Electrospun carbon nanofiber@CoS$_2$ core/sheath hybrid as an efficient all-pH hydrogen evolution electrocatalyst†

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Developing all-pH and efficient non-noble-metal electrocatalysts for the hydrogen evolution reaction (HER) remains a great challenge in science nowadays. In this work, carbon nanofiber (CNF)@CoS$_2$ hybrids with a hierarchical core/sheath structure have been successfully fabricated via a combination of electrospinning and hydrothermal methods for use as HER electrocatalysts under all-pH media. The CoS$_2$ nanoparticles are uniformly attached onto the three-dimensional (3D) CNF framework, and have different morphologies on adjusting the concentration of the Co and S precursors. This shows that the morphology of the electrocatalytically active CoS$_2$ has a great impact on the HER performance, which can be attributed to the difference in the exposure of the effective catalytic sites. Moreover, the CNF@CoS$_2$ hybrids distinctly exhibit superior HER activities compared to that of pure CoS$_2$ agglomerates, which highlights the importance of the 3D carrier with a nanostructure to increase the number of exposed electrocatalytic sites. When applied as electrocatalysts operating at all-pH values, the CNF@CoS$_2$ hybrids exhibit remarkable HER properties, with a low onset potential of $\approx 40$ mV, a small Tafel slope of 66.8 mV per decade and a large current density (10.0 mA cm$^{-2}$ at $\eta = 110$ mV) in acid medium, along with a low onset potential of $\approx 130$ mV in both alkaline and neutral conditions. Apart from the facile preparation of an efficient all-pH electrocatalyst, this work highlights the impacts of morphology and nanostructure of the catalyst on the HER activity, which is universally applicable in the field of catalysis.

1. Introduction

Due to the serious air pollution induced by overusing fossil fuels, it has become a matter of utmost urgency to search for clean and carbon-neutral energy sources to meet the growing power appetite of the world.$^1$ Hydrogen is a universally recognized energy carrier, which can store solar and electric energy via formation of a chemical bond between two hydrogen atoms.$^2$ Electr...
chalcogen atoms, e.g., NiS$_2$, CoS$_2$ and CoSe$_2$. The application of these TMDs in HER catalysts is inspired by their similar structure and composition to hydrogenases, whose catalytic activity is comparable to that of platinum. Indeed, a large number of studies have reported that these first-row TMDs are highly HER active electrocatalysts, which even possess advantages over 2D layered TMDs like WS$_2$ and MoS$_2$. In particular, cobalt based dichalcogenides are emerging as attractive HER electrocatalysts owing to their low free energy barrier for the adsorption of H atoms on Co atoms, which has been further proved using experimental studies. For instance, Cui’s group prepared a series of first-row TMDs (FeS$_2$, CoS$_2$, NiS$_2$, CoSe$_2$, et al.) through sulfurization/selenization of e-beam evaporated metal films, and among them, cobalt dichalcogenides presented superior HER activity relative to the others. Apart from their intrinsic HER catalytic activity, a lot of highly conductive carbon materials with large surface areas have been involved in constructing composite electrocatalysts with outstanding performance. CoS$_2$ nanoparticles grown on carbon fiber paper were successfully prepared via a two-step reaction, and the obtained hybrid exhibited excellent HER activity in acid electrolyte, achieving a superhigh current density of 100 mA cm$^{-2}$ at a low overpotential of 180 mV, along with exceptional durability. Peng et al. fabricated a 3D CoS$_2$/reduced graphene oxide (rGO)–carbon nanotube (CNT) nanocomposite film with a nanoporous structure and high electrical conductivity, which exhibited a low overpotential of 142 mV at a current density of 10 mA cm$^{-2}$. In addition to hybridization, tuning the morphology of electrocatalysts is also an effective pathway to improve their HER performance. Hollow cobalt sulfide nanoprisms were synthesized using a microwave-assisted synthetic route, exhibiting a superior catalytic performance to those prepared using a traditional solvothermal method, due to their higher surface area with larger porosity, thus facilitating access to the catalytically active sites. Jin’s group reported the controllable preparation of CoS$_2$ materials with three morphologies, i.e., films, microwires and nanowires, and further demonstrated that the micro- and nanostructured CoS$_2$ could increase the electrocatalytic surface area and promote the release of the generated hydrogen bubbles, compared with that of CoS$_2$ film in 0.5 M H$_2$SO$_4$ solution.

Moreover, it is worth mentioning that cobalt based dichalcogenides not only possess a remarkable HER performance in acid media, but also have great catalytic potential in alkaline and neutral conditions. As is well known, most works based on noble-metal-free electrocatalysts mainly focus on their acid-stable HER activities for proton exchange technology. However, commercial alkaline electrolysis technology and microbial electrolysis cells also require effective catalysts for use in alkaline and neutral electrolytes. As a result, it is of great significance to develop efficient HER electrocatalysts over the whole pH range. However, only limited success has been achieved up to now.

Herein, carbon nanofiber@CoS$_2$ (CNF@CoS$_2$) hybrids with a core/sheath structure have been faciley fabricated through a combination of electrospinning and hydrothermal methods, and utilized as efficient HER electrocatalysts across the pH range. By adjusting the concentration of the Co and S precursors, CNF@CoS$_2$ hybrids can form CoS$_2$ nanoparticles with different morphologies, thus having a great influence on their electrocatalytic performance. It can be found that angular shaped CoS$_2$, distributed uniformly with a relatively small size on the CNF carrier, exhibits the best HER performance, which can be attributed to its nanoscale structure with more exposed active sites compared with the others. Benefiting from the synergistic effects between the electroactive material and the carrier, the CNF@CoS$_2$ hybrid exhibits an outstanding HER performance, with a low onset potential of 40 mV, a small Tafel slope of 66.8 mV per decade as well as a large current density (10.0 mA cm$^{-2}$ at $\eta = 110$ mV) in acid medium. Notably, it also works well in alkaline and neutral conditions, both with an onset potential of $\sim$130 mV, indicating its great potential as an all-pH and efficient hydrogen-evolution electrocatalyst.

2. Experimental

2.1. Materials

Polyacrylonitrile (PAN, $M_w = 150\,000$ g mol$^{-1}$) was purchased from Sigma-Aldrich. CoCl$_2$·6H$_2$O and thiourea (CS(NH$_2$)$_2$) were purchased from J&K Chemical, with a purity of 99%. N,N-Dimethylformamide (DMF) and ethanol were obtained from Shanghai Chemical Reagent Company. Deionized water was used throughout all the experiments.

2.2. Preparation of electrospun CNF nanofiber membranes

The preparation process of the CNF@CoS$_2$ nanofiber membranes was schematically shown in Scheme 1. PAN nanofiber membranes were produced through a simple single-nozzle electrospinning technique. Typically, 1.0 g PAN was dissolved overnight in 10 mL DMF with vigorous stirring, forming a viscous homogenous solution. Then, it was sucked into a 5 mL viscous homogenous solution. Then, it was sucked into a 5 mL
syringe equipped with a stainless spinneret, which was positioned 18 cm away from the collector. When a high voltage of 20 kV was applied to the system, PAN nanofibers were generated and collected with a feeding rate of 0.25 mm min⁻¹ as self-standing membrane. The pre-oxidation process was operated under 250 °C for 2 h with a heating rate of 1 °C min⁻¹ in an air atmosphere to obtain oxidized-PAN (o-PAN) nanofiber membranes. Afterwards, they were carbonized in a nitrogen flow at 800 °C for 2 h to prepare the CNF membranes.

The CNF@CoS₂ core/sheath hybrid membranes were fabricated with a one-step hydrothermal method. First, the prepared CNF membranes were immersed in a bath of HNO₃/H₂O (1 : 1 v/v) for 6 h to increase their hydrophilicity, and then washed with deionized water several times. After that, certain amounts of CoCl₂·6H₂O (i.e., 0.3, 1.0, and 3.0 mmol) and CS(NH₂)₂ (i.e., 1.0, 5.0, and 15.0 mmol, respectively) were dissolved in 36 mL deionized water with sonication for a few minutes. The solution was transferred to a 100 mL Teflon-lined stainless steel autoclave with a piece of acid-treated CNF membrane (2 × 2 cm²) immersed in the reaction solution. The autoclave was sealed and maintained at 180 °C for 17 h, and then cooled down to room temperature. Finally, the obtained hybrid membrane was rinsed with deionized water and dried at 70 °C for 6 h. The products were named CNF@CoS₂-1, CNF@CoS₂-3 and CNF@CoS₂-9, respectively. Pure CoS₂ was also synthesized via the same procedure without the addition of CNF membrane.

2.3. Characterization

The morphology of the products was observed using a field emission scanning electron microscope (FESEM, Ultra 55, Zeiss) at an acceleration voltage of 5 kV. X-ray diffraction (XRD) experiments were conducted on an X'Pert Pro X-ray diffractometer with Cu Kα radiation (λ = 0.1542 nm) under a voltage of 40 kV and a current of 40 mA. Thermogravimetric analysis (Pyris 1 TGA) was performed under an air flow from 100 °C to 900 °C at a heating rate of 20 °C min⁻¹. X-ray photoelectron spectroscopy (XPS) analyses were made with a VG ESCALAB 220i-XL device and all XPS spectra were corrected using the C 1s line at 284.6 eV.

2.4. Electrochemical measurements

Electrochemical tests were performed in a standard three-electrode setup using a CHI 660D electrochemical workstation (Shanghai Chenhua Instrument Co., China), where a graphite rod was used as the counter electrode, a saturated calomel electrode (SCE) as the reference electrode and a modified glassy carbon electrode (GCE) as the working electrode. A piece of pre-cut CNF@CoS₂ core/sheath hybrid membrane was fixed onto the GCE using Nafion solution as a conductive binder, with a loading amount of 0.6–0.8 mg in a square shape of 3 × 3 mm². Linear sweep voltammetry (LSV) was measured with a scan rate of 2 mV s⁻¹ in 0.5 M H₂SO₄, 1.0 M KOH solution and 1.0 M potassium phosphate buffer solution (PBS). All of the potentials were calibrated to the reversible hydrogen electrode (RHE) by adding a value of (0.241 + 0.059 × pH) V.

Tafel plots were obtained by plotting the polarization curves as the overpotential (η) versus log current density (log j) to assess the HER kinetics. The linear portions of the Tafel plots fitted well to the Tafel equation (η = b log(j) + a, where b is the Tafel slope). The double-layer capacitance (Cdl) of the various products can be calculated from the cyclic voltammograms in the region of 0.15–0.25 V vs. RHE. By plotting Δj at 0.2 V vs. RHE against the scan rate, the slope is twice Cdl. Electrochemical impedance spectroscopic (EIS) measurements were carried out under various overpotentials in the frequency range from 10⁶ Hz to 10⁻² Hz with an amplitude of 5 mV. The electrocatalytic stability of the products was evaluated using the current-time response under a fixed overpotential for a period of time.

3. Results and discussion

3.1. Morphology and structure of CNF@CoS₂ core/sheath hybrid membranes

Electrospinning is a low-cost, scalable and versatile method to prepare nanofiber membranes with tunable nanostructures, large specific surface areas and good structural stability. As shown in Fig. S1† the electrospun CNFs possess an average diameter of 300–400 nm and have smooth surfaces without defects. Such a randomly arranged structure forms an interconnected network, which can be utilized as an ideal carrier for the attachment of electrochemically active materials. In this work, CoS₂ nanoparticles were decorated onto the CNF membranes using a one-step hydrothermal process. It can be clearly seen from Fig. 1A–C that all the nanofibers are uniformly covered with CoS₂ nanoparticles in this overall view. However, as observed from images at high magnification (Fig. 1D–F), the morphology and size of the attached CoS₂ nanoparticles show a great change on increasing the concentration of CoCl₂·6H₂O and CS(NH₂)₂. For CNF@CoS₂-1, CoS₂ nanoparticles have an angular shape with a random polygonal structure. Furthermore, the average diameter of the hybrid nanofibers is rather large in size, up to nearly 1 μm. As shown in Fig. 1E, the CoS₂ nanoparticles in CNF@CoS₂-3 have a much smaller size compared with that in CNF@CoS₂-1, along with a more homogeneous dimension. Such a nano-scaled structure with relatively small and even size may increase the effective electrocatalytic surface area, thus exerting a positive effect on the HER performance. EDS mapping images of the CNF@CoS₂-3 hybrid (Fig. S2†) indicate the uniform distribution of Co and S elements, verifying the uniform growth of CoS₂ on the CNF carrier. With an excessive amount of cobalt salt, the CoS₂ particle size becomes even smaller for the CNF@CoS₂-9 hybrid without such distinct angular edges (Fig. 1C, F and S3†). In addition, the CoS₂ nanoparticles are so small (around 50–100 nm in diameter) that some part of them inevitably accumulate densely, which could impede effective contact between the electrolyte and the electrocatalytic active materials. For pure CoS₂ without CNF carrier (Fig. S4†), the CoS₂ particles self-aggregate to form large agglomerates,
indicating the importance of the CNF carrier to effectively disperse CoS₂ nanoparticles.

The surface chemical states of CNF@CoS₂-3 hybrid were characterized using XPS as shown in Fig. 2A–C. The survey spectrum confirms the coexistence of C, Co, O and S elements within the hybrid without any detectable impurity. The high-resolution Co 2p spectrum exhibits two strong peaks located at 778.7 and 793.8 eV, corresponding to Co 2p₃/₂ and Co 2p₁/₂, respectively. In the high resolution S 2p spectrum, peaks at 162.9 eV and 164.1 eV are observed, referring to S 2p₃/₂ and S 2p₁/₂ of the divalent sulfide ions (S²⁻), respectively. For the XRD patterns in Fig. 2D, the CNF membrane displays a broad peak ranging from 20° to 30°, implying its amorphous structure. Both pure CoS₂ and CNF@CoS₂-3 display diffraction peaks at 2θ = 28.3°, 32.7°, 36.7°, 40.3°, 46.8° and 55.5°, which can be well indexed to the (111), (200), (210), (211), (220) and (311) planes of the cubic phase of CoS₂ (JCPDS 41-1471), respectively. All the above results further demonstrate the successful preparation of the CNF@CoS₂ hybrids.

The loading amount of CoS₂ in different hybrids was studied using TGA analysis, as shown in Fig. 3. There exist several stages of weight change for pure CoS₂, which correspond to different phase changes. When performed in an air flow, CoS₂ begins to lose sulfur at 430 °C, which subsequently transforms to its stable form of metal sulfate. The products collected after thermal treatment at 700 °C and 900 °C were further investigated using XRD, as shown in Fig. S5.† The diffraction peaks at 2θ = 31.6°, 37.2°, 45.0°, 56.1°, 59.7° and 65.7° can be assigned to the (220), (311), (400), (422), (511) and (440) crystal planes of Co₃O₄ (JCPDS 42-1467), respectively.
indicating that CoS$_2$ has been oxidized completely at 900 °C. In contrast, although the product at 700 °C has some characteristic diffraction peaks of Co$_3$O$_4$, the peak intensities are far weaker than those at 900 °C. Furthermore, some additional peaks appear at $2\theta = 25.2^\circ$, $34.7^\circ$, which correspond to the planes of CoSO$_4$ (JCPDS 03-0843). As a result, when calcined in air, CoS$_2$ first loses sulfur, then changes to cobalt sulphate, and finally turns into pure Co$_3$O$_4$ at 900 °C. The TGA curve of CNF displays a huge weight loss from 430 °C to 680 °C, which results from its burning in air. As a result, the TGA process for the CNF@CoS$_2$ hybrids combines the decomposition of CNF with the transformation of CoS$_2$. To our surprise, it was found that the three CNF@CoS$_2$ hybrids had the same residue weight at 900 °C, demonstrating that the loading ratios of the electroactive CoS$_2$ remained almost the same, about 74.8 wt% in the hybrids (see ESI† for calculation details). This result indicates that the concentration of precursor has a great influence on the morphology instead of the loading amount.

3.2. All-pH-value HER performance of CNF@CoS$_2$ hybrids

The electrocatalytic activity of the CNF@CoS$_2$ hybrids was first investigated in an acid medium of 0.5 M H$_2$SO$_4$ solution with a three-electrode setup at a scan rate of 2 mV s$^{-1}$. The LSV curves for the CNF@CoS$_2$ hybrids with different morphologies are shown in Fig. 4A. Two key points are utilized to evaluate the performance of the HER electrocatalysts from the LSV curves. One lies in the overpotential ($\eta$), which is the difference between the applied and thermodynamic potential for a given electrochemical reaction. The other point refers to the electrochemical current density, indicating the output of the electrocatalytically evolved hydrogen. Naturally, an excellent HER catalyst is capable of generating a large current density under a low overpotential, i.e., more hydrogen can be produced using less energy if the reaction is undertaken using a well-performing electrocatalyst. Therefore, it can be deduced from Fig. 4A that among all the CNF@CoS$_2$ hybrids, the CNF@CoS$_2$-3 hybrid displays the best electrocatalytic activity, with an onset overpotential of 40 mV. The other two LSV curves are more or less negatively shifted compared with that of the CNF@CoS$_2$-3 hybrid, implying an inferior HER performance. Furthermore, the CNF@CoS$_2$-3 hybrid also has an apparent current density ($j$) of 10.0 mA cm$^{-2}$ at a low overpotential ($\eta$) of 110 mV, compared with 166 mV required for CNF@CoS$_2$-1 and 143 mV for the CNF@CoS$_2$-9 hybrid. A possible reason for the difference in their electrocatalytic activities can be speculated from the morphological changes discussed above. Compared to the other two hybrids, the CNF@CoS$_2$-3 hybrid possesses an optimal hierarchical structure with uniformly distributed CoS$_2$ nanoparticles of a relatively smaller size without severe agglomeration, which may offer a large contact area between electrolyte and catalyst, thus facilitating ion/electron transfer, leading to the best HER performance. This hypothesis was further confirmed from the electrochemical double-layer capacitance ($C_{dl}$) and electrochemical impedance spectra (EIS). $C_{dl}$ measurements were carried out using cyclic voltammetry (CV) tests to estimate the effective surface area of the solid–liquid interface. It can be observed from Fig. 4B and C that the CNF@CoS$_2$-3 hybrid exhibits a larger $C_{dl}$ of 17.5 mF cm$^{-2}$ than those of its counterparts (Fig. S6†), indicating a higher

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**Fig. 4**  (A) LSV polarization curves for the CNF@CoS$_2$-1, CNF@CoS$_2$-3 and CNF@CoS$_2$-9 hybrid modified GCE in 0.5 M H$_2$SO$_4$ solution. (B) CV curves of the CNF@CoS$_2$-3 hybrid in the region of 0.15–0.25 V vs. RHE. (C) Plot showing the extraction of the double layer capacitance ($C_{dl}$) for the CNF@CoS$_2$-3 hybrid at 0.2 V. (D) Nyquist plots of the CNF@CoS$_2$-1, CNF@CoS$_2$-3 and CNF@CoS$_2$-9 hybrids at an overpotential of 0.17 V.
exposure of the electrocatalytic sites, thus contributing to its excellent HER performance. Fig. 4D displays the Nyquist plots of the different CNF@CoS$_2$ hybrids obtained under the same overpotential of 170 mV. $R_{\text{ct}}$ represents the charge transfer resistance, which corresponds to the semicircle of the Nyquist plots. It can be clearly observed that CNF@CoS$_2$-3 shows the lowest $R_{\text{ct}}$ of 9.9 $\Omega$, in comparison with 12.5 $\Omega$ for CNF@CoS$_2$-9 and 22.9 $\Omega$ for the CNF@CoS$_2$-1 hybrid. Measured under the same proton reduction potential, a lower $R_{\text{ct}}$ value reflects a more effective electron transfer in the CNF@CoS$_2$-3 hybrid, leading to more efficient hydrogen evolution. EIS responses of the CNF@CoS$_2$-3 hybrid at various overpotentials were also tested, as shown in Fig. S7.† $R_{\text{ct}}$ of 28.9 $\Omega$, 9.9 $\Omega$ and 6.4 $\Omega$ under overpotentials of 120, 170 and 220 mV were found, respectively. This reveals that with increasing overpotential, more efficient hydrogen generation takes place, along with a more rapid electron transfer.

Further comparison of the HER performance between pure CNF, CoS$_2$, the CNF@CoS$_2$-3 hybrid and commercially available Pt catalyst was also made. As shown in Fig. 5A, the Pt catalyst exhibits a superb HER activity, with a near zero onset potential and a large current density, while pure CNF hardly displays any catalytic performance. Although pure CoS$_2$ agglomerates exhibit a non-negligible electrocatalytic performance, the onset potential and current density are far inferior to those of the CNF@CoS$_2$-3 hybrid, which is due to its serious aggregation inhibiting the efficient exposure of its electrocatalytic sites. As confirmed from the analysis of $C_{\text{dl}}$, the catalytically active surface area of CNF@CoS$_2$-3 (17.5 mF cm$^{-2}$ in Fig. 4B and C) is over ten times larger than that of pure CoS$_2$ (1.3 mF cm$^{-2}$ in Fig. 5B and C). Such a huge difference between the hybrid and pure electrocatalytic materials strongly supports the importance of constructing a hierarchical structure with exposed HER active sites on a nanoscale size. The corresponding Tafel plots are shown in Fig. 5D. As is well known, the reaction mechanism of a catalyst can be deduced from its Tafel slope, which is determined from the inherent properties of the catalyst. For CNF@CoS$_2$-3 measured in 0.5 M H$_2$SO$_4$, the Tafel slope is calculated to be 66.8 mV per decade, suggesting that the HER activity occurs through a Volmer–Heyrovsky mechanism, in which fast discharge of a proton takes place, followed by a rate-limiting electrochemical recombination with an additional proton. Moreover, it should be noted that the Tafel slope of the CNF@CoS$_2$-3 hybrid is lower than that of pure CoS$_2$ (86.8 mV per decade), indicating a higher HER efficiency. This result can be further explained using the $R_{\text{ct}}$ from the EIS analysis, as shown in Fig. 5E. On the one hand, the semicircle of pure CoS$_2$ is far larger than that of the CNF@CoS$_2$-3 hybrid, demonstrating a higher charge transfer resistance. On the other hand, the solution resistance ($R_s$), a value that can be obtained from the intersection of the curves at the real axis in the range of the high frequency region, displays the same tendency (8.2 $\Omega$ for the CNF@CoS$_2$-3 hybrid and 15.6 $\Omega$ for pure CoS$_2$). In terms of the cycling stability in 0.5 M H$_2$SO$_4$ solution, the CNF@CoS$_2$-3 hybrid displays a slight fluctuation in HER performance, with the response current first increasing over 2000 s, and then decreasing later (Fig. 5F). Consequently, the morphology of the CNF@CoS$_2$-3 hybrid after cycling for 2000 s was characterized, as shown in Fig. S8.† It can be observed that no

![Fig. 5](image-url)
obvious change occurs in the morphology after cycling for 2000 s. The increase in HER activity may be attributed to the activation process at the beginning, and this similar phenomenon has been reported in previous studies. For the later decrease in current density, a possible reason may be due to the partial loss of the electrocatalytically active sites of the CNF@CoS$_2$-3 hybrid during the violent bubble accumulation and release process, which can be further proved from its morphology after the whole cycling test (Fig. S9†). However, it is worth mentioning that despite the partial collapse, the CoS$_2$ nanoparticles still generally well-covered the surface of the CNF nanofibers after the cycling test, demonstrating that the CNF carrier plays an important role in stabilizing the structure. In contrast, pure CoS$_2$ particles display a huge decline in electrocatalytic activity (Fig. S10†). All the above results clearly verify that the HER performance of the hybrid materials is better than that of the pure aggregate, demonstrating the rational design of the hybrid product, whose electrocatalytic performance is also remarkable when compared to some previous studies in acid media, as listed in Table 1.

The HER performance of the CNF@CoS$_2$-3 hybrid was further examined in 1.0 M KOH solution and 1.0 M PBS solution to evaluate its activity as an all-pH-range electrocatalyst (Fig. 6). In alkaline medium, the CNF@CoS$_2$-3 hybrid exhibits an onset overpotential of 130 mV, accompanied by significant hydrogen evolution ($i = 10.0$ mA cm$^{-2}$) at a voltage of $-207$ mV. Furthermore, the Tafel slope of the CNF@CoS$_2$-3 hybrid (113.3 mV per decade) is also relatively smaller than that of pure CoS$_2$ (172.5 mV per decade) in KOH solution, implying a faster HER process with higher efficiency. When applied in neutral conditions, it could be easily found that all the electrocatalysts have a much poorer activity than those in acid and alkaline electrolytes. The CNF@CoS$_2$-3 hybrid catalyst exhibits an onset potential of $-130$ mV, a Tafel slope of 163.7 mV per decade, and a current density ($i$) of 10.0 mA cm$^{-2}$ at an overpotential ($\eta$) of 360 mV. In addition, the working stability is another important criterion when making an excellent electrocatalyst. Fig. 6C and F depict the current–time ($I$–$t$) curves of the CNF@CoS$_2$-3 hybrid catalyzed HER under a constant overpotential of 0.27 V in KOH and |Inorg. Chem. Front., 2016, 3, 1280–1288| This journal is © the Partner Organisations 2016 Published on 16 August 2016. Downloaded on 03/12/2016 10:31:57.

![Fig. 6](image-url) (A, D) LSV polarization curves for different material modified GCE in 1.0 M KOH and 1.0 M PBS. (B, E) Tafel plots for Pt, CoS$_2$, the CNF@CoS$_2$-3 hybrid modified GCE in 1.0 M KOH and 1.0 M PBS. (C, F) Time dependence of the current density for the CNF@CoS$_2$-3 hybrid modified GCE recorded at $-0.27$ V and $-0.50$ V versus RHE in 1.0 M KOH and 1.0 M PBS, respectively.

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$\eta_{10}$: overpotential ($\eta$) for current density ($j$) of 10.0 mA cm$^{-2}$; $\eta_{20}$: $\eta$ for $j = 20.0$ mA cm$^{-2}$; $\eta_{100}$: $\eta$ for $j = 100.0$ mA cm$^{-2}$.
0.50 V in PBS, respectively, which also exhibit good cycling performance. Moreover, taking further insight from the i-t curves obtained using different electrolytes, it can be seen that the serrated waves of the i-t curve obtained in PBS solution are more violent than those in H_2SO_4 and KOH solutions, indicating that the CNF@CoS_2:3 hybrid is less effective in driving H_2 (g) bubble release in neutral media, which can be attributed to different HER mechanisms in different electrolyte media. In addition, it is speculated that these accumulated H_2 (g) bubbles can generate an effective protective layer separating the electrocatalyst from the neutral electrolyte, thus periodically hindering the dissolution or collapse of the catalyst, leading to an enhanced stability compared with that in acid and alkaline electrolytes.

4. Conclusions

In summary, CNF@CoS_2 core/sheath hybrids with different morphologies were prepared through a combination of electrospinning and hydrothermal methods for use as HER electrocatalysts in all-pH-value conditions. A hierarchical structure is facilely constructed with nanosized CoS_2 particles evenly distributed on the CNFs, thus effectively preventing self-agglomeration of electroactive CoS_2. In addition, the attached CoS_2 nanoparticles display different morphologies on tuning the precursor concentration, but their loading amounts remain the same. The differences in morphology have a huge impact on the electrocatalytic performance, which can be attributed to the differences in the exposure of the efficient electroactive sites. Owing to the synergistic effects between the 3D CNF carrier and the electrocatalytically active CoS_2 nanoparticles, the optimal CNF@CoS_2 hybrid exhibits an excellent all-pH HER performance, with a low onset potential of −40 mV, a small Tafel slope of 66.8 mV per decade and a large current density (10.0 mA cm\(^{-2}\)) at \(\eta = 110\) mV in acid media, as well as remarkable activities in alkaline and neutral media.

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Notes and references