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# A Typha Angustifolia-like MoS<sub>2</sub>/carbon nanofiber composite for high performance Li-S batteries

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A Typha Angustifolia-like MoS<sub>2</sub>/carbon nanofiber composite as both the chemically trapping agent and redox conversion catalyst for lithium polysulfides have been successfully synthesized via a simple hydrothermal method. By **applying** the Typha Angustifolia-like MoS<sub>2</sub>/carbon nanofiber as the interlayer for pure sulfur cathode, the cycling performance and coulombic efficiency have been improved significantly, **which the capacity degradation is only 0.09% per cycle and the coulombic efficiency can reach as high as 99%.**

**Keyword:** Li-S batteries, MoS<sub>2</sub>/BCF, electrocatalysis, chemically trapping, polysulfides

## INTRODUCTION

Lithium-sulfur (Li-S) batteries attracted considerable interests due to its high energy density (2600 Wh/kg) as well as the cathode material, sulfur, is cost-effective, natural abundant, and environment friendly (Gu and Lai, 2019). However, Li-S batteries are plagued with various challenges. Among those, the serious lithium polysulfides (LiPSs) shuttling, inducing large capacity degradation, severe polarization, sluggish reaction kinetics and inefficient self-discharge, is one of the most significant issues (Liu et al., 2019; Xu et al., 2019).

In view of such a serious situation, tremendous efforts have been made to suppress polysulfides shuttling through physical confinements and chemical adsorption by construction various kinds of nanostructures, such as the nonpolar porous carbon (Rehman et al., 2016; Guo et al., 2018), graphene (Yin et al., 2016), carbon nanotubes (Yang et al., 2018), as well as the polar metal oxides (Gu et al., 2016; Song et al., 2018), metal sulfides (He et al., 2019; Lin et al., 2019), metal carbide (Chen et al., 2018; Dong et al., 2018; Song et al., 2019), metal nitride (Jiao et al., 2019; Wang et al., 2019), etc. Accordingly, the LiPSs shuttling has alleviated to some extent. While recently, researchers focused on the electrocatalysis of reducing sulfur to LiPSs and oxidizing  $\text{Li}_2\text{S}_2/\text{Li}_2\text{S}$  to LiPSs even to sulfur during the charge-discharge process, which is important for achieving high reversible capacity and coulombic efficiency.

By applying the electrocatalysis concept of enhancing the redox reactions of **polysulfides**, increasing numbers of catalysts suitable for redox conversion of lithium **polysulfides** have been reported (Jeong et al., 2017; Liu et al., 2018; Hao et al., 2019; He et al., 2019; Jiao et al., 2019; Lin et al., 2019; Yuan et al., 2019).

In this work, we synthesized a new 1D **nanostructure**, Typha Angustifolia-like MoS<sub>2</sub>/carbon nanofiber composite as both the chemical **trapping agent** and redox conversion catalyst for **LiPSs** to enhance the sulfur cathode performances. The sulfur cathode with the MoS<sub>2</sub>/carbon nanofiber interlayer illustrate an initial capacity as high as 926.1 mAh/g at charge-**discharge** current of 0.5 C, even after 300 cycles, a reversible capacity of 661.5 mAh/g could maintain.

## **EXPERIMENTAL**

### **Materials Preparation**

Bamboo carbon fiber (BCF) preparation: the bamboo stick was immersed in 8M KOH solution and hydrothermal reaction for 12 h. Then the resulted bamboo fiber was dried and annealed at 800 °C for 2 h under Ar atmosphere. Finally, the BCF was obtained by washing with distilled water and drying overnight.

BCF/MoS<sub>2</sub> preparation: 114 mg Ammonium molybdate tetrahydrate ((NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>•4H<sub>2</sub>O) and stoichiometric overdose thiourea were dissolved in 60 mL distilled water, then 40 mg BCF dispersed in the

mixture solution by ultrasonication. Next the solution was transferred into the Teflon **autoclave** and reacted for 12 h at 200 °C. At that time, a black composite was obtained. After washing by the distilled water and ethanol and drying, the composite was annealed in H<sub>2</sub>/N<sub>2</sub> (5% volume percent of H<sub>2</sub>) atmosphere at 800 °C for 1 h to obtain the **finally** Typha Angustifolia-like BCF/MoS<sub>2</sub> composites.

### **Materials Characterizations**

The samples' structures were characterized by X-ray diffraction (XRD, (Model LabX-6000, Shimadzu, Japan) and JSM-7001F scanning electron microscope (SEM) (JEOL, Japan).

### **Electrochemical Measurements**

S, carbon black and polyvinylidene fluoride (analytical reagent, Sigma-Aldrich) in a weight ratio of 80:10:10, were mixed with solvent of 1-methyl-2-pyrrolidinone (analytical reagent, Sigma-Aldrich). After stirring for 12h, the electrode slurry was obtained. Then the slurry was pasted on the Aluminium foil via the blade-coating method. After drying at 60 °C in a vacuum oven overnight, the electrode was cut into wafers with a size of 0.5 cm<sup>2</sup> and a weight of approximately 1.5 mg. **The interlayer was made by BCF/MoS<sub>2</sub>, carbon black and polytetrafluoroethylene in a weight ratio of 80:10:10 with solvent of 1-methyl-2-pyrrolidinone to form a flexible film.** After drying at 60 °C in a vacuum oven overnight, the **film** was cut into wafers with a **diameter of**

11 mm, thickness of 150  $\mu\text{m}$  and a weight of approximately 1.2 mg.

Then batteries were assembled in a glove box (Vigor, China), using the lithium metal as the counter electrode, polypropylene (Celgard 2300) as the separator, and 1 M lithium bis(trifluoromethane)sulfonimide (LiTFSI) in 1,3-dioxolane/1,2-dimethoxyethane (DOL/DME) (1:1, v/v) containing 0.2 M  $\text{LiNO}_3$  as the electrolyte. And the BCF/ $\text{MoS}_2$  wafer could be placed between the separator and the electrode as the interlayer during the battery assembling process. Finally, the charge and discharge performances of the coin cells were tested with a LAND CT-2001A instrument (Wuhan, China) and the cyclic voltammetry (CV) curves were performed on a CHI 660D electrochemical workstation (CHI Instrument, Shanghai, China), both which the potential range was controlled between 1.5 and 3.0 V at room temperature. And the capacities were calculated based on the sulfur mass. Additionally, the electrode impedance spectrums (EIS) were tested on CHI 660E (frequency range from 100 kHz and 10 mHz).

## RESULTS AND DISCUSSIONS

Firstly, the XRD was used to examine crystallization structure of the synthesized product. As shown in Figure 1, The BCF/ $\text{MoS}_2$  has been successfully synthesized by using a simple hydrothermal method. From the XRD spectrum of BCF, there is a wide peak around at 2 theta of  $23^\circ$ , which belongs to the partial graphitization of carbon, implying the good

conductivity of BCF(Gu et al., 2015). While from the spectrum of BCF/MoS<sub>2</sub>, the peak belonging to the graphitization carbon has been covered by other strong peaks. All these peaks could be ascribed to the MoS<sub>2</sub>, and the crystal phase could match well with the MoS<sub>2</sub> stand PDF card (37-1492).

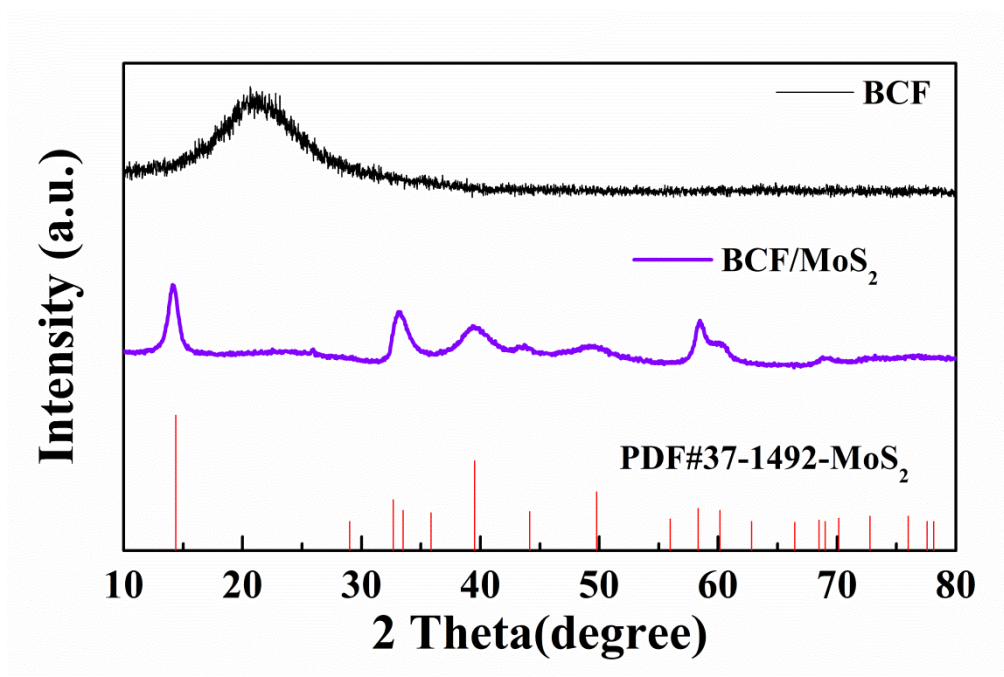


Figure 1 XRD spectra of BCF, BCF/MoS<sub>2</sub> and Stand XRD spectrum of MoS<sub>2</sub>.

Following the morphology information of BCF and BCF/MoS<sub>2</sub> have been investigated by SEM. As shown in Figure 2a, the bamboo carbon with **unique** fiber structure has successfully synthesized. While in Figure 2b and 2c, **the BCF as a core, and the MoS<sub>2</sub> grown in the direction of the nanofiber line as a shell has been observed.** Such a unique one-dimensional structure is very much like *Typha Angustifolia* as shown in Figure 2d.

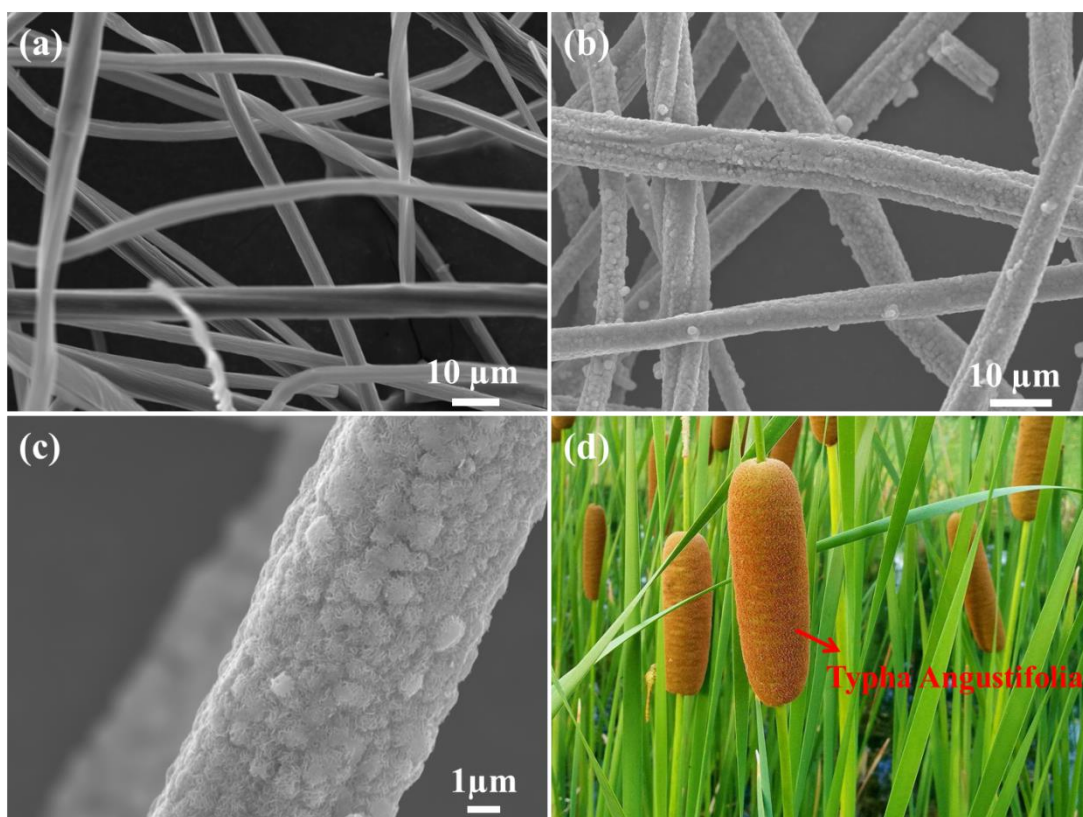


Figure 2 SEM images of (a) bamboo carbon fiber, (b-c) BCF/MoS<sub>2</sub> and (d) the digital photo of *Typha Angustifolia*.

Then the electrochemical performances of the sulfur cathode with and without the BCF/MoS<sub>2</sub> interlayer have been investigated. As shown in Figure 3a, there are two obvious and stable redox peaks for the sulfur cathode with BCF/MoS<sub>2</sub> interlayer. While in Figure 3b, pure sulfur electrode (BCF/MoS<sub>2</sub> interlayer) illustrate deformed and widened redox peaks in the CV curves, which suggests a sluggish kinetic process(Li et al., 2017;Liu et al., 2018). Comparing the peak potentials (Figure 3c) during the redox reactions, it is evident that the sulfur cathode with BCF/MoS<sub>2</sub> interlayer shows higher reduction potential and lower oxidation potential than that without BCF/MoS<sub>2</sub> interlayer, indicating that the BCF/MoS<sub>2</sub> interlayer **significantly** lowers the electrode



polarization(Gu et al., 2015;Wang et al., 2018;He et al., 2019). This can be attributed to the catalysis effect of MoS<sub>2</sub> on the oxidation/reduction of lithium polysulfides/Li<sub>2</sub>S(Wang et al., 2018;He et al., 2019). In terms of the onset potentials shown in Figure 3d, **the** onset potential of the sulfur cathode with BCF/MoS<sub>2</sub> interlayer in the oxidation reaction is  $\approx 2.23$  V, compared with  $\approx 2.21$  V for the pure sulfur cathode without BCF/MoS<sub>2</sub> interlayer. With respect to the reduction reaction, the onset potentials for sulfur cathode with BCF/MoS<sub>2</sub> interlayer are  $\approx 2.42$  and  $\approx 2.12$  V, compared with  $\approx 2.4$  and  $\approx 2.1$  V for the pure sulfur cathode without BCF/MoS<sub>2</sub> interlayer, which are lower by  $\approx 20$  mV. These results demonstrate that by inserting a conductive BCF/MoS<sub>2</sub> interlayer, the redox kinetics are accelerated and the polarization losses are also **significantly** reduced for the Li-S battery.(Gu et al., 2015;Li et al., 2017;He et al., 2019)

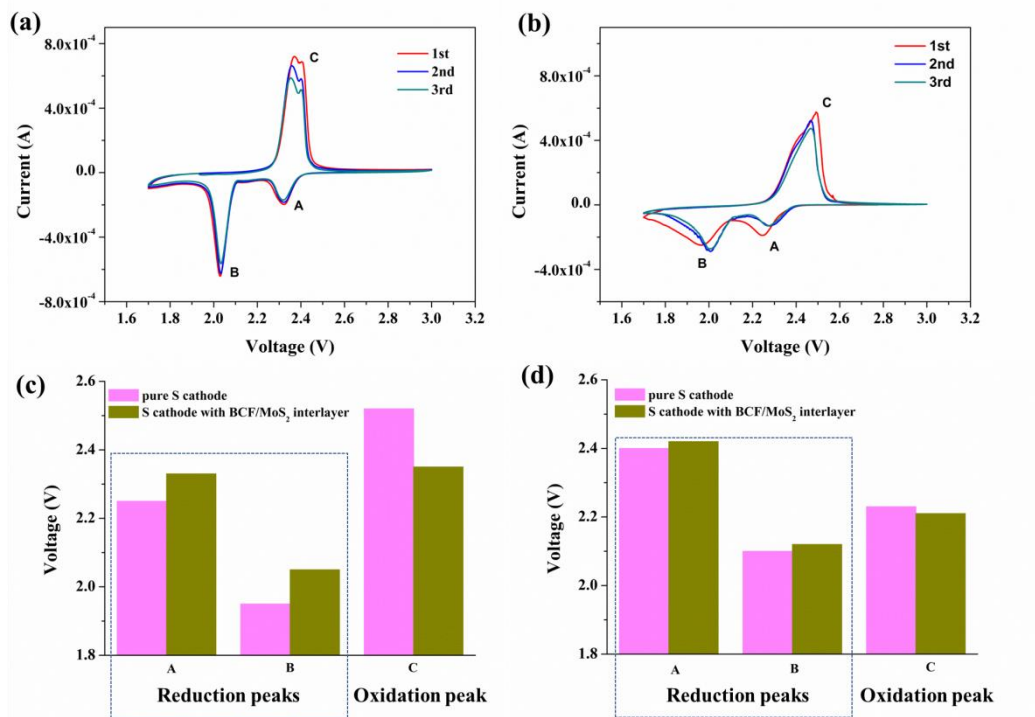


Figure 3 Kinetics of electrochemical reactions in Li-S batteries. CV test of (a) sulfur cathode with BCF/MoS<sub>2</sub> interlayer and (b) pure sulfur electrode. Corresponding (c) peak potentials and (d) onset potentials of the sulfur cathodes with and without BCF/MoS<sub>2</sub> interlayer from the first CV cycle in (a) and (b).

Finally, we carried out the long cycling performances and rate capabilities of the sulfur cathode with and without BCF/MoS<sub>2</sub> interlayer. As shown in Figure 4a, the sulfur cathode with interlayer shows a high initial specific capacity of 926.1 mAh/g. After cycling 300 cycles, it can still maintain a high reversible capacity of 661.5 mAh/g, and the capacity degradation rate is only 0.09% per cycle. However, the pure sulfur cathode without BCF/MoS<sub>2</sub> interlayer, only demonstrates an initial capacity of 510 mAh/g and extremely low reversible capacity of 56.3 mAh/g after 300 cycles. By contrast, the initial average discharge capacity of the pure sulfur cathode without interlayer is  $\approx$ 400 mAh/g

lower than the sulfur cathode with BCF/MoS<sub>2</sub> interlayer, indicating **significant** dissolution and loss of **LiPSs** into the electrolyte during the initial cycles, and such severe dissolution and loss continues throughout the whole charge and discharge process because the ultimate reversible capacity is also extremely low. **Additionally, from figure 4b, it is clearly observed that the sulfur cathode with BCF/MoS<sub>2</sub> interlayer shows far better rate capabilities compared to the one without BCF/MoS<sub>2</sub> interlayer. Even the charge-discharge current increases to 2 C, a reversible capacity of around 456 mAh/g could still be reserved, and after the current switch to low density of 0.2 C, a recoverable capacity of approximately 900 mAh/g could reach. Therefore, the BCF/MoS<sub>2</sub> is highly effective as polysulfide immobilizers to enhance the cycling life and rate capabilities(Gu et al., 2015).**

**What's more,** it can be observed that the sulfur cathode with BCF/MoS<sub>2</sub> interlayer demonstrate an excellent coulombic efficiency approximately 99%, but the sulfur cathode without interlayer shows an obvious weaker coulombic efficiency, particularly in the tens of cycles ahead. The coulombic efficiency results indicates that BCF/MoS<sub>2</sub> as electrocatalyst could significantly accelerate the redox reaction in Li-S batteries and improve the coulombic efficiency(Gu et al., 2015;Jeong et al., 2017;Wang et al., 2018).

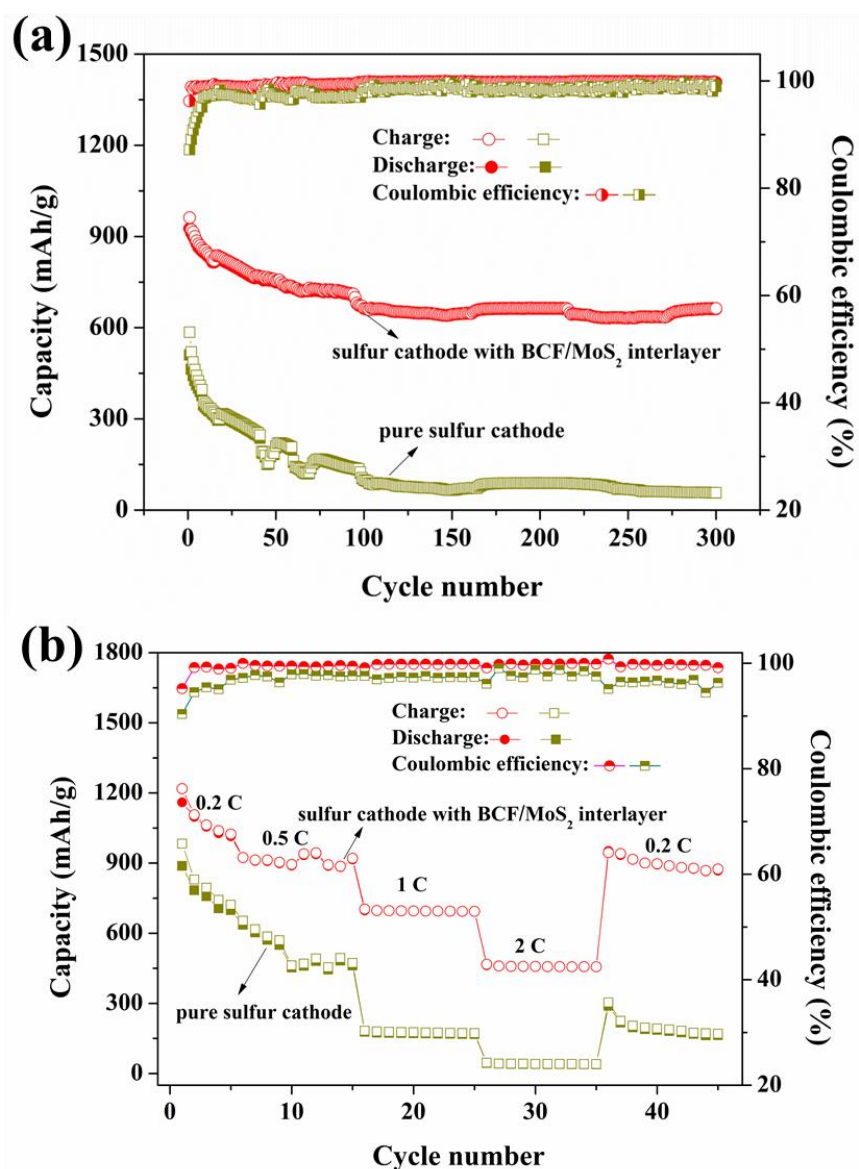


Figure 4 (a) The long-cycling performances of sulfur cathode with and without BCF/MoS<sub>2</sub> interlayer at charge-discharge current of 0.5 C, (b) the rate capabilities of sulfur cathode with and without BCF/MoS<sub>2</sub> interlayer.

## CONCLUSIONS

In summary, the Typha Angustifolia-like MoS<sub>2</sub>/carbon nanofiber composites has been successfully employed as the interlayer in Li-S batteries. The BCF/MoS<sub>2</sub> interlayer bestows Li-S batteries excellent long-term cycle stability (only 0.09% capacity fade per cycle) and high

coulombic efficiency (99%) even when the sulfur content is as high as 65% in the electrode. The exceptional performance can be attributed to (i) the resultant conductive fiber networks, providing conductive skeletons for the electrons transfer; (ii) **abundant** gaps and pores to store the sulfur; (iii) polar MoS<sub>2</sub> shell chemically trapping the LiPSs as well as catalyzing the LiPSs redox reaction. **Therefore, the unique Typha Angustifolia-like MoS<sub>2</sub>/carbon nanofiber interlayer has shed a light for high-performance Li-S batteries development.**

#### **AUTHOR CONTRIBUTIONS**

All authors listed have made a substantial, direct and intellectual contribution to the work, and approved it for publication.

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