Characterization of Sn₄P₃–Carbon Composite Films for Lithium-Ion Battery Anode Fabricated by Aerosol Deposition

Toki Moritaka, Yuh Yamashita, Tomohiro Tojo, Ryoji Inada, and Yoji Sakurai

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Abstract: We fabricated tin phosphide–carbon (Sn₄P₃/C) composite film by aerosol deposition (AD) and investigated its electrochemical performance for a lithium-ion battery anode. Sn₄P₃/C composite powders prepared by a ball milling was used as raw material and deposited onto a stainless steel substrate to form the composite film via impact consolidation. The Sn₄P₃/C composite film fabricated by AD showed much better electrochemical performance than the Sn₄P₃ film without complexing carbon. Although both films showed initial discharge (Li⁺ extraction) capacities of approximately 1000 mAh g⁻¹, Sn₄P₃/C films retained higher reversible capacity above 700 mAh g⁻¹ after 100 cycles of charge and discharge processes while the capacity of Sn₄P₃ film rapidly degraded with cycling. In addition, by controlling the potential window in galvanostatic testing, Sn₄P₃/C composite film retained the reversible capacity of 380 mAh g⁻¹ even after 400 cycles. The complexed carbon works not only as a buffer to suppress the collapse of electrodes by large volume change of Sn₄P₃ in charge and discharge reactions but also as an electronic conduction path among the atomized active material particles in the film.

Keywords: tin phosphide; carbon; composite film; aerosol deposition; lithium-ion battery; anode

1. Introduction

Li-ion batteries (LiBs) are widely used as a power source for portable electronic devices, and recently have attracted much attention as a large-scale power source for electric vehicles and plug-in hybrid electric vehicles. In order to achieve advanced LiBs with higher energy density, development of anode materials with higher capacity is indispensable. Graphite with a theoretical capacity of 372 mAh g⁻¹ is commonly used as an anode for LiBs, while lithium alloys such as Li–Si and Li–Sn with a higher theoretical capacity (Li₄₄Si: 4200 mAh g⁻¹, Li₄₄Sn: 990 mAh g⁻¹) have been extensively studied [1–3]. However, they result in poor cycling stability due to a large volume change during charge and discharge reactions. In order to improve the cycling stability, various composite materials including metal oxides, multiphase alloys and intermetallic compounds have been studied as alternatives to graphite anode for LiBs [3–7]. These materials show much higher capacities than graphite and improved cycling performance compared to lithium alloy materials. The enhancement of cycling stability in these Li-alloy-based materials attributed to an inactive matrix [4]. Li-alloy-based materials form an inactive matrix during cycling and this matrix is expected to suppress the volume change of the alloying reaction, and keeps the electrode particles mechanically connected together resulting in a reversible alloying reaction.

Tin phosphide Sn₄P₃ (theoretical gravimetric capacity = 1255 mAh g⁻¹) is known as one of the high capacity alloy-based anode materials for LiBs [8–11]. Sn₄P₃ has a layered structure (space group: R-3m)
suitable for lithium insertion and high intrinsic electronic conductivity at room temperature. In addition, \( \text{Sn}_4\text{P}_3 \) forms \( \text{Sn} \) and \( \text{Li}_3\text{P} \) in the lithium insertion reaction. \( \text{Li}_3\text{P} \) has high ionic conductivity \[12\] and would act as a matrix for suppressing the volume change during the alloying reaction. According to these features, application of a \( \text{Sn}_4\text{P}_3 \) anode for high-capacity sulfide-based solid-state batteries has been also demonstrated \[13\].

As reported in the literature \[8,9\], \( \text{Sn}_4\text{P}_3 \) shows initial reversible capacity as high as 900 mAh g\(^{-1}\) and by controlling the electrical potential window in galvanostatic charge and discharge testing, it maintains a reversible capacity above 400 mAh g\(^{-1}\) after 50 cycles. Reducing the size and morphology of \( \text{Sn}_4\text{P}_3 \) particles \[14,15\] and doping of a small amount of Fe \[16\] and Mn \[17\] into \( \text{Sn}_4\text{P}_3 \) are also effective for further improvement of the cycling stability. Moreover, complexing the carbon materials with nano-structured \( \text{Sn}_4\text{P}_3 \) particles significantly enhances both the rate performance and cycling stability \[18–25\]. The complexed carbon behaves as the buffer for the volume change of active material particles during charge and discharge reaction and maintains the electric conduction between the particles.

In general, the electrodes used in actual batteries are fabricated by coating a slurry composed of electrode active materials on metallic foils and contain conducting carbon additives and binders. For the case of carbon complexed alloy-based anodes, the weight fraction of active materials in an electrode becomes small (less than ~70%) due to significant amounts of conducting additives and binders. Consequently, gravimetric specific capacity calculated by the total mass of the electrode (including carbon additives and binders) is reduced significantly. To address this issue, we are focusing on the aerosol deposition (AD) method \[26–28\] as an electrode fabrication process, which uses impact consolidation for ceramic particles at room temperature. This method is known as a fabrication process of various functional ceramic films at room temperature. By controlling the size and morphology of the base powder material, the film fabricated by AD has a dense structure made of nanocrystalline particles, and the structural and physical properties are similar to the base powder material. Moreover, adhesion strength between the film formed by AD and the substrate is high without adding binders \[26,27\]. To date, several works for the application of AD to rechargeable battery materials have been reported. The electrochemical performance for film-shaped electrodes of Si alloy or composite \[29,30\], tin-phosphide with different compositions \[31\], transition metal oxides \[32–40\] formed on a metal and a ceramic-based solid electrolyte substrate have been studied to verify the feasibility of AD. Moreover, as-deposited solid electrolyte films show a moderate \( \text{Li}^+ \) conductivity of \( 10^{-7} \text{ to } 10^{-5} \text{ S cm}^{-1} \) at room temperature \[41–45\].

In this work, we fabricated \( \text{Sn}_4\text{P}_3 \)-carbon (\( \text{Sn}_4\text{P}_3/C \)) composite films on a stainless steel substrate by AD and the electrochemical performance of the LiB anode was evaluated. \( \text{Sn}_4\text{P}_3/C \) composite powder was prepared by ball milling and used as a raw material to form the composite film via impact consolidation. The influence of complexed carbon on the cycling stability of both the microstructure and reversible capacity was examined.

2. Materials and Methods

2.1. Fabrication and Characterization of \( \text{Sn}_4\text{P}_3/C \) Composite Powders

\( \text{Sn}_4\text{P}_3 \) powder was prepared using a simple mechanochemical synthesis with a planetary ball-milling \[8,9,13\]. Sn (99%, Kojundo Chemical Laboratory, Saitama, Japan) and red P (99.9%, Kojundo Chemical Laboratory, Saitama, Japan) powders were used as starting materials. Stoichiometric amounts of the starting materials (10 g) were put into a ZrO\(_2\) vessel (45 mL) with ZrO\(_2\) balls that were 10 mm in diameter (100 g) and reacted in a planetary ball milling apparatus (Nagao System, Planet M2-3F, Kawasaki, Japan) with a fixed rotation speed of 350 rpm for 8 h under an Argon atmosphere.

It is known that controlling the particle size of raw powder is important for film fabrication by AD \[14–16,28,30,32\], and we could not form \( \text{Sn}_4\text{P}_3 \) film by AD with as-synthesized powder. In order to prepare \( \text{Sn}_4\text{P}_3 \) powders suitable for AD, as-synthesized \( \text{Sn}_4\text{P}_3 \) powder (~10 g) was put into a ZrO\(_2\) vessel with ethanol (30 mL) and ZrO\(_2\) balls with diameters of 1 mm (30 g) and 2 mm (100 g) and then pulverized by a planetary ball-milling at 350 rpm and 24 h. After the pulverization, the obtained \( \text{Sn}_4\text{P}_3 \)
powder and acetylene black (AB) were mixed with a weight ratio of Sn₄P₃:AB = 8:2. The mixture (~5 g) was put into a ZrO₂ vessel again with ZrO₂ balls with diameters of 5 mm (50 g) and 10 mm (100 g), and Sn₄P₃/C composite powder was prepared by a planetary ball-milling at 350 rpm for 24 h.

The crystal phase of as-synthesized Sn₄P₃, ball-milled Sn₄P₃ and Sn₄P₃/C powder was evaluated by X-ray Diffractometer (XRD; Rigaku, MultiFlex, Tokyo, Japan) using Cu Ka radiation (λ = 0.15418 nm), with a measurement range 2θ of 5°–90° and a step interval of 0.002°. Field emission scanning electron microscopy (FE-SEM; Hitachi High-Technologies, SU8000 Type II, Tokyo, Japan) was used to observe the size and morphology for all powder samples. Energy dispersive X-ray (EDX) analysis was also performed using FE-SEM, to observe Sn₄P₃/C particles and the corresponding distribution of Sn, P and C elements.

### 2.2. Fabrication and Characterization of Sn₄P₃/C Composite Films by AD

As shown in the literature [34,40,42], an AD apparatus consists of a carrier gas supplying system, an aerosol chamber, a deposition chamber equipped with a motored X-Y-Z stage and a nozzle with a thin rectangular-shaped orifice with the cross-sectional size of 10 mm × 0.5 mm. Sn₄P₃/C powder was used as a raw material for fabricating Sn₄P₃/C composite film by AD. A carrier nitrogen (N₂) gas flows out from the gas supply system to the aerosol chamber. In the aerosol chamber, the powder is dispersed into the carrier gas to make an aerosol. Using a pressure difference between the evacuated deposition chamber and the carrier gas system, the aerosol flows into the deposition chamber through a nozzle and is sprayed onto an SUS316L stainless steel substrate. The deposition area was masked into a circular shape 8 mm in diameter. The deposition chamber was evacuated to a low vacuum state at approximately 20 Pa and deposition was carried out for 20–30 min. During the deposition process, the stage was moved uni-axially with a back-and-forth motion length of 50 mm and a speed of 10 mm s⁻¹. Based on the results in our previous works [34,38,40,42,44], the distance between the substrate and nozzle tip was set to 10 mm and the mass flow of the N₂ carrier was fixed at 20 L min⁻¹.

The crystal phase of the Sn₄P₃/C composite film was evaluated by XRD using Cu Ka radiation (λ = 0.15418 nm), with a measurement range 2θ of 5°–90° and a step interval of 0.002°. Microstructure observation of composite films was carried out by using FE-SEM, and EDX analysis was also performed to observe the distribution of Sn, P and C elements in the composite film.

The electrochemical properties of as-deposited Sn₄P₃/C films were evaluated by using a two-electrode set up. A Sn₄P₃/C film on an SUS316L substrate was used as a working electrode, where as a single lithium foil served as both a counter and a reference electrode. The electrolyte solution was 1 mol L⁻¹ LiPF₆ in a mixture of ethylene carbonate (EC) and dimethyl carbonate (DMC) with a volume ratio of 1:1 (Kishida Chemical Co., Ltd., Osaka, Japan). Together with a separator (Celgard, Celgard 3501, Tokyo, Japan), these components were assembled in a CR2032 coin-type cell. The cell assembly was carried out in a dry Argon-filled glove box. The cells were charged and discharged over the cell voltage ranges of 0 to 0.75, 1, 1.25 and 2.5 V at a fixed current density (per total mass of composite film) of 50 mA g⁻¹ and 25 °C using the Battery Test System (TOSCAT-3100, TOYO-SYSTEM, Iwaki, Japan). After the cycling test, the cells were disassembled in a dry Argon-filled glove box and the microstructure of the Sn₄P₃/C composite films was observed using FE-SEM. Before the observations, cycled Sn₄P₃/C films were cleaned with DMC to eliminate residual Li salt therein.

### 3. Results and Discussion

#### 3.1. Crystal Phase and Microstructure of Sn₄P₃/C Composite Powder and Film

The XRD data for as-synthesized Sn₄P₃, ball-milled Sn₄P₃ and Sn₄P₃/C composite powders are summarized in Figure 1. The pattern for Sn₄P₃ (JCPDS No. 03-066-0017) is also shown as the reference. It is confirmed that peak patterns for all sample powders agree well with the reference, suggesting that any structural changes did not occur in the ball-milling and carbon-complexing process. No peaks from carbon were detected in the Sn₄P₃/C powder because complexed carbon with Sn₄P₃ has amorphous structure.
Figure 2 shows scanning electron microscopy (SEM) images for as-synthesized Sn₄P₃, ball-milled Sn₄P₃ and Sn₄P₃/C composite powders. As-synthesized powder consists of agglomerated particles with a size of 1–5 μm (Figure 2a). After ball-milling, the particle size reduces to 0.5–1.5 μm (Figure 2b). On the other hand, after complexing carbon (acetylene black (AB)) with Sn₄P₃ by ball-milling, the particle size is not changed remarkably (Figure 2c).

![XRD patterns](image)

**Figure 1.** Comparison of X-ray diffraction (XRD) patterns for as-synthesized Sn₄P₃ powder, ball-milled Sn₄P₃ powder and Sn₄P₃/C composite powder.

![SEM images](image)

**Figure 2.** Scanning electron microscopy (SEM) images for (a) as-synthesized Sn₄P₃ powder, (b) ball-milled Sn₄P₃ powder and (c) Sn₄P₃/C composite powder.

A higher magnified SEM image of Sn₄P₃/C particles and elementary distributions for Sn, P and C in an observation area are shown in Figure 3. As shown in Figure 3a, the sample powder looks like agglomerated particles with a particle size of about 0.5–2 μm. Sn and P show similar distribution in an observed area and were detected on agglomerated particles, and interestingly, C is also distributed along the particle shape. This suggests that carbon (AB) particles are complexed successfully with Sn₄P₃ particles by a simple ball-milling process.
deformed or fractured particles. Moreover, Sn, P and C are distributed uniformly in an observed area, suggesting that carbon is included successfully in the film.

**Figure 3.** (a) SEM image of Sn$_4$P$_3$/C composite powder and corresponding element mapping for Sn, P and C are shown in (b), (c) and (d).

Figure 4 shows the XRD pattern and cross-sectional SEM image of the Sn$_4$P$_3$/C composite film formed on an SUS316L substrate by AD. Note that the peak intensity for the composite film is one order lower than the raw powder. Together with the peaks from the substrate, the peaks from the Sn$_4$P$_3$ phase are clearly confirmed but become broader compared to raw powder. No other secondary phases are formed during film fabrication by AD. The thickness of the composite film is confirmed to be 2.5–3 µm. An SEM image and elementary distributions for Sn, P and C for the broader surface of the Sn$_4$P$_3$/C composite film are shown in Figure 5. It can be seen that the film is composed of deformed or fractured particles. Moreover, Sn, P and C are distributed uniformly in an observed area, suggesting that carbon is included successfully in the film.

**Figure 4.** (a) XRD patterns for the Sn$_4$P$_3$/C composite powder and film formed on an SUS316L substrate and (b) cross-sectional SEM image for the Sn$_4$P$_3$/C composite film formed on an SUS316L substrate.
with ball-milled Sn.

3.2. Electrochemical Performance of Sn Film and retains a reversible capacity above 700 mAh g$^{-1}$ in the SUS316L substrate. Therefore, the rapid degradation of Sn coin-type cell after testing and confirmed that the majority of the film was collapsed and peeled off.

$m (Figure S1). As can be seen, the capacity fading in the Sn film electrode without complexing carbon. The Sn film was fabricated by AD with ball-milled Sn$_3$P$_3$ powder (Figure 2b) on an SUS316L substrate, with a thickness of approximately 2 µm (Figure S1). As can be seen, the capacity fading in the Sn$_3$P$_3$ film with cycling is relatively fast and reduces to less than 10 mAh g$^{-1}$ at the 80th cycle. We checked the Sn$_3$P$_3$ film taken from a disassembled coin-type cell after testing and confirmed that the majority of the film was collapsed and peeled off from the SUS316L substrate. Therefore, the rapid degradation of Sn$_3$P$_3$ films with cycling is caused by the mechanical damage due to the large volume change of the Sn$_3$P$_3$ film in charge and discharge reactions. On the other hand, the Sn$_4$P$_3$/C composite film shows much better cycling stability than the Sn$_3$P$_3$ film and retains a reversible capacity above 700 mAh g$^{-1}$ even at the 100th cycle, indicating that the

Figure 5. (a) SEM image of the broader surface of the Sn$_4$P$_3$/C composite film. Corresponding element mapping for Sn, P and C are shown in (b), (c) and (d).

3.2. Electrochemical Performance of Sn$_4$P$_3$ Film Electrodes

Next, we discuss the electrochemical performance for Sn$_4$P$_3$/C composite film as a LiB anode. Figure 6 shows the galvanostatic charge (Li$^+$ insertion) and discharge (Li$^+$ extraction) curves at different cycle numbers for the Sn$_4$P$_3$/C film electrode in a coin-type cell. The cell voltage window is from 0 to 2.5 V. At the first cycle, charge capacity reaches 1750 mAh g$^{-1}$, which is much higher than the theoretical capacity (1255 mAh g$^{-1}$) of Sn$_3$P$_3$, while discharge capacity is confirmed to be 1200 mAh g$^{-1}$. Coulombic efficiency at the first cycle is 69%, but irreversibility in charge and discharge reaction is greatly reduced after the second cycle and the Coulombic efficiency retains 96–98%. This suggests that the charge capacity in the first cycle includes the contribution of side reactions such as the decomposition of an organic liquid electrolyte at a lower cell voltage to form Li$^+$ conducting solid-electrolyte interphase (SEI) on the electrode surface. The reversible capacity decreases monotonically with cycling but retains a high capacity of 800 mAh g$^{-1}$ at the 50th cycle. The averaged operation potential is approximately 0.7 V and step-like profiles are confirmed in both charge and discharge processes, which is consistent with the results reported in the literature [8–11].

Cycling stability for the Sn$_4$P$_3$/C composite film electrode is shown in Figure 7, together with the data for the Sn$_4$P$_3$ film electrode without complexing carbon. The Sn$_4$P$_3$ film was fabricated by AD with ball-milled Sn$_4$P$_3$ powder (Figure 2b) on an SUS316L substrate, with a thickness of approximately 2 µm (Figure S1). As can be seen, the capacity fading in the Sn$_4$P$_3$ film with cycling is relatively fast and reduces to less than 10 mAh g$^{-1}$ at the 80th cycle. We checked the Sn$_4$P$_3$ film taken from a disassembled coin-type cell after testing and confirmed that the majority of the film was collapsed and peeled off from the SUS316L substrate. Therefore, the rapid degradation of Sn$_4$P$_3$ films with cycling is caused by the mechanical damage due to the large volume change of the Sn$_4$P$_3$ film in charge and discharge reactions. On the other hand, the Sn$_4$P$_3$/C composite film shows much better cycling stability than the Sn$_4$P$_3$ film and retains a reversible capacity above 700 mAh g$^{-1}$ even at the 100th cycle, indicating that the
complexed carbon is effective at enhancing the cycling performance of Sn₄P₃. However, the capacity fading is accelerated by further cycling above 100 cycles and the reversible capacity is reduced to 300 mAh g⁻¹ at the 200th cycle. It is worth noting that we also fabricated a Sn₄P₃/C composite film with a lower carbon content (weight ratio Sn₄P₃:C = 9:1) and evaluated the electrochemical performance, but the degradation with cycling becomes more significant and the reversible capacity at 100 cycles was reduced to 500 mAh g⁻¹ (Figure S2).

![Graph](image_url)

**Figure 6.** Galvanostatic charge and discharge curves at different cycle numbers for the Sn₄P₃/C composite film electrode at 25 °C and 50 mA g⁻¹. Cell voltage window for galvanostatic cycling is 0–2.5 V.

![Graph](image_url)

**Figure 7.** Comparison of cycling stability of charge and discharge capacities for the Sn₄P₃/C composite film and the Sn₄P₃ film electrodes. Cell voltage window for galvanostatic cycling is 0–2.5 V.

As shown in Figure 7, the degradation mode of the Sn₄P₃/C composite film in galvanostatic cycling seems to be classified into three processes: (1) Rapid degradation below 15 cycles, (2) gradual degradation from 15 to 100 cycles, (3) accelerated degradation after 100 cycles. In order to further examine the degradation process, differential capacities dQ/dV (Q: capacity (mAh), V: cell voltage (V)) are calculated and plotted against a cell voltage in Figure 8. Based on the examination of the electrochemical reaction mechanism for Sn₄P₃ anode characterized by ex-situ XRD and X-ray absorption
spectroscopy (XAS) analysis in the literature [8], the reactions occurring in the charge process are considered as follows: (A) Li\(^+\) insertion into the layered structure of Sn\(_4\)P\(_3\) (at ~0.85 V), (B) Formation of LiP and partial transformation from LiP to Li\(_3\)P (at ~0.65 V), (C) and (D) Allooying of Sn with Li to form Li\(_x\)Sn alloy (at ~0.5 V and ~0.3 V) and (E) Formation of Li\(_3\)P and Li\(_x\)Sn (at <0.25 V). On the other hand, reactions in the discharge process are considered as follows: (F) and (G) Dealloying reaction of Li\(_x\)Sn (at ~0.45 V and ~0.65 V), (H) Dealloying reaction of Li\(_x\)Sn and Li extraction to Li\(_3\)P to form LiP (at ~0.75 V) and (I) Li extraction from Li\(_x\)P (at >0.8 V). Labels (A)–(H) for these expected reactions are also plotted in a graph.

![Differential capacity dQ/dV curves](image)

**Figure 8.** Differential capacity dQ/dV (Q: capacity, V: cell voltage) curves for the Sn\(_4\)P\(_3\)/C composite film calculated from the data for galvanostatic cycling test.

In the range of 20 cycles or less, the contributions of (A) and (B) in charge and (I) in discharge decrease remarkably with cycling, while the reactivity of Sn for (D) and Li\(_x\)Sn for (G) seems to be activated with cycling. After discharging, Sn\(_4\)P\(_3\) is not completely formed reversibly but amorphous Sn and P are formed [8]. Moreover, it is demonstrated that the reversibility of the P ↔ LiP reaction is not good due to the large volume change and poor conductivity of the LiP phase [8,9,46,47], resulting in rapid capacity fading of the Sn\(_4\)P\(_3\)/C composite film during the initial 20 cycles. The dQ/dV profiles at the 20th and 100th cycle are similar but the peak intensity for (B), (C) and (D) in charge and (E), (G) and (H) in discharge decreases with the cycle progress. At the 150th cycle, these specific peaks in dQ/dV profile become smaller and broader compared to the profile at the 100th cycle.

For further examination, we fabricated another Sn\(_4\)P\(_3\)/C composite film to observe the change of the microstructure during the galvanostatic cycling. After the galvanostatic testing with different cycles, we took out the film electrodes of the disassembled cells and confirmed that all the films were not delaminated from the SUS316L substrates (see insets in Figure 9). SEM images for Sn\(_4\)P\(_3\)/C composite film after the first, 100th, 120th and 160th cycle are summarized in Figure 9. It is worth noting that capacity fading behavior with cycling for all Sn\(_4\)P\(_3\)/C films is nearly the same as the data shown in Figure 7. After the first cycle (Figure 9a), it can be seen that the asperities on the surface of the composite film are clearly increased compared to the as-deposited film (Figure 5a). This is caused by the large volume expansion and contraction in the charge and discharge reactions of active material. Such structural change is repeated during cycling and induces the gradual capacity fading with cycling. At the 100th cycle (Figure 9b), generation of many small cracks and agglomeration of the particles are confirmed and become significant with further cycling (Figure 9c,d). The former breaks the electrical conduction path and the latter reduces to electrochemical utilization of active materials in
the composite film, resulting in acceleration of the degradation of reversible capacities after 100 cycles as shown in Figure 7.

![SEM images of Sn₄P₃/C composite films](image)

**Figure 9.** SEM images for the broader surface of Sn₄P₃/C composite films taken out of disassembled cells: (a) after the first cycle, (b) after the 100th cycle, (c) after the 120th cycle and (d) after the 160th cycle. Insets in (b), (c) and (d) are the photo images of films with different cycle numbers.

The improvement of the cycling stability of the Sn₄P₃ anode has been demonstrated by controlling the cell voltage window in the literature but the cycle numbers were limited to only 50 [8,9], so we investigated the long-term cycling stability for Sn₄P₃/C composite films at different cell potential windows of 0–0.75 V, 0–1 V and 0–1.25 V. The galvanostatic charge and discharge curves at the 20th cycle and cycling performance for Sn₄P₃/C composite films tested at different cell potential windows are shown in Figures 10 and 11. Although the reversible capacities at the 20th cycle reduce monotonically with decreasing cell voltage for discharge from 1.25 to 0.75 V, the cycling stability is dramatically improved. The film tested at 0–0.75 V shows a reversible capacity of 380 mAh g⁻¹ at the 400th cycle and the capacity retention reaches 80%. The film electrodes tested at 0–1 V and 0–1.25 V also show higher reversible capacities of 400 and 500 mAh g⁻¹ at the 200th cycle than the film tested at 0–2.5 V (300 mAh g⁻¹, see Figure 7), but the capacity fading is accelerated with further cycling. Reversible capacities at the 400th cycle for the films tested at 0–1 V and 0–1.25 V are only 250 and 110 mAh g⁻¹, respectively. By limiting the potential for discharge below 0.75 V, the extraction reaction of Li from LiₓP with a larger volume change is greatly suppressed, which could contribute to better cycling stability.
After 40 cycles, the film tested at 0–1.25 V is 97% after 40 cycles. As shown in Figure 13a, no delamination of the film from the SUS316L substrate was seen after cycling at 0–1.25 V. In addition, the structural change of the film cycled at 0–0.75 V is less than that for the film cycled at 0–1.25 V, which contributes to the better cycling performance. For the film cycled at 0–1.25 V with the lowest capacity retention (Figure 13b), delamination of many 

Figure 10. Comparison of galvanostatic charge (Li⁺ insertion) and discharge (Li⁺ extraction) curves at the 20th cycle for the Sn₄P₃/C composite film electrode tested at different cell voltage windows.

Figure 11. Cycling stability of charge and discharge capacities for the Sn₄P₃/C composite film tested at different cell voltage windows.

In Figure 12, the Coulombic efficiencies for Sn₄P₃/C composite films tested at different cell voltage windows are plotted against the cycle numbers. It can be seen that the efficiency for the initial several cycles becomes lower by reducing the cell voltage for discharging (Li extraction) from 1.25 to 0.75 V. This is mainly attributed to the reduction of the Li⁺ extraction reaction from Li₄P that occurred at a cell voltage above 0.8 V. After 20 cycles, the efficiencies for all films reached approximately 95%. After 40 cycles, the film tested at 0–0.75 V shows an efficiency as high as 97–98% and maintains it stably in whole measurement range. This is consistent with the good cycling stability of this film (Figure 11). On the other hand, the films tested at 0–1 V and 0–1.25 V show a slightly lower efficiency of 96–97% after 40 cycles.

For further examination, we took the Sn₄P₃/C composite film electrodes out of the disassembled cells after cycling at different cell voltage windows and observed their microstructures by SEM (Figure 13). As shown in Figure 13a, no delamination of the film from the SUS316L substrate was observed after cycling at 0–0.75 V. In addition, the structural change of the film cycled at 0–0.75 V is less than that for the film cycled at 0–1.25 V, which contributes to the better cycling performance. For the film cycled at 0–1.25 V with the lowest capacity retention (Figure 13b), delamination of many
parts of the film from the substrate is confirmed and the large transversal cracks were generated at the location without peeling.

![Coulombic efficiency graph](image)

**Figure 12.** Coulombic efficiencies plotted against the cycle numbers for Sn$_4$P$_3$/C composite films tested at different cell voltage windows.

**Figure 13.** SEM images for the broader surface of Sn$_4$P$_3$/C composite films taken out of disassembled cells: (a) cycled at 0–0.75 V and (b) cycled at 0–1.25 V. Insets are the photo images of films.

Lastly, we compare the electrochemical performance of some Sn$_4$P$_3$ anode materials for LiBs reported in the literature [8,9,11,14–17,20–24], which is listed in Table 1. It is worth noting that our current results reported in this paper are not the top performance for Sn$_4$P$_3$ anode materials for LiBs. As reported in [21], Sn$_4$P$_3$/C nanospheres synthesized by carbonization/reduction and phosphorization of SnO$_2$–GCP (glucose-derived, carbon-rich polysaccharide) nanospheres showed an outstanding rate performance and cycling stability. These nanospheres can also be applicable for ultra-stable anode materials for sodium-ion batteries (SiBs). However, as mentioned above, the electrodes with these Sn$_4$P$_3$/C composite anode materials used in batteries are fabricated by a slurry coating process with a large amount of carbon additives and binders [20–24], resulting in the decrease of the fraction of Sn$_4$P$_3$ in the electrode. For the Sn$_4$P$_3$/C composite anode listed in Table 1, gravimetric capacities calculated by the total mass of electrode (including both carbon additives and binders) are 20–30% lower than the listed values. On the other hand, Sn$_4$P$_3$/C composite film electrodes formed by AD do not contain other carbon additives and binders, resulting in higher gravimetric capacity for the whole
electrode. There is room to improve the electrochemical performance of the size and content of carbon materials for Sn₄P₃/C powders used for AD. We are now trying to optimize complexed carbon content and increase the composite film thickness and the progress will be reported in a forthcoming paper.

**Table 1.** Comparison of electrochemical performance of some Sn₄P₃ anode materials for Li-ion batteries (LiB). AD: aerosol deposition.

| Samples                                      | Current Density/mA g⁻¹ | Cycle Numbers | Specific Capacity/mAh g⁻¹ | References |
|----------------------------------------------|------------------------|---------------|---------------------------|------------|
| Sn₄P₃                                        | 100                    | 50            | 370                       | [8]        |
| Sn₄+xP₃                                      | 100                    | 50            | 530 (x = 1)               | [9]        |
| Sn₄P₃ film by pulsed laser deposition (PLD)  | 0.2 mA cm⁻²            | 10            | 553                       | [11]       |
| Fe doped Sn₄P₃                              | 100                    | 100           | 420                       | [14]       |
| Mn doped Sn₄P₃                              | 100                    | 150           | 488                       | [15]       |
| Sn₄P₃                                       | 100                    | 20            | 261                       | [16]       |
| Sn₄P₃                                       | 100                    | 300           | 442                       | [17]       |
| Sn₄P₃/graphite                              | 100                    | 100           | 651                       | [20]       |
| Sn₄P₃/C nanosphere                          | 2000                   | 50            | 1050                      |            |
| Sn₄P₃/SnO₂–C                                | 400                    | 200           | 733                       | [22]       |
| Sn₄P₃/hollow graphene sphere                 | 100                    | 100           | 606                       | [23]       |
| Sn₄P₃/N doped C                             | 100                    | 120           | 718                       | [24]       |
| Sn₄P₃/C film by AD                          | 50                     | 400           | 380                       | This work  |

4. Conclusions

Sn₄P₃/C composite film was successfully fabricated by the AD method and its electrochemical performance for a lithium-ion battery anode was examined. The Sn₄P₃/C composite film fabricated by AD showed much better electrochemical performance than the Sn₄P₃ film without complexing carbon. Although both films showed initial discharge (Li⁺ extraction) capacities of approximately 900–1000 mAh g⁻¹, Sn₄P₃/C films retained the higher reversible capacity above 700 mAh g⁻¹ after 100 cycles of charge and discharge processes while the capacity of the Sn₄P₃ film rapidly degraded with cycling. Precise control of the potential window in galvanostatic testing of the Sn₄P₃/C composite film results in remarkable improvement in the cycling performance. We obtained a reversible capacity of approximately 400 mAh g⁻¹ after 400 cycles by controlling the cell potential window, which is mainly attributed to the suppression of structural change of the film electrode during the cycling.

Supplementary Materials: The following are available online at http://www.mdpi.com/2079-4991/9/7/1032/s1, Figure S1: (a) SEM image of broader surface (left) and transverse cross section (right) of the Sn₄P₃ film fabricated by AD and (b) Galvanostatic charge and discharge curves for the Sn₄P₃ film, Figure S2: (a) Galvanostatic charge and discharge curves for the Sn₄P₃/C composite film (Sn₄P₃:AB = 9:1 in weight) and (b) Comparison of cycling stability for Sn₄P₃/C composite films with different carbon content.

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