Pomegranate-Like Structured Si@SiOₓ Composites With High-Capacity for Lithium-Ion Batteries

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Silicon anodes with an extremely high theoretical specific capacity of 4,200 mAh g⁻¹ have been considered as one of the most promising anode materials for next-generation lithium-ion batteries. However, the large volume expansion during lithiation hinders its practical application. In this work, pomegranate-like Si@SiOₓ composites were prepared using a simple spray drying process, during which silicon nanoparticles reacted with oxygen and generated SiOₓ on the surface. The thickness of the SiOₓ layer was tuned by adjusting the drying temperature. In the unique architecture, the SiOₓ which serves as the protection layer and the void space in pomegranate-like structure could alleviate the volume expansion during repeated lithium insertion/extra ction. As a lithium-ion battery anode, pomegranate-like Si@SiOₓ composites dried at 180°C delivered a high specific capacity of 1746.5 mAh g⁻¹ after 300 cycles at 500 mA g⁻¹.

Keywords: lithium-ion battery, anode, silicon, pomegranate-like structured, spray drying

INTRODUCTION

Lithium-ion batteries (LIBs) have been widely used in portable electronics, electric vehicles, and large-scale energy storage systems owing to their high energy density, long cycling stability, and environmental friendliness (Armand and Tarascon, 2008; Yu et al., 2018; Zuo et al., 2019; Kwon et al., 2020; Li et al., 2020; Wang K. et al., 2020). Nowadays, the traditional commercial anode graphite (372 mAh g⁻¹) is not able to meet increasing demands for higher energy density. In this context, a Si anode with an ultra-high theory specific capacity (4,200 mAh g⁻¹) and lower work potential (< 0.5 V vs. Li⁺/Li) have emerged as an alternative (Szczech and Jin, 2011; Zhuang et al., 2017; Zuo et al., 2017; Fang et al., 2020a). However, during the lithiation process, the Si anode undergoes large volume expansion (~320%, Li₂₂Si₃), which leads to active material pulverizing and splitting away from the current. A solid electrolyte interface (SEI) for a Si anode is unstable, resulting in fracture and regeneration during the charge/discharge process. The SEI layer continuously grows and irreversibly consumes lithium-ions in the electrolyte, leading to inferior lithium-ion kinetics and poor electrochemical performance. Moreover, the Si anode is a semiconductor with poor conductivity, which is not conducive to delivering capacity (Zheng et al., 2019; Zuo et al., 2019; Kwon et al., 2020).
Many efforts have been made to solve these limitations. Studies have shown that nanostructures including 0D nanoparticles (Xu Y. et al., 2017), 1D nanowires/nanotubes (Ge et al., 2012; Kennedy et al., 2017), 2D nanofilms, and 3D porous structures (Yang et al., 2015; Xu et al., 2017; Chen et al., 2018; Mu et al., 2019; Fang et al., 2020b) can significantly reduce the volume effect and shorten the diffusion distance of lithium-ion. Combining Si with other matrices is also an efficient alternative to alleviate the volume expansion stress (Cheng et al., 2018; Jo et al., 2019; Tao et al., 2019). Mu et al. used self-assembly and magnesiothermic reduction to prepare a pomegranate-like core-shell structured Si@C composite (Mu et al., 2019). Benefiting from the void space in the special architecture, the Si@C anode delivered a reversible capacity of 1,494 mAh g\(^{-1}\) after 100 cycles at 100 mA g\(^{-1}\), corresponding to an outstanding capacity retention of 97% (He et al., 2017). In another work, Kim et al. creatively obtained Si/SiO\(_x\) from a sol-gel reaction using trithoxysilane as the precursor. After coating it with a uniform carbon film, it maintained a discharge capacity of 740 mAh g\(^{-1}\) after 100 cycles at 200 mA g\(^{-1}\). This satisfactory performance is mainly due to the amorphous SiO\(_x\) matrix, which can alleviate the volume expansion during the charge/discharge process (Park et al., 2014). During the lithiation process, SiO\(_x\) reacts with Li\(^{+}\) to generate Li\(_2\)O and a series of lithium silicates, which favors the construction of a stable SEI (Song et al., 2013; Wang et al., 2014; Liu et al., 2019).

\[
\begin{align*}
\text{SiO}_x + 2x\text{Li}^{+} + 2xe^- & \rightarrow \text{Si} + x\text{Li}_2O \quad (1) \\
\text{SiO}_x + y\text{Li}^{+} + ye^- & \rightarrow \text{Li}_y\text{SiO}_x \quad (2)
\end{align*}
\]

Based on the above discussion, by structural design and introducing SiO\(_x\) (silicon oxide, 0<x<2) as the protection layer, the volume expansion stress of a Si anode can be effectively alleviated. In this study, we developed pomegranate-like Si/SiO\(_x\) second particles by a one-step spray drying process, during which silicon nanoparticles reacted with oxygen and produced a uniform SiO\(_x\) protection layer. The unique pomegranate-like structure also contained a number of voids that could further maintain structural stability during cycling. We believe that this unique pomegranate-like Si/SiO\(_x\) could provide excellent electrochemical performance in LIBs.

**EXPERIMENTAL SECTION**

**Synthesis of Pomegranate-Like Structured Si@SiO\(_x\) Composites**

Pomegranate-like structured Si@SiO\(_x\) composites were fabricated through simple spray drying using nano-silicon as the precursor. Typically, 3 g of silicon nanoparticles (SiNPs, 20–120 nm, Guangzhou Hongwu Material Technology Co., Ltd) was added to 300 mL of deionized water and sonicated for 30 min. The homogeneous suspension was then pumped into the spray dryer at a speed of 10 mL min\(^{-1}\) and dehydrated using hot air. During the drying process, the fresh surface of the SiNPs reacted with oxygen in the hot air. By adjusting the drying temperature to 150, 180, and 210°C, the thickness of the SiO\(_x\) layer was tuned, and the Si@SiO\(_x\) composites were denoted as pSi-150, pSi-180, and pSi-210, respectively.

**Materials Characterization**

The morphology and structure of the prepared Si@SiO\(_x\) were observed using scanning electron microscopy (SEM, JEOL JSM-7500F) and transmission of electron microscopy (TEM, Tecnai G2 F20 S-TWIN). The crystal structure of Si@SiO\(_x\) was characterized by X-ray diffraction (XRD) using X’pert MPD DY1219 diffractometer (copper K\(_\alpha\) source, 40 kV, 20 mA), and the 29 angle was from 15 to 85° with scanning steps of 0.02°/s. The elemental distribution was analyzed by energy dispersive X-ray spectroscopy (EDS) attached to SEM. X-ray photoelectron spectroscopy (XPS) was performed on a PHI 5,600 physical electronic device. Raman spectra were recorded on LabRAM HR with a laser wavelength of 532 nm, and Fourier transform infrared spectroscopy (FT-IR) was conducted on KBr pellets. The specific surface area and pore size distribution of the sample were measured using nitrogen adsorption-desorption measurements on a Kubo-X1000 spectrometer.

**Electrochemical Measurement**

The CR2032 coin cell was used to test the electrochemical performance of the Si@SiO\(_x\) composites. First, active material pSi composite, conductive agent SP, and binders including sodium carboxymethyl cellulose (CMC), and styrene-butadiene rubber (SBR) were mixed at a mass ratio of 80:10:4:6, respectively. Then, the mixture was ground for 1 h in an agate mortar and coated on copper foil (Wason Copper Foil Co., Ltd). After drying in a vacuum oven at 60°C for 1 h, the copper with anode material was cut into pieces with a diameter of 12 mm and finally dried at 105°C for 12 h in a vacuum oven. Lithium metal was used as the counter electrode. 1 M LiPF\(_6\) was dissolved in the solution of ethylene carbonate (EC), dimethyl carbonate (DMC), and ethyl methyl carbonate (EMC) in a volume ratio of 1:1:1 as electrolyte. In addition, 2% fluoroethylene carbonate (FEC) was added as the electrolyte additive. Eventually, the coin cell was assembled in an Ar-filled glove box (Chengdu Derek Co., Ltd). The cycling performance and rate capacity of the newly prepared cells were tested on the Neware 4000s in a potential window from 0.01 to 2 V vs. Li\(^{+}/\text{Li}\). Cyclic voltammetry (CV) and electrochemical impedance spectroscopy (EIS) tests were performed on an electrochemical workstation (Multi Autolab M204, Metrohm). The CV scan region was 0.001–3 V with a sweep speed of 0.1 mV s\(^{-1}\) and the EIS measurement ranged from 10 mHz to 100 kHz with an amplitude of 5 mV.

**RESULTS AND DISCUSSION**

The preparation process of the pomegranate-like structured Si@SiO\(_x\) composite is illustrated in Figure 1. First, Si nanoparticles were dispersed in an aqueous solution with stirring and sonication. Subsequently, the homogeneous slurry was spray dried at various temperatures, during which the
surface of the SiNPs was partly oxidized to SiO$_x$ and rearranged to the pomegranate-like structured material.

As shown in Figure 2a, commercial SiNPs (20–120 nm) disorderly aggregate. After the spray drying process, SiNPs rearranged into the pomegranate-like structured second particles (Figures 2b,c, 1–7 µm, the particle size distribution maps are shown in Supplementary Figure 1). The EDS elemental mapping images in Figures 2d,e indicate that Si and O elements are uniformly distributed in the pSi-180 composite. The O element comes from the SiO$_x$ coating layer, formed when SiNPs were in contact with hot air. During the lithiation process, SiO$_x$ reacted with lithium-ions and generated Li$_2$O and a series of lithium silicates, which could partly relieve the volume expansion (Wang et al., 2014).

In TEM images, the unique pomegranate-like structure is also observed (Figure 3a) and pores (bright area) are displayed in the composite, which could accommodate the volume expansion of a Si anode and contribute to the permeation of lithium-ion. Figures 3b–e, Supplementary Figure 2 shows the morphology of SiNPs and the pomegranate-like composites formed under
different temperatures. The SiO$_x$ layer can be observed on the surface of SiNPs, with thicknesses of 2, 5, 10, and $\sim$15 nm for SiNPs, pSi-150, pSi-180, and pSi-210, respectively. With increasing temperature, the SiO$_x$ coating layer becomes thicker. Figure 3f displays the selected area electron diffraction (SEAD) of the pSi-180. The four diffraction rings correspond to (111), (220), (311), and (400) crystal planes of Si, indicating that SiO$_x$ was formed on the surface and SiNPs remains in the crystalline state inside.

The crystal phase states of the SiNPs, pSi-150, pSi-180, and pSi-210 were further analyzed by XRD (Figure 4A). The peaks at 28.4, 47.3, 56.1, 69.1, and 76.4° correspond to (111), (220), (311), and (400) crystal planes of Si, indicating that SiO$_x$ was formed on the surface and SiNPs remains in the crystalline state inside.
(311), (400), and (331) crystal planes of Si (PDF#77-2107), which is consistent with the SEAD result in Figure 3f. After the spray drying process, the peak positions of pSi-150, pSi-180, and pSi-210 are the same as those of SiNPs, indicating that the main crystal structure of SiNPs did not change. However, due to the formation of SiO$_x$ on the surface, the peak intensity decreased with increasing drying temperature. According to the FT-IR spectra in Figure 4B, symmetric, and asymmetric Si-O-Si stretching vibrations are observed at 1,101 and 878 cm$^{-1}$, respectively. As expected, the intensity of the two peaks of pSi-180 is higher than that of SiNPs due to the increasing amount of SiO$_x$. The Raman spectra of SiNPs and pSi-180 are shown in

![FIGURE 5](image1.png) ![FIGURE 6](image2.png)

**FIGURE 5** | XPS spectra of SiNPs (A), and pSi-180 (B); High resolution XPS spectra of Si 2p for SiNPs (C), and pSi-180 (D).

**FIGURE 6** | Nitrogen adsorption-desorption isotherms of (A) SiNPs, and (B) pSi-180.
Supplementary Figure 3. The peak at 514 cm\(^{-1}\), which belongs to Si-Si, has a decreased intensity after drying at 180\(^\circ\)C due to the formation of SiO\(_x\).

XPS was used to further investigate the chemical composition and state of SiNPs and pSi-180. The peaks at 532, 150, and 99 eV are associated with O 1s, Si 2s, and Si 2p in Figures 5A,B. After the spray drying process, the strength of the O 1s peak increased. Figures 5C,D show the high-resolution XPS spectra of Si 2p for SiNPs and pSi-180, respectively. The peaks at 98.7 and 102.1 eV correspond to Si\(^0\) and Si\(^{2+}\), respectively. After the spray drying process, a new peak appears at the higher binding energy (103.0 eV), which is consistent with Si\(^{4+}\). The amount of SiO\(_x\) on the surface increased from 9.1 to 59.4% (calculated by CasaXPS), which could improve the cycling performance of the Si@SiO\(_x\) anode (Zhang X. L. et al., 2019).

Figure 6 shows the N\(_2\) adsorption-desorption isotherms of SiNPs and pSi-180. According to the Brunauer, Emmett, and Teller (BET) model, the specific surface area of SiNPs and pSi-180 was 33.5 and 35.0 m\(^2\) g\(^{-1}\), respectively. The improvement of the specific surface area was mainly due to the SiO\(_x\) on the surface of SiNPs, which could accommodate the agglomeration among SiNPs (Zheng et al., 2018). Based on the Barret, Joyner, and Halenda (BJH) model, the pore volume also increased after the formation of the pomegranate-like structure (0.15 to 0.29 cc g\(^{-1}\)). During the spray drying process, SiNPs rearranged to the pomegranate-like structure, which could effectively alleviate the volume expansion stress.
during the repeated lithiation/delithiation process. Furthermore, the unique structure is conducive to the penetration of electrolyte and the diffusion of lithium-ion (Yu et al., 2018).

To investigate the lithium-ion storage in the pomegranate-like structured Si@SiO\textsubscript{x} anode, CV was performed at different rates. As shown in Figures 7A,C, the peak at \( \sim 0.17 \) V in the reduction process are ascribed to the alloying process between Li\textsuperscript{+} and Si (Si\textsubscript{12}Si\textsubscript{5}). The anodic peaks at 0.33 V and 0.53 V are related to the reversible process (Li\textsubscript{22}Si\textsubscript{5} amorphous Si). Although the CV curves deliver a similar shape, the anodic peak intensity of pSi-180 is higher than that of SiNPs. According to the Sevcik equation (Equation 3), the lithium-ion diffusion coefficient (\( D_{Li^{+}} \)) for SiNPs (Figure 7B) and pSi-180 (Figure 7D) is \( 6.0 \times 10^{-10} \) and \( 1.4 \times 10^{-9} \) cm\textsuperscript{2}s\textsuperscript{-1}, respectively, indicating...
that the pomegranate-like structured anode is conducive to the diffusion of lithium-ion (Feng et al., 2019; Luo et al., 2019; Mu et al., 2019).

\[ I_p = 2.69 \times 10^5 n^{3/2} A D_{Li^+}^{1/2} v^{1/2} C_0 \]  (3)

where n is the number of reaction electrons, A is the specific surface area of the electrode, C is the concentration of lithium-ion.

To better understand the role of SiO\(_x\) and the pomegranate-like structure, EIS measurements were conducted between 10 mHz and 100 kHz. As shown in Figure 7E, Nyquist plots of SiNPs and pSi-180 consist of two parts before cycling: a semicircle in the high-frequency region corresponding to the charge transfer resistance (R\(_{ct}\)) and an oblique line in the low frequency region associated with the lithium-ion diffusion impedance in the electrode (W\(_o\)). The point of intersection between the semicircle and the X-axis in the high region represents the contact resistance (R\(_s\)). According to the equivalent circuit shown in Supplementary Figure 4, the calculated R\(_{ct}\) and R\(_s\) for SiNPs decreased after spray drying at 180°C (R\(_{ct}\): 818.7 Ω → 235.2 Ω; R\(_s\): 6.0 Ω → 0.8 Ω). According to Equation 5, the slope in Figure 7F represents the Warburg impedance coefficient (σ\(_w\)), whose square is inversely proportional to D\(_{Li^+}\). Hence, the D\(_{Li^+}\) of pSi-180 is slightly higher than that of SiNPs (calculated through EIS), which is in accordance with the result calculated through CV curves with different scan rates. The pomegranate-like structure could enhance the contact area between the active materials and electrolyte, thereby improving the charge transfer resistance and lithium-ion diffusion rate (Li et al., 2018; Luo et al.,

| Anode material | Reversible specific | Current | References |
|----------------|---------------------|---------|------------|
| Pomegranate-like Si@C | 1,204 mAh g\(^{-1}\) (100th) | 0.5 A g\(^{-1}\) | Mu et al., 2019 |
| Pomegranate-like structure | 581 mAh g\(^{-1}\) (200th) | 0.2 A g\(^{-1}\) | Shen et al., 2017 |
| Pomegranate-like Si/N-G | 1,100 mAh g\(^{-1}\) (150th) | 0.1 A g\(^{-1}\) | Lin et al., 2016 |
| Pomegranate-like SiO\(_x\)/C | 1,024 mAh g\(^{-1}\) (200th) | 0.5 A g\(^{-1}\) | Yu et al., 2018 |
| Core-bishell Si@SiO\(_x\)@TiO\(_2\)-δ | 650 mAh g\(^{-1}\) (200th) | 0.2 A g\(^{-1}\) | Hu et al., 2019 |
| Si-SiO\(_x\)-C | 1,035 mAh g\(^{-1}\) (200th) | 0.1 A g\(^{-1}\) | Lee et al., 2017 |
| Hollow B-Si/SiO\(_x\)@VN/PC | 1,237 mAh g\(^{-1}\) (300th) | 0.5 A g\(^{-1}\) | Zhang X. L. et al., 2019 |
| Pomegranate-like Si@SiO\(_x\) | 1,747 mAh g\(^{-1}\) (300th) | 0.5 A g\(^{-1}\) | This work |

**Figure 9** | Differential capacity (dQ/dV) plots for Si (A), pSi-150 (B), pSi-180 (C), and pSi-210 (D) after 50 cycles at 200 mA g\(^{-1}\).
where \( R \) is the gas constant, \( T \) is the ambient temperature, \( n \) is the number of reaction electrons, \( F \) is the Faraday constant, \( A \) is the specific surface area of the electrode, \( C \) is the concentration of lithium-ion, and \( \sigma_w \) is the Warburg impedance coefficient.

Figures 8A–D shows the voltage-capacity curves of SiNPs, pSi-150, pSi-180, and pSi-210 at 100 mA g\(^{-1}\) for the first cycle, and 200 mA g\(^{-1}\) for the following cycles. In the first cycle, the charge/discharge specific capacity was 3179.0/418 8.7, 3031.2/4103.6, 2891.2/3781.6, and 2644.6/3459.5 mAh g\(^{-1}\), for the four samples, respectively. And the Coulombic efficiency was 75.9, 73.8, 76.5, and 76.4%, respectively. With the increase in the drying temperature, the initial reversible specific capacity decreased for the surface of SiNPs, which was slightly oxidized to SiO\(_x\). In addition, Supplementary Figure 5 compares the voltage-capacity curves of the above samples in the first cycle, and it is observed that they exhibit similar charge/discharge plateaus. However, with increasing spray drying temperature, the voltage difference between charge and discharge plateaus become larger due to the passivation of SiO\(_x\) on the surface (Zheng et al., 2018). In the second cycle, the reversible specific capacity was 2385.4, 2525.7, 2656.9, and 2378.5 mAh g\(^{-1}\) for SiNPs, pSi-150, pSi-180, and pSi-210, respectively. After 50 cycles, the reversible specific capacity was 1148.9, 2054.5, 2394.9, and 1984.7 mAh g\(^{-1}\), respectively. The 50th capacity retention of the above samples...
was 48.2, 81.3, 90.1, and 83.4%, respectively. With the aid of the SiO$_x$ buffer layer and the unique pomegranate-like structure, the cycling stability was significantly improved.

**Figure 8E** shows the long cycling performance of pSi-180 at 500 mA g$^{-1}$. After 100 cycles, the pSi-180 electrode delivers a high reversible specific capacity of 1786.3 mAh g$^{-1}$. In the following cycles, pSi-180 remained stable and after 400 cycles it still presented a reversible specific capacity of 1642.9 mAh g$^{-1}$ and remarkable cycling stability (0.09% decay per cycle). The rate capability was also tested at various current densities between 100 mA g$^{-1}$ and 2,000 mA g$^{-1}$, shown in **Supplementary Figure 6**. Reversible specific capacities of 2853.9, 1786.3, 1519.8, and 1247.7 mAh g$^{-1}$ were obtained at 100, 500, 1,000, and 2,000 mA g$^{-1}$, respectively. When the current density returned to 100 mA g$^{-1}$ from the high rate, pSi-180 delivered 2686.7 mAh g$^{-1}$, corresponding to a capacity recovery of 94.1%.

As shown in **Figure 9**, the dQ/dV curves display a similar shape after cycling, which is consistent with the CV results. After introducing SiO$_x$ and the formation of the pomegranate-like structure, the peak intensity of the structured composites increased compared to that of SiNPs, indicating better electrode activity. Additionally, the polarization potential difference between anodic and cathodic peaks for SiNPs, pSi-150, pSi-180, and pSi-210 after 50 cycles was 0.20, 0.14, 0.11, and 0.12 V, respectively. After the spray drying process, pomegranate-like structured composites suffered less polarization and delivered better lithium-ion lithiation/delithiation kinetics (Zhang Y. et al., 2019).

**CONCLUSION**

The pomegranate-like structured Si@SiO$_x$ was successfully prepared through a one-step spray drying process. By controlling the inlet air temperature, the thickness and oxidation degree of SiNPs could be tuned. In the series of the pomegranate-like Si@SiO$_x$ composites, pSi-180 delivered a high discharge capacity of 1,747 mAh g$^{-1}$ and outstanding long-cycling performance (0.13% loss per cycle) after 300 cycles at 500 mA g$^{-1}$. In addition, pSi-180 presented a reversible capacity of 1519.8 mAh g$^{-1}$ at 2 A g$^{-1}$. This excellent electrochemical property comes from the SiO$_x$ coating layer and the unique pomegranate-like structure, which can accommodate volume expansion during the repeated charge/discharge process. In this work, a one-step spray drying technology was used, which is conducive to the large-scale production of a Si anode. This method could be used in other nanoparticles to prepare pomegranate-like structured materials.

**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the article/Supplementary Material.

**AUTHOR CONTRIBUTIONS**

JL is responsible for conceptualization, methodology, and wrote the main manuscript text. WL, YQ, ZX, and MQ contributed immensely to the writing and discussion of the final manuscript. GP and YR are responsible for project administration and funding acquisition. In addition, they all approved the final manuscript for publication. All authors contributed to the article and approved the submitted version.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2020.00666/full#supplementary-material
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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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