Hierarchical NiS/N-doped carbon composite hollow spheres with excellent supercapacitor performance†

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Designing electrode materials with high specific capacitance is crucial for further improving the energy storage performance of supercapacitors. Nickel sulfide (NiS) nanosheets anchored on the surface of N-doped hollow carbon spheres (NiS/NHCS) were rationally designed and synthesized *via* a multistep transformation approach. Uniform nickel silicate nanosheets were first deposited on the surface of NHCSs, followed by chemical conversion into NiS nanosheets. The NiS/NHCS composite shows excellent electrochemical properties as an electrode material for supercapacitors. The NiS/NHCS electrode can deliver a high specific capacitance of 1150 F g⁻¹ at 1 A g⁻¹ and show outstanding cycling performance with a retention rate of 76% over 4000 cycles, which are much higher than those of pure NiS hollow spheres (NiS-HS) (400 F g⁻¹ and 63%, respectively). Moreover, a hybrid supercapacitor assembled with NiS/NHCS as the cathode and an activated carbon electrode as the anode delivers a high energy density of 38.3 W h kg⁻¹ at a power density of 160 W kg⁻¹ and has an excellent cycle performance with the retention rate of 96% after 5000 cycles. The exceptional electrochemical performance of the NiS/NHCS composite indicates its potential application in high-performance supercapacitors.

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Introduction

Energy storage and conversion devices, in particular rechargeable batteries and supercapacitors, have attracted increasing attention for the better utilization of renewable resources.^{1–5} Supercapacitors have been considered as one of the most promising device candidates due to their intriguing characteristics of fast charge/discharge capability, outstanding cycle life and high power density.^{6–10} Nonetheless, the relatively low energy density of supercapacitors compared with rechargeable batteries still requires significant improvement. Enhancing the specific capacitance of the electrode materials is one of the compelling approaches to solve this issue.

Recently, transition metal sulfides have been extensively investigated as novel electrode materials for supercapacitors,¹¹ due to their high theoretical specific capacitance, excellent

electrical conductivity, and mechanical and thermal stability, including nickel sulfides (NiS,¹² Ni₂S₃ (ref. 13) and Ni₇S₆ (ref. 14)), cobalt sulfides (Co₃S₄ (ref. 15)), manganese sulfides (MnS¹⁶), and binary metal sulfides (NiCo₂S₄ (ref. 17) and MnCo₂S₄ (ref. 18)). Among them, nickel sulfide has received special interest. For instance, NiS box-in-box hollow structures reported by Yu *et al.*¹⁹ delivered a high specific capacitance of 668 F g⁻¹ at the current density of 1 A g⁻¹ and distinguished rate performance (71% capacitance retention at 20 A g⁻¹). Yang *et al.*²⁰ synthesized hierarchical flower-like nickel sulfide which exhibited a specific capacitance of 972.3 F g⁻¹ at 2 A g⁻¹ and excellent cycling stability. Zhu *et al.*²¹ fabricated NiS hollow spheres *via* a facile template-engaged transformation approach, which showed high specific capacitances of 927–583 F g⁻¹ at different current densities of 4.08–10.2 A g⁻¹. These studies on nickel sulfide materials indicate that these unique nanostructures are able to improve the electrochemical performance for supercapacitors.

On the other hand, the growth and anchoring of functional nanomaterials on carbonaceous materials, such as graphene,^{22–25} carbon nanotubes²⁶ and carbon fiber,^{18,27} is an effective method to improve the cycling performance and rate properties of supercapacitors.²⁸ Hollow carbon spheres (HCSs) have been regarded as an alternative substrate for preparing electrode material due to the following advantages: internal cavities serving as ion reservoirs and rapidly providing electrolyte ions for the electrochemical reactions, a carbon shell with

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good conductivity facilitating electron transport,²⁹ and a large surface area favoring the deposition of active materials. It can therefore be expected that a thin and porous layer of active material grown on the HCS surface could exhibit enhanced energy storage capacity and rate capacity.

Herein, we reported the design and fabrication of porous NiS nanosheets grown on N-doped hollow carbon spheres (NHCSs) *via* an efficient template-engaged method. The first step involves fabricating N-doped carbon spheres@SiO₂ (N-carbon/SiO₂) with a core-shell structure, which serves as the template for the growth of nickel silicate (NiSi) nanosheets on the surface of NHCSs in the second step, yielding NiSi/NHCS hollow structures. Finally, the NiSi/NHCS hollow structure is transformed into NHCS-supported NiS nanosheets (NiS/NHCS) by hydrothermal sulfidation treatment. When used as an electrode material for supercapacitors, the NiS/NHCS composite with a hierarchically hollow structure exhibits better electrochemical properties than pure NiS hollow spheres (NiS-HS).

Results and discussion

The synthesis process of the hierarchical NiS/NHCS composite with a hollow structure is illustrated in Fig. 1. First, N-carbon/SiO₂ with uniform sizes (Fig. S1a and b†) is synthesized *via* a one-step process involving the hydrolysis of tetraethoxysilane (TEOS) and the polymerization of dopamine, followed by the carbonization of the polydopamine/SiO₂ spheres. Then nickel silicate nanosheets are *in situ* formed on the NHCS surface *via* a facile hydrothermal treatment in dilute ammonia solution, where NHCSs and SiO₂ cores serve as templates and the self-sacrificial substrate, respectively (step I). In this process, ammonia can not only produce an alkaline condition but also provide NH₃ to coordinate with Ni²⁺.³⁰ Ammonium chloride is a necessary additive to mediate the dissociation of ammonia and inhibit the formation of nickel hydroxide. Silicate anions are released from the dissolution of the SiO₂ core under alkaline conditions at high temperature. The reaction between the silicate ions and Ni²⁺ leads to the nucleation of NiSi, which could preferentially deposit on the carbon shell. Continuous dissolution of the SiO₂ core supplies more silicate anions for the growth of NiSi nanosheets. When there are not enough dissolved Ni²⁺ ions, an outer shell composed of nickel silicate nanosheets on the NHCS surface and the remaining SiO₂ core

enclosed within form a rattle-like structure (NiSi/NHCS/SiO₂). As the reaction continues, the SiO₂ core is fully dissolved and a hollow nanostructure of NiSi/NHCS is eventually fabricated. Finally, the as-prepared NiSi/NHCS is transformed into NiS/NHCS by a solution-based sulfidation approach with Na₂S as the S²⁻ source (step II).

The microstructures of reaction products for each stage were characterized by transmission electron microscopy (TEM). The N-carbon/SiO₂ (Fig. 2a and e) solid spheres had a uniform diameter of 310 nm. For NiSi/NHCS/SiO₂ obtained after reaction for 2 h (Fig. 2b and f), nickel silicate nanosheets are closely grown on the surface of NHCSs, and a gap between the NiSi/NHCS shell and SiO₂ core can be clearly observed. After 12 h of reaction, NiSi/NHCS (Fig. 2c and g) shows a hollow structure with a 400 nm outer diameter and shell thickness of 50 nm, and this structure is maintained in the final product NiS/NHCS (Fig. 2d and h). The hollow structure combining the NiS nanosheet shell and NHCSs can provide abundant electrochemically active sites and a large electrolyte-electrode contact area for a pseudocapacitive reaction. Meanwhile, the N-doped carbon core can boost the electron transfer, which can improve the rate capacity of supercapacitors. Additionally, the aggregation of electroactive particles and a pronounced volume variation during charge/discharge processes can be effectively inhibited due to the hollow carbon cavities.

Field-emission scanning electron microscope (FESEM) images (Fig. 3a and b) confirmed that the surface of both NiSi/NHCS and NiS/NHCS is assembled by using numerous nanosheets, and the morphology was not altered during sulfidation. The NiSi hollow sphere (NiSi-HS) (Fig. S2†) and NiS-HS (Fig. 3c) are also composed of interconnected nanosheets, and TEM images of NiS-HS (Fig. 3d) further confirm the hierarchical hollow structure. The average diameter of NiS-HS is about 380 nm, which is smaller than that of NiS/NHCS, mainly due to the absence of the NHCS core. The N₂ adsorption-desorption isotherms (Fig. S3a†) of NiS/NHCS and NiS-HS can be classified as type IV with an overlap of type H2 and H3 hysteresis loops, which shows the existence of ink-bottle mesopores within the NiS nanosheets or porous NHCSs and slit-like mesopores formed between NiS nanosheets, respectively.³¹ NiS/NHCS and NiS-HS have hierarchical pore structures according to the wide pore size distribution (Fig. S3b†), which is beneficial for the electrolyte ion penetration.³² Especially, the BET specific surface area and pore volume of NiS/NHCS (106 m² g⁻¹ and 0.30 cm³ g⁻¹) are higher than those of NiS-HS (86 m² g⁻¹ and 0.15 cm³ g⁻¹), which can provide more electroactive sites for participating in the faradaic redox reactions, leading to improved electrochemical performance.

The crystal structures of NHCSs, NiSi/NHCS and NiS/NHCS were characterized by the XRD patterns (Fig. 4a). The peak at 23.5° can be assigned to graphite carbon sheets of NHCSs. In the XRD spectrum of NiSi/NHCS, the weak peaks located at 35.5° and 60.7° are indexed to the (200) and (060) planes of monoclinic nickel hydrosilicate, respectively.²¹ In the XRD spectrum of NiS/NHCS, the peaks at 19.4°, 30.3°, 32.7°, 35.9°, 49.4°, 53.5° and 59.8° can be indexed to (110), (101), (300), (021), (410), (401) and (012) crystal planes of NiS (JCPDF #86-2281).

Fig. 1 Scheme for the formation of nickel sulfide (NiS) nanosheets supported on the NHCS surface by a novel conversion route: (I) growth of nickel silicate (NiSi) nanosheets on the NHCS surface with the silica core simultaneously eliminated; (II) chemical conversion of nickel silicate into NiS to form NiS/NHCS.

Fig. 2 TEM images of N-carbon/SiO₂ (a and e), NiSi/NHCS/SiO₂ (b and f), NiSi/NHCS (c and g) and NiS/NHCS (d and h).

From the Raman spectra of NHCSs and NiS/NHCS (Fig. 4b), typical D and G bands at 1358 and 1596 cm⁻¹ were observed, further confirming the existence of carbon in the NiS/NHCS composite. The relatively high intensity of the D band demonstrates the existence of substantial defects, mainly resulting from nitrogen dopants and contaminants.³³ The peak at

578 cm⁻¹ in the spectrum of NiS/NHCS is attributed to the F_{2g} model of Ni-S.³⁴

The surface elemental composition and chemical state of N-carbon/SiO₂ and NiS/NHCS were analyzed by X-ray photoelectron spectroscopy (XPS). The survey XPS spectrum of N-carbon/SiO₂ (Fig. S4a†) demonstrates the presence of C, N, O

Fig. 3 FESEM images of NiSi/NHCS (a), NiS/NHCS (b) and NiS-HS (c) and the TEM image of NiS-HS (d). The scale bars of the insets are 100 nm.

Fig. 4 (a) XRD patterns of NHCSs, NiS/NHCS and NiS/NHCS. (b) Raman spectra of NHCSs and NiS/NHCS. High-resolution XPS spectra of Ni 2p before and after charging (c) and S 2p (d) for NiS/NHCS.

Fig. 5 Cyclic voltammetry curves (a), GCD curves (b), specific capacitance at various current densities (c) and Nyquist plots (d) of NiS/NHCS and NiS-HS electrodes.

and Si elements. Particularly, the N signal confirms the successful introduction of N into the carbon shell. The C 1s peak in the high-resolution spectrum (Fig. S4b†) can be deconvoluted into four peaks at 284.6, 285.6, 287.2 and 289.0 eV, corresponding to sp^3 -hybridized C-C/C=C, C-O/C-N, C=O and -COO bonds.²⁹ The high-resolution N 1s spectrum (Fig. S4c†) can be deconvoluted into four peaks at 398.3, 400.0, 400.9 and 403.5 eV, corresponding to pyridinic-N, pyrrolic-N, quaternary-N and N-oxide. The N element can not only increase the conductivity of carbon materials, but also facilitate electrolyte penetration with a feasible pathway. The survey XPS spectrum of NiS/NHCS (Fig. S5†) shows peaks for Ni, S, C, N and O elements, where the signal for O can be attributed to the oxygen-containing functional group in the NHCS core and sulfur oxides in the NiS shell.^{33,35,37} The high-resolution XPS spectrum of Ni 2p (Fig. 4c) shows two characteristic peaks at 852.1 and 869.7 eV and two shake-up satellites at 858.0 and 876.1 eV, which can be attributed to Ni 2p_{3/2} and Ni 2p_{1/2} orbitals of Ni²⁺.³⁶ Furthermore, the XPS peak of S 2p (Fig. 4d) is deconvoluted into two binding energy peaks at 162.5 and 163.8 eV, which correspond to S 2p_{3/2} and S 2p_{1/2} orbitals of S²⁻, respectively. Another peak at 168.3 eV belongs to sulfur oxide from the oxidation of S with residual O₂.^{37,38}

The electrochemical performance of NiS/NHCS and NiS-HS was investigated by using a three-electrode configuration in

2.0 M KOH aqueous solution. The cyclic voltammetry (CV) curves of both samples (Fig. 5a) exhibit a couple of faradaic redox peaks, which can be assigned to the reversible faradaic redox process of Ni³⁺/Ni²⁺.²⁹ The high-resolution XPS spectra of Ni 2p for the NiS/NHCS after charging was used to prove the Ni valence change in the charging/discharging process. The two evident peaks appearing at 854.8 and 872.4 eV can be ascribed to Ni³⁺, indicating the occurrence of the redox reaction. It is obviously seen that the integrated CV area of NiS/NHCS is much larger than that of NiS-HS, indicating that NiS/NHCS possesses better capacitive properties. In the galvanostatic charge/discharge (GCD) curve of NiS/NHCS (Fig. 5b), the discharging time matches well with the charging time, revealing a high reversibility of the pseudocapacitive reaction at the surface of the electrodes. And the potential plateaus correspond to the redox peaks of the CV curves. Notably, the discharge time of NiS/NHCS is much longer than that of NiS-HS, indicating the markedly improved specific capacitance due to the introduction of NHCSs as the conductive substrate.

The GCD curves of NiS/NHCS and NiS-HS electrodes were measured at various current densities (Fig. S7a and c†) to evaluate the rate performance (Fig. 5c). The NiS/NHCS electrode outperforms NiS-HS over the entire current density range tested. For example, the NiS/NHCS electrode shows a high specific capacitance of 1150 F g⁻¹ at 1 A g⁻¹ and a capacitance

Fig. 6 (a) Configuration and illustration of the energy storage mechanism for the NiS/NHCS||AC device. (b) CV curves of AC and NiS/NHCS from -1.0 to 0 V and 0 to 0.5 V, respectively, at 10 mV s⁻¹ in a three-electrode system. (c) CV curves of the as-fabricated NiS/NHCS||AC device in various operation voltages at a scan rate of 10 mV s⁻¹.

retention rate of 52.2% at the high current density of 20 A g^{-1} , which are much higher than those of the NiS-HS electrode (400 F g^{-1} at 1 A g^{-1} and 45.6% at 20 A g^{-1}). The N-doped carbon hollow core uniformly deposited with the NiS nanosheets can promote the electron transfer in the NiS/NHCS electrode, leading to enhanced rate performance. The mechanism for the excellent capacitive properties of the NiS/NHCS electrode was further investigated by electrochemical impedance spectroscopy (Fig. 5d). In the high frequency region, there is a pseudo-semicircle in the Nyquist plot of NiS-HS while the semicircle is negligible for NiS/NHCS, indicating that NiS/NHCS exhibits a smaller charge-transfer resistance than NiS-HS. In the low frequency region, the slope for the NiS/NHCS electrode is larger than that for NiS-HS, suggesting that NiS/NHCS not only possesses lower diffusive resistance but offers faster ion transport. To evaluate the durability of the electrode materials, the long-term cycling performance of NiS/NHCS and NiS-HS were tested by GCD measurements up to 4000 cycles at 5 A g^{-1} . 76% of the specific capacitance of NiS/NHCS was retained after 4000 cycles compared to 63% retention for NiS-HS (Fig. S8†).

To further evaluate the NiS/NHCS electrode for practical application, a hybrid supercapacitor (HSC) device was assembled using the NiS/NHCS electrode as the cathode and the active carbon (AC) electrode as the anode in KOH aqueous electrolyte (Fig. 6a). Fig. 6b shows the CV curves of NiS/NHCS and AC electrodes at a scan rate of 10 mV s^{-1} with a stable potential range of -1 to 0 V and 0 to 0.5 V in the three-electrode system,

respectively. The energy storage mechanism of the NiS/NHCS||AC device is depicted in Fig. 6a. The NiS/NHCS cathode can store energy by an oxidation reaction of NiS with the incorporation of OH^- during the charging process while the AC anode can accumulate electrons by cation (*e.g.* K^+) adsorption.³⁹ In the charging process, the NiS/NHCS cathode and AC anode can produce an electric potential from 0 to 0.5 V and from -1 to 0 V , respectively, endowing the HSC device with a broad operating voltage window of 1.5 V . In the discharging process, the NiS/NHCS cathode and AC anode can release energy *via* the reduction reaction and desorption of ions. The employed AC electrode exhibits a specific capacitance of 280 F g^{-1} at the current density of 1 A g^{-1} (Fig. S9†). According to the charge balance theory between the anode and cathode in the HSC device, the optimal mass loading ratio for the NiS/NHCS and the AC electrode is $m_{(\text{NiS/NHCS})} : m_{(\text{AC})} = 0.54 : 1$, while that for the NiS-HS and the AC electrode is $m_{(\text{NiS-HS})} : m_{(\text{AC})} = 1.4 : 1$. Utilizing the different potential window of the NiS/NHCS and the AC electrode, the operating voltage of the fabricated HSC device is expected to be 1.5 V (Fig. 6c and d).

As depicted in Fig. 7a and b, the integrated area of the CV curve of NiS/NHCS||AC was much larger than that of NiS-HS||AC, while the discharging time of the former is also much and longer than that of the latter, indicating that the specific capacitance of NiS/NHCS||AC is much larger than that of NiS-HS||AC (Fig. 7c). The specific capacitance of the NiS/NHCS||AC exhibited a maximum specific capacitance of 120 F g^{-1} at the

Fig. 7 (a) CV curves, (b) GCD curves, (c) current–density dependence of the specific capacitance and (d) Ragone plots of NiS/NHCS||AC and NiS-HS||AC. The inset of part d shows two LEDs powered by two NiS/NHCS||AC devices in series.

current density of 0.2 A g^{-1} and maintained a capacitance of 46 F g^{-1} at 5 A g^{-1} , while those of NiS-HS||AC were 75.3 F g^{-1} at 0.2 A g^{-1} and 28.1 F g^{-1} at 5 A g^{-1} . In contrast to the single NiS/NHCS electrode in the three-electrode system, the GCD curves of the HSC devices (Fig. S10a and S11c†) exhibited a better linear feature, demonstrating that the electron release from the NiS/NHCS redox reaction in the cathode has been well matched with the electron accumulation by cation adsorption in the AC anode.

Energy density and power density are two important factors for evaluating the electrochemical performance for a hybrid supercapacitor device. It can be observed from the Ragone plot (Fig. 7d) that the NiS/NHCS||AC device demonstrates a high energy density of 38.3 W h kg^{-1} at a power density of 160 W kg^{-1} with a charging time of 1150 s (Fig. S10a†). It is worth noting that, even when charging for only 25 s , the HSC can still possess an energy density of 13.5 W h kg^{-1} at a high power density of 4000 W kg^{-1} . By contrast, the NiS-HS||AC device shows energy densities of 24.8 W h kg^{-1} at 150 W kg^{-1} and 8.8 W h kg^{-1} at 3944 W kg^{-1} . To manifest the practical operating performance of the NiS/NHCS||AC device, two LEDs (with lowest working voltage of $\sim 2.2 \text{ V}$) can be powered by two HSC devices connected in series ($\sim 1.4 \text{ mg}$ active materials in total for each), with red light emission lasting for 110 s . Fig. S12† shows the long-term cycling property of the NiS/NHCS||AC at a current density of 3 A g^{-1} . Remarkably, the NiS/NHCS||AC exhibited excellent cycling stability and still delivered 96% of its initial capacitance after 5000 cycles. After 5000 cycles, NiS nanosheets were still anchored on the surface of NHCSs (Fig. S13†), indicating that the NiS/NHCS composite possessed a stable chemical structure.

The excellent supercapacitive performance of NiS/NHCS can be attributed to several aspects. On the one hand, the NiS/NHCS composite with a novel hierarchical architecture can facilitate the diffusion of electrolyte ions and supply sufficient electroactive sites for redox reactions. On the other hand, the NHCS core can markedly increase the electron transfer efficiency in the electrode material and alleviate the aggregation of NiS nanosheets during the charge/discharge process.^{40–42} Finally, the NHCSs as the backbone could support the NiS nanosheets stably during the charge/discharge process, which is beneficial for the cycling stability. Hence, the NiS/NHCS composite can be regarded as a promising candidate material for high-performance supercapacitors, due to its high specific capacitance and outstanding stability.

Conclusion

A hierarchical NiS/NHCS composite with enhanced supercapacitive performance can be rationally designed and fabricated by a sacrificial template method. The NiS/NHCS composite exhibits a specific capacitance of 1150 F g^{-1} at 1 A g^{-1} and 600 F g^{-1} at the high current density of 20 A g^{-1} . And the hybrid supercapacitor device assembled with NiS/NHCS as the cathode and an activated carbon electrode as the anode delivers a high energy density of 38.3 W h kg^{-1} at a power density of 160 W kg^{-1} and has an excellent cycle performance with the retention rate of 96% after 5000 cycles. Moreover, the

strategy reported herein can be applied for synthesizing other hierarchical carbon-metal sulfide composite materials for energy conversion and storage applications.

Experimental section

Synthesis of N-doped carbon/SiO₂ spheres

N-doped carbon/SiO₂ (N-carbon/SiO₂) spheres were synthesized by a facile one-step method involving the formation of spherical SiO₂ cores and the coating of polydopamine. Briefly, a mixture containing 12 mL of anhydrous ethanol, 0.5 mL of ammonia (25%) and 40 mL of distilled water was vigorously stirred for 10 min . One milliliter of tetraethoxysilane (TEOS) was added to the mixture dropwise, and the stirring continued for an additional 15 min . Subsequently, 4 mL of dopamine hydrochloride (DA) aqueous solution (50 mg mL^{-1}) was added to the above solution and kept stirring for 32 h . Brown polydopamine/SiO₂ (PDA/SiO₂) spheres were collected after centrifugation and washed with purified water three times. N-carbon/SiO₂ spheres were obtained after calcination at 800°C for 2 h under a nitrogen (N₂) atmosphere.

Synthesis of the NiSi/NHCS composite

For preparation of NiSi/NHCS, 40 mg of N-carbon/SiO₂ was dispersed in 40 mL of water by sonication for 0.5 h , followed by the addition of NiCl₂ (0.5 mmol), NH₄Cl (5 mmol) and NH₃·H₂O (0.5 mL) under mild stirring. After stirring for 0.5 h , the mixture suspension was transferred into a Teflon-lined autoclave and maintained at 105°C . After reaction for 2 h and 12 h , NiSi/NHCS/SiO₂ and NiSi/NHCS composites were obtained, respectively.

Synthesis of NiS/NHCS

40 mg of NiSi/NHCS and 100 mg of polyvinylpyrrolidone (PVP, K-30) were added to 60 mL of distilled water and sonicated for 0.5 h . Subsequently, 600 mg of Na₂S·9H₂O was added during stirring, and the suspension was transferred to a 100 mL Teflon-lined autoclave and heated at 160°C for 12 h . The product was obtained after washing with purified water and ethanol several times and annealing at 300°C for 2 h in a N₂ atmosphere. For comparison, NiS hollow spheres (NiS-HS) were fabricated under the same conditions as those of the synthesis of NiS/NHCS, except that SiO₂ spheres were used instead of the N-carbon/SiO₂.

Materials characterization

Powder X-ray diffraction (XRD) patterns of the samples were recorded on a Rigaku X-ray diffractometer with Cu K_α radiation ($\lambda = 0.15418 \text{ nm}$). The morphologies were examined with a field emission scanning electron microscope (FESEM, JSM-7500) and transmission electron microscope (TEM, Titan G2). Nitrogen adsorption-desorption experiments were performed on a Micromeritics ASAP 2020 nitrogen adsorption apparatus to measure the specific surface area and porosity of the samples. The X-ray photoelectron spectroscopy (XPS) was conducted on a Thermo ESCALAB 250. Raman spectrum analysis was

performed by using a Renishaw Raman microscope under $\lambda_{\text{exc}} = 633$ nm laser excitation.

Electrochemical measurements

The working electrodes were obtained by brushing a mixture of active materials (NiS/NHCS, NiS-HS or AC), acetylene black and polyvinylidene fluoride (mass ratio 8 : 1 : 1) in N-methyl pyrrolidone (NMP) solvent onto nickel foam and air-drying at 100 °C overnight. The mass loading of the active material for individual electrodes was 1.0 mg cm⁻². A three-electrode system configuration was used to test the electrochemical performance of individual electrodes, where a platinum foil served as the counter electrode and a Ag/AgCl electrode served as the reference electrode. Cyclic voltammetry (CV), galvanostatic charge/discharge (GCD) and electrochemical impedance spectroscopy (EIS) were carried out on a electrochemical workstation (CHI 760E) in 2.0 M KOH electrolyte. Moreover, the ACS device was assembled with NiS/NHCS as the cathode and AC as the anode to measure its potential for practical application.

Conflicts of interest

There are no conflicts to declare.

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