



Cite this: *Nanoscale*, 2018, **10**, 3777

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

Marc Walter,^{a,b} Simon Doswald,^{a,b} Frank Krumeich,^a Meng He,^{a,b} Roland Widmer,^c Nicholas P. Stadie^{†‡} and Maksym V. Kovalenko^{*,a,b}

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.

Received 30th September 2017,
Accepted 31st January 2018

DOI: 10.1039/c7nr07309g

rscl.li/nanoscale

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-

^aDepartment of Chemistry and Applied Biosciences, ETH Zürich – Swiss Federal Institute of Technology Zürich, Vladimir Prelog Weg 1, 8093 Zürich, Switzerland. E-mail: mvkovalenko@ethz.ch

^bEmpa-Swiss Federal Laboratories for Materials Science and Technology, Laboratory for thin films and photovoltaics, Überlandstrasse 129, 8600 Dübendorf, Switzerland

^cEmpa-Swiss Federal Laboratories for Materials Science and Technology, Nanotech@surfaces Laboratory, Überlandstrasse 129, 8600 Dübendorf, Switzerland

†Electronic supplementary information (ESI) available. See DOI: 10.1039/c7nr07309g

‡Present address: Department of Chemistry and Biochemistry, Montana State University, 59717 Montana, USA.



phous CoSn_2O_x 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_2O_x 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_2O_x NPs” is used for oxidized and mostly amorphous NPs (with $<5 \text{ nm}$ crystal domain sizes).

Experimental

Colloidal synthesis of Sn and Co NPs

To prepare Sn NPs, a solution of NaBH_4 (96 mmol, 98%, ACBR) in anhydrous 1-methyl-2-pyrrolidone (NMP, 85 mL, 99.5%, Fisher BioReagents) was heated to $60 \text{ }^\circ\text{C}$ under flowing N_2 . Then, a solution of SnCl_2 (1 mmol, $\geq 99\%$, 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 $150 \text{ }^\circ\text{C}$ under flowing N_2 , followed by the injection of CoCl_2 (8 mmol, $\geq 98\%$, 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_2O_x 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_2O_x 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 $80 \text{ }^\circ\text{C}$ 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 ($\text{O}_2 < 0.1 \text{ ppm}$, $\text{H}_2\text{O} < 0.1 \text{ ppm}$) using elemental lithium as the counter electrode for half-cells and LiCoO_2 on Al foil (MTI Corporation, $\sim 20 \text{ mg cm}^{-2}$) 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 multi-channel 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- $\text{K}\alpha_1$ irradiation, $\lambda = 1.540598 \text{ \AA}$). X-ray photoelectron spectroscopy (XPS) measurements were carried out in normal emission using a monochromatized Al $\text{K}\alpha$ 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 NPs⁶⁰ based on the borohydride reduction of metal chlorides in NMP. In contrast to the reduction reactions of SbCl_3 ⁶⁰ and SnCl_2 which exhibited suitable kinetics at $60 \text{ }^\circ\text{C}$, a higher temperature of $150 \text{ }^\circ\text{C}$ 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), $a = 5.831 \text{ \AA}$, $c = 3.182 \text{ \AA}$, 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.† 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



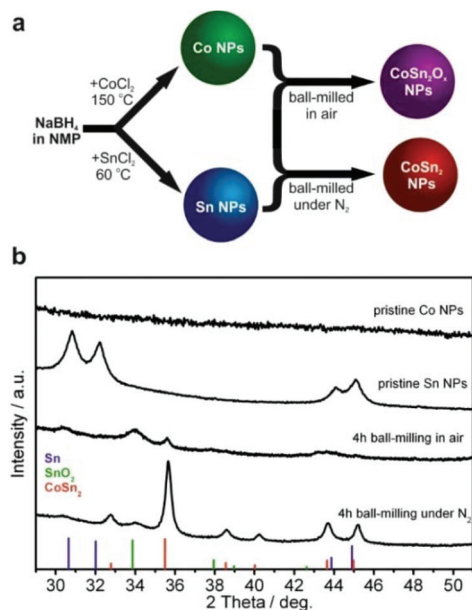


Fig. 1 (a) Schematic of the preparation of Co–Sn NPs. (b) X-ray diffraction (XRD) patterns of colloiddally synthesized Co and Sn NPs and mechanochemically prepared CoSn₂O_x and crystalline CoSn₂ NPs. Reference patterns: tetragonal SnO₂, space group *P42/mnm* (136), *a* = 4.7391 Å, *c* = 3.1869 Å, ICDD PDF entry 00-077-0448; tetragonal CoSn₂, space group *I4/mcm* (140), *a* = 6.363 Å, *c* = 5.456 Å, ICDD PDF entry 00-025-0256.

XRD reflections corresponding to SnO₂ and CoSn₂. For simplicity, the resulting product is denoted as CoSn₂O_x NPs throughout this work. In agreement with the XRD results (Fig. 1b), *d*-spacing values deduced from HR-TEM measurements of CoSn₂O_x 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 CoSn₂O_x 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, CoSn₂O_x NPs and CoSn₂ NPs, Li-ion half-cells were assembled. Electrodes were composed of NPs, carbon black (CB), and

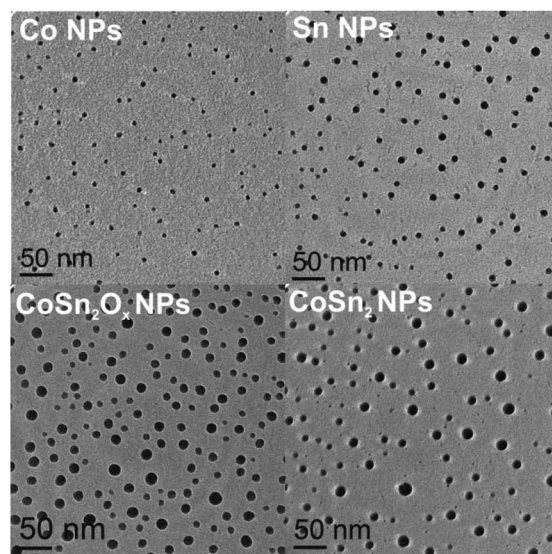


Fig. 2 Transmission electron microscopy (TEM) images of colloiddally synthesized Co and Sn NPs and mechanochemically prepared CoSn₂O_x and CoSn₂ NPs.

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 CoSn₂O_x 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 → Li_{4.4}Sn conversion reaction (theoretical capacity: 992 mA h g⁻¹), a current density

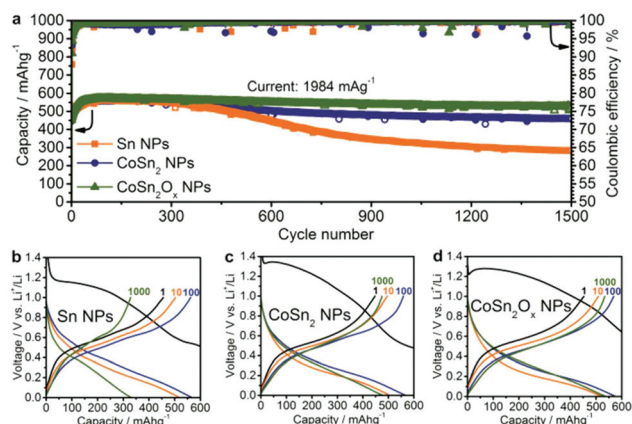


Fig. 3 (a) Cycling stability measurements of Sn NPs, CoSn₂ NPs and CoSn₂O_x NPs in lithium-ion half-cells at a current of 1984 mA g⁻¹ within the potential range of 0.005–1.0 V. Galvanostatic charge/discharge curves for (b) Sn NPs, (c) CoSn₂ NPs, and (d) CoSn₂O_x NPs after a number of cycles corresponding to the half-cells in (a) (1st discharge curves with full capacity range are shown on Fig. S8a–c†).



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 CoSn₂ 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 only the 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, CoSn₂O_x NPs, and CoSn₂ 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, CoSn₂ NPs retain a capacity of 462 mA h g⁻¹ after 1500 cycles (corresponding to a reduction of 18%). CoSn₂O_x NPs exhibit even 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 Li₂O as a byproduct.

With such a high cycling stability, the CoSn₂O_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.^{8,47–54,67–72} 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†). 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 CoSn₂ NPs and CoSn₂O_x NPs in terms of cycling stability might be attributed to the fact that CoSn₂O_x NPs are highly oxidized. The surface oxides are likely converted into Li₂O 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, CoSn₂ NPs, and CoSn₂O_x NPs are equally low, with an average value of ~0.5 V vs. Li⁺/Li (Fig. 3b–d).

Apart from by the mechanical ball-milling of Co and Sn NPs, crystalline CoSn₂ 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 SnCl₂/CoCl₂ into a solution of NaBH₄ 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 Co²⁺, and the resulting CoSn₂ 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 colloiddally synthesized CoSn₂ 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 CoSn₂ 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, Li-ion 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 CoSn₂O_x NPs, the slight difference in capacity compared to CoSn₂ 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, KS6) 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 LiCoO₂ as the cathode were assembled (Fig. 5). Based on the results of the half-cell experiments, CoSn₂O_x NPs were selected for primary investigation in full-cell experiments.

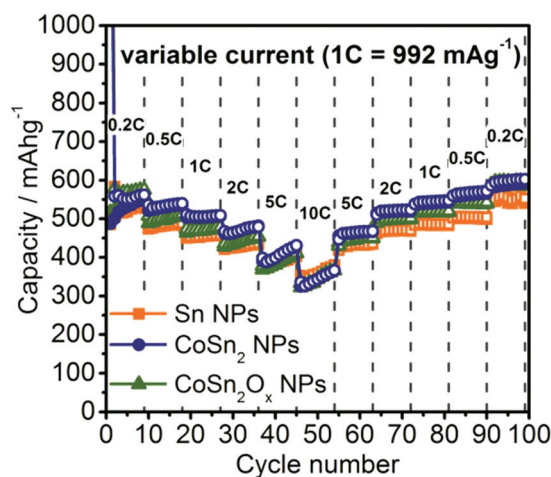


Fig. 4 Rate capability tests for Sn, CoSn₂, and CoSn₂O_x NPs in lithium-ion half-cells within the potential range of 0.005–1.0 V.



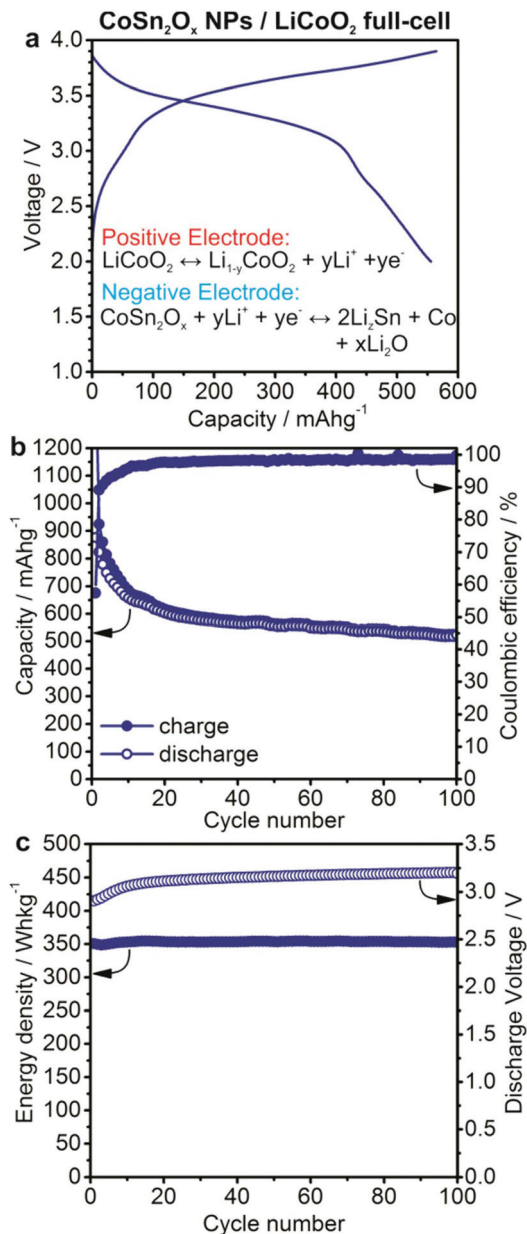


Fig. 5 Electrochemical performance of a lithium-ion full-cell battery with CoSn₂O_x NPs as the anode and LiCoO₂ as the cathode material. (a) Galvanostatic charge/discharge curves during the 50th cycle (see also Fig. S15†). (b) Capacity retention and coulombic efficiency. (c) Specific energy density and discharge voltage. Cells were cycled at a current of 500 mA g⁻¹ in the potential range of 2.0–3.9 V. Specific capacities and currents correspond to the mass of CoSn₂O_x NPs.

Herein, all specific capacities and currents correspond to the mass of CoSn₂O_x NPs. Full-cells of CoSn₂O_x/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, CoSn₂O_x 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 CoSn₂O_x/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), CoSn₂O_x 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, CoSn₂O_x 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, CoSn₂O_x 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 CoSn₂O_x. Considering that high rates were used in both half- and full-cell experiments, the CoSn₂O_x 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.

Acknowledgements

This work was financially supported by ETH Zürich (Grant No. ETH-56 12-2), the Competence Center for Energy and Mobility (CEEM, project SLIB), the Swiss Federal Commission for Technology and Innovation (CTI-Project No. 14698.2 PFIW-IW)



and the CTI Swiss Competence Centers for Energy Research (SCCER, Heat and Electricity Storage). Electron microscopy was carried out at the Scientific Center for Optical and Electron Microscopy (ScopeM) at ETH Zurich and at the Empa Electron Microscopy Center.

Notes and references

- J. Cabana, L. Monconduit, D. Larcher and M. R. Palacín, *Adv. Mater.*, 2010, **22**, E170–E192.
- N. Nitta and G. Yushin, *Part. Part. Syst. Charact.*, 2014, **31**, 317–336.
- P. G. Bruce, B. Scrosati and J.-M. Tarascon, *Angew. Chem., Int. Ed.*, 2008, **47**, 2930–2946.
- M. R. Palacín, *Chem. Soc. Rev.*, 2009, **38**, 2565–2575.
- A. Magasinski, P. Dixon, B. Hertzberg, A. Kvit, J. Ayala and G. Yushin, *Nat. Mater.*, 2010, **9**, 353–358.
- C. K. Chan, R. N. Patel, M. J. O'Connell, B. A. Korgel and Y. Cui, *ACS Nano*, 2010, **4**, 1443–1450.
- M. F. Oszajca, M. I. Bodnarchuk and M. V. Kovalenko, *Chem. Mater.*, 2014, **26**, 5422–5432.
- K. Kravchyk, L. Protesescu, M. I. Bodnarchuk, F. Krumeich, M. Yarema, M. Walter, C. Guntlin and M. V. Kovalenko, *J. Am. Chem. Soc.*, 2013, **135**, 4199–4202.
- N. Liu, Z. Lu, J. Zhao, M. T. McDowell, H.-W. Lee, W. Zhao and Y. Cui, *Nat. Nanotechnol.*, 2014, **9**, 187–192.
- A. Jahel, C. M. Ghimbeu, L. Monconduit and C. Vix-Guterl, *Adv. Energy Mater.*, 2014, **4**, 1400025.
- J. Liu, Y. Wen, P. A. van Aken, J. Maier and Y. Yu, *Nano Lett.*, 2014, **14**, 6387–6392.
- M. He, M. Walter, K. V. Kravchyk, R. Erni, R. Widmer and M. V. Kovalenko, *Nanoscale*, 2015, **7**, 455–459.
- M. He, K. Kravchyk, M. Walter and M. V. Kovalenko, *Nano Lett.*, 2014, **14**, 1255–1262.
- C. C. Li, Q. H. Li, L. B. Chen and T. H. Wang, *J. Mater. Chem.*, 2011, **21**, 11867–11872.
- C. Guo, L. Wang, Y. Zhu, D. Wang, Q. Yang and Y. Qian, *Nanoscale*, 2015, 10123–10129.
- G. Zeng, N. Shi, M. Hess, X. Chen, W. Cheng, T. Fan and M. Niederberger, *ACS Nano*, 2015, **9**, 4227–4235.
- J. Zhang, K. Wang, Q. Xu, Y. Zhou, F. Cheng and S. Guo, *ACS Nano*, 2015, **9**, 3369–3376.
- X.-Y. Yu, H. Hu, Y. Wang, H. Chen and X. W. Lou, *Angew. Chem., Int. Ed.*, 2015, 7395–7398.
- J. Zhou, J. Qin, X. Zhang, C. Shi, E. Liu, J. Li, N. Zhao and C. He, *ACS Nano*, 2015, **9**, 3837–3848.
- H. Hwang, H. Kim and J. Cho, *Nano Lett.*, 2011, **11**, 4826–4830.
- J. S. Cho, Y. J. Hong and Y. C. Kang, *ACS Nano*, 2015, **9**, 4026–4035.
- L. Wang, J. Liang, Y. Zhu, T. Mei, X. Zhang, Q. Yang and Y. Qian, *Nanoscale*, 2013, **5**, 3627–3631.
- J. B. Cook, E. Detsi, Y. Liu, Y.-L. Liang, H.-S. Kim, X. Petrissans, B. Dunn and S. H. Tolbert, *ACS Appl. Mater. Interfaces*, 2017, **9**, 293–303.
- Y. Jin, Y. Tan, X. Hu, B. Zhu, Q. Zheng, Z. Zhang, G. Zhu, Q. Yu, Z. Jin and J. Zhu, *ACS Appl. Mater. Interfaces*, 2017, **9**, 15388–15393.
- H. Zhang, X. Huang, O. Noonan, L. Zhou and C. Yu, *Adv. Funct. Mater.*, 2017, **27**, 1606023.
- H. Ying and W.-Q. Han, *Adv. Sci.*, 2017, **4**, 1700298.
- A. D. W. Todd, R. E. Mar and J. R. Dahn, *J. Electrochem. Soc.*, 2007, **154**, A597–A604.
- S.-I. Lee, S. Yoon, C.-M. Park, J.-M. Lee, H. Kim, D. Im, S.-G. Doo and H.-J. Sohn, *Electrochim. Acta*, 2008, **54**, 364–369.
- P. P. Ferguson, A. D. W. Todd and J. R. Dahn, *Electrochem. Commun.*, 2008, **10**, 25–31.
- P. P. Ferguson, M. L. Martine, A. E. George and J. R. Dahn, *J. Power Sources*, 2009, **194**, 794–800.
- Z. Chen, J. Qian, X. Ai, Y. Cao and H. Yang, *J. Power Sources*, 2009, **189**, 730–732.
- X.-Y. Fan, F.-S. Ke, G.-Z. Wei, L. Huang and S.-G. Sun, *J. Alloys Compd.*, 2009, **476**, 70–73.
- L. Huang, J.-S. Cai, Y. He, F.-S. Ke and S.-G. Sun, *Electrochem. Commun.*, 2009, **11**, 950–953.
- F. Nacimiento, R. Alcántara and J. L. Tirado, *J. Electrochem. Soc.*, 2010, **157**, A666–A671.
- J. He, H. Zhao, J. Wang, J. Wang and J. Chen, *J. Alloys Compd.*, 2010, **508**, 629–635.
- X.-L. Wang, W.-Q. Han, J. Chen and J. Graetz, *ACS Appl. Mater. Interfaces*, 2010, **2**, 1548–1551.
- F. Nacimiento, P. Lavela, J. Tirado and J. Jiménez-Mateos, *J. Solid State Electrochem.*, 2012, **16**, 953–962.
- G. Ferrara, L. Damen, C. Arbizzani, R. Inguanta, S. Piazza, C. Sunseri and M. Mastragostino, *J. Power Sources*, 2011, **196**, 1469–1473.
- Z. Du and S. Zhang, *J. Phys. Chem. C*, 2011, **115**, 23603–23609.
- L.-J. Xue, Y.-F. Xu, L. Huang, F.-S. Ke, Y. He, Y.-X. Wang, G.-Z. Wei, J.-T. Li and S.-G. Sun, *Electrochim. Acta*, 2011, **56**, 5979–5987.
- M.-Y. Li, C.-L. Liu, M.-R. Shi and W.-S. Dong, *Electrochim. Acta*, 2011, **56**, 3023–3028.
- D.-H. Nam, R.-H. Kim, C.-L. Lee and H.-S. Kwon, *J. Electrochem. Soc.*, 2012, **159**, A1822–A1826.
- G. Ferrara, C. Arbizzani, L. Damen, M. Guidotti, M. Lazzari, F. G. Vergottini, R. Inguanta, S. Piazza, C. Sunseri and M. Mastragostino, *J. Power Sources*, 2012, **211**, 103–107.
- R. M. Gnanamuthu, Y. N. Jo and C. W. Lee, *J. Alloys Compd.*, 2013, **564**, 95–99.
- J. R. Gonzalez, F. Nacimiento, R. Alcántara, G. F. Ortiz and J. L. Tirado, *CrystEngComm*, 2013, **15**, 9196–9202.
- X. Liu, J. Xie, H. Zhao, P. Lv, K. Wang, Z. Feng and K. Świerczek, *Solid State Ionics*, 2015, **269**, 86–92.
- C. Wu, J. Maier and Y. Yu, *Adv. Funct. Mater.*, 2015, **25**, 3488–3496.
- B.-O. Jang, S.-H. Park and W.-J. Lee, *J. Alloys Compd.*, 2013, **574**, 325–330.



- 49 X. Li, X. He, Y. Xu, L. Huang, J. Li, S. Sun and J. Zhao, *J. Mater. Chem. A*, 2015, **3**, 3794–3800.
- 50 N. Mahmood, C. Zhang, F. Liu, J. Zhu and Y. Hou, *ACS Nano*, 2013, **7**, 10307–10318.
- 51 N. Mahmood, J. Zhu, S. Rehman, Q. Li and Y. Hou, *Nano Lett.*, 2015, **15**, 755–765.
- 52 G. O. Park, J. Yoon, J. K. Shon, Y. S. Choi, J. G. Won, S. B. Park, K. H. Kim, H. Kim, W.-S. Yoon and J. M. Kim, *Adv. Funct. Mater.*, 2016, **26**, 2800–2808.
- 53 X. Shi, H. Song, A. Li, X. Chen, J. Zhou and Z. Ma, *J. Mater. Chem. A*, 2017, **5**, 5873–5879.
- 54 J. Zhu, D. Wang, T. Liu and C. Guo, *Electrochim. Acta*, 2014, **125**, 347–353.
- 55 N. Tamura, Y. Kato, A. Mikami, M. Kamino, S. Matsuta and S. Fujitani, *J. Electrochem. Soc.*, 2006, **153**, A1626–A1632.
- 56 C. M. Ionica-Bousquet, P. E. Lippens, L. Aldon, J. Olivier-Fourcade and J. C. Jumas, *Chem. Mater.*, 2006, **18**, 6442–6447.
- 57 Y. Xu, M. Zhou and Y. Lei, *Adv. Energy Mater.*, 2016, **6**, 1502514.
- 58 W. Li, X. Sun and Y. Yu, *Small Methods*, 2017, **1**, 1600037.
- 59 B. Zhang, X.-S. Li, C.-L. Liu, Z.-H. Liu and W.-S. Dong, *RSC Adv.*, 2015, **5**, 53586–53591.
- 60 M. Walter, R. Erni and M. V. Kovalenko, *Sci. Rep.*, 2015, **5**, 8418.
- 61 V. Etacheri, O. Haik, Y. Goffer, G. A. Roberts, I. C. Stefan, R. Fasching and D. Aurbach, *Langmuir*, 2011, **28**, 965–976.
- 62 S. Komaba, T. Ishikawa, N. Yabuuchi, W. Murata, A. Ito and Y. Ohsawa, *ACS Appl. Mater. Interfaces*, 2011, **3**, 4165–4168.
- 63 A. M. Chockla, K. C. Klavetter, C. B. Mullins and B. A. Korgel, *Chem. Mater.*, 2012, **24**, 3738–3745.
- 64 L. Ji, M. Gu, Y. Shao, X. Li, M. H. Engelhard, B. W. Arey, W. Wang, Z. Nie, J. Xiao, C. Wang, J.-G. Zhang and J. Liu, *Adv. Mater.*, 2014, **26**, 2901–2908.
- 65 Z. Yang, A. A. Gewirth and L. Trahey, *ACS Appl. Mater. Interfaces*, 2015, **7**, 6557–6566.
- 66 I. A. Courtney and J. R. Dahn, *J. Electrochem. Soc.*, 1997, **144**, 2045–2052.
- 67 X. Huang, S. Cui, J. Chang, P. B. Hallac, C. R. Fell, Y. Luo, B. Metz, J. Jiang, P. T. Hurley and J. Chen, *Angew. Chem., Int. Ed.*, 2015, **54**, 1490–1493.
- 68 N. Zhang, Q. Zhao, X. Han, J. Yang and J. Chen, *Nanoscale*, 2014, **6**, 2827–2832.
- 69 Z. Zhu, S. Wang, J. Du, Q. Jin, T. Zhang, F. Cheng and J. Chen, *Nano Lett.*, 2014, **14**, 153–157.
- 70 J. Liu, Y. Wen, P. A. van Aken, J. Maier and Y. Yu, *Nano Lett.*, 2014, **14**, 6387–6392.
- 71 Y. Xu, Q. Liu, Y. Zhu, Y. Liu, A. Langrock, M. R. Zachariah and C. Wang, *Nano Lett.*, 2013, **13**, 470–474.
- 72 J. Hwang, S. H. Woo, J. Shim, C. Jo, K. T. Lee and J. Lee, *ACS Nano*, 2013, **7**, 1036–1044.
- 73 S. R. Sivakkumar, J. Y. Nerkar and A. G. Pandolfo, *Electrochim. Acta*, 2010, **55**, 3330–3335.
- 74 F. Ding, W. Xu, D. Choi, W. Wang, X. Li, M. H. Engelhard, X. Chen, Z. Yang and J.-G. Zhang, *J. Mater. Chem.*, 2012, **22**, 12745–12751.
- 75 P. Gevorkian, *Large-Scale Solar Power Systems: Construction and Economics*, Cambridge University Press, 2012.
- 76 M. N. Obrovac and V. L. Chevrier, *Chem. Rev.*, 2014, **114**, 11444–11502.
- 77 D. Foster, J. Wolfenstine, J. Read and J. L. Allen, *Performance of Sony's Alloy Based Li-Ion Battery*, DTIC Document, 2008.

