

Evaluation of Metal Phosphide Nanocrystals as Anode Materials for Na-ion Batteries

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Abstract: Sodium-ion batteries (SIBs) are potential low-cost alternatives to lithium-ion batteries (LIBs) because of the much greater natural abundance of sodium salts. However, developing high-performance electrode materials for SIBs is a challenging task, especially due to the ~50% larger ionic radius of the Na⁺ ion compared to Li⁺, leading to vastly different electrochemical behavior. Metal phosphides such as FeP, CoP, NiP₂, and CuP₂ remain unexplored as electrode materials for SIBs, despite their high theoretical charge storage capacities of 900–1300 mAh g⁻¹. Here we report on the synthesis of metal phosphide nanocrystals (NCs) and discuss their electrochemical properties as anode materials for SIBs, as well as for LIBs. We also compare the electrochemical characteristics of phosphides with their corresponding sulfides, using the environmentally benign iron compounds, FeP and FeS₂, as a case study. We show that despite the appealing initial charge storage capacities of up to 1200 mAh g⁻¹, enabled by effective nanosizing of the active electrode materials, further work toward optimization of the electrode/electrolyte pair is needed to improve the electrochemical performance upon cycling.

Keywords: Anode materials · Li-ion batteries · Na-ion batteries · Nanocrystals · Synthesis

1. Introduction

Lithium-ion batteries (LIBs) have become the battery technology of choice for applications demanding high energy and power densities, such as portable electronics and electric vehicles, and also show great promise for the large-scale grid storage of electricity. Yet, the irregular geographic distribution and relatively low natural abundance of lithium salts raise doubts as to the future security and cost of supply. In this regard, conceptually identical sodium-ion batteries (SIBs) are a favorable alternative due to the much greater abundance (by a factor of 10³) and therefore lower price of sodium salts.^[1] However, the seemingly simple replacement

of the Li⁺ ion with its 50% larger group I neighbor has drastic consequences for the resulting electrochemistry. For instance, both silicon and graphite, which are well-known anode materials with outstanding lithium ion storage properties, show negligible capacities for sodium ions.^[2] Extensive research toward new electrode materials is needed to advance the development of high-performance SIBs.

Of all possible anode materials for SIBs, red phosphorus (P) is probably the most appealing candidate due to its low cost, nontoxicity and, most importantly, extremely high sodium capacity (2596 mAh g⁻¹ for P ↔ Na₃P, the highest Na⁺ capacity known) at a low desodiation potential (~0.6 V vs. Na⁺/Na). However, similarly to other alloying/conversion type materials, P suffers from massive volume changes during sodiation/desodiation ($\Delta V = 291\%$, by molar volume) leading to the mechanical disintegration of the electrodes and therefore rapid capacity fading due to loss of electrical contact. The other main disadvantage of P is its relatively low electronic conductivity, causing slow reaction kinetics. Although noticeable progress has been demonstrated for P-based SIB anodes,^[3] typically very large amounts of conductive carbons are used to provide sufficient conductivity as well as mechanical stability of the electrodes, and often high capacities with good cycling stability can only be achieved at low charge/discharge currents of ~100 mA g⁻¹ (~0.05 C).

In this study, we were intrigued by the possibility of addressing the aforementioned issues facing P-based SIB anodes by using metal phosphide nanocrystals (NCs) as the active material. Generally, nanostructured materials often show improved electrochemical performance over their bulk counterparts due to mitigation of the effects caused by volumetric changes and improved ionic and electronic conductivities upon homogeneous mixing with conductive carbon additives.^[4] Moreover, metallic inclusions, which form *in situ* upon electrochemical conversion of the transition metal phosphide to alkali metal phosphide, are also expected to improve the electronic connectivity within the electrode. Despite the additional mass of the transition metal, theoretical specific charge-storage capacities of metal phosphides are still extremely high (900–1300 mAh g⁻¹), surpassing all of the main alternatives to P such as Sn (847 mAh g⁻¹) and Sb (660 mAh g⁻¹). Herein, we present the sodium and also lithium storage properties of highly uniform FeP, CoP, NiP₂ and CuP₂ NCs prepared *via* colloidal synthesis methods. It should be noted that, with the exception of a recent study on FeP,^[5] this is the first report on the electrochemical performance of such metal phosphides in SIBs. All of the phosphide NCs investigated in this work show high charge-storage capacities, close to the theoretically expected values. In comparison to the corresponding metal sulphide NCs, the phosphides

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exhibit lower desodiation potentials and are hence better suited as SIB anode materials, but suffer from very fast capacity loss upon cycling. Further work on the optimized formulation of the electrodes and the selection of suitable electrolytes and electrolyte additives is needed to improve long-term cycling stability.

2. Experimental

2.1 Synthesis of Metal Phosphide Nanocrystals

2.1.1 FeP Nanowires (NWs)

In a typical experiment, 2.5 g tri-n-octylphosphine oxide (TOPO, 99%, Strem) and 3 mL tri-n-octylphosphine (TOP, ≥97%, Strem), previously dried at 100 °C under vacuum for 1 h, were heated to 300 °C under Ar. At 300 °C, 0.5 mL of Fe stock solution, prepared by mixing 1 mL TOP and 0.25 mL Fe(CO)₅ (99.99%, Strem), was injected into the TOP/TOPO mixture. After 30 min, a second injection of 0.5 mL of stock solution was carried out. The reaction was stopped after an additional 30 min. FeP NWs were precipitated by adding hexane and ethanol, separated by centrifugation, and re-dispersed in chloroform containing 1 wt% oleic acid. The second precipitation was induced by adding ethanol. After centrifugation, the FeP nanowires were re-dispersed in chloroform and stored under ambient conditions.

2.1.2 NiP₂ NCs

In a typical experiment, 4.5 mL octadecene (ODE, 90%, Sigma-Aldrich), 6.4 mL oleylamine (OLA, 95%, Strem) and 0.25 g (1 mmol) nickel(II) acetylacetonate (≥98%, Merck) were dried at 110 °C under vacuum for 1 h to remove water and low-boiling point impurities. Then, 2 mL of TOP were added to the flask under Ar and the reaction mixture was heated to 320 °C and held at this temperature for 1 h. The flask was cooled to 200 °C by flowing air and then 105 mg (3.4 mmol) of red phosphorous (≥97%, Sigma-Aldrich) were added. The reaction mixture was then heated again to 330 °C and held at this temperature for 22 h. NiP₂ NCs were precipitated twice by adding chloroform and ethanol, separated by centrifugation, and re-dispersed again in chloroform. After centrifugation, the NiP₂ NPs were re-dispersed in chloroform and stored under ambient conditions.

2.1.3 CoP NCs

In a typical experiment, 4.5 mL ODE, 6.4 mL OLA and 0.25 g (1 mmol) cobalt(II) acetylacetonate (≥98%, Merck) were dried at 110 °C under vacuum for 1 h to remove water and low-boiling point impurities.

Then, 2 mL of TOP were added to the flask under Ar and the reaction mixture was heated to 320 °C for 65 min. The flask was cooled to 200 °C by flowing air and then 105 mg (3.4 mmol) of red phosphorous were added. Then the reaction mixture was heated again to 330 °C and held at this temperature for 22 h. CoP NCs were isolated and purified identically to the NiP₂ NCs above.

2.1.4 CuP₂ NCs

In a typical experiment, 4.5 mL ODE, 6.4 mL OLA and 0.262 g (1 mmol) copper(II) acetylacetonate (≥97%, Sigma-Aldrich) were dried at 110 °C under vacuum for 1 h to remove water and low-boiling point impurities. Then, 2 mL of TOP were added to the flask under Ar atmosphere and the reaction mixture was heated to 320 °C for 75 min. The flask was cooled to 200 °C by flowing air and then 200 mg (6.4 mmol) of red phosphorous were added. The reaction mixture was then heated again to 330 °C and was held at this temperature for 22 h. CuP₂ NCs were isolated and purified identically to the NiP₂ NCs above.

2.2 Characterization of Metal Phosphide Nanocrystals

Transmission electron microscopy (TEM) was performed using a JEOL JEM-2200FS instrument operated at 200 kV, using carbon-coated Cu grids as substrates (Ted-Pella). Powder X-ray diffraction (XRD) was measured using a STOE STADI P diffractometer (with Cu-Kα₁ irradiation, λ = 1.540598 Å).

2.3 Electrode Preparation, Cell Assembly and Electrochemical Measurements

In order to evaluate the electrochemical properties of FeP, CoP, NiP₂ and CuP₂ NCs, Na-ion and Li-ion half-cells were assembled. Prior to electrode preparation, organic ligands were removed from the surface of the NCs by stirring them in a 1 M solution of hydrazine in acetonitrile for 2 h at room temperature, as is commonly performed for colloidal quantum dots.^[6] Electrodes were prepared by mixing the respective metal phosphide NCs (63.75 wt%) with carbon black (21.25 wt%, TIMCAL), carboxymethylcellulose (CMC, 15 wt%) and water as a solvent using a planetary ball-mill at 500 rpm for 1 h. The aqueous slurries were coated onto Cu current collectors, which were dried at 80 °C under vacuum overnight prior to cell assembly. For electrochemical testing, coin cells with elemental Na or Li were assembled in an Ar-filled glovebox (O₂ < 0.1 ppm, H₂O < 0.1 ppm) using either 1 M NaClO₄ in propylene carbonate (PC) with 10% fluoroethylene carbonate (FEC) or 1 M LiPF₆ in a 1:1 mixture of ethylene carbonate (EC)

and dimethyl carbonate (DMC) with 3% FEC. FEC was added to the electrolyte in both the Li and Na coin cells to improve capacity retention.^[3c,7] All electrochemical tests were carried out at room temperature and the capacities were reported relative to the mass of the metal phosphide NCs.

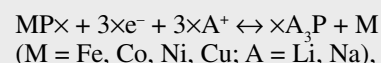
3. Results and Discussion

3.1 Synthesis and Characterization of Metal Phosphide Nanocrystals

FeP NWs were synthesized according to the procedure reported by Qian *et al.*^[8] In order to obtain NiP₂ NCs, a two-step procedure was developed. First, Ni₂P NCs were synthesized according to a known protocol reported by Popczun *et al.*^[9] We then added a second step: conversion of the as-prepared Ni₂P NCs into NiP₂ NCs by adding red P to the reaction mixture, followed by heating at 330 °C for 22 h. Analogously, this two-step approach was also applied in the synthesis of CuP₂ and CoP NCs, simply by replacing nickel(II) acetylacetonate with the respective copper or cobalt salt (for details, see the Experimental section). Fig. 1 summarizes the characterization of the metal phosphide NCs obtained by these methods. FeP NCs were on average ~300 nm in length and ~7 nm in width. CoP, NiP₂ and CuP₂ NCs exhibited diameters of 25, 10 and 60 nm, respectively. All materials showed phase-pure XRD patterns, indexed according to the standard ICSD files for these compounds.

3.2 Electrochemical Performance of Metal Phosphide Nanocrystals

Fig. 2 shows the electrochemical performance of the metal phosphide NCs in Na-ion and Li-ion half-cells. Na-ion cells were cycled at a current rate of 100 mA g⁻¹ in the potential range of 0.02–2.5 V. For Li-ion cells, current rates of 300 mA g⁻¹ and a potential range of 0.02–2.0 V were used. Assuming the formation of Na₃P or Li₃P via the general conversion reaction



the metal phosphides FeP, CoP, NiP₂ and CuP₂ have theoretical capacities of 926, 894, 1333 and 1282 mAh g⁻¹, respectively. In close agreement, CuP₂ indeed showed the highest capacity in the first cycle. However, the capacities of all studied materials rapidly faded during cycling. The compounds with higher P content, NiP₂ and CuP₂, showed higher initial capacities but poorer capacity retention. Namely, for CuP₂ NCs the charge capacity decreased from 1140 mAh g⁻¹ to 570 mAh g⁻¹ within the first 16 cycles. For the FeP, CoP and

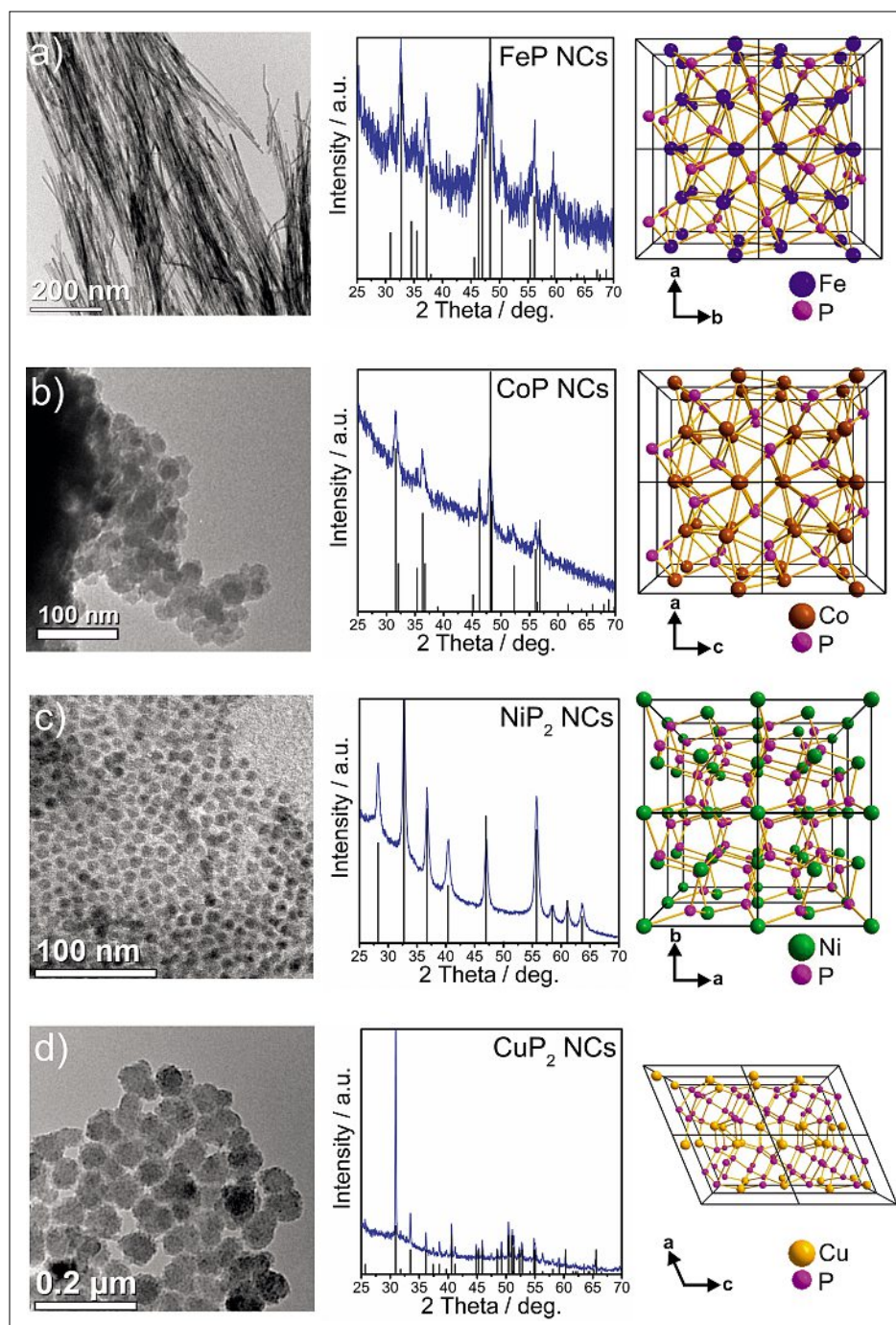


Fig. 1. Characterization of metal phosphide NCs. Transmission electron microscopy (TEM) images, X-ray diffraction (XRD) patterns and schematic representations of the crystal structures (from left to right) of FeP, CoP, NiP_2 and CuP_2 NCs (a–d). The XRD patterns are indexed according to the ICSD database: to orthorhombic FeP (PDF No.: 00-071-2262, space group Pna21 (33), $a = 5.193 \text{ \AA}$, $b = 5.792 \text{ \AA}$, $c = 3.099 \text{ \AA}$), orthorhombic CoP (PDF No.: 00-029-0497, space group Pnma (62), $a = 5.077 \text{ \AA}$, $b = 3.281 \text{ \AA}$, $c = 5.587 \text{ \AA}$), cubic NiP_2 (PDF No.: 00-073-0436, space group Pa3 (205), $a = 5.4706 \text{ \AA}$) and monoclinic CuP_2 (PDF No.: 00-076-1190, space group P21/c (14), $a = 5.8004 \text{ \AA}$, $b = 4.8063 \text{ \AA}$, $c = 7.5263 \text{ \AA}$, $\beta = 112.7^\circ$).

NiP_2 NCs, the capacities fell to below 600 mAh g^{-1} after just the first 10 cycles, and faded to less than 400 mAh g^{-1} during subsequent cycling. Similar observations were made when testing the metal phosphide NCs in Li-ion half-cells (Fig. 2b). Due to the high surface area of the nanosized materials, low coulombic efficiencies (20–70%) were obtained for the first cycle due to the irreversible decomposition of the electrolyte forming the solid electrolyte

interface (SEI) (see Figs 2c and 2d). Notably, rather poor coulombic efficiencies of 92–95% for Na-ion and 96–98% for Li-ion cells were obtained during subsequent cycles, indicating continuous deterioration and reformation of the SEI caused by pulverization of the electrode material.

Figs 2e and 2f show the galvanostatic charge and discharge voltage profiles for the first cycle for all tested metal phosphide NCs. For sodium ion storage, a desodiation

plateau can be identified in all cases at $\sim 0.6 \text{ V}$ vs. Na^+/Na , which is at the same potential as reported for the electrochemical reaction of red P with Na.^[3e] This implies that metal phosphides rather convert into elemental P and that cycling proceeds mainly by the reaction $\text{P} + 3\text{e}^- + 3\text{Na}^+ \leftrightarrow \text{Na}_3\text{P}$, as has been suggested for FeP.^[5] In Li-ion half-cells, the majority of delithiation occurs at a potential of more than 1.0 V showing that metal phosphides are generally better suited as SIB anode materials due to lower voltages of desodiation.

3.3 Comparison of FeP and FeS_2 NCs as Anode Materials for Na-ion Batteries

Clearly, from the prospects of low cost and low toxicity, iron-based sodium storage electrode materials are the most interesting candidates, in particular when the other chemical constituents of the compound comprise equally abundant elements such as phosphorus and sulfur. Hence, iron sulfides can be seen as a main alternative to phosphides. Similar difficulties with capacity fading might occur for FeS_2 (pyrite) due to its large ($\sim 280\%$) volume expansion upon Na_2S formation.^[10] In order to compare the electrochemical performance of Fe phosphides and sulfides, we synthesized pyrite FeS_2 NCs with sizes from 50–100 nm and tested them under the same conditions as the FeP NCs. The synthesis, characterization and electrochemical properties of FeS_2 NCs have been detailed in our recent report.^[11] Assuming the formation of Na_2S , FeS_2 NCs possess a theoretical maximum capacity of 894 mAh g^{-1} , similar to the value for FeP (926 mAh g^{-1}). However, as can be seen in Fig. 3, the electrochemical performance of FeS_2 and FeP NCs is in fact very different. Whereas FeP NCs show rapid capacity fading upon cycling, FeS_2 NCs exhibit stable capacities of $\geq 800 \text{ mAh g}^{-1}$ (near the theoretical value), clearly demonstrating that the identity of the anion in a conversion-type electrode material plays a critical role in determining its electrochemical properties. The only relevant previous investigation of FeP as a SIB anode material is the recent report by Li *et al.*:^[5] in that work, anodes prepared by ball-milling FeP showed much faster capacity fading, from 460 mAh g^{-1} to $\sim 200 \text{ mAh g}^{-1}$ within 40 cycles at a current of 50 mA g^{-1} . Compared to FeP NCs, the only obvious drawback of FeS_2 NCs is the higher desodiation potential (Figs 3b and 3c), that is, however, well compensated by good capacity retention.

4. Conclusion

In conclusion, we have prepared NCs of FeP, CoP, NiP_2 and CuP_2 using colloidal

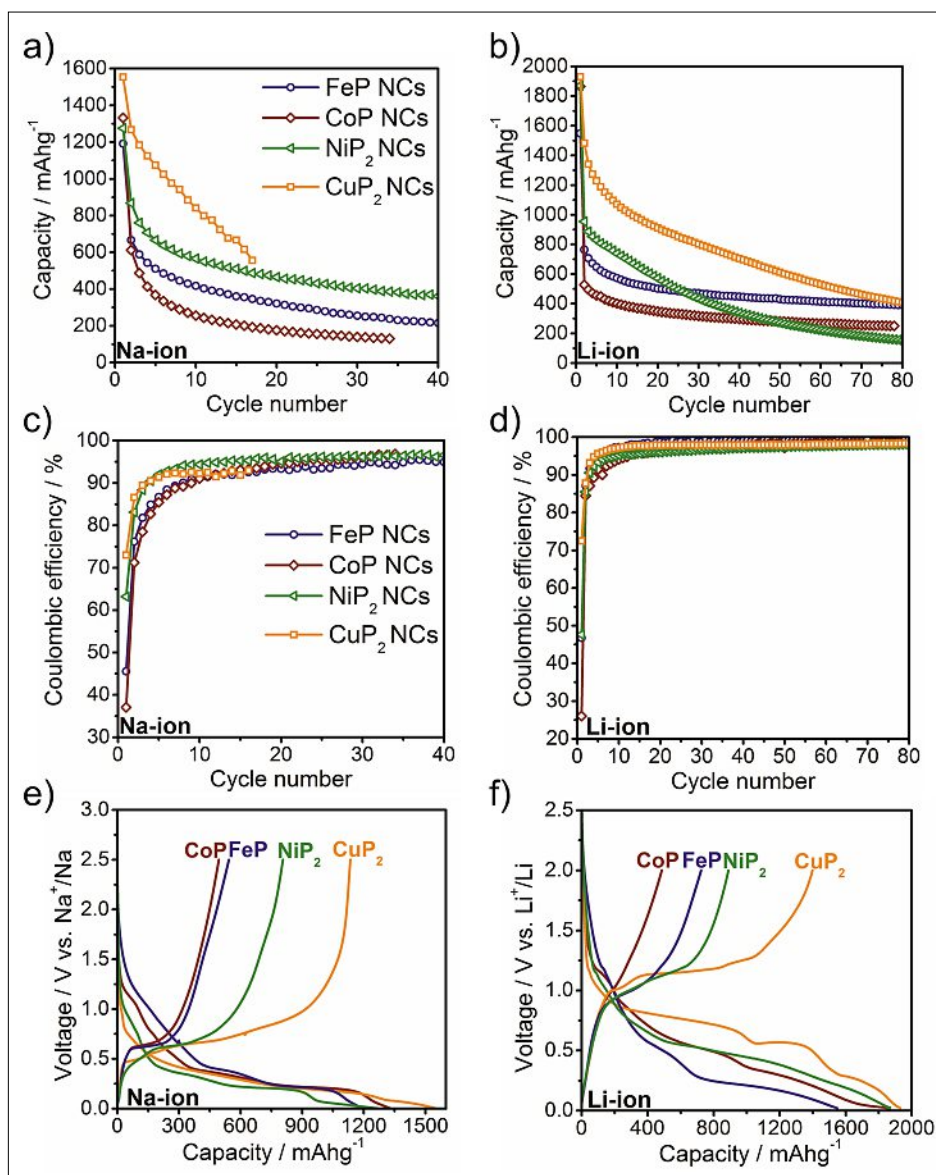


Fig. 2. Electrochemical performance of metal phosphide nanocrystals. Galvanostatic cycling of metal phosphide NCs in Na-ion (a) and Li-ion half-cells (b) with the respective coulombic efficiency plots (c, d). Galvanostatic charge and discharge curves for the first cycle for Na-ion (e) and Li-ion (f) half-cells. Electrodes were composed of 63.75% metal phosphide NCs, 21.25% CB and 15% CMC. 1 M NaClO₄ in PC with 10% FEC served as the electrolyte for Na-ion and 1 M LiPF₆ in EC:DMC (1:1) for Li-ion half-cells. Galvanostatic cycling tests were carried out with a current of 100 mA g⁻¹ in the potential range of 0.02–2.5 V for Na-ion and 300 mA g⁻¹ in the potential range of 0.02–2.0 V for Li-ion half-cells.

dal synthesis methods. Motivated by their high theoretical capacities, the low electrochemical potentials needed for the anodic side of the battery, and the high natural abundance of their constituting elements, we explored the potential of these nanomaterials as SIB anode materials. We also compared their sodium storage properties with that of lithium. We find that with conventionally formulated electrodes (by mixing with amorphous carbon and water-soluble binders), metal phosphide NCs deliver high sodium capacities but exhibit low capacity retention upon cycling. The direct comparison of FeP and FeS₂ as SIB anode materials indicates the much better cyclability of the latter. We thus conclude

that metal phosphides are inherently less stable than their corresponding sulfides and this instability is a combined effect of large volumetric changes and phosphorus-specific processes, such as the reactivity of Na₃P towards the electrolyte.^[3c] Further progress towards high performance phosphide-based electrodes is expected to result from smart electrode engineering by designing secondary structures in which the metal phosphide NPs are encapsulated into conductive carbons,^[12] thereby eliminating the direct large-area contact with the electrolyte. Furthermore, a sensible choice of the electrolyte and electrolyte additives might enable the higher stability of the SEI layer in future studies.

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- [1] a) R. Van Noorden, *Nature* **2014**, 507, 26; b) V. Palomares, P. Serras, I. Villaluenga, K. B. Hueso, J. Carretero-Gonzalez, T. Rojo, *Energy Environ. Sci.* **2012**, 5, 5884; c) M. D. Slater, D. Kim, E. Lee, C. S. Johnson, *Adv. Funct. Mater.* **2013**, 23, 947; d) H. Pan, Y.-S. Hu, L. Chen, *Energy Environ. Sci.* **2013**, 6, 2338; e) N. Yabuuchi, K. Kubota, M. Dahbi, S. Komaba, *Chem. Rev.* **2014**, 114, 11636.
- [2] a) P. Ge, *Solid State Ionics* **1988**, 28, 1172; b) S. Komaba, Y. Matsuura, T. Ishikawa, N. Yabuuchi, W. Murata, S. Kuze, *Electrochem. Commun.* **2012**, 21, 65; c) L. D. Ellis, B. N. Wilkes, T. D. Hatchard, M. N. Obrovac, *J. Electrochem. Soc.* **2014**, 161, A416.
- [3] a) J. Qian, X. Wu, Y. Cao, X. Ai, H. Yang, *Angew. Chem. Int. Ed.* **2013**, 52, 4633; b) Y. Kim, Y. Park, A. Choi, N.-S. Choi, J. Kim, J. Lee, J. H. Ryu, S. M. Oh, K. T. Lee, *Adv. Mater.* **2013**, 25, 3045; c) N. Yabuuchi, Y. Matsuura, T. Ishikawa, S. Kuze, J.-Y. Son, Y.-T. Cui, H. Oji, S. Komaba, *ChemElectroChem* **2014**, 1, 580; d) W.-J. Li, S.-L. Chou, J.-Z. Wang, H.-K. Liu, S.-X. Dou, *Nano Lett.* **2013**, 13, 5480; e) M. Walter, R. Erni, M. V. Kovalenko, *Sci. Rep.* **2015**, 5; f) J. Song, Z. Yu, M. L. Gordin, S. Hu, R. Yi, D. Tang, T. Walter, M. Regula, D. Choi, X. Li, A. Manivannan, D. Wang, *Nano Lett.* **2014**, 14, 6329; g) Y. Zhu, Y. Wen, X. Fan, T. Gao, F. Han, C. Luo, S.-C. Liou, C. Wang, *ACS Nano* **2015**, 9, 3254.
- [4] a) M. F. Oszejka, M. I. Bodnarchuk, M. V. Kovalenko, *Chem. Mater.* **2014**, 26, 5422; b) N. Liu, Z. Lu, J. Zhao, M. T. McDowell, H.-W. Lee, W. Zhao, Y. Cui, *Nat. Nanotechnol.* **2014**, 9, 187; c) A. Magasinski, P. Dixon, B. Hertzberg, A. Kvit, J. Ayala, G. Yushin, *Nat. Mater.* **2010**, 9, 353; d) C. K. Chan, R. N. Patel, M. J. O'Connell, B. A. Korgel, Y. Cui, *ACS Nano* **2010**, 4, 1443; e) P. G. Bruce, B. Scrosati, J.-M. Tarascon, *Angew. Chem. Int. Ed.* **2008**, 47, 2930; f) M. R. Palacin, *Chem. Soc. Rev.* **2009**, 38, 2565; g) K. Kravchyk, L. Protesescu, M. I. Bodnarchuk, F. Krumeich, M. Yarema, M. Walter, C. Guntlin, M. V. Kovalenko, *J. Am. Chem. Soc.* **2013**, 135, 4199; h) M. He, K. Kravchyk, M. Walter, M. V. Kovalenko, *Nano Lett.* **2014**, 14, 1255; i) M. He, M. Walter, K. V. Kravchyk, R. Erni, R. Widmer, M. V. Kovalenko, *Nanoscale* **2015**, 7, 455; j) J. Liu, Y. Wen, P. A. van Aken, J. Maier, Y. Yu, *Nano Lett.* **2014**, 14, 6387; k) A. Jägel, C. M. Ghimbeu, L. Monconduit, C. Vix-Guterl, *Adv. Energy Mater.* **2014**, 4, DOI: 10.1002/aenm.201400025.
- [5] W.-J. Li, S.-L. Chou, J.-Z. Wang, H.-K. Liu, S.-X. Dou, *Chem. Commun.* **2015**, 51, 3682.
- [6] D. V. Talapin, C. B. Murray, *Science* **2005**, 310, 86.
- [7] a) A. M. Chockla, K. C. Klavetter, C. B. Mullins, B. A. Korgel, *Chem. Mater.* **2012**, 24, 3738; b) V. Etacheri, O. Haik, Y. Goffer, G. A. Roberts, I. C. Stefan, R. Fasching, D. Aurbach, *Langmuir* **2011**, 28, 965; c) S. Komaba, T. Ishikawa, N. Yabuuchi, W. Murata, A. Ito, Y. Ohsawa, *ACS Appl. Mater. Interfaces* **2011**, 3, 4165; d) L. O.

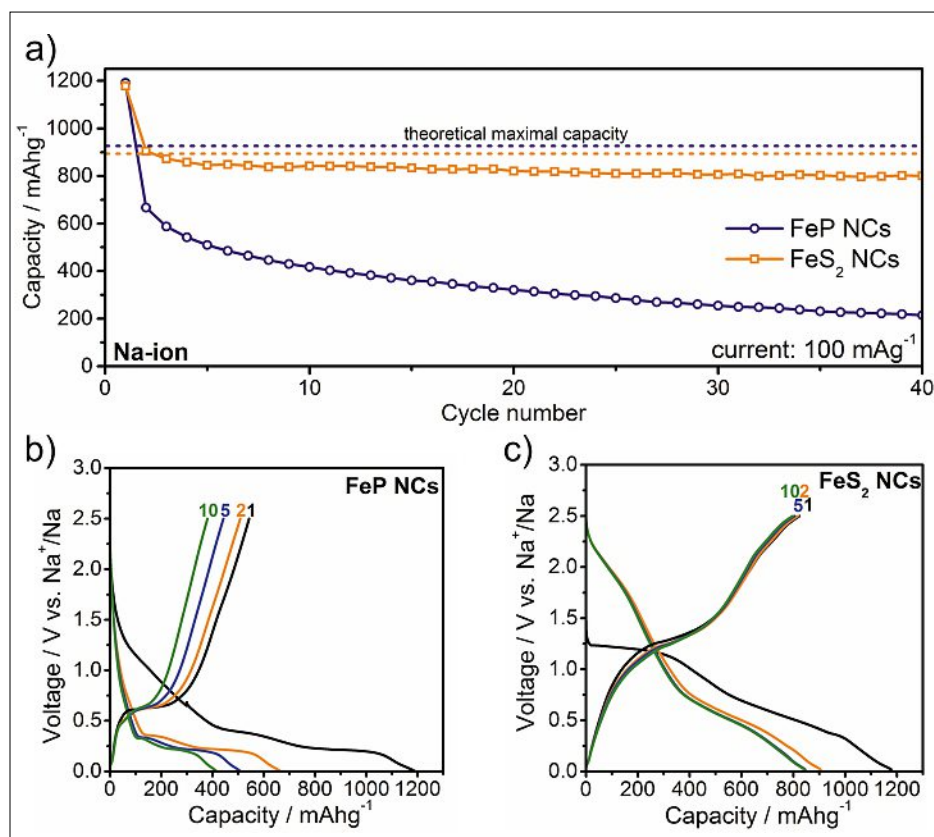


Fig. 3. Comparison of the electrochemical performance of FeP and FeS₂ NCs. Galvanostatic cycling of FeS₂ and FeP in Na-ion half-cells (a) and the respective charge/discharge curves (b, c). Galvanostatic cycling tests were carried out with a current of 100 mA g⁻¹ in the potential range of 0.02–2.5 V.

- Vogt, M. El Kazzi, E. Jämstorp Berg, S. Pérez Villar, P. Novák, C. Villevieille, *Chem. Mater.* **2015**, 27, 1210.
- [8] C. Qian, F. Kim, L. Ma, F. Tsui, P. Yang, J. Liu, *J. Am. Chem. Soc.* **2004**, 126, 1195.
- [9] E. J. Popczun, J. R. McKone, C. G. Read, A. J. Biacchi, A. M. Wiltrout, N. S. Lewis, R. E. Schaak, *J. Am. Chem. Soc.* **2013**, 135, 9267.
- [10] F. Klein, B. Jache, A. Bhide, P. Adelhelm, *Phys. Chem. Chem. Phys.* **2013**, 15, 15876.
- [11] M. Walter, T. Zund, M. Kovalenko, *Nanoscale* **2015**, 7, 9158.
- [12] a) X. Ji, K. T. Lee, L. F. Nazar, *Nat. Mater.* **2009**, 8, 500; b) X. W. Lou, C. M. Li, L. A. Archer, *Adv. Mater.* **2009**, 21, 2536; c) H. Zhang, X. Yu, P. V. Braun, *Nat. Nanotechnol.* **2011**, 6, 277; d) Y. Yu, L. Gu, C. Wang, A. Dhanabalan, P. A. van Aken, J. Maier, *Angew. Chem. Int. Ed.* **2009**, 48, 6485; e) N. Liu, H. Wu, M. T. McDowell, Y. Yao, C. Wang, Y. Cui, *Nano Lett.* **2012**, 12, 3315.