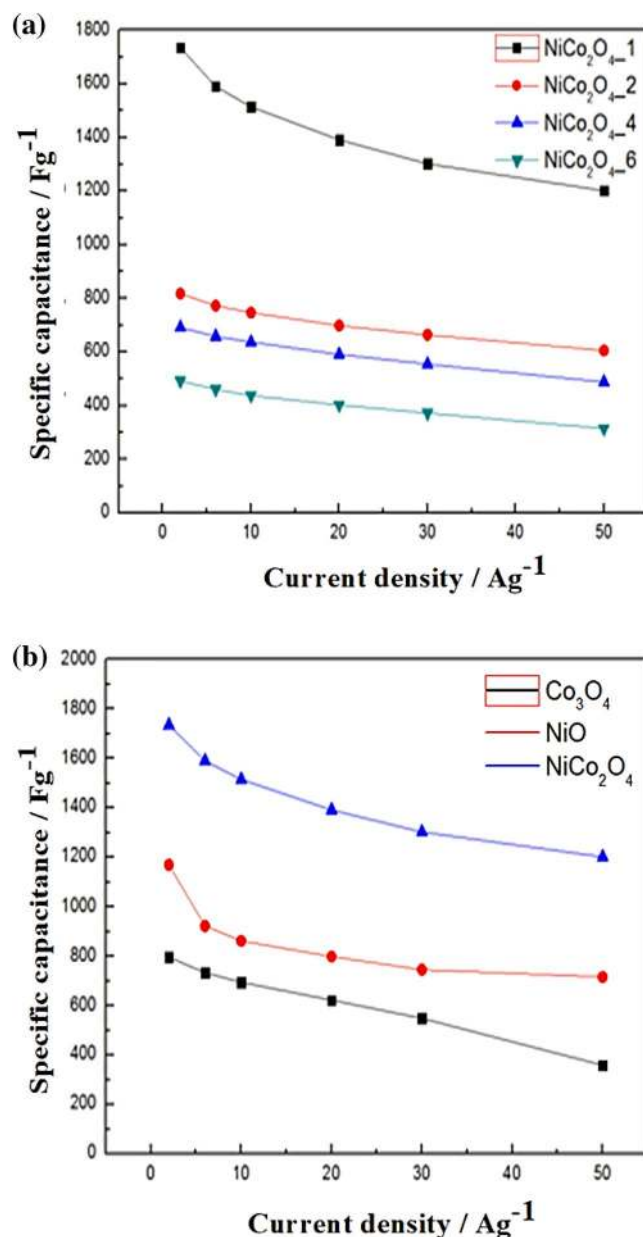


Figure 8. (a) Specific capacitance of NiCo₂O₄ with different numbers of electrodeposition cycles at different CDs. (b) Specific capacitance of NiCo₂O₄, NiO, and Co₃O₄ at different CDs.

$$E = 1/2(C \times \Delta V^2) \quad (2)$$

$$P = 3600 (E/\Delta t) \quad (3)$$

where ΔV , C , and Δt are the potential drop during discharge, specific capacitance derived from the charge–discharge calculations, and full discharge duration, respectively.

Figure 12 depicts the power and energy density plot (Ragone plot) of the devices at different charge–discharge rates. The calculated energy densities of the NiCo₂O₄/CNT ASC were 42.25, 41.43, 29.25, 23.73, and 19.16 W h kg⁻¹ at power densities of 298.79, 596.66, 1196.52, 2966.29, and 5796.04 W kg⁻¹, respectively. These findings indicate that the ASC device attained a greater energy density than reported devices, such as colloidal quantum dots/NiCo₂O₄/activated carbon (AC) (27.8 W h kg⁻¹ at 128 W kg⁻¹)^{48,49}, NiCo₂O₄-reduced graphene oxide/AC (23.32 W h kg⁻¹ at 324.9 kW kg⁻¹)⁵⁰, Ni_xCo_{1-x} LDH–zinc tin oxide/AC (23.7 W h kg⁻¹ at 284.2 W kg⁻¹)⁵¹, and CoO@polypyrrole/AC (43.5 W h kg⁻¹ at 87.5 W kg⁻¹)⁵².

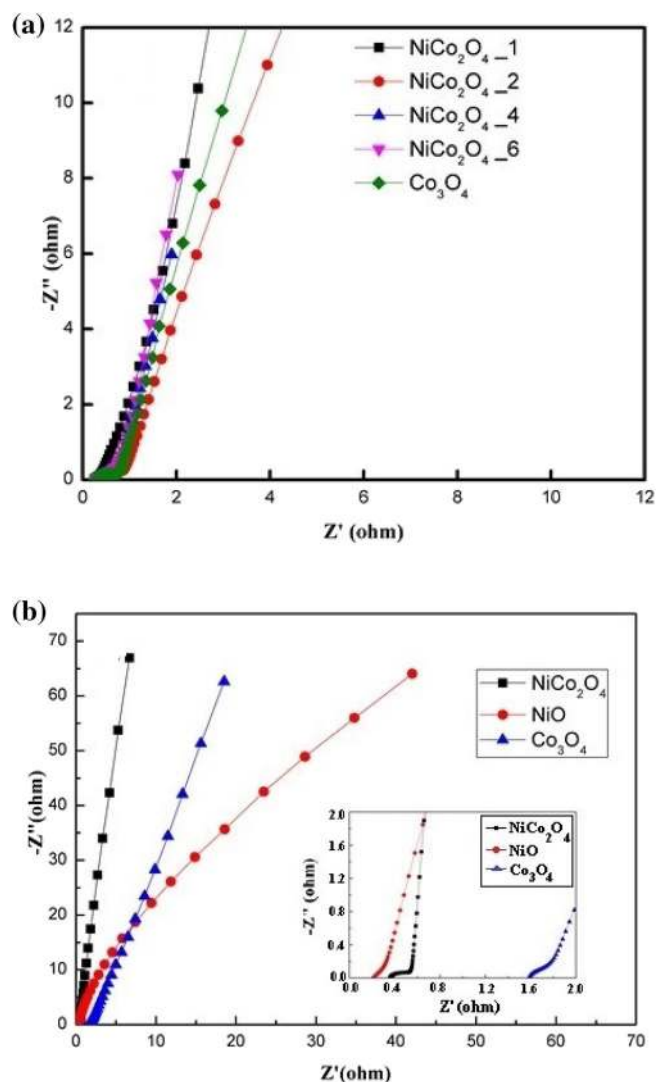


Figure 9. Nyquist impedance plots of (a) NiCo_2O_4 with different numbers of deposition cycles and (b) NiCo_2O_4 , NiO , and Co_3O_4 .

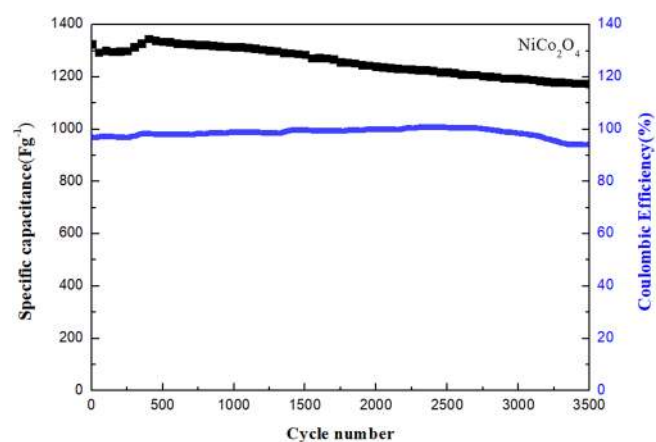


Figure 10. Cycling stability and Coulombic efficiency of the NiCo_2O_4 electrode at a CD of 30 A g^{-1} for 3500 cycles.

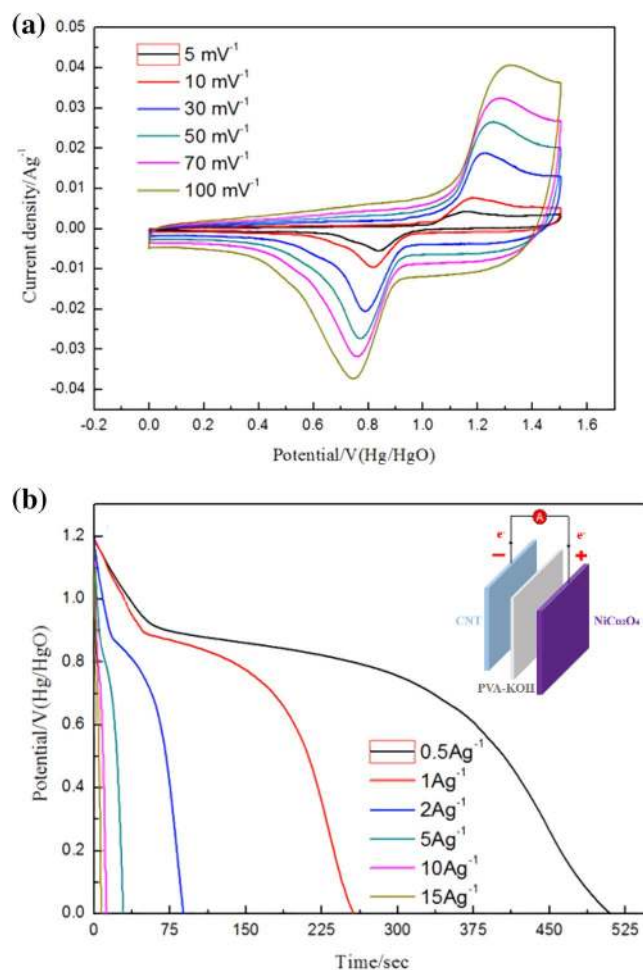


Figure 11. (a) CV curves of the ASC device. (b) Charge–discharge curves of the ASC device. The embedded picture is a schematic of the ASC device.

Conclusions

In summary, NiCo₂O₄ was synthesised on nickel foam via electrodeposition. NiCo₂O₄ was found to have superior specific capacitances of 1734.9 and 1201.8 F g⁻¹ at CDs of 2 and 50 A g⁻¹, respectively, with great cycling stability (only 12.7% loss after 3500 cycles). Furthermore, a high performance solid-state ASC was built utilising NiCo₂O₄ and CNTs as the anode and cathode, respectively, and the solid-state polyvinyl alcohol–KOH gel as an electrolyte. A specific capacitance of 212.47 F g⁻¹ was attained at a 0.5 A g⁻¹ CD. Moreover, the ASC exhibited a high energy density (42.25 W h kg⁻¹) at a 298.79 W kg⁻¹ power density and a high power density (5,796.04 W kg⁻¹) at a 19.16 W h kg⁻¹ energy density.

Materials and methods

Preparation of nickel cobaltite. The electrochemical deposition was carried out at ambient temperature to synthesize NiCo₂O₄ nanosheets onto nickel foam in a 3-electrode cell in which nickel foam, saturated calomel, and Pt foil utilized as working, reference, and counter electrodes, respectively. Nickel–Cobalt layered double hydroxide precursor was deposited over nickel foam in an aqueous mixed electrolyte of 2 mM cobalt nitrate and 1 mM nickel nitrate using a ZIVE SP2 electrochemical workstation.

Versatile numbers of cycles like 1, 2, 4, and 6 cycles were selected for electrodeposition of working electrodes with the potential of –1.2–0.5 V (vs. SCE). The nickel foams were ultrasonically cleaned and rinsed three times with distilled water and ethanol after electrodeposition and then dried at ambient temperature. Then, the electrodeposited working electrodes was placed in a muffle furnace and annealed for two hours at 300 °C, to transform the coating into interconnected mesoporous NiCo₂O₄ nanosheets; it was carefully weighed after annealing (Fig. 1). The obtained NiCo₂O₄ specimens are called as NiCo₂O₄-*n* (where *n* is the number of cycles, *n* = 1, 2, 4, and 6), and henceforth “NiCo₂O₄” denotes the *n* = 1 sample when NiCo₂O₄ is compared with other materials.

Preparation of flexible solid electrolyte. The solution-casting technique was used to prepare the potassium hydroxide and PVA polymer electrolyte. 2 g of polyvinyl alcohol was added into 30 ml of double-distilled

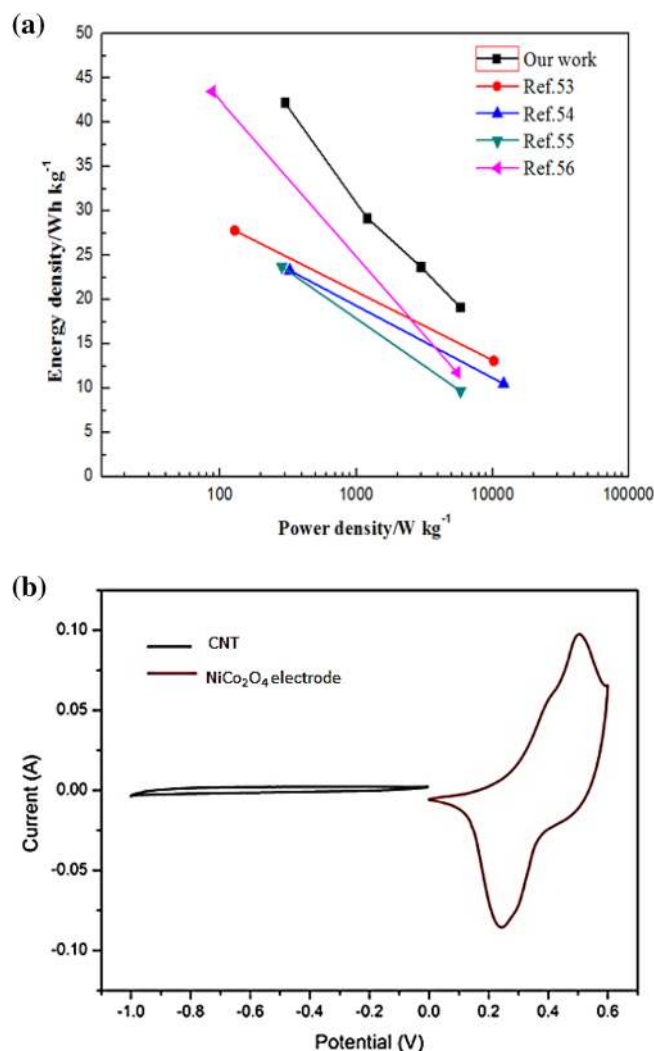


Figure 12. (a) Power and energy density plot (Ragone plot) of the current ASC device at different charge–discharge rates compared with the literature. (b) CV curves of the CNT and NiCo₂O₄ electrodes in a three-electrode system at a scan rate of 5 mV s⁻¹.

water and stirred for three hours at 80 °C. Then 6 M potassium hydroxide was mixed the prepared solution with agitation at room temperature for 3 h. After dissolution was complete, the final mixture was constantly agitated until a homogeneous viscous product was obtained. Finally, to attain a jelly electrolyte, the mixture was transferred to an oven (vacuum) and kept at 70 °C overnight.

Characterization of NiCo₂O₄. The morphology and microstructure of specimens were examined by FESEM (FESEM, LEO-1550) equipped with EDS at a 5 kV applied voltage. XRD tests were carried out (Bruker D8 Advance X-ray diffractometer) with Cu K- α radiation ($\lambda = 0.154056$ nm) at 30 mA and 40 kV. The speed of scanning was 5° min⁻¹ with 0.02° steps. Elemental mapping (FEI Talos microscope operating at a 200 kV accelerating voltage), high-angle annular dark-field scanning, and TEM (HAADF-STEM-EDS) were used to characterize the specimens. Thermo VG Escalab 250 photoelectron spectrometer was used for XPS analysis. The pore structures were evaluated by N₂ adsorption at 77 K using volumetric equipment (Quantachrome AS-1-MP) after pre-evacuation for 2 h at 423 K while maintaining a base pressure of 10⁻⁴ Pa.

Electrochemical characterizations. The electrochemical characterizations containing electrochemical impedance spectroscopy (EIS), galvanostatic charge–discharge (GCD), and cyclic voltammetry (CV) were carried out in a conventional three-electrode configuration at ambient temperature in which the NiCo₂O₄ on Ni foam electrode was utilized as the working electrode, and a Hg/HgO and Pt foil were the reference and counter electrodes. The electrochemical characterizations were done using a 6 M potassium hydroxide solution with the aid of ZIVE SP2 workstation (10 μ Hz–4 MHz). The voltage range was from 0 to 0.6 V vs. Hg/HgO for the NiCo₂O₄ electrode. To examine the energy storage performance of the working electrode for practical applica-

tion, the anode electrode material was nickel cobalt oxide, and the cathode electrode material was carbon nanotubes (CNTs). The voltage range was from 0 to 1.5 V vs. Hg/HgO for the CNT electrode.

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Author contributions

R.X., X.C., L.H. conceived the main idea. F.M., F.J., E.Z., S.B., K.S.H., and K.N.H. performed all the calculation work. All authors analyzed the results and wrote the paper.

Competing interests

The authors declare no competing interests.

Additional information

Correspondence and requests for materials should be addressed to E.Z. or S.B.

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