

Cite this: *J. Mater. Chem. A*, 2019, 7, 7553

Hierarchical MoS₂–carbon porous nanorods towards atomic interfacial engineering for high-performance lithium storage†

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Hierarchical nanostructures have attracted considerable attention for rechargeable battery systems since they combine the benefits of size effects induced by nanoscaling with the integrity of bulk materials. Despite significant progress, the hierarchical structures reported so far are designed only down to the nanoscale. To improve the battery performance, downsizing the designed building blocks of the hierarchical structure to smaller scales (molecular or even atomic level) is essential. This novel concept has been realized in a MoS₂/C composite system, where MoS₂ and N-doped carbon molecular layers are alternately stacked to form nanosheet building blocks, which are further assembled into a porous nanorod structure. This hierarchical heterostructure converts the guiding principle of sub-nanoscale engineering into practice, aiming at increasing the interfaces between MoS₂ and carbon towards the largest possible molecular contact level. The resultant MoS₂/N-doped carbon porous nanorods (MoS₂/NC-PNR) electrode exhibits outstanding performances in lithium-ion batteries including high initial discharge capacity of ~1300 mA h g⁻¹, cycling stability for 700 cycles and excellent rate performance (443 mA h g⁻¹ at 10C). The outstanding performance of the MoS₂/NC-PNR superstructure illustrates the enormous potential of the hierarchically designed 2D compounds from molecular layer level, which could be extended to other layered materials.

Received 20th December 2018
Accepted 1st March 2019

DOI: 10.1039/c8ta12293h

rsc.li/materials-a

Introduction

Molybdenum disulfide (MoS₂) is one of the most stable and multifunctional members of the transition metal dichalcogenide family and hence promises a wide range of applications, including hydrogen evolution,^{1,2} solar cells^{3,4} and rechargeable

batteries.^{5–8} Its tuneable layered structure and high theoretical lithium storage capacity of 670 mA h g⁻¹ render it a great candidate as anode material in Li-ion batteries (LIBs).^{9,10} However, the semi-conductive nature and large volume changes during the conversion reaction process, which cause disintegration of the structure, lead to very limited cycling stability and poor rate capability.¹¹ Furthermore, the involved redox couple Li₂S/S suffers from the shuttle effect of polysulfides, which causes the loss of active material, significantly reducing the specific capacities as well.¹²

Rational design of nanoscale, hierarchical MoS₂ composite structures has been proven to be an effective route to tackle the abovementioned problems.^{13–16} A hierarchical structure both exploits the size effects of the bare nanoscaled building blocks and maintains the integrity of the bulk material to some extent. On the one hand, the Li⁺ diffusion pathways are shortened due to the size effect. On the other hand, a relatively loose arrangement of the individual building blocks can facilitate electrochemical reactions by providing increased contact area between active material and electrolyte.¹⁷ In this context, an adequate backbone structure is needed to guarantee the stability of the composite by buffering mechanical strain during electrochemical cycling. In case of MoS₂-based composites, usually carbon layers are introduced to increase the overall

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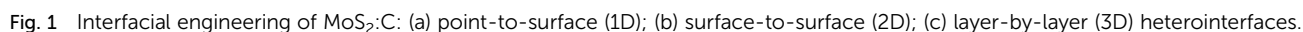
† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ta12293h

In this work, we apply a facile bottom-up method to synthesize $\text{MoS}_2/\text{N-doped carbon porous nanorods}$ ($\text{MoS}_2/\text{NC-PNR}$), targeting the hierarchical superstructure to achieve high performance lithium storage. In the reported material, similar hexagonal structures enable alternating stacking of MoS_2 molecular layers and N-doped carbon layers, which promotes the largest possible contact interfaces between the active material (MoS_2 layers) and the buffer layers (carbon). In addition, such ordered stacking yields an expanded interlayer distance as compared to bulk hexagonal MoS_2 (2H- MoS_2). In this architecture, the superior properties of both MoS_2 and N-doped carbon are fully maintained while a synergetic effect due to a molecular layer level homogeneous distribution is introduced, potentially leading to ultra-fast electron and ion transfer.²⁸ Furthermore, the interoverlapped MoS_2/C nanosheets serve as building blocks to construct a hierarchical rod-like structure which is robust enough to endure the repeated de-/lithiation processes and offers a large number of electrochemically active sites as well. These merits endow $\text{MoS}_2/\text{NC-PNR}$ as anode material in LIB with greatly enhanced lithium storage performance. Importantly, this materials design strategy has great potential to be extended to other 2D material systems, thereby realizing high performance hierarchical materials for other energy related applications.

Results and discussion

Synthesis of the MoS₂/NC-PNR

As illustrated in Fig. 2, the MoS₂/NC-PNR superstructure was prepared *via* a modified template method. Firstly, α -MoO₃ nanorods with around several μ m long and 200 nm in diameter (see SEM images in Fig. S1a†) were mixed into a dopamine (DOPA)/TRIS solution to form a Mo(VI) – DOPA complex (MDC), which served as self-sacrifice template. The formation of this complex is indicated by the initially colourless DOPA/TRIS solution changing to a light brown suspension, immediately after the addition of the α -MoO₃ nanorods (see Fig. S1c†). During complexation of the MDC, the morphology (Fig. S1b†) and orthorhombic crystal structure (Fig. S1d†) of the α -MoO₃ nanorod template are maintained. However, the UV-vis spectra of α -MoO₃, DOPA and MDC reveal a different electronic



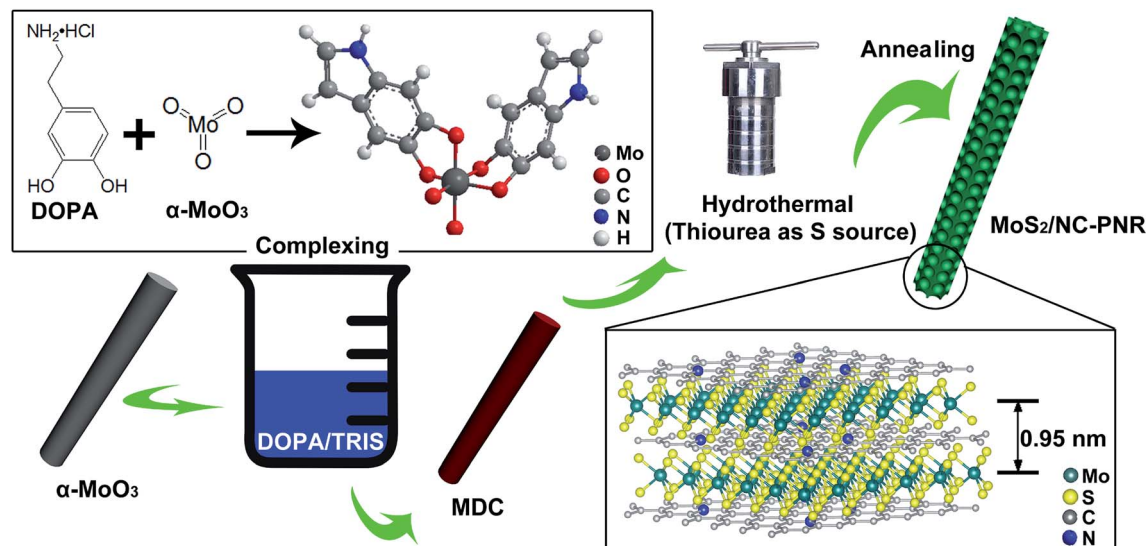


Fig. 2 Schematic illustration of MoS₂/NC-PNR formation.

structure in MDC compared to α -MoO₃ due to different chemical environments. In Fig. 3a, a broad absorption peak at about 315 nm, which corresponds to bulk-type molybdate species,^{29,30} is observed in the α -MoO₃ spectrum but disappeared in the MDC spectrum. In the latter spectrum, an absorption peak at 280 nm presents in the same position as that of the DOPA spectrum, which originates from the L_a-L_b coincident transition of DOPA molecules³¹ due to an excess amount of DOPA in the complexation suspension. High-resolution N 1s spectra of DOPA (Fig. 3b) and MDC (Fig. 3c), collected using X-ray photoemission spectroscopy (XPS), show the presence of different amines in the compounds. In case of DOPA, two species at binding energies of 401.8 eV and 399.2 eV contribute to the signal and are assigned to primary and secondary amine, respectively.³² Although the latter indicates the polymerization of DOPA to a certain degree, the main peak at 401.8 eV confirms that the dominant species is monomeric. In the N 1s spectrum of MDC, four separate peaks at 401.7 eV, 399.5 eV, 398.6 eV, and

396.5 eV are observed, which are attributed to primary, secondary and tertiary amine, and molybdenum (Mo 3p_{3/2}), respectively. The large share of tertiary amine, which is associated with tautomeric species of the intermediates, can be explained by electron donation from DOPA to Mo(vi), resulting in reduced electron density in the DOPA part.³³ These results confirm the formation of the MDC complex as an essential intermediate to finally synthesize the layer-by-layer structures.

In order to obtain the final product, the MDC complex was hydrothermally treated with thiourea as the sulfur source followed by an annealing procedure. In the early stage, self-polymerization of DOPA is supposed to take place with the emission of NH₃ from thiourea which forms a mild basic environment (pH = 10.2 after hydrothermal reaction).³⁴ In the meantime, Mo(vi) in the MDC complex is reduced to MoS₂ with the help of H₂S, growing *in situ* on the surface of polydopamine (PDA). Thus, the hydrothermal reaction yields MoS₂/polydopamine porous nanorods (MoS₂/PDA-PNR), which were then

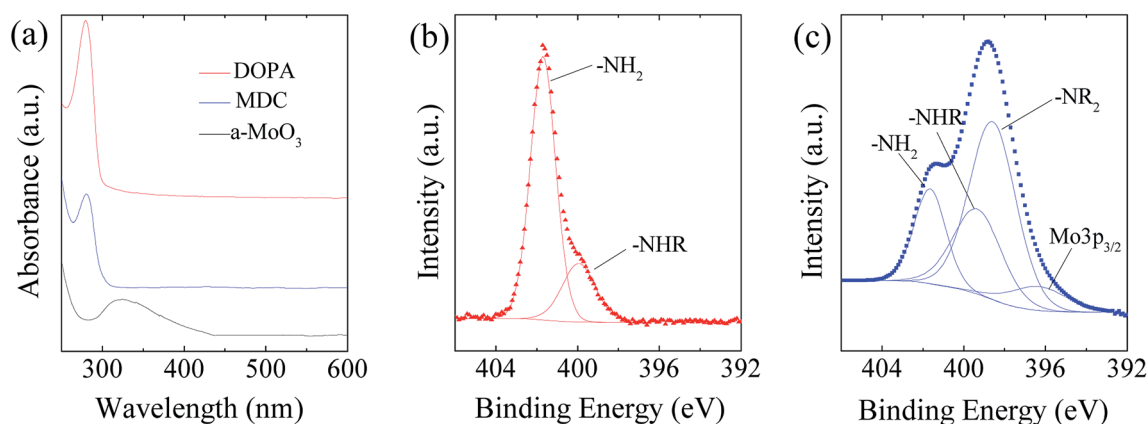


Fig. 3 (a) UV-vis spectra of α -MoO₃, DOPA and MDC; spectra are offset for clarity. (b and c) XPS N 1s spectra of DOPA and MDC, respectively. -NH₂, -NHR and -NR₂ indicate primary, secondary and tertiary amines.

sintered at 900 °C under Ar atmosphere to prepare the final product MoS₂/NC-PNR. Altogether, this procedure results in the formation of a layer-by-layer MoS₂/NC structure while on the larger length scale the porous nanorod morphology is maintained.

Characterization of the interoverlapped structure

Analysis of the chemical composition of the resultant MoS₂/NC-PNR superstructure was carried out by means of XPS, displayed in Fig. 4a and S2.† The survey spectrum (Fig. S2a†) shows distinct peaks of Mo, S, C, N and O in which the molar ratio of S : Mo equals 2.40 ± 0.01 . The discrepancy to the stoichiometry of MoS₂ can be explained by surface defects of the nanocomposite.³⁵ A high-resolution scan of the Mo 3d range (Fig. 4a left) shows a doublet at 229.2 eV and 232.7 eV, which is attributed to the Mo⁴⁺ in MoS₂. A minor contribution from MoO_x is also observed. The S 2p spectrum (Fig. S2b†) consists of a single doublet at binding energies of 162.3 eV and 163.6 eV, which relates to the S 2p_{3/2} and 2p_{1/2} orbitals of S²⁻.³⁶ In the C 1s spectrum (Fig. 4a mid), a dominant C=C signal at 284.5 eV and traces of a C=O one at 287.4 eV are observed, indicating low oxygen content in the carbon layers. Furthermore, the C-N species at 285.8 eV accounts for 14.1% of all carbon, revealing the N-rich nature of the carbon component.³⁷ The N 1s

spectrum (Fig. 4a right) is composed of a Mo 2p_{3/2} peak and three different N species, *i.e.*, graphitic, pyrrolic, and pyridinic, suggesting a successful doping of N into the carbon layer.³⁸ The Raman shift spectrum of MoS₂/NC-PNR in Fig. 4b further confirms the chemical composition of the hierarchical superstructure. At lower wave numbers, the peaks present at 383.43 and 404.71 cm⁻¹ correspond to the in-plane E_{2g}¹ and out of plane A_{1g} mode of MoS₂, respectively.³⁹ It is worth noting that the A_{1g} peak is more pronounced than the E_{2g}¹ peak, indicating a more edge-terminated and less layered structure.⁴⁰ Actually, the frequency difference of the two peaks $\Delta k = 21.28 \text{ cm}^{-1}$ is close to the value of single layer MoS₂ reported in the literature³⁹ ($20.2\text{--}21.2 \text{ cm}^{-1}$), suggesting that single layer MoS₂ is featured in the MoS₂/NC-PNR composite. At higher wave numbers, characteristic peaks at 1369 and 1598 cm⁻¹ correspond to disordered carbon (D-band) and ordered graphitic carbon (G-band), respectively, with a peak intensity ratio of approximately 1 : 1. The pronounced D-band indicates a defect rich nature of the N-doped carbon layers.

The TGA data (Fig. 4d) of both MoS₂/NC-PNR and bulk MoS₂ illustrate oxidation of MoS₂ to MoO₃, accompanied by a theoretical mass loss of 10.1 wt%. The bigger total mass loss in case of MoS₂/NC-PNR confirms a significant carbon content, which is determined to 25.8 wt%, assuming complete C combustion.

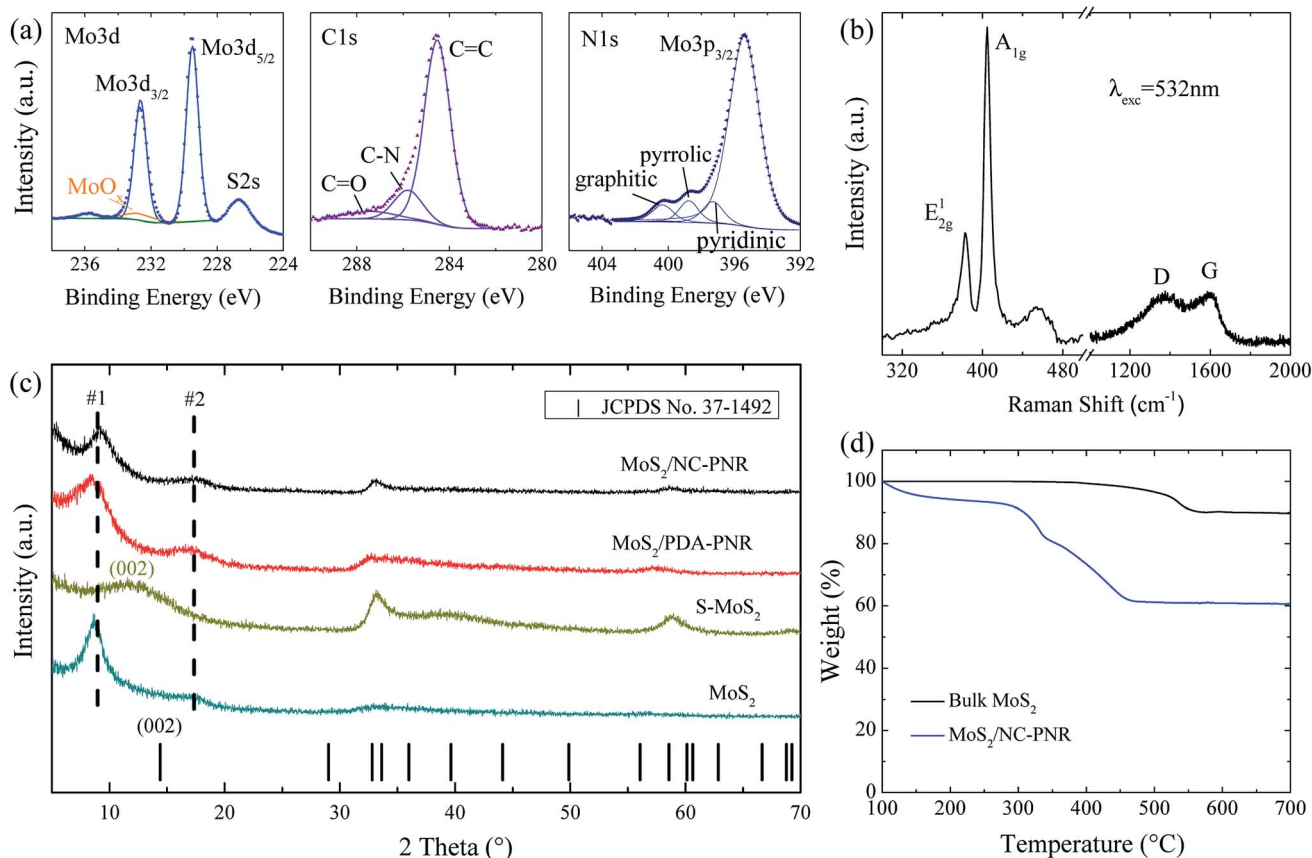


Fig. 4 (a) XPS Mo 3d, C 1s and N 1s spectrum of MoS₂/NC-PNR; (b) Raman spectrum of the MoS₂/NC-PNR excited at 532 nm; (c) XRD patterns of MoS₂/NC-PNR, MoS₂/PDA-PNR, S-MoS₂ and MoS₂. Vertical ticks display the standard Bragg peaks of MoS₂ (JCPDS no. 37-1492); (d) TGA curve of MoS₂/NC-PNR and of bulk MoS₂.



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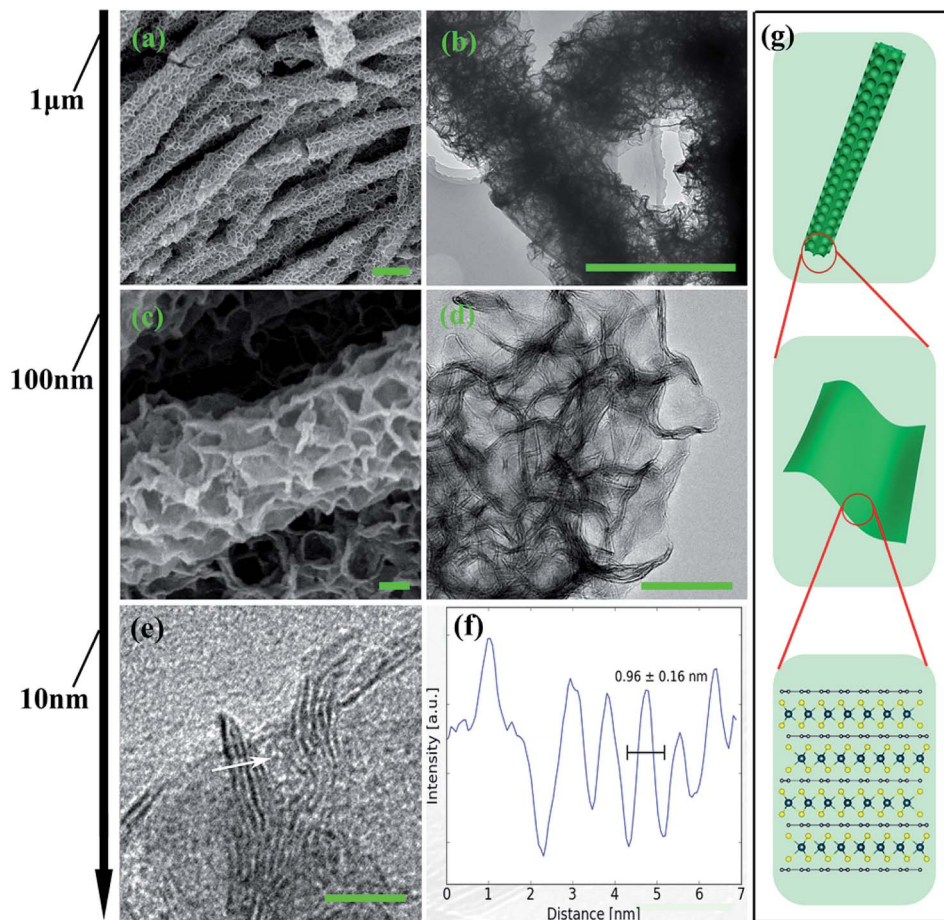


Fig. 5 Structural organisation of MoS₂/NC-PNR on several length scales: (a, c) SEM images (scale bar 1 μm), (b, d) TEM images (scale bar 100 nm), (e) HR-TEM images (scale bar 10 nm), (f) line scan of the white arrow in (e) and (g) schematic illustration of the hierarchical structure.

and a moderate shift of the oxidation peak up to 2.31 V in cycle 10. Upon continued cycling, further electrochemical activity is identified by means of several reductive and oxidative features in the voltage ranges of 0.2–1.6 V (cathodic) and 1.1–1.9 V (anodic). They may, *i.e.*, originate from the de-/lithiation of an amorphous Mo/Li₂S matrix, including the re-formation of MoS_x. The N-doped carbon in the superstructure can also accommodate Li⁺, which is indicated by a distinct stable reduction feature at 0.01 V.

The charge–discharge profile of a MoS₂/NC-PNR-based electrode upon galvanostatic cycling at a current density of 0.5C (1C = 670 mA g^{−1}) is shown in Fig. 7b. In agreement with the CV curves, there are two plateaus at ~1.0 and ~0.7 V during the first discharge process referring to the different lithium storage processes, while only one prominent plateau appears at ~2.2 V during the first charge process. From the second cycle on, the dominant plateaus at 2.0/2.25 V as well as the slopes below 2.0 V, corresponding to the Li₂S/S redox pair, show quite reversible behavior, which is superior to usually reported Li–S batteries.⁴⁶ The observed high reversibility may be promoted by the N-doped carbon layers and by the appearance of conductive Mo nanoparticles.¹² The stacked N-doped carbon layers could offer physical confinement to the active species and provide

more electroactive sites for the strong chemisorption of lithium polysulfides. The Mo nanoparticles derived from the MoS₂ reduction boost the electrochemical reaction, further hindering the polysulfide shuttle effect. As a result, the corresponding cycling performance in Fig. 7c presents a stable discharge capacity of ~800 mA h g^{−1} for up to 150 cycles after initial decay from 1086 mA h g^{−1}. The initial capacity loss is mainly attributed to the partially irreversible conversion reaction and the formation of solid-electrolyte interphase (SEI).⁴⁷ The durable cycling capability of the hierarchical superstructure is highlighted by the comparison with a S-MoS₂ electrode, whose discharge capacities drop noticeably after 40 cycles before stabilizing at around underwhelming 30 mA h g^{−1}. The poor cycling stability of S-MoS₂ can be attributed to structural changes of the active MoS₂ material as well as the shuttle effect of *in situ* formed (poly)sulfide from the 2nd cycle on.⁴⁸ However, the MoS₂/NC-PNR superstructure design specifically aims at maintaining the high specific capacities by maximizing the interfacial contact area between active material and interoverlapped carbon, which may buffer the mechanical strain to the largest extent possible.²³ In the SEM images of electrodes after 150 cycles (Fig. S5†), the rod-like morphology of MoS₂/NC-PNR is partially preserved, while in case of the S-MoS₂ material only



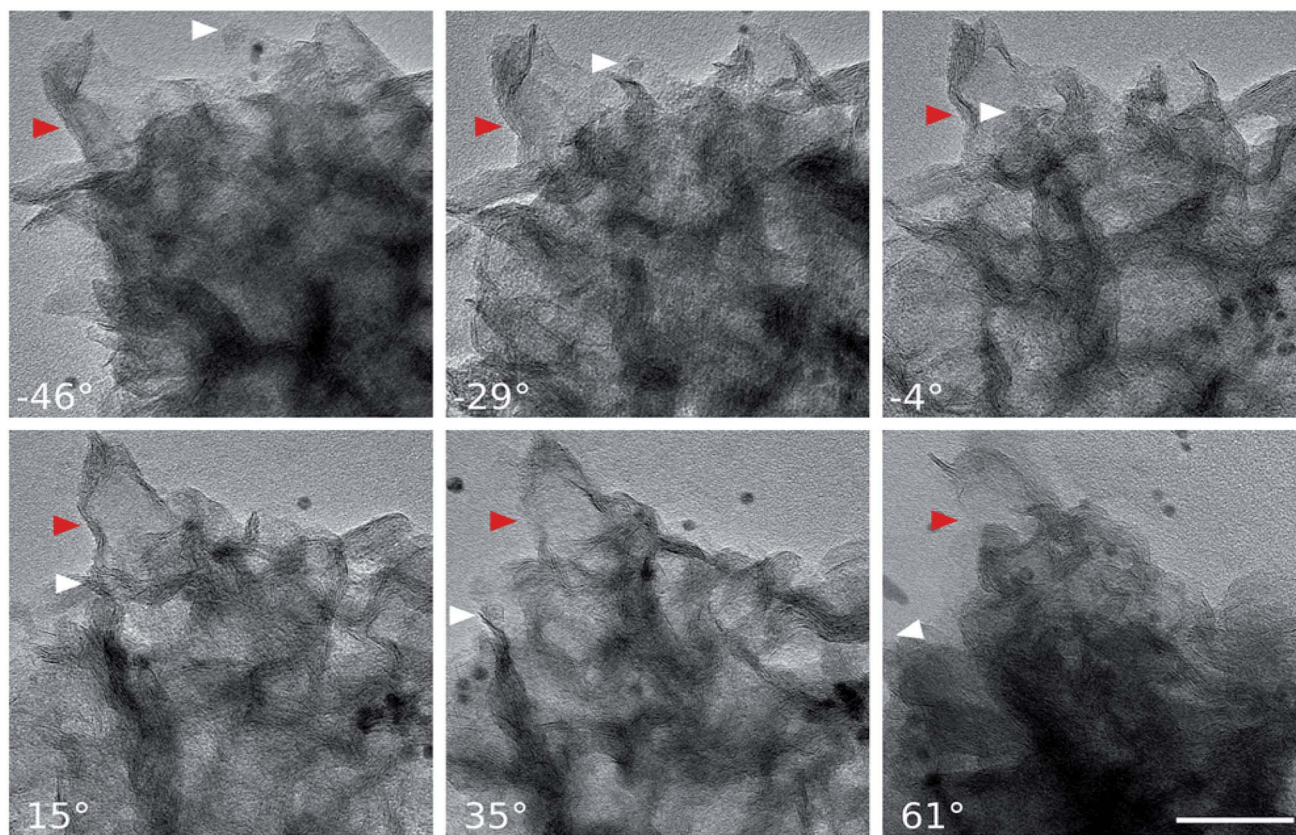


Fig. 6 HR-TEM images: selected frames from a movie (Movie S1†) showing a tilt series. The arrows indicate the same part of the nanorod as it is being tilted. While -46° to 15° show the lines of the stacking, indicated by the red arrow, the sample is tilted in such a manner at 35° and 61° , that the projection goes straight through the layers (scale bar: 50 nm).

irregular nanoparticles remained. Further post mortem characterizations of the $\text{MoS}_2/\text{NC-PNR}$ electrode demonstrate the buffer function of the N-doped carbon layers to the amorphous products. In Fig. S6a,† the XRD patterns of the $\text{MoS}_2/\text{NC-PNR}$ electrode after cycling suggest completely non-crystallized feature of the MoS_2 structure. The HAADF-STEM image of the cycled electrode and the corresponding SAED patterns in Fig. S7a–c† also present a disordered structure of the Mo species. In the SAED patterns, the only weak diffraction rings can be indexed as disordered graphitic carbon. The EDX together with the STEM images (Fig. S7d–f†) indicate a homogeneous Mo distribution in the carbon matrix. All the results highlight the significant role of carbon in maintaining the structural integrity of the electrode. Therefore, the $\text{MoS}_2/\text{NC-PNR}$ electrode exhibits an ultra-long cycling lifetime of 700 cycles at a current of 2C, delivering a stable capacity of $\sim 520 \text{ mA h g}^{-1}$ as shown in Fig. 7e. In addition, the N-doped carbon layers compensate the intrinsic low conductivity of MoS_2 in $\text{MoS}_2/\text{NC-PNR}$, allowing faster electron transfer during the dis-/charge process.⁴⁹ As a consequence, the $\text{MoS}_2/\text{NC-PNR}$ electrode shows much improved rate capability as compared to the S- MoS_2 electrode (Fig. 7d). The former exhibits an initial discharge capacity of 1294 mA h g^{-1} at 0.1C, stabilizing at 810, 720, 636, 575, and 507 mA h g^{-1} upon increasing the current density to 0.2, 0.5, 1, 2, and 5C, respectively. All of the values are

higher than that of the latter. Especially at high current density of 10C, the $\text{MoS}_2/\text{NC-PNR}$ electrode presents a large reversible capacity of 443 mA h g^{-1} , which is twice the value of the S- MoS_2 electrode.

In order to clarify the enhanced electrochemical kinetics of the $\text{MoS}_2/\text{NC-PNR}$ electrode, further CV measurements and electrochemical impedance spectroscopy (EIS) have been carried out. The CV curves at different scan rates (see Fig. 8a,) show similar shape, which indicates excellent adaptability even at large scan rates. The peak current (I) and the scan rate (ν) obey a power law $I = a\nu^b$, from which both the capacitive and the diffusion controlled contributions of the charge storage process can be evaluated.⁵⁰ Fitting the data in Fig. 8b yields that the parameter b equals to 1.02, 0.86, 0.77 and 0.88 for the cathodic and anodic peaks (C1, C2, A1, A2), respectively. The values suggest that capacitive processes (mainly the redox pseudocapacitive contribution) are dominant in the associated electrochemical reactions. The exact (pseudo)capacitive and diffusion controlled contributions can be assessed by using the equation $I = k_1\nu + k_2\nu^{1/2}$ (see Fig. S8†).⁵¹ In Fig. 8c, the (pseudo)capacitive contribution is proportional to the scan rate, reaching 85% at 2 mV s^{-1} . The high (pseudo)capacitive contribution suggests fast redox reactions in the $\text{MoS}_2/\text{NC-PNR}$ electrode, which is straightforwardly attributed to the effective contact between MoS_2 and N-doped carbon as well as the porous nanorods



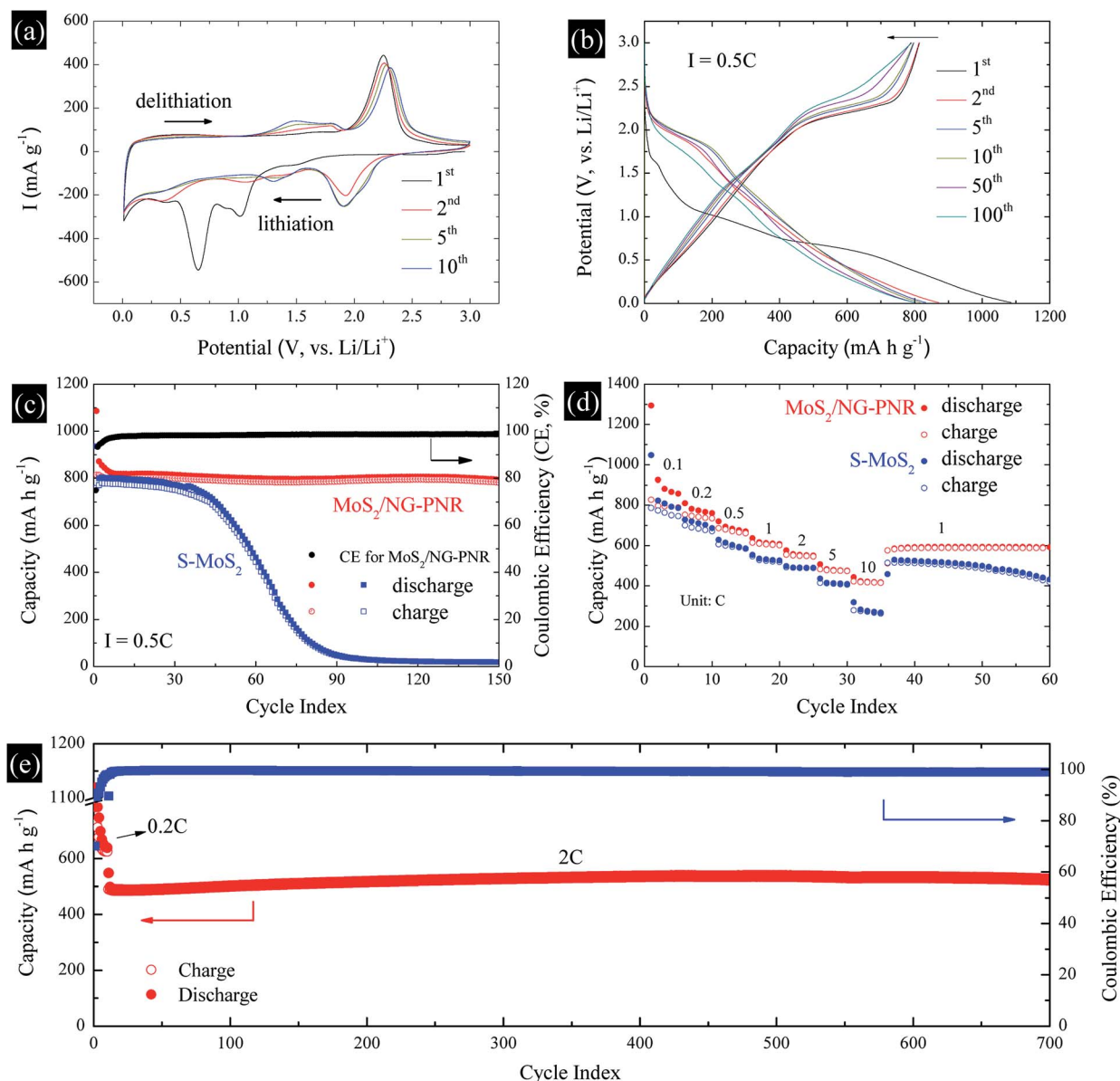


Fig. 7 Electrochemical performance of MoS₂/NC-PNR: (a) CV, (b) GCPL measurements at specific cycles, (c) cycling stability at 0.5C and the comparison with S-MoS₂, (d) rate capability and (e) long-term cycling stability at 2C (activated at 0.2C for 10 cycles).

architecture. This conclusion is further supported by EIS. In the Nyquist plots obtained at open circuit voltage (OCV) in Fig. 8d, both MoS₂/NC-PNR- and S-MoS₂-based electrodes show a depressed semi-circle in the high to medium frequency range as well as a sloping behavior in the low frequency range. The diameter of the semicircle, which correlates with the charge transfer resistance (R_2 in the corresponding equivalent circuit as shown in the inset of Fig. 8d), is only half the value in MoS₂/NC-PNR ($183 \pm 2 \Omega$) compared to S-MoS₂ ($342 \pm 1 \Omega$). Upon cycling, R_2 decreases abruptly in the MoS₂/NC-PNR electrode (Fig. S9†) which can be attributed to irreversible structural changes during the initial lithiation. Interestingly, the MoS₂/NC-PNR electrode exhibits few changes in subsequent cycles, which also holds for the low frequency domain, revealing a stable and reversible de/lithiation process. These results

imply not only improved conductivity of the pristine MoS₂/NC-PNR compound, but also more efficient ion transport pathways through the N-doped carbon layers of the superstructure.

The abovementioned results clearly imply that the hierarchical MoS₂/C composite with well-designed molecular contact interface ensures an enhanced electrochemical performance. This conclusion is further confirmed by comparing the present results with our previous work.²⁵ In ref. 27, we have reported C@MoS₂@C sandwiched hollow spheres with a surface-to-surface contact between MoS₂ and carbon layers. Interestingly, the carbon content in C@MoS₂@C sandwiched hollow spheres and the MoS₂/NC-PNR superstructures is almost identical (~26%). With the same chemical composition, the major differences between the two materials are the interfacial contact and morphological architecture. Due to its much more effective



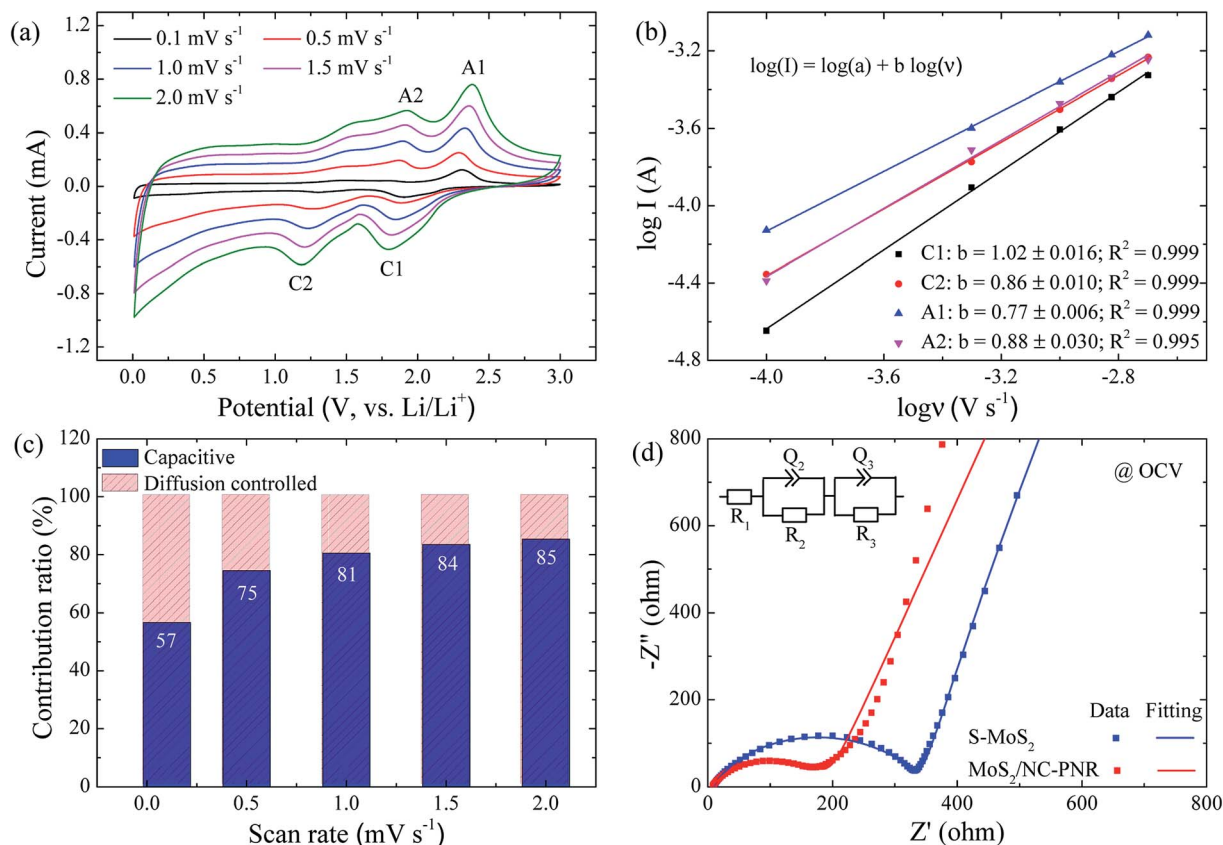


Fig. 8 Kinetic analysis of MoS₂/NC-PNR as LIB anode: (a) CV curves at different scan rate; (b) current response versus scan rate (log *I* vs. log *v*) plots at each redox peak; (c) capacitive and diffusion controlled contributions to the electrochemical processes at corresponding scan rate; the number labels the percentile capacitive contributions; (d) Nyquist plots and the fitting curves of MoS₂/NC-PNR and S-MoS₂ electrodes. The inset shows the equivalent circuit.

interfacial contact (layer-by-layer contact), the MoS₂/NC-PNR electrode exhibits well enhanced electrochemical performance as compared to the C@MoS₂@C electrode including higher capacities of ~800 mA h g⁻¹ vs. ~640 mA h g⁻¹ at 0.5C; longer cycling stability up to 700 cycles at 2C vs. 200 cycles at 1C; and a better rate capability of 507 mA h g⁻¹ vs. 382 mA h g⁻¹ at 5C. Actually, these values are comparable or even better than most of recently published MoS₂-based hierarchical structures as seen in Table S1† while our approach employs an advantageous environmentally friendly synthetic route and exploits a low cost, easy access carbon source. In general, the reported performances show that the rational structure design of nanocomposites is indispensable to their application potential and the present work provides a route to realize such a hierarchical structure down to the molecular layer level.

Conclusions

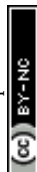
In summary, we report on a hierarchically structured nanocomposite with optimized atomic interfacial contact for application in lithium ion batteries. The MoS₂/NC-PNR superstructure has been prepared by means of a one-pot synthesis approach. The alternating stacking of the MoS₂ layers with intercalated N-doped carbon layers provides

a maximum specific contact area between the active material and the buffer layers, effectively alleviating the loss of active material due to pulverization of MoS₂ or dissolution of polysulfides. In addition, the buffer layers also act as a conductive matrix, enabling faster Li-ion and electron transfer during the de-/lithiation processes. At a larger length scale, the rod-like porous structure offers large specific surface area and abundant active sites for electrochemical reactions which also contributes to the dominant capacitive charge storage behaviour. As a result, the interfacially engineered MoS₂/NC-PNR superstructure exhibits a high specific capacity of 1294 mA h g⁻¹ at 0.1C, stable cycling performance up to 700 cycles at 2C, and a high rate capability of 443 mA h g⁻¹ at 10C. The ordered MoS₂/NC-PNR superstructure with well-defined interfaces is capable of buffering huge mechanical strain and offering efficient electron and ion transport pathways, which indicates a new way of rationally designing other hierarchical functional materials.

Experimental methods

Synthesis of α-MoO₃ nanorods

α-MoO₃ nanorods were prepared through a hydrothermal method according to the literature.⁵² 1000 mg ammonium



molybdate tetrahydrate (AMT, Sigma-Aldrich, 81.0–83.0% MoO₃ basis) was dissolved in 34 mL deionized water. 6 mL nitric acid (65 wt%) was added dropwise before stirring the AMT solution for 15 min. After stirring for another 15 min, the mixture was transferred to a 50 mL Teflon autoclave, heating at 180 °C for 24 h. The milky white precipitate was washed and collected for further use.

Synthesis of MoS₂/NC-PNR superstructure and S-MoS₂

MoS₂/NC-PNR was produced by adding α -MoO₃ nanorods into a dopamine/Tris solution and then stirring for 24 h. Then the precipitate was centrifuged and added into a thiourea solution under vigorous stirring. The homogeneously dispersed suspension was hydrothermally treated at 200 °C for 24 h. After the reaction, the black precipitate was washed with distilled water and ethanol for at least 3 times before vacuum drying at 80 °C. Finally, the black powder was sintered at 900 °C for 3 h under Ar atmosphere. For comparison, pure MoS₂ nanosheets were synthesized by using the same α -MoO₃ nanorods and same sulfur source at the same hydrothermal conditions and subsequent thermal treatment. The pure MoS₂ samples before and after sintering are labeled as MoS₂ and S-MoS₂, respectively.

Characterization

The morphology and microstructure of the sample was characterized by scanning electron microscopy (SEM, ZEISS Leo 1530) and transmission electron microscopy (TEM, JEM 2100F). TEM tilt-series images were acquired using a ZEISS Libra 200 MC Cs-Corr DMU Kronos with 200 kV acceleration voltage at highest magnification (pixel size: 0.578 Å) from –68° to +64° in 1° steps. 5 images were acquired at each step with exposure times of 1000 ms, which were later aligned and summed to increase resolution. The tilt-series was then aligned using the software provided by the IMOD package.^{53–55} For TEM measurement, the sample (powder) was dispersed in ethanol : water (2 : 1) mixture at 3 mg mL^{–1}. The dispersion was ultrasonicated for 30 minutes and then centrifuged at 14 800 rpm for 60 seconds. 1 μ L of the material that had collected at the bottom of the tube was then mixed with 30 μ L of water. 2.5 μ L of this mixture was dropped onto a copper TEM grid with carbon-coated Formvar film. Before the sample could dry, 2.5 μ L of a mixture of water and gold fiducials (6 nm in size), used for image alignment, was dropped onto both sides of the TEM grid. After waiting for one minute, excess liquid was carefully removed with filter paper.

X-Ray powder diffraction (XRD) was performed in Bragg–Brentano geometry (Bruker-AXS D8 ADVANCE ECO) applying Cu-K α_1 radiation ($\lambda = 1.54056$ Å). The step size $\Delta 2\theta$ was 0.02° with a scan speed of 0.5° min^{–1}. X-ray photoemission spectroscopy (XPS) was carried out in an ESCALAB 250Xi ultra-high vacuum system with an Al K α radiation source ($h\nu = 1486.6$ eV), 900 μ m spot size and 20 eV pass energy. Nitrogen physisorption measurements were performed at 77 K with a Gemini V (Micromeritics, Norcross, GA) after degassing the sample at 120 °C for 2 h. Brunauer–Emmett–Teller (BET) analysis from the amount of N₂ absorbed at various relative vapor pressures (six points

0.05 < P/P_0 < 0.3, nitrogen molecular cross-sectional area = 0.162 nm²) was used to determine the surface area. Barrett–Joyner–Halenda (BJH) analysis was done on the adsorption isotherms in order to obtain the pore-size distribution. Raman spectroscopy was performed on bulk powder samples using a Renishaw InVia microscope with 532 nm excitation lasers in air under ambient conditions. The laser power was kept below 100 μ W to avoid decomposition. Measurements on minimum five different spots were performed and showed the samples to be highly homogeneous under these measurement conditions. The spectra were baseline corrected and averaged after acquisition. UV-vis absorption spectra were measured with a Jasco UV-660 spectrophotometer in the range from 250 to 600 nm. The absorption of the substrate was subtracted as baseline correction. Thermogravimetric analysis was performed by using a TGA/DSC1 STARE System (Mettler Toledo) at a heating rate of 10 °C min^{–1} in air.

Electrochemical measurement

Electrochemical studies were carried out using Swagelok-type cells.⁵⁶ For both the pure MoS₂ and the MoS₂/NC-PNR electrodes, the active material, carbon black (SuperP, Timcal) and polyvinylidene fluoride (PVDF, Sigma-Aldrich, 99%) binder were mixed in anhydrous 1-methyl-2-pyrrolidinone (NMP, Sigma-Aldrich, 99%) with a weight ratio of 7 : 2 : 1. The slurry was pasted on a circular Cu plate (~10 mm in diameter) with a mass loading of 1 mg cm^{–2}, dried overnight under vacuum at 80 °C and pressed. The two-electrode Swagelok-type cells were assembled in an Ar atmosphere glove box using lithium foil as counter electrode and 1 M LiPF₆ in a 1 : 1 mixture of ethylene carbonate and dimethyl carbonate as liquid electrolyte (Merck LP30). Cyclic voltammetry and galvanostatic cycling of the cells were performed at 25 °C between 0.01 and 3.0 V *versus* Li/Li⁺ at various scan/current rates using a VMP3 multichannel potentiostat (Bio-Logic SAS). Electrochemical impedance spectroscopy (EIS) was carried out also by using the VMP3 multichannel potentiostat in the frequency range of 100 kHz to 0.1 Hz. The EIS data were analyzed with the help of the Z Fit function of the EC-Lab software (Bio-Logic).

Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors thank I. Glass for experimental support. Financial support by the CleanTech-Initiative of the Baden-Württemberg-Stiftung (Project CT3 Nanostorage) and by the IMPRS-QD is gratefully acknowledged. Z. L. acknowledges financial support



by the Chinese Scholarship Council, the Excellence Initiative of the German Federal Government, and by the Götze foundation.

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