Oxidized Co–Sn nanoparticles as long-lasting anode materials for lithium-ion batteries†

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Herein, we present the synthesis and systematic comparison of Sn- and Co–Sn-based nanoparticles (NPs) as anode materials for lithium-ion batteries. These nanomaterials were produced via inexpensive routes combining wet chemical synthesis and dry mechanochemical methods (ball milling). We demonstrate that oxidized, nearly amorphous CoSn2O x NPs, in contrast to highly crystalline Sn and CoSn2 NPs, exhibit high cycling stability over 1500 cycles, retaining a capacity of 525 mA h g−1 (92% of the initial capacity) at a high current density of 1982 mA g−1. Moreover, when cycled in full-cell configuration with LiCoO2 as the cathode, such CoSn2O x NPs deliver an average anodic capacity of 576 mA h g−1 over 100 cycles at a current of 500 mA g−1, with an average discharge voltage of 3.14 V.

Introduction

Despite an intensive research effort to develop new materials for rechargeable lithium-ion batteries (LIBs) over the past four decades, graphite remains the most common anode material in commercial devices. This is a surprising circumstance considering the fact that graphite possesses relatively low gravimetric and volumetric capacities (372 mA h g−1 and 820 mA h cm−3, respectively) compared to a large number of both alloying- (e.g., Si, Ge, and Sn) and conversion-type materials (e.g., Fe3O4, MoS2, and SnSb).1,2 The primary reasons for this are the rapid decline in capacity of many of these alternative anode materials due to massive volume changes during cycling (e.g., ΔV = 100–300%) causing mechanical disintegration of the electrodes, and/or due to irreversible reactions during charge and discharge. It has been demonstrated in a variety of case studies that these issues can be mitigated by using nanostructuring strategies in the synthesis of the active anode materials.3–26 Nevertheless, the commercialization of such high-capacity alloying- and conversion-type anodes has remained hampered for several reasons. In conversion-type anodes, a major fraction of the capacity is often obtained at potentials above 1.0 V vs. Li+/Li, reducing the voltage of the full-cell and thus the energy density of the corresponding battery. Secondly, the synthesis of nanostructured electrodes is often elaborate and therefore costly. Tin remains among the few materials which are realistic candidates to replace graphite in commercial LIBs, due to its high gravimetric and volumetric capacities (992 mA h g−1 and ~7300 mA h cm−3), low delithiation potential (0.2–1 V), high electrical conductivity, low cost and natural abundance. In fact, anodes comprising amorphous Sn–Co–C nanocomposites have indeed been employed in commercial rechargeable batteries (e.g., Nexelion™, Sony Corp., Japan) since at least 2005. The success of this product has triggered further intensive research on Co–Sn-based anodes for LIBs.27–54 The superior cycling stability of many Co–Sn-based materials over elemental Sn has been attributed to two factors: (1) the effective buffering of volume changes occurring during the lithiation of Sn by the inactive Co matrix and (2) the prevention of aggregation of Sn particles during delithiation due to the preferred formation of intermetallic Co–Sn phases.28,55–58 Notable recent examples of nanostructured Co–Sn-based materials include, for instance, CoSnO3 NPs in a graphene network47 and Sn–Co NPs encapsulated in carbon spheres,59 which both exhibit stable capacities over ≥100 cycles in a wide potential range of up to 3.0 V vs. Li+/Li.

In this report, we present a systematic comparison of well-defined Co–Sn nanoparticle (NP) based materials as anodes for LIBs. We introduce a simple synthetic procedure which can be used to prepare either amorphous Co NPs or crystalline Sn NPs, and subsequently demonstrate their conversion into crystalline non-oxidized CoSn2 NPs or oxidized, nearly amor-
pous CoSn$_2$O$_2$ NPs by simple mechanochemical methods. Half-cell electrochemical experiments were carried out to compare pure Sn NPs to both Co–Sn NP systems, indicating superior characteristics of the CoSn$_2$NPs. Finally, full-cell experiments were carried out on this most promising system using LiCoO$_2$ as the cathode; stable cycling with anodic capacities of 576 mA h g$^{-1}$ for 100 cycles could be demonstrated at a current of 500 mA g$^{-1}$, with an average discharge voltage of 3.14 V. Note: In the work described herein, “CoSn$_2$ NPs” is used to refer to the highly crystalline, non-oxidized CoSn$_2$ material (seen by X-ray diffraction to be phase-pure), while the term “CoSn$_2$O$_2$ NPs” is used for oxidized and mostly amorphous NPs (with $<$5 nm crystal domain sizes).

Experimental

Colloidal synthesis of Sn and Co NPs

To prepare Sn NPs, a solution of NaBH$_4$ (96 mmol, 98%, ACB) in anhydrous 1-methyl-2-pyrrolidone (NMP, 85 mL, 99.5%, Fisher BioReagents) was heated to 60 °C under flowing N$_2$. Then, a solution of SnCl$_2$ (1 mmol, ≥99%, Alfa Aesar) in anhydrous NMP (3 mL) was injected and the reaction mixture was immediately cooled to room temperature using a water-ice bath. To prepare Co NPs, a NaBH$_4$ solution in anhydrous NMP (32 mmol in 15 mL) was heated to 150 °C under flowing N$_2$, followed by the injection of CoCl$_2$ (8 mmol, ≥98%, Sigma-Aldrich) and immediate cooling to room temperature using a water-ice bath. The obtained Co NPs and Sn NPs were purified by washing once with dimethyl sulfoxide and then two times with water after separation from the supernatant by centrifugation. Finally, the reaction product was dried under vacuum at room temperature.

Synthesis of crystalline CoSn$_2$ NPs and oxidized CoSn$_2$O$_2$ NPs

For the preparation of Co–Sn based NPs, Sn NPs (1.4 mmol) were ball-milled for 4 hours with Co NPs (0.7 mmol) at a frequency of 30 s$^{-1}$ using a Fritsch Pulverisette 23 mill (10 mL ZrO$_2$ vessel, loaded with two 10 mm ZrO$_2$ balls). In order to prepare crystalline CoSn$_2$ NPs, the vessel was loaded and sealed under nitrogen atmosphere. Oxidized CoSn$_2$O$_2$ NPs were obtained when the starting Co and Sn NPs were loaded in air.

Electrode fabrication, cell assembly and electrochemical measurements

The following battery components were used: carbon black (CB, Super C65, TIMCAL), carboxymethylcellulose (CMC, Grade: 2200, Daicel Fine Chem Ltd), fluoroethylene carbonate (FEC, Solvay, battery grade), 1 M LiPF$_6$ in ethylene carbonate : dimethyl carbonate (EC : DMC, 1 : 1 by volume, Merck, battery grade), glass microfiber separator (GF/D, Whatman), and Cu foil (9 µm, MTI Corporation). For electrode preparation, aqueous slurries were prepared by mixing the respective NPs (64 wt%) with CB (21 wt%), CMC (15 wt%) and water using a planetary ball-mill (Fritsch Pulverisette 7 mill), and then coated onto Cu current collectors. The electrodes were then dried at 80 °C under vacuum for 12 h prior to assembling the cells. Material loadings were 0.5 mg cm$^{-2}$ for half-cell and 1 mg cm$^{-2}$ for full-cell experiments. Electrochemical measurements were conducted in air-tight coin-type cells assembled in an Ar-filled glove box (O$_2$ < 0.1 ppm, H$_2$O < 0.1 ppm) using elemental lithium as the counter electrode for half-cells and LiCoO$_2$ on Al foil (MTI Corporation, ~20 mg cm$^{-2}$) as the cathode in full-cells. Glass microfiber was used as the separator in all cases. A standard solution of 1 M LiPF$_6$ in EC : DMC with 3% FEC was used as the electrolyte. Galvanostatic cycling tests were performed at room temperature on a MPG2 multichannel workstation (BioLogic). Anodic capacities were determined corresponding to the mass of the Co–Sn material in both half- and full-cell experiments.

Materials characterization

Transmission electron microscopy (TEM) images were obtained with a Philips CM30 microscope operated at 300 kV, using carbon-coated Cu grids as substrates (Ted-Pella). Energy-dispersive X-ray spectroscopy (EDX) measurements were carried out using a NanoSEM 230 scanning electron microscope. Scanning transmission electron microscopy (STEM) images and EDX spectroscopy maps were collected on a FEI Talos F200X microscope operated at 200 kV. Powder X-ray diffraction (XRD) was measured on a STOE STADI P diffractometer (Cu-Ko irradiation, $\lambda = 1.540598$ Å). X-ray photoelectron spectroscopy (XPS) measurements were carried out in normal emission using a monochromatized Al Ka X-ray radiation source and a Scienta R3000 display analyzer.

Results and discussion

Inexpensive routes to Co–Sn-based NPs were derived by combining the wet colloidal synthesis of unary metallic Co and Sn NPs with dry mechanochemical reactions between them (Fig. 1a), conducted in or without the presence of air. Unary metallic NPs of Co (amorphous) and Sn (crystalline) were synthesized by an extension of the method that we previously reported for Sb NPs$^{60}$ based on the borohydride reduction of metal chlorides in NMP. In contrast to the reduction reactions of SbCl$_3$ and SnCl$_2$ which exhibited suitable kinetics at 60 °C, a higher temperature of 150 °C was required for CoCl$_2$. The resulting Co NPs were amorphous (Fig. 1b, see also Fig. S1 and S2a and b†), whereas the Sn NPs were highly crystalline in the form of β-Sn (Fig. 1b, indexed as phase-pure tetragonal Sn, space group I41/amd (141), $a = 5.831$ Å, $c = 3.182$ Å, ICDD PDF entry no.: 00-004-0673). High-resolution transmission electron microscopy (HR-TEM) image of Sn NP along with selected area electron diffraction (SAED) and d-spacing is shown on the Fig. S2c and d.$^7$ Mixtures of Co and Sn NPs (molar ratio 1:2) were then ball-milled either in air or under a nitrogen atmosphere with the intention of intimately mixing and alloying these materials and to study the effects of oxidation. Synthesis in air yielded a largely amorphous/poorly crystalline product, with broad and weak
XRD reflections corresponding to SnO₂ and CoSn₂. For simplicity, the resulting product is denoted as CoSn₂O₅ through-out this work. In agreement with the XRD results (Fig. 1b), d-spacing values deduced from HR-TEM measurements of CoSn₂O₅ NPs indicate the presence of small crystalline domains composed of CoSn₂ and SnO₂ (Fig. S3†). Elemental mapping with energy-dispersive X-ray spectroscopy in scanning TEM (EDX-STEM) studies confirms oxidation throughout the material, with a rather homogeneous distribution of Co, Sn, and O (Fig. S4†). As follows from XPS spectra (see Fig. S5†), both cobalt and tin are completely oxidized on the surface, most probably forming Co(OH)₂ and SnO₂. In addition, minor amount of metallic tin was observed in agreement with XRD results. In comparison with tin, cobalt is hardly detectable by XPS. Ball-milling under inert conditions, however, resulted in the formation of highly crystalline CoSn₂ NPs (Fig. 1, see also Fig. S2e and f†), with only a minor content of SnO₂. In all cases, the NPs obtained were small: 4–7 nm for Co NPs, 5–10 nm for Sn NPs and 6–20 nm for CoSn₂O₅ NPs and CoSn₂ NPs (Fig. 2 and see also Fig. S6†). Importantly, as control experiments, attempts to synthesize Co–Sn nanomaterials with commercial microcrystalline powders of Co and Sn did not result in the formation of the CoSn₂ phase (Fig. S7†).

Lithium-ion half-cell experiments
In order to compare the electrochemical properties of Sn, CoSn₂O₅, NPs and CoSn₂ NPs, Li-ion half-cells were assembled. Electrodes were composed of NPs, carbon black (CB), and CMC binder (in a ratio of 64:21:15 by weight), and were tested against elemental lithium using 1 M LiPF₆ in EC: DMC. FEC was added to the electrolyte to improve cycling stability by forming a stable SEI layer.⁶¹–⁶⁵ All specific capacities and currents presented herein correspond to the combined mass of Sn and Co, excluding CB and CMC. Fig. 3 shows the capacity retention performance of CoSn₂O₅ NPs compared to Sn NPs over 1500 cycles at a high current density of 1984 mA g⁻¹ in the potential range of 0.005–1.0 V vs. Li⁺/Li.

Since lithium is reversibly stored in all of the systems studied herein primarily via the Sn → Li₄.₄Sn conversion reaction (theoretical capacity: 992 mA h g⁻¹), a current density...
of 992 mA g\(^{-1}\) was designated as the 1C rate.\(^{66}\) Assuming that Co does not contribute to any Li-ion capacity, the theoretical capacity of CoSn\(^2\) is 795 mA h g\(^{-1}\). During galvanostatic cycling of the half-cells, the upper cut-off potential was limited to 1.0 V in order to include the upper region relevant to subsequent full-cell experiments. As can be seen in Fig. 3, at a high current density of 1984 mA g\(^{-1}\) (“2C”), Sn NPs, CoSn\(^2\)O\(_x\) NPs, and CoSn\(^2\) NPs initially show similar capacities of 550–600 mA h g\(^{-1}\), with the highest values typically reached after 50–100 cycles. The increase of the capacities in the initial cycles might be attributed to restructuring processes in the electrode, which lower the resistivity and hence lead to a higher utilization of the capacity. Upon extended cycling, Sn NPs exhibit significant capacity fading within 400 cycles, while Co–Sn-based NPs show a much better retention of their capacity. Specifically, CoSn\(^2\) NPs retain a capacity of 462 mA h g\(^{-1}\) after 1500 cycles (corresponding to a reduction of 18%). CoSn\(^2\)O\(_x\) NPs even exhibit better cyclability, retaining a capacity of 525 mA h g\(^{-1}\) (a reduction of 8%) after 1500 cycles. It should be noted that for all three systems, the average coulombic efficiency (CE) is 99.6% during cycling. First-cycle CE is, however, as low as ~30%, which can be attributed to the formation of solid electrolyte interface (SEI) and the reduction of surface oxides to their corresponding metals with Li\(_2\)O as a byproduct.

With such a high cycling stability, the CoSn\(^2\)O\(_x\) NPs in this work compare favorably to the majority of recently investigated Sn-based Li-ion anode materials (see Table S1 for a detailed comparison), especially when considering that the cycling performed herein was restricted to potentials below 1.0 V vs. Li\(^+\)/Li in order to obtain values that are truly relevant for full-cells.\(^{5,47–54,67–72}\) This superior cycling stability of NPs incorporating Co and Sn compared to pure Sn NCs might be attributable to two effects. Firstly, due to the fact that it does not form lithium alloys, Co can serve as an inactive matrix during cycling and therefore buffer the volume changes caused by the lithiation/delithiation of Sn. Secondly, the presence of Co can prevent Sn NCs from aggregating upon delithiation and therefore further improve the retention of the starting structure of the active anode material.\(^{28,55,56}\) Further, cyclic voltammetry (CV) measurements indicate a third possible effect: lithiation/delithiation occurs through fewer (most likely amorphous) phase transitions for Co–Sn-based NPs compared to Sn NCs (Fig. S9–11 at). It is well known that pure Sn forms a multitude of intermediate crystalline phases during lithiation, leading to increased anisotropic strain in the particles during cycling and therefore lower cycling stability.

The difference between CoSn\(^2\) NPs and CoSn\(^2\)O\(_x\) NPs in terms of cycling stability might be attributed to the fact that CoSn\(^2\)O\(_x\) NPs are highly oxidized. The surface oxides are likely converted into Li\(_2\)O during the first cycle(s), which is also known to inhibit the sintering of Sn domains during cycling.\(^{66}\) It should be noted that the average delithiation potentials for Sn NCs, CoSn\(^2\) NPs, and CoSn\(^2\)O\(_x\) NPs are equally low, with an average value of ~0.5 V vs. Li\(^+\)/Li (Fig. 3b–d).

Apart from the mechanical ball-milling of Co and Sn NPs, crystalline CoSn\(^2\) NPs can also be synthesized by the same colloidal method as used herein to synthesize the pure Co and Sn NPs by simultaneous injection of SnCl\(_2\)/CoCl\(_2\) into a solution of NaBH\(_4\) in NMP. However, such a synthesis strategy suffers from an imbalance between the reactivities of the two metal chlorides. A high temperature of 150 °C is necessary to reduce Co\(^{2+}\), and the resulting CoSn\(^2\) NPs are larger than 20 nm, which is significantly larger than the Co–Sn based NPs prepared by ball-milling (see Fig. S12†). As a result of the larger size of colloidally synthesized CoSn\(^2\) NPs, much poorer capacity retention was observed: the capacity reduced to <200 mA h g\(^{-1}\) after 100 cycles (see Fig. S13†).

To evaluate the rate capability of the Co–Sn based NPs developed in this work, galvanostatic cycling was performed at current rates between 0.2C to 10C (Fig. 4, 1C = 992 mA g\(^{-1}\)). Due to the similar particle sizes of these systems, and therefore similar reaction kinetics in all cases, comparable rate capabilities were observed. The only exception was at currents of 0.5C–2C where Sn NPs showed ~50 mA h g\(^{-1}\) lower capacities compared to CoSn\(^2\) NPs. Even at rates as high as 10C, all three materials still retained a capacity of ~350 mA h g\(^{-1}\).

Interestingly, it was observed that at such high currents, lithium capacities increased during cycling, resulting in the same or even higher capacities during the stepwise decrease of the rate back to 0.2C. Especially for CoSn\(^2\)O\(_x\) NPs, the slight difference in capacity compared to CoSn\(^2\) NPs initially observed at rates of 0.5C–2C becomes fully diminished during cycling. For comparison, graphite is known to exhibit much poorer rate capability,\(^{73,74}\) which was confirmed in this work by control experiments using graphite anodes (TIMCAL, K86) tested under identical conditions (Fig. S14†).

**Lithium-ion full-cell experiments**

In order to investigate the applicability of Co–Sn-based NPs as anode materials for commercial batteries, anode-limited full-cells using LiCoO\(_2\) as the cathode were assembled (Fig. 5). Based on the results of the half-cell experiments, CoSn\(^2\)O\(_x\) NPs were selected for primary investigation in full-cell experiments.

**Fig. 4** Rate capability tests for Sn, CoSn\(^2\), and CoSn\(^2\)O\(_x\) NPs in lithium-ion half-cells within the potential range of 0.005–1.0 V.
Herein, all specific capacities and currents correspond to the mass of CoSn₂Oₓ NPs. Full-cells of CoSn₂Oₓ/LiCoO₂ were cycled galvanostatically at a current of 500 mA g⁻¹ in the potential range of 2.0–3.9 V. Under such conditions, CoSn₂Oₓ NPs exhibit an average capacity of 576 mA h g⁻¹ upon extended cycling, similar to the values observed in half-cell experiments. Neglecting the excess of cathode material used in this work, the theoretical charge storage capacity of the cell is estimated as 112.6 mA h g⁻¹ based on: \( C_{\text{cell}} = C_{\text{anode}}C_{\text{cathode}}/(C_{\text{anode}} + C_{\text{cathode}}) \). Taking the average discharge voltage of 3.14 V into account, the resulting average specific energy density for the CoSn₂Oₓ/LiCoO₂ cell is 353 W h kg⁻¹ and, most importantly, it remains stable at this value for 100 cycles (see Fig. 5c). This specific energy density is comparable to that of the state-of-the-art graphite/LiCoO₂ system (∼360 W h kg⁻¹ based on 140 mA h g⁻¹/3.7 V vs. Li⁺/Li for LiCoO₂ and 372 mA h g⁻¹/0.15 V vs. Li⁺/Li for graphite).⁷⁵ However, considering the much higher density of bulk β-Sn (7.3 g cm⁻³) and Co (8.9 g cm⁻³) compared to graphite (2.2 g cm⁻³), CoSn₂Oₓ NPs can theoretically exhibit improved volumetric energy densities by up to ∼40% (see also Table S2†). For a large variety of portable electronic devices, the importance of volumetric energy density is greater than that of gravimetric energy density.⁷⁶ For comparison, the improvement of the volumetric energy density of the Sony Nexelion™ battery is 20% over identical cells using graphite anodes.⁷⁷

Conclusions

Co and Sn NPs with diameters of ≤10 nm were synthesized via the simple reduction of their respective chlorides using NaBH₄ in NMP, and were subsequently converted into intermetallic Co–Sn NPs by ball-milling. The resulting nanostructured materials can be seen as well-defined model systems, suited for investigating the effects of crystallinity and composition on electrochemical properties upon lithiation/delithiation cycling. Despite the fact that Sn and CoSn₂ NPs show good cycling stability for several hundred cycles, CoSn₂Oₓ NPs show the most outstanding retention of capacity, losing only 8% of their initial capacity over 1500 cycles at 1984 mA g⁻¹. In addition, in lithium-ion full-cell experiments with LiCoO₂ as the cathode material, CoSn₂Oₓ NPs provide capacities of on average 576 mA h g⁻¹ with an average discharge voltage of 3.14 V. This system, therefore, exhibits stable specific energy densities comparable to state-of-the-art LIBs based on graphite, and potentially much higher volumetric energy densities due to the higher density of CoSn₂Oₓ. Considering that high rates were used in both half- and full-cell experiments, the CoSn₂Oₓ NPs presented herein also offer a potential improvement in power density over cells assembled using conventional graphite anodes and also over Sony’s commercialized Nexelion™ battery which has been designed specifically for low power (e.g., camcorder) devices.⁷⁷

Conflicts of interest

There are no conflicts to declare.

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