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Enhanced Electrochemical Performance of α -Fe $_2$ O $_3$ Grains Grafted onto TiO $_2$ -Carbon Nanofibers via a Vapor-Solid Reaction as Anode Materials for Li-Ion Batteries

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Abstract

α-Fe₂O₃ grains grafted onto TiO₂/carbon nanofibers (CNFs) for use as anode materials in lithium-ion batteries have been successfully fabricated by electrospinning and vapor-solid reaction (VSR). Scanning electron microscopy (SEM), transmission electron microscopy (TEM) and N₂ adsorption-desorption isotherms reveal that the ultrafine α-Fe₂O₃ nanoparticles were formed on the TiO₂/CNFs and have uniform dispersion along the fiber direction. The VSR approach could retard nucleation, thus making TiO₂/CNFs with small Fe₂O₃ grains grafting (approximately 5 nm in diameter). The TiO₂/CNFs are capable of buffering the large volume variation of α-Fe₂O₃ during cycling and preventing electrode pulverization and aggregation, as well as providing sufficiently large interstitial space within the crystallographic structure to host Li ions. The electrochemical properties of the composite electrodes were tested by galvanostatic cycling at both constant and variable current rates. The composite delivers both good rate capability under an uprated current density of 1000 mA g⁻¹ and especially enhanced cycle stability (~600 mAh g⁻¹ after 200 cycles at a current density of 1000 mA g⁻¹). The super electrochemical performance is attributed to a synergetic effect between α-Fe₂O₃ and TiO₂-CNFs as well as the threedimensional (3D) network, which contributes to greatly enhanced diffusion kinetics and structural stability for lithium-ion batteries. This VSR approach can be extended to other hierarchical metal oxide nanostructures for favorable applications in

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