

Engineering artificial solid electrolyte interface via carbon coating and VC additive for SnO₂ thin-film lithium-ion anodes

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Lithium-ion microbatteries (LIBs) are energy storage systems that can power at microscale. They should have a high energy density and show durable cycling behavior [1]. Tin dioxide (SnO₂) is of particular interest as an anode material, as it has a high theoretical capacity of 1494 mAhg⁻¹, which significantly exceeds the performance of traditional graphite anodes. Furthermore, SnO₂ is an accessible, inexpensive, and environmentally friendly material, making it promising for the creation of next-generation environmentally friendly microbatteries [2]. However, the huge volume changes during lithium ion insertion/extraction led to capacity decay due to materials pulverization and solid electrolyte interface (SEI) redormation.

In this study, we attempted to build stable SEI layer, which can prolong cycle life of SnO₂ anode. SnO₂ thin-film anodes were obtained by RF magnetron sputtering, followed by thermal treatment to increase crystallinity and enhance surface morphology. First, the film surface was coated with carbon in order to prevent the initial side reactions on the SnO₂ surface. Secondly, the vinylene carbonate (VC) additive was added to electrolyte to create more smooth inbreakable SEI layer.

Electrochemical performance was evaluated using cyclic voltammetry and galvanostatic charge-discharge measurements conducted at a current rate of 0.1 C in the voltage range of 0.01-3.0 V. A comparative study was conducted to evaluate the electrochemical properties of C-coated SnO₂ anodes in two types of electrolytes: a bare 1 M LiPF₆ in a mixture of EC:DEC:EMC (1:1:1 by volume) and the same electrolyte mixture but with the addition of 5 wt.% VC. Thus, the electrode was covered by carbon and then additional VC-based layer during SEI formation in the initial cycles. The obtained data indicate a significant influence of the each layers on the behavior of the anode.

Figure 1 shows SEM images of the surface of thin films. The pure SnO₂ sample showed a porous surface with visible

microcracks, caused by thermal treatment. In comparison, the C-coated SnO₂ had a morphology that is more dense and even, with a smoother surface and fewer cracks, indicating increased structural stability.

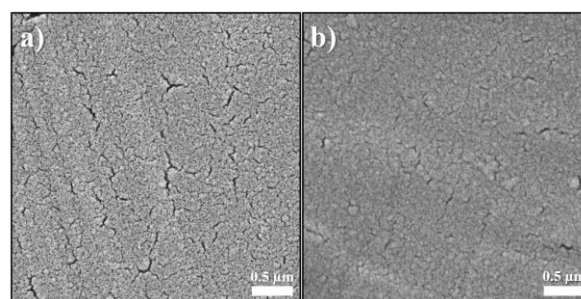


Figure 1. SEM surface images of SnO₂ (a) pure SnO₂, (b) C-coated SnO₂ thin film.

The presented electrode material exhibits outstanding stability and capacity retention, making it a promising candidate for next-generation energy storage systems.

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References

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