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Charge compensation in a layered van der Waals NiPS₃ host through various cationic intercalations†

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The layered structure of van der Waals compounds enables facile insertion of guest species between layers, resulting in material multifunctionality through easily modifying its physical properties. Isostructural MPS₃ compounds with 3d transition metal cations such as Mn, Fe, Co and Ni can serve as hosts for relatively small alkali metals as well as larger organic molecules. NiPS₃ is the most exotic representative among them, because despite 30 years of intense research, its electronic structure still evokes numerous questions, not to mention the electronic structure of intercalated NiPS₃. There are two possibilities for electron transfer in semiconducting NiPS₃ upon insertion of electron-donating species, either to a discrete Ni atomic level, or to a molecular level of the (P₂S₆)⁴⁻ unit. We performed a systematic structural and spectroscopic study of NiPS₃ upon electrochemical intercalation of Li, Na and 1-ethyl-3-methylimidazolium (EMIM) cations. Up to 0.5 Li or 0.5 Na per NiPS₃ formula unit can be inserted into free octahedral spaces in the interlayers without visible changes in the diffraction pattern of the host. In contrast, more than 1 EMIM per NiPS₃ unit can be intercalated between host layers leading to a significant interlayer distance expansion from 6.33 Å to 11.3 Å. The charge compensation was found to be different for the three intercalants: upon Li insertion, the electron density increases on the (P₂S₆)⁴⁻ unit and Ni remains redox-inactive, while intercalation of Na leads to reduction of Ni. In contrast, uptake of larger EMIM cations does not result in any changes in Ni, S and P K-edge near edge XANES spectra of NiPS₃ and results in only very little change in their extended X-ray absorption fine structure spectra. It is likely that there is an electrochemical reduction of EMIM cations to heterocyclic carbenes with their possible dimerization. The impact on magnetization of Li and EMIM intercalation was also studied.

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1. Introduction

Layered transition metal phosphorus trisulfides (M₂P₂S₆ or MPS₃ where M = Ni, Mn, Fe *etc.*) have been of interest towards opto-electronic applications due to their versatile anisotropic properties¹ and the possibility of novel ground states.^{2,3} In the recent decades, they have also been found to be promising for electrocatalytic applications.⁴ The family of MPS₃ compounds

crystallizes in a CdCl₂-type structure with the monoclinic space group *C2/m*, as shown in Fig. 1a. In the layers, transition metal ions M, usually in the oxidation state 2+, are octahedrally surrounded by S anions, which are interconnected *via* P cations. Since there are P–P bonds in the structure, the correct presentation of the chemical formula corresponds to M₂P₂S₆ showing (P₂S₆)⁴⁻ units. There are competing inter- and intra-layer magnetic interactions in the structure. The relatively weaker van der Waals interaction between magnetic cations in neighboring layers can be controlled through change of the interlayer distance. Since it is not always feasible to synthesize the desired structures, post-mortem intercalation of cations may allow tuning of the structure.

Intercalation of alkali ions in MPS₃ and the corresponding structural and electronic changes have been extensively studied, especially with Li ions.^{5–10} Due to the limited absorption capacity of lithium by MPS₃ and the modest influence on the physical properties, lithium is only suitable for controlled tuning of the magnetism to a certain extent.^{5,11,12} It is understood that the small alkali ions occupy two types of octahedral

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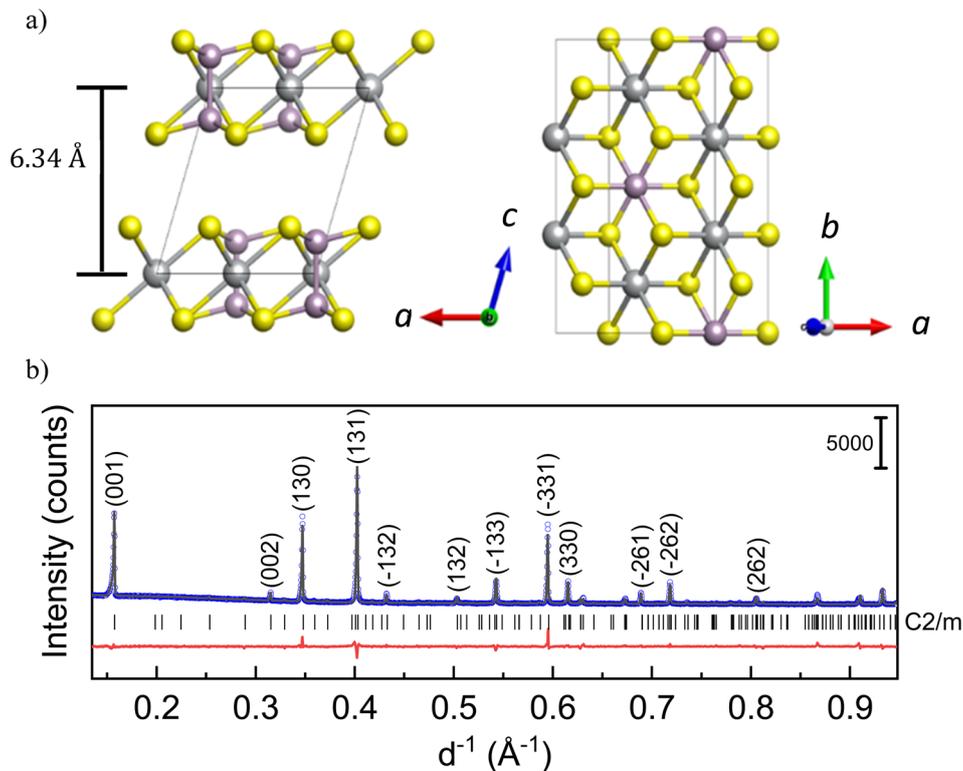


Fig. 1 (a) Crystal structure of MPS_3 compounds with $\text{M} = \text{Ni}$, Mn , and Fe ; left – along the c -axis, and right – in the ab -plane. Metal cations (grey spheres) are octahedrally coordinated by S anions (yellow spheres). P cations (violet) are 4-fold-coordinated by 3S and 1P atoms, resulting in $(\text{P}_2\text{S}_6)^{4-}$ units. (b) X-ray powder diffraction pattern of NiPS_3 with the observed (blue dots) and calculated (black solid line) curves together with their difference curve (red solid line) and Bragg positions given as black vertical lines.

voids present within the layers, without affecting the interlayer spacing.¹¹ An extended Li intercalation results in structure decomposition into mainly metal–sulfur and metal–phosphorus compounds likely due to the high reactivity of alkali ions with chalcogens.^{5–10} However, there is still a discrepancy in the literature regarding the electrochemical potential for intercalation as well as the composition of an intercalated compound with structural stability. Moreover, the charge compensation mechanism upon intercalation is still not completely understood. Therefore, as an alternative, intercalation of large cations or organic molecules, leading to detectable changes in the crystal structure, is also of interest for these compounds. For example, insertion of complex ammonium cations with different organic groups such as tetrabutyl ammonium (TBA^+), tetrapropyl ammonium (TPA^+) and cetyltrimethyl ammonium (CTA^+) were demonstrated in NiPS_3 ,^{13,14} while MnPS_3 is known to be a host for poly(phenylene vinylene) and for pyridine.^{15,16}

In the family of transition metal phosphorus trisulfides, different aspects of intercalation have been comprehensively studied in NiPS_3 .^{9,14,17,18} This compound attracts a special attention due to a big difference in the mechanistic behavior of organic molecule intercalation compared to that of other MPS_3 systems regarding the charge compensation. For instance, it was reported that there is a strong tendency in MnPS_3 and FePS_3 for cation exchange, along with the creation of M^{2+} vacancies during intercalation.^{15,16,19–21} In contrast, no direct intercalation

of pyridine was observed for NiPS_3 , although metal-to-ligand interaction would be of a similar bond strength.²² Pattayil *et al.* suggested that a possible reason for this is the difference in the crystal field energy, which stabilizes Ni in the crystal structure,²² thereby impeding the pyridine intercalation. Similarly, no reaction was reported with 1,10-phenanthroline on NiPS_3 .¹⁸ It should be mentioned that intercalation into NiPS_3 was only possible either by Na -ion mediation (first Na intercalation, followed by cation replacement) or by NiPS_3 doping with other transition metal cations such as Fe^{2+} .^{18,23} During the electrochemical intercalation process with TBA^+ , a decreasing Raman band associated with octahedrally coordinated Ni^{2+} cations was observed, giving an indication of the reduction of nickel.¹⁴ Furthermore, it has been confirmed through electron energy loss spectroscopy (EELS) analysis that both Ni and P were reduced upon more than 1 Li intercalation per NiPS_3 formula unit.¹⁰

Summing up, a solution for a controllable tuning of the interlayer spacing in MPS_3 did not exist until now. The understanding of the charge compensation mechanism is in the nascent stage. Even for the intensively studied lithium intercalation into NiPS_3 , to the best of our knowledge, no structural monitoring *via operando* diffraction methods has been published to date.

Therefore, in the present work we focused on the intercalation principles of NiPS_3 with regard to small alkali metal cations



Li and Na as classical intercalation cations, along with a big organic cation. We aimed to understand structural changes in NiPS₃ and the redox activity of each element upon intercalation, and to compare these with those of other MPS₃ compounds. For this, we synthesized NiPS₃ powder of high quality (see Fig. 1b) and performed a detailed *operando* X-ray diffraction and X-ray absorption spectroscopy study on Li and Na intercalation. We further presented successful organic cation intercalation in NiPS₃ using 1-ethyl-3-methylimidazolium trifluoromethylsulfonylimide [EMIM]⁺[TFSI]⁻ ionic liquid (IL). ILs are notable for their wide electrochemical stability window.²⁴ EMIM-TFSI is one of the more commonly used ionic liquids, being in the liquid state at room temperature.^{25–27} More importantly, EMIM-TFSI has a high conductivity (0.96 S m⁻¹) and low viscosity (24.1 mPa s), which is favorable for electrochemical intercalation.²⁸ Detailed *ex situ* and *operando* characterization was performed to study the structure and morphology of the intercalated materials, towards understanding the mechanism and nature of charge storage upon intercalation in NiPS₃.

2. Experimental section

2.1 Synthesis of pristine NiPS₃

The NiPS₃ sample was synthesized directly from the elements *via* a solid-state reaction. Ni (powder, Alfa Aesar, 99.8%), P (powder, Alfa Aesar, 99.999%), and S (powder, Alfa Aesar, 99.999%) were pressed into pellets (1 cm diameter at approximately 30 kN using a hydraulic press) in stoichiometric ratio and sealed in a quartz tube (12 mm inner diameter and 3 mm wall thickness) under 300 mbar Ar, after evacuation. After an initial heat treatment at 573 K for 24 h, the sample was heated to 973 K and held for 12 days. The heating rate was 50 K h⁻¹. After this, the tube was cooled down to room temperature at the rate of 300 K h⁻¹ and opened under an argon atmosphere. The NiPS₃ structure was confirmed by powder X-ray diffraction measurements (Fig. 1b).

2.2 X-ray diffraction (XRD) measurements

Lab-based powder XRD was performed in transmission geometry on a STOE Stadi P diffractometer with a curved Ge(111) crystal monochromator and Cu K_{α1} radiation ($\lambda = 1.54056 \text{ \AA}$), equipped with a Dectris single-strip Mythen 1K detector. In order to avoid air exposure, the samples were sealed using Kapton tape inside an Ar-filled glovebox. The NiPS₃ samples after cation intercalation for *ex situ* measurements were washed with acetonitrile and dried at 323 K under vacuum to remove all organic residues. All handling was undertaken inside an Ar-filled glovebox.

Operando XRD measurements were performed in the lab on a STOE Stadi P diffractometer using Mo K_{α1} radiation ($\lambda = 0.70926 \text{ \AA}$), at the synchrotron facilities DESY, Hamburg, Germany (P02.1 beamline, $\lambda = 0.2074 \text{ \AA}$) and ALBA, Barcelona, Spain (MSPD, $\lambda = 0.41273 \text{ \AA}$; and NOTOS, $\lambda = 0.6894 \text{ \AA}$, beamlines). All samples for *operando* measurements were prepared in special coin cells with fused silica or Kapton windows. This

allows measurements in the transmission mode with simultaneous control of the potential of the cell. A multi-sample holder for parallel *operando* measurements was used.²⁹ XRD analysis was performed by the Rietveld method using JANA2006 software.³⁰

Elemental analysis was performed by inductively coupled plasma – optical emissions spectroscopy (ICP-OES) on an iCAP 6500 Duo View from ThermoFisher Scientific.

2.3 Electrochemical tests

A BioLogic VMP3 potentiostat was used for all electrochemical studies, including galvanostatic cycling with potential limitation (GCPL), cyclic voltammetry (CV), and the galvanostatic intermittent titration technique (GITT). For electrode preparation, the NiPS₃ powder was mixed with Super P carbon (BASF) as a conductive agent, and polytetrafluoroethylene (PTFE) binder (Aldrich) for better contact with the current collector. The mixture, in a 75 : 20 : 5 weight ratio, was pressed on a copper mesh with an average loading of 11 mg cm⁻². The electrodes were assembled in Swagelok-type cells, using lithium chips (12 mm diameter and 250 μm thickness, Chemetall), platinum mesh (12 mm diameter, 0.25 mm thickness, 99.9% purity, Alfa Aesar) or sodium metal (Alfa Aesar, 99.95%), which was rolled out into flat disks, as a counter electrode. Two glass fiber separators (Whatman GF/D, GE) were applied and soaked with electrolyte. Intercalation of alkali cations was performed with a LP30 electrolyte containing 1 M LiPF₆ in ethylene carbonate (EC, BASF)/dimethyl carbonate (DMC, BASF) in a 1 : 1 ratio (LP30 Selectilyte, BASF), or with a home-made electrolyte composed of 1 M NaPF₆ (abcr, 99%) dissolved in an EC/DMC mixture (1 : 1). For intercalation of organic species, the ionic liquid EMIM-TFSI from Sigma Aldrich was used as the electrolyte. Electrodes and cells were built in an Ar-filled glove box (H₂O < 1 ppm, O₂ < 1 ppm) to avoid contact with oxygen and water. All cells were operated in a climate chamber at room temperature. For diffusion coefficient determination, the galvanostatic intermittent titration technique (GITT) with a constant current pulse of 0.1C (1C corresponds to the current needed for intercalation of 1 EMIM ion per NiPS₃ formula unit in 1 h) was applied, followed by an open circuit phase for several hours. The steps were repeated until the EMIM content exceeded 4 times the formula unit.

2.4 X-ray absorption spectroscopy (XAS) studies

Operando X-ray absorption spectroscopy measurements at the Ni K-edge were carried out at the DESY Synchrotron (P64 PETRA III beamline, Hamburg, Germany) and at the ALBA synchrotron (NOTOS beamline, Barcelona, Spain). The NOTOS beamline enables a combination of XAS-XRD measurements. The samples were assembled in electrochemical coin cells with Kapton windows. A multi-sample holder for parallel *operando* measurements was used.²⁹ The measurements were performed in both transmission and fluorescence mode. *Ex situ* XAS measurements of the S, P and Ni K-edges were carried out at the BESSY II synchrotron (HiKE endstation located at the KMC-1 bending magnet beamline, Berlin, Germany).^{31,32}



In case of the P and S K-edge, the incident X-rays were monochromatized using a Si (111) double-crystal monochromator and focused by means of a piezo refocusing capillary to a size of $100\ \mu\text{m} \times 100\ \mu\text{m}$ (H \times V), if the beam is perpendicular to the sample. For the Ni K-edge, the Si (422) crystal pair was used. The X-ray absorption near edge structure (XANES) was collected in partial fluorescence yield (PFY) mode at a grazing angle of 45° with a Bruker XFlash 4010 fluorescence silicon drift detector (SDD). The incoming photon energy was calibrated using Au 4f spectrum of clean Au foil. The obtained data were processed with Athena software, including background subtraction and normalization.³³

2.5 Magnetometry

Magnetization measurements were performed on a superconducting quantum interference device (SQUID) magnetometer (MPMS, Quantum Design) using powder samples. The magnetization was measured as a function of temperature at an applied magnetic field of 2 T in the field-cooled (FC) and zero-field-cooled (ZFC) mode, in the temperature range of 5 to 330 K.

2.6 Single crystal X-ray diffraction (SCXRD)

SCXRD data acquisition was accomplished on a Bruker D8 Venture (Mo $K\alpha$, $\lambda = 0.71073\ \text{\AA}$) equipped with a PHOTON 100 CMOS detector. The measurement was performed at room temperature. Indexing was performed using APEX3 software.³⁴ Data integration and absorption corrections were performed using the SAINT and SADABS^{34,35} software, respectively. The crystal structure was solved by dual-space methods implemented in the SHELXT³⁶ program and refined by the full-matrix least-squares method on F^2 with SHELXL.³⁷

2.7 High-resolution transmission electron microscopy (HRTEM) and electron diffraction (ED)

HRTEM, ED and energy-dispersive X-ray (EDX) spectroscopy measurements were performed with an aberration-corrected FEI Titan 80–300 electron microscope (ThermoFisher, USA) equipped with an in-column EDX detector. Diffraction patterns were treated in the kinematical approximation.

2.8 Raman spectroscopy

Raman measurements were performed using a S&I Monovista CRS+ confocal Raman spectrometer at 405 nm or 514 nm, with a 2400 grating. The laser was adjusted and focused through a Nikon 20 \times objective (N.A. 0.35; WD = 20 mm) to give an incident laser power of approximately 0.5–5 mW, depending on the sample being measured. Spectra were recorded for a total accumulation time of 75–500 s, depending on the sensitivity and spectral quality of the sample. The peak positions were corrected with respect to the Raman spectrum of a toluene–acetonitrile mixture. Baseline correction and spectral analysis were performed using Qpipi (a Matlab-based software for spectral analysis) and OriginPro.

3. Results

3.1 Structural evolution of NiPS₃ upon alkali metal intercalation

According to the literature reports, NiPS₃ can incorporate up to 1.5 Li per formula unit at a very low current density close to the open circuit voltage (OCV) regime.³⁸ The pristine structure of NiPS₃ contains two different octahedral gaps in the van der Waals interlayer, into which alkali-metal cations Li and Na can

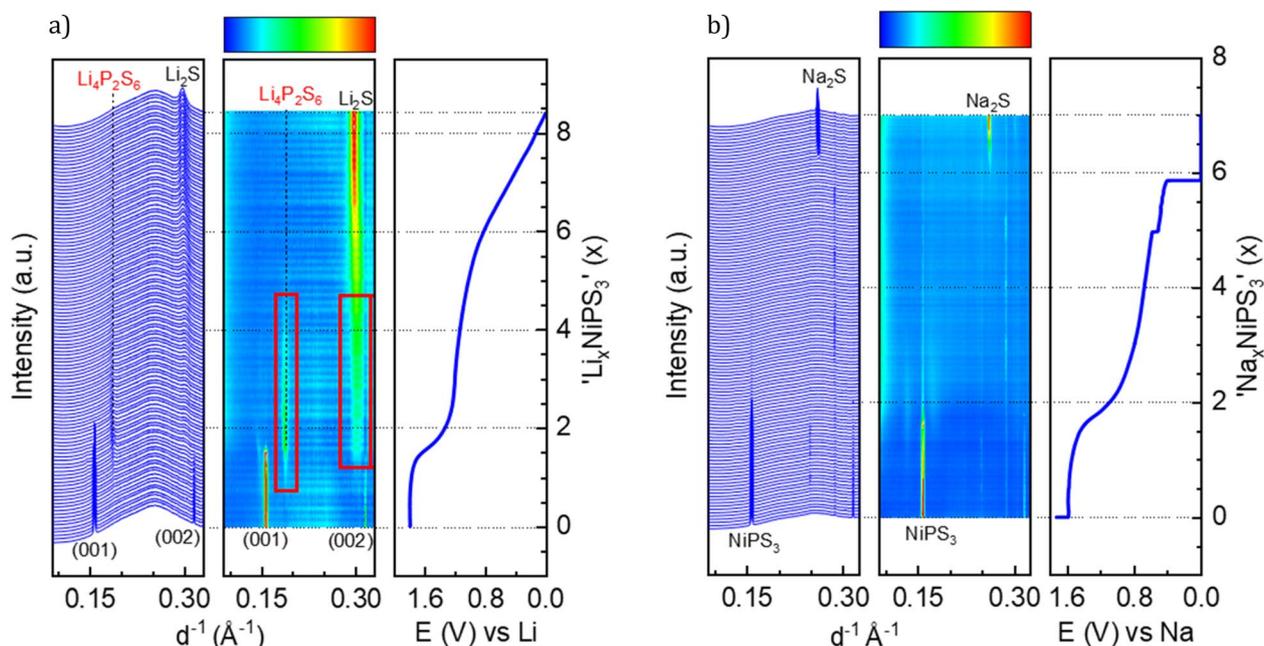


Fig. 2 Waterfall plot, contour plot and potential profile from *operando* XRD. (a) Li_xNiPS_3 , with a current density of $71\ \text{mA g}^{-1}$; red boxes highlight the intermediate phase, and (b) Na_xNiPS_3 with a current density of $61\ \text{mA g}^{-1}$.



potentially be intercalated without affecting the interlayer distance.³⁸ 1/3 of the octahedral spaces belong to the 2*d* Wyckoff position and 2/3 to the 4*h* positions. However, the intercalation behavior of MPS₃ (M – transition metal cation Fe, Mn or Ni) compounds in electrochemical cells can depend on the applied current density: a higher current abruptly decreases the cell potential, thus influencing the Li-ion mobility and structural changes.^{38,39}

We studied Li and Na intercalation behavior in NiPS₃ at current densities significantly exceeding the OCV regime (current densities of 60 to 70 mA g⁻¹). Fig. 2 shows *operando* XRD data collected during intercalation of (a) Li and (b) Na cations into NiPS₃, in the form of a waterfall plot and a contour plot for better visualization, including the corresponding potential curve of an *operando* electrochemical cell.

At the beginning of Li-intercalation up to $x(\text{Li}) = 0.5$ (Li_{0.5}-NiPS₃ composition), Bragg reflections in the diffraction patterns remain unaffected, likely due to insertion into free octahedral spaces.⁴⁰ The second step up to $x(\text{Li}) \sim 1.6$ is accompanied by a continuous decrease in reflection intensities of the parent compound, until they completely disappeared, suggesting structural degradation. This behavior, clearly confirmed by evolution of the characteristic (001) and (002) reflections of NiPS₃ shown in Fig. 2a, is in line with the DFT calculations of Choi *et al.*, which show a significant structural distortion at $x(\text{Li}) = 0.875$ induced by Li intercalation.¹⁰ The authors also observed some changes in the near surrounding of nickel cations starting from Li_{0.8}NiPS₃, using pair distribution function (PDF) analysis. Evolution of reflections corresponding to Li₂S is observed starting from $x = 1.6$. This marks the end of the decomposition process of NiPS₃ according to the reaction $9\text{Li} + \text{NiPS}_3 \rightarrow 3\text{Li}_2\text{S} + \text{Li}_3\text{P} + \text{Ni}$.⁴¹ The fact that no reflections of elemental nickel and lithium phosphide were observed could be due to an amorphous structure and/or a very small crystallite size. Additionally, we could also identify Li₄P₂S₆ as an intermediate phase, based on a set of reflections developing at 0.187 and 0.301 Å⁻¹.⁴² This phase is observed at a Li-content of $0.8 < x < 4.5$ as a pathway for decomposition of the parent structure to Li₂S. The Li₄P₂S₆ phase is not stable during the subsequent lithiation process, as demonstrated by the disappearance of the reflection at 0.187 Å⁻¹ and a slight shift of the reflection at 0.301 Å⁻¹.

Choi *et al.*,¹⁰ based on the results of XPS and Raman measurements, also concluded a multistep reduction process of Ni²⁺, which includes partially reduced inorganic intermediates such as Ni_xPS_y and Li_xPS_y, that eventually leads to Li₂S as the final decomposition product. Formation of Li₂S was also observed by Foot *et al.*³⁸ during intercalation at a low current density for $x > 1.6$.

In a comparable study of Brec *et al.*,⁴⁰ performed at a lower current rate, two separate plateaus are visible in the range of $0 < x(\text{Li}) < 1.5$ and above 2.25 V *vs.* Li⁺/Li (in contrast to the first plateau at 1.75 V *vs.* Li⁺/Li in our case). These plateaus were attributed to two stages of intercalation arising from filling two different Wyckoff positions. Under those conditions close to the equilibrium state, some metastable, partially ordered phases can be formed, which are not detectable when the current density is high. Note that the

authors³⁹ measured single crystals in the presence of a different electrolyte, which can slightly impact the total potential *vs.* Li⁺/Li.

This brought us to the conclusion that the high galvanostatic currents applied to NiPS₃, result in aggravate filling of interlayers in the structure probably due to a kinetic reason, causing an abrupt potential drop that leads to structural decomposition of NiPS₃, since other phases become thermodynamically more stable at such a low cell potential. Therefore, we observed the loss of crystallinity of the initial compound and the formation of lithium decomposition products at much lower states of lithiation than reported in the literature.⁴⁰ It should be mentioned that in contrast to Foot *et al.*,³⁸ who detected Li₂S above $x > 1.6$, the final transformation in our experiments is delayed. After delithiation, some reflection shifts are visible, indicating re-oxidation of Li₂S, but in contrast to Choi *et al.* no crystalline NiPS₃ could be observed.¹⁰

A similar multistep decomposition process of NiPS₃ with the formation of the Na₂S end product was observed during galvanostatic sodium intercalation. The end of the first plateau at $x(\text{Na}) = 1.8$ in the galvanostatic curve coincides with the disappearance of pristine NiPS₃ reflections (Fig. 2b). However, in contrast to lithium, no crystalline intermediate products could be detected during structural decomposition. Moreover, the reflections of Na₂S are sharper than of its Li counterpart suggesting a higher crystallinity of Na₂S. However, the peak evolution occurs at significantly higher $x(\text{Na}) > 6$ values, especially in a constant voltage profile.

3.2 Redox changes in NiPS₃ upon alkali metal intercalation

Furthermore, we performed *operando* XAS measurements at the Ni K-edge upon Li and Na insertion into NiPS₃, which provide direct information about the valence state and local surrounding of the absorbing atom.⁴³ Fig. 3a shows the evolution of the Ni K-edge spectra recorded *operando* during electrochemical Li-intercalation, which was limited to 1 Li per NiPS₃ formula unit, together with the NiO reference spectrum. The edge positions of the NiO reference material and of NiPS₃ are different. The difference arises mostly from various chemical surroundings of Ni and different covalencies of the chemical bonds, which can be qualitatively estimated from Pauling electronegativity values. Although Ni is octahedrally coordinated in both NiO and NiPS₃, the more electronegative oxygen atoms (3.44 Pauling units) in ionic Ni–O bonds strongly attract the electron density from Ni, making it more difficult to remove core level electrons and causing a shift to a higher edge energy, in comparison to more covalent Ni–S bonds with S having an electronegativity of 2.589 Pauling units.

According to the work of Brec and Ouvrard,⁴⁴ the extended X-ray absorption fine structure (EXAFS) analysis of chemically lithiated Li_xNiPS₃ samples points to pronounced migration of Ni cations from the octahedral to tetrahedral sulfur surrounding their simultaneous reduction. Therefore, as per the work of Brec and Ouvrard,⁴⁴ 50% of Ni exists in the zero oxidation state in sulfur tetrahedra in the Li₁Ni_{0.5}⁰Ni_{0.5}²⁺PS₃



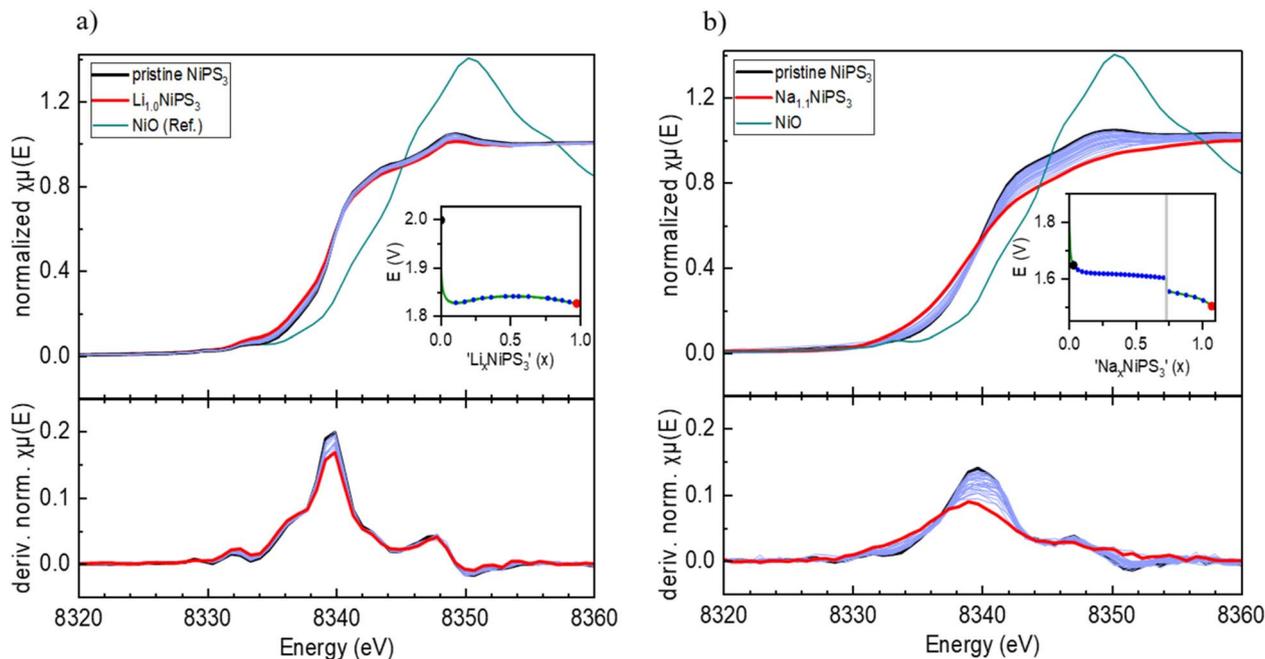


Fig. 3 Operando XAS spectra of alkali ion intercalation into NiPS₃: (a) Ni K-edge and $d\mu(E)/dE$ plots of Li_xNiPS₃ ($0 \leq x \leq 1$). (b) Ni K-edge and $d\mu(E)/dE$ plots of Na_xNiPS₃ ($0 \leq x \leq 1.1$). Insets show the corresponding potential profiles and the specific points of measurement.

composition. The Ni–S distance for the first Ni coordination sphere is reported to decrease from 2.45 Å (NiS₆-octahedra) to 2.30 Å (NiS₄-tetrahedra).

However, the reported Ni-reduction and Ni-diffusion in Li_x-NiPS₃ are in contradiction to our *operando* XAS measurements. According to Fig. 3a, showing Ni K-edge spectra of Li_xNiPS₃ also as $d\mu(E)/dE$ derivative plots, there are no visible changes in the spectra for compositions in the range of $0 \leq x(\text{Li}) \leq 1$. As is well known, the pre-edge of transition metals in the K-edge spectrum originates from a $1s \rightarrow 3d$ electron transition and depends, among other factors, on the symmetry and the coordination number of the central atom.⁴³ In agreement with the dipole selection rules, only a quadrupole transition is allowed, hence showing either zero or a low intensity in the octahedral environment.⁴³ In the case of tetrahedral surrounding, the p- and d-orbitals of Ni overlap and can hybridize, thus enabling a dipole transition from the $1s$ to a hybridized p-orbital.⁴⁵ This should strongly enhance the pre-edge intensity with the pronounced Li-insertion. However, such an enhancement could not be detected in Li-containing compositions. There are also no visible shifts in the main edge and post-edge regions of the Ni K-edge spectra, indicating no change in the coordination sphere and Ni oxidation state. Therefore, we conclude that Ni is redox-inactive during lithiation of Li_xNiPS₃.

Similarly, sodium intercalation into NiPS₃ (Fig. 3b) was studied up to $x(\text{Na}) = 1.1$. The normalized plots show many more changes upon intercalation in comparison to lithium. Here, the maximum in the derivative plots is shifted from 8340 eV to 8339 eV, showing the beginning of partial Ni reduction, in agreement with the work of van Dinter *et al.*⁹ The changes in the post-edge region with increasing sodium content are significant as well. Although no structural changes were

observed till $x(\text{Na}) = 1.5$, the larger size of the Na-ion likely leads to local distortions in the structure and the corresponding partial redox.

In order to further explore interlayer modification, intercalation with a larger organic cation was undertaken. In contrast to the literature where organic salts are used in solvent media, an ionic-liquid mediated approach was used without the need for additional component(s). EMIM-TFSI was selected and the corresponding intercalation of an EMIM cation is presented next.

3.3 Intercalation of EMIM and co-intercalation of (EMIM,Li)

Prior to galvanostatic EMIM insertion experiments, the electrochemical stability window of EMIM-TFSI was probed using a symmetric Pt–Pt two-electrode cell setup at room temperature (Fig. 4a). Cyclic voltammetry measurements, performed between -5 V and 1 V at a low scan rate of 0.1 mV s^{-1} , show an irreversible redox process starting below -4.1 V, which lies outside the working potential range of EMIM intercalation between -2.4 V and -3.9 V vs. Pt.

Insertion of EMIM cations from the EMIM-TFSI electrolyte was performed using either metallic Li or Pt as a counter electrode. In the case of the using a Li-electrode, co-insertion of Li during the long intercalation time is expected and the system is referred to as EMIM-Li. A galvanostatic profile of NiPS₃ during insertion, using EMIM-TFSI electrolyte and a lithium electrode is shown in Fig. 4b along with the data for pure Li intercalation for comparison. The potential curve for EMIM-Li shows two distinct plateaus at 1.8 V and 1.2 V, similar to that of Li intercalation. However, the extent of the first plateau is significantly less, ending at around $x(\text{Li}) \sim 0.6$.



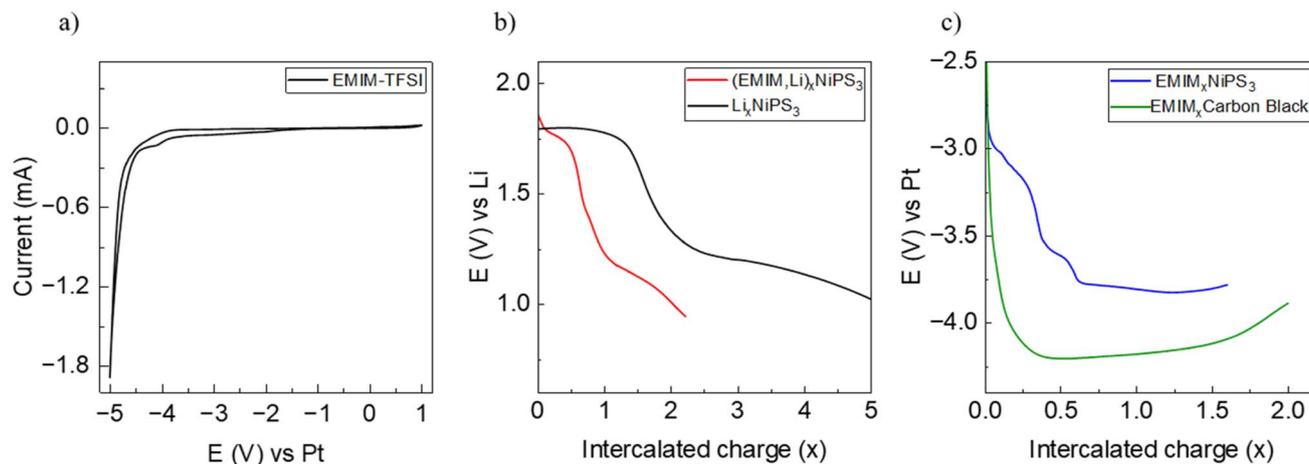


Fig. 4 (a) Cyclic voltammetry of EMIM-TFSI in a symmetric Pt–Pt cell, (b) galvanostatic discharge of NiPS₃ with Li (black), (EMIM,Li) (red), and (c) EMIM-only (blue), and galvanostatic discharge of a carbon black material in EMIM-TFSI vs. Pt (green).

Fig. 4c shows the electrochemical EMIM intercalation against the Pt counter electrode. The overall potential difference is due to the Pt–Li difference in the galvanic series. The reference measurement (green line) shows the galvanostatic curve of the cell with carbon black as a working electrode. However, since the potential drops below -4.1 V vs. Pt, the observed curve most likely represents decomposition of the electrolyte. An initial plateau is seen at ~ -3.3 V, which gradually shifts to a higher potential during intercalation likely due to a reduction in the overpotential of the material with continued intercalation.

3.4 Crystal and electronic structure changes

Structural changes in NiPS₃ during reaction with EMIM-TFSI were monitored by *operando* XRD measurements using EMIM-TFSI electrolyte. Fig. 5 shows the XRD data where Li and Pt metals were used as counter electrodes. Here, a new crystalline phase is formed during the insertion of EMIM-cations in cells with both Li and Pt counter electrodes. In the case of (EMIM,Li) co-intercalation (Fig. 5b), the conversion represents a two-phase process with subsequent disappearance of the pristine phase, which is characterized by the first galvanostatic plateau up to $x = 0.6$, and growing reflections of a new phase. The most pronounced change between two phases is related to the (00 l) reflections, see for example a clear shift of the (001) reflection from 0.1578 \AA^{-1} in pristine NiPS₃ to 0.0885 \AA^{-1} in the new phase (Fig. 5b). The newly formed phase has the (001') reflection at much smaller reciprocal lattice lengths, signaling an expansion of the NiPS₃ interlayers. This phase transformation is completed with the EMIM_{0.6}NiPS₃ composition at the end of the first plateau in the potential profile. During further intercalation, no other changes occur in the lattice parameters. Only a fading of the intensity of the (001') reflection is seen for $x > \sim 1.4$, likely as a result of continued Li intercalation. The intercalation of pure EMIM ions into NiPS₃ is similar to the co-intercalation process (Fig. 5a) with the evolution of a new phase. In contrast to the loss of pristine NiPS₃ reflections in alkali

metal intercalation (Fig. 2) and (EMIM-Li), with pure EMIM intercalation the reflections of the initial compound are merely faded until an effective inserted charge of $\sim 2.4x$. Although a new phase is formed as well, the conversion process is not completed even after reaching a composition of EMIM_{2.4}NiPS₃. In addition, the new (001') reflection is at a slightly higher $1/d$ position of 0.1010 \AA^{-1} , indicating less expansion along the c -axis. Its intensity does not appear to fade and, therefore, one can conclude that there is no visible decomposition here, in contrast to the co-intercalation.

For comparison, Fig. 5c shows the X-ray powder diffractograms of pristine NiPS₃ and the materials after intercalation, corresponding to the (EMIM,Li)_{0.5}NiPS₃ and EMIM_{0.5}NiPS₃ compositions. Additional reflections and an increased background of the intercalated samples originate from *operando* cell components like the copper current collector and ionic liquid. The increase in the lattice parameter c , which is related to the interlayer distance in the ab -plane, from 6.62 \AA to 10.65 \AA , defined by using the Le Bail analysis method on the assumption of the same symmetry of the new phase (see Fig. S2 in the ESI†), is in agreement with a common observation for intercalation of organic ions into transition metal phosphorus trisulfides.^{13,19,21,46} For example, tetrabutylammonium (TBA⁺) intercalation into NiPS₃ was demonstrated by Choi *et al.* with an increased c -parameter of 14.5 – 15.3 \AA .¹⁰ The variation is a result of various crystallographic phases with different orientations of the organic molecule between the layers. In contrast, all EMIM_{0.5}NiPS₃ reflections could be assigned to only one crystallographic phase. Interestingly, a similar reaction between TBA⁺ and NiPS₃ was performed by Mi *et al.*, demonstrating an interlayer distance of only 11.46 \AA .¹³ This may be an effect of different amounts of intercalation, leading to the assumption that this is a stepwise process, which is supported by the work of Chen *et al.*¹⁹ They investigated the reaction between FePS₃ and 1,10-phenanthroline after certain intercalation intervals with *ex situ* XRD, and also observed different stages resulting from changes in the intercalant orientation. Changes in oxidation



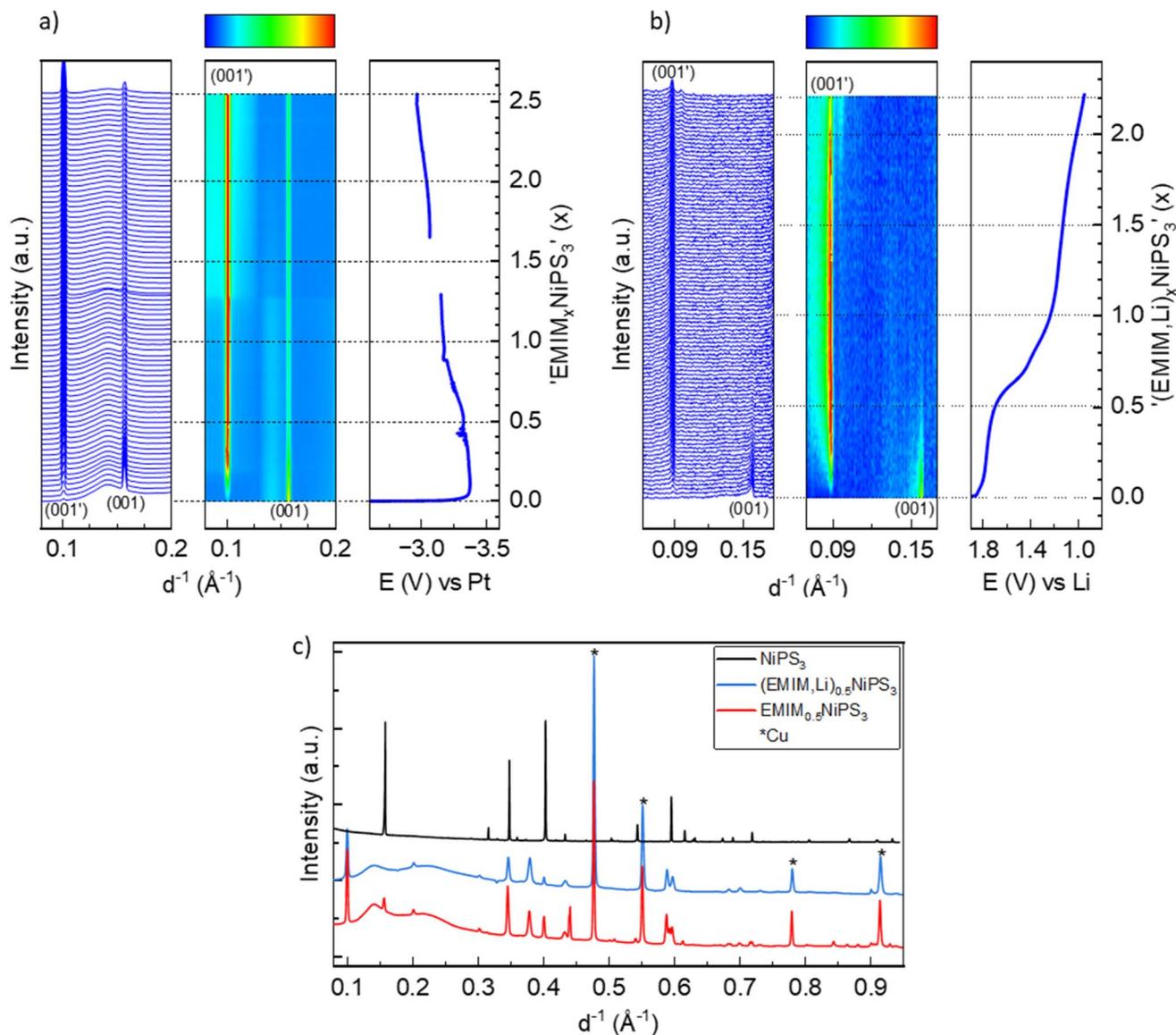


Fig. 5 (a) Waterfall plot, contour plot and potential profile from *operando* XRD of NiPS₃ with EMIM-TFSI vs. Pt, and vs. Li (b). (c) XRD of the pristine (black line) and intercalated phases corresponding to the (EMIM,Li)_{0.5}NiPS₃ (blue line) and EMIM_{0.5}NiPS₃ (red line) composition.

states and the local structure upon EMIM intercalation were monitored with *operando* Ni K-edge XAS, and *ex situ* P and S K-edge XAS measurements, as shown in Fig. 6 and 7.

Similar to Li-insertion into NiPS₃, the Ni K-edge spectra of intercalated (EMIM,Li)_xNiPS₃ and (EMIM)_xNiPS₃ resemble each other on increasing the *x*-value, which is clearly seen in the first derivative graphs of the normalized absorption coefficient (Fig. 6).

Here we would like to make a note of caution. As recently reported,⁴⁷ exposure of battery components such as electrolytes or electrodes to X-ray radiation in XAS experiments can induce some material damage, leading to incorrect interpretation of the obtained results. For example, it was shown that reactivity of the InSb/Al electrode in a Mg-battery with organic electrolytes is significantly retarded during the *operando* synchrotron X-ray absorption measurement, and the spectra evolution occurred

only after a relaxation time, where the beam was powered off.⁴⁷ To exclude such an effect in our case, we undertook multiple measurements to avoid a misinterpretation of data. First, an *operando* combined XAS/XRD experiment was performed at the NOTOS beamline (ALBA, Spain) enabling simultaneous monitoring of the crystal and electronic structures of NiPS₃ upon EMIM intercalation. The electrochemical cell design including electrode thickness and material density was almost the same as those in the XAS experiments at the P64 Petra III beamline (DESY, Germany). The maximal flux on the sample at NOTOS corresponds to 1×10^{11} Ph s⁻¹ in the energy range of 3d metals, and 1×10^{13} Ph s⁻¹ at the P64 beamline. At NOTOS, the Ni K-edge spectra were recorded with an exposure time of 10 min, which was only half of the exposure time at P64. In addition, *ex situ* Ni K-edge measurements were performed at HiKE/KMC-1 (BESSY II, Germany) on (EMIM)_xNiPS₃ samples at various



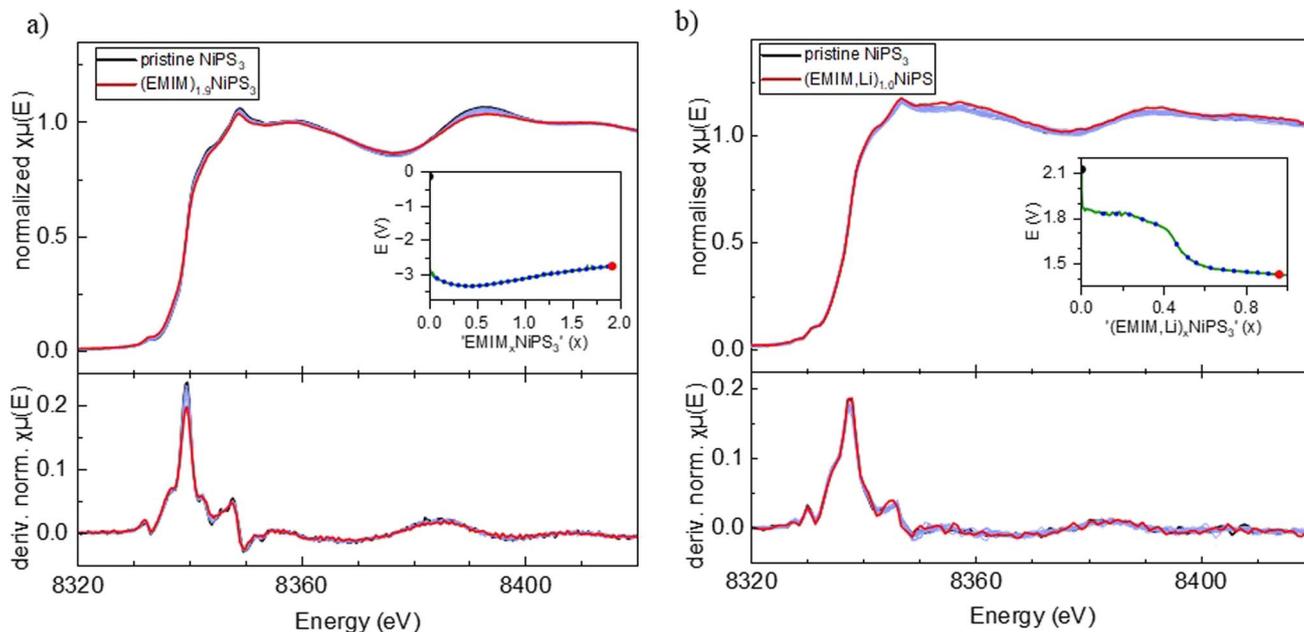


Fig. 6 Operando Ni K-edge XAS data. (a) EMIM cation intercalated NiPS₃ including the first derivative of the normalized absorption coefficient. (b) EMIM and Li co-intercalated NiPS₃ including the first derivative of the normalized absorption coefficient. Insets show the corresponding potential profiles and the specific points of measurement.

intercalation stages. In the EMIM intercalation experiment at NOTOS, the two-phase reaction mechanism was confirmed *operando* XRD while the observed changes in the Ni K-edge spectra were close to zero, in agreement with the *operando* results from the P64 beamline of PETRA III and *ex situ* results from the HiKE/KMC-1 beamline of BESSY II. The uniformity of all Ni K-edge spectra for various (EMIM)_xNiPS₃ samples recorded at different X-ray source facilities, unambiguously confirms Ni-inertness during either Li intercalation, (EMIM-Li) co-intercalation or only EMIM intercalation.

In contrast, changes in the P and S K-edge spectra of pristine and intercalated materials are more noticeable (Fig. 7). The P K-edge spectra of (EMIM,Li)-co-intercalated and (EMIM)-intercalated samples differ from each other, pointing to different electronic processes upon Li and EMIM insertion into NiPS₃. The near-edge region of the P K-edge spectrum of NiPS₃ consists of two peaks. The energy position of the corresponding adsorption edge was set as the maximum of the first derivative of the normalized absorption coefficient, corresponding to the inflection point.⁴⁸ Based on the “rigid-band electronic model”,⁴⁹ the low-energy shoulder at 2145.5 eV might correspond to the transition from the P 1s orbital to the antibonding $\sigma^*(\text{P-P})$ orbital of the (P₂S₆)⁴⁻ entity, whereas the maximum at 2147.2 eV probably reflects the transition P 1s → antibonding $\sigma^*(\text{P-S})$ orbital, since the $\sigma^*(\text{P-P})$ level is assumed to be low-lying in energy.⁵⁰ Upon co-intercalation of EMIM- and Li-cations, the intensity of the first peak noticeably decreases, indicating a decreasing amount of unoccupied states. A natural interpretation of this observation is that these states become occupied by electrons from the intercalated Li or EMIM. This is consistent with the fact that the first shoulder continuously loses

intensity upon (EMIM,Li) intercalation. At the same time, the peak at 2147.5 eV shifts slightly to lower energy and becomes significantly broader, pointing to an increased electron density at P (Fig. 7). There are also noticeable changes in the near-edge region at higher energy.

Generally, phosphorus has a narrow range of absorption edge energies for species with various oxidation states, which makes the procedure of conventional determination of oxidation numbers difficult: for example, the energy difference is about 2.5 eV for compounds with the formal oxidation state varying from 0 (elemental phosphorus) to P⁵⁺ in Na₄P₂O₇.⁵¹ Furthermore, with increasing Li-content in the samples, a new peak at a much lower energy of 2143.2 eV arises for both Li intercalation and (EMIM-Li) co-intercalation, which is in line with the onset of structural decomposition of NiPS₃ at the pronounced intercalation confirmed by our *operando* XRD studies in both.

In the case of pure EMIM intercalation (Fig. 7b), the P K-edge spectra of samples with various amounts of EMIM show less changes. The positions of two main peaks in the derivative plots at 2145.5 eV and 2147.2 eV do not change with the EMIM content. However, the maximum of the white line systematically grows with a higher EMIM content, in contrast to that of the Li-containing samples. The near-edge structures at higher energies also remain almost constant. We conclude that pure EMIM intercalation, in contrast to Li intercalation and (EMIM-Li) intercalation, does not affect the electronic structure and the crystal structure around P.

We also recorded spectra of P₂S₅ as a reference material for phosphorus and sulfur. In both NiPS₃ and P₂S₅ compounds, P-atoms have a distorted tetrahedral surrounding of three S-atoms with bond lengths varying between 2.02 and 2.10 Å.



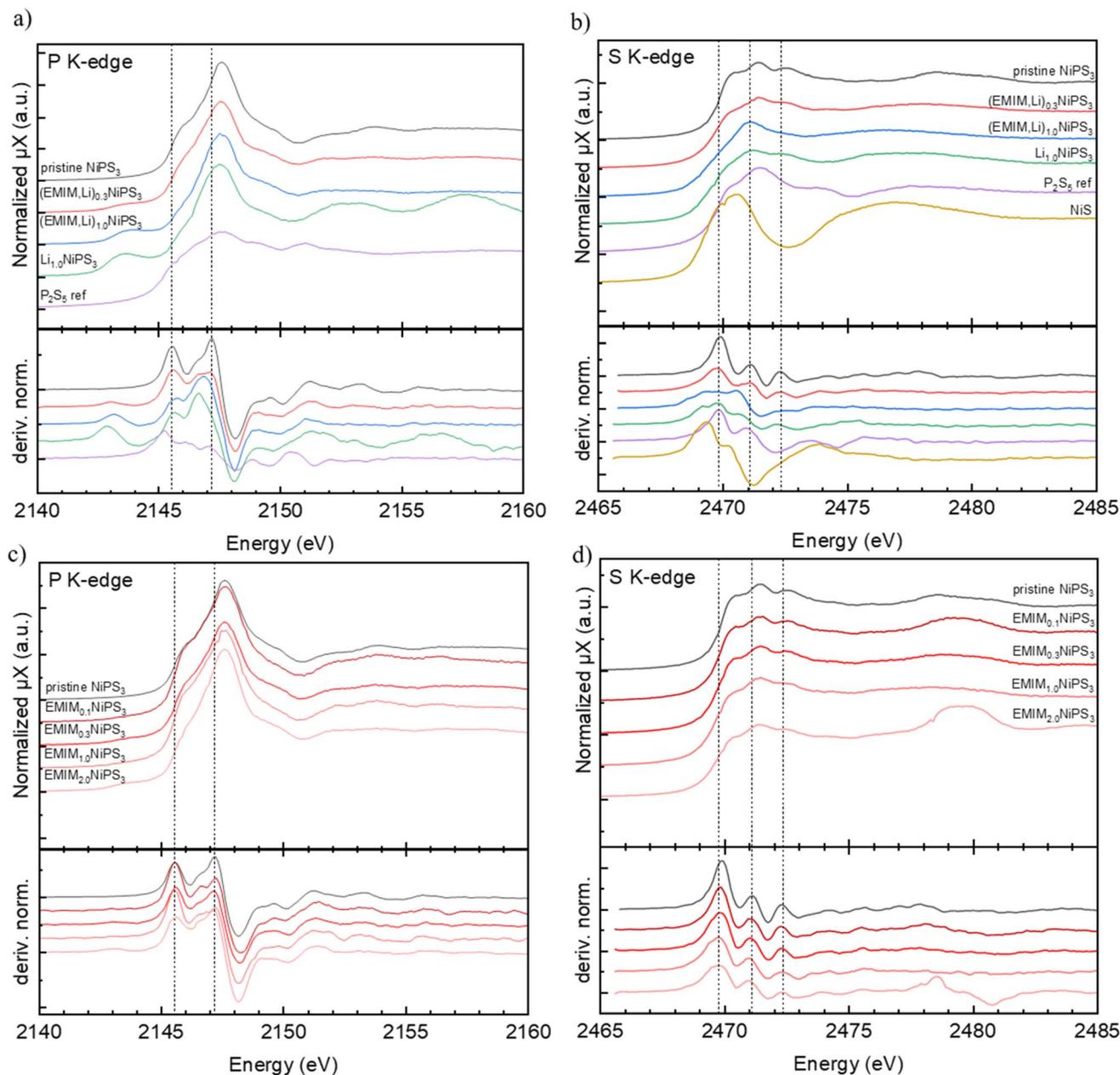


Fig. 7 *Ex situ* XAS data, (a) P K-edge of pristine, EMIM and Li co-intercalated NiPS₃ and a P₂S₅ reference material including the first derivative of the normalized absorption coefficient. (b) S K-edge of pristine, EMIM and Li co-intercalated NiPS₃ and a P₂S₅ reference material including the first derivative of the normalized absorption coefficient. (c) P K-edge of pristine and EMIM intercalated NiPS₃. (d) S K-edge of pristine and EMIM intercalated NiPS₃. Data has been plotted with y-axis offsets for better visualization.

The fourth bond in the tetrahedron is either shorter like the P=S double bond of 1.91 Å in P₂S₅, or longer like the P-P single bond of 2.15 Å in NiPS₃. Despite the similarity in the structural surrounding, the P near-edge spectra of P₂S₅ and NiPS₃ are rather different. The first maximum in the derivative plot of the normalized absorption coefficient for P₂S₅ is shifted to a lower energy of 2145.05 eV, compared to 2145.5 eV in NiPS₃.

Considering the S K-edge spectra, the changes in the case of (EMIM,Li) co-intercalation are again much more pronounced than those after EMIM intercalation (Fig. 7c and d). The near-edge region in NiPS₃ is composed of three well-distinguished

peaks, which correspond to a transition of S 1s to various possible empty levels such as hybridized S 3p – Ni 3d orbitals and a $\sigma^*(\text{P-S})$ orbital of the (P₂S₆)⁴⁻ unit.⁵² The assignment of peaks to certain transitions in the case of S K-edge spectra is not straightforward without supporting theoretical calculations, since the electronic structure of pristine NiPS₃ around the Fermi level has not been well-understood until now.⁵³ The first two peaks in the derivative plots of NiPS₃ with energy positions at 2469.8 eV and 2471.1 eV are very close to spectral features of the P₂S₅ reference.



After (EMIM,Li) co-intercalation (Fig. 7a and b), a new pre-edge feature at 2469.0 eV appears in the S K-edge spectra of (EMIM,Li)_xNiPS₃ samples. Simultaneously, the derivative peak at 2471.1 eV moved to a lower energy of 2470.5 eV. The third peak at 2472.3 eV remained nearly constant. Similar to the P K-edge, the intensity of the first shoulder at 2470 eV decreases upon (EMIM,Li) intercalation. These states are understood to be occupied by electrons donated by Li. The overall shift to lower energies reflects the increasing electron density. After intercalation of EMIM (Fig. 7c and d), only a broadening of the first maximum in the derivative plots could be detected, and the other peaks remained unchanged. In particular, the intensity of the first shoulder does not decrease.

Therefore, our spectroscopic experiments showed noticeable changes in the electronic structure of S and P in layered NiPS₃ upon Li and EMIM co-intercalation, while Ni cations were not affected at all. In particular, our results show that the first unoccupied states of S and P are filled by electrons. Hence, the (P₂S₆)⁴⁻ sublattice absorbs the electrons donated by Li-intercalation. The changes are much more pronounced upon insertion of small Li-cations compared to bigger EMIM molecules. EMIM intercalation appears to be completely ineffective in impacting the electronic structure. In the case of Na intercalation, Ni reduction was observed, pointing to a conventional redox reaction mechanism. This is in accordance with the NiPS₃ structure decomposition detected by *operando* XRD measurements.

3.5 Elemental analysis

To address the open question regarding the charge compensation mechanism(s), elemental analysis by ICP-OES was performed before and after intercalation. It is known from other 2D MPS₃ materials that the transition metal can leave the host structure during intercalation, for example during intercalation of *N*-alkylamine in FePS₃.⁵⁴ The mass fractions of Ni and P were determined in both the untreated sample and two intercalated NiPS₃ compounds, after EMIM intercalation with Li and Pt counter electrodes for $x = 1$ charge transfer. The Ni/P molar ratios for the pristine, EMIM-intercalated and (EMIM,Li) co-

intercalated samples were found to be rather similar and close to 1, confirming no elemental losses due to dissolution or secondary reactions.

Additionally, using ICP-OES the amount of lithium in (EMIM,Li)₁NiPS₃ (intercalation *vs.* the Li electrode) was determined to be 0.59 Li per NiPS₃ unit. This shows the preferential intercalation of Li ions over EMIM ions, even when the electrolyte did not contain any Li in the beginning. Furthermore, in the case of pure EMIM cation intercalation, the effective EMIM content in the material was evaluated to be 0.74 ± 0.07 , based on the mass fraction of Ni and P in EMIM₁NiPS₃ (1EMIM from the galvanostatic experiment) using ICP-OES. This intercalated amount is less than the calculated introduced charge through GCPL. This difference could be due to the unavoidable washing step to remove excess of ionic liquid, which might also remove surface-adsorbed EMIM ions and/or portions of carbon and the binder. Another possibility may be other redox reactions occurring in parallel to intercalation.

3.6 Cation mobility in intercalated NiPS₃

Based on the chemical analysis, a co-intercalation process of Li and EMIM into NiPS₃ occurs in the cell with a Li-anode and EMIM-TFSI electrolyte, pointing to unequal rates of Li and EMIM intercalation kinetics. We studied cation mobility in NiPS₃ during the intercalation process in the EMIM-TFSI electrolyte with Li or Pt as a counter electrode using the GITT. In the case of (EMIM,Li) co-intercalation, determination of only an average diffusion coefficient for both cations is possible. Starting from $2 \times 10^{-9} \text{ cm}^2 \text{ s}^{-1}$, the average $D_{\text{EMIM,Li}}$ diffusion coefficient shows a small maximum during the potential decrease from 1.9 to 1.7 V *vs.* Li⁺/Li, followed by a drop of several orders of magnitude, reaching a nearly constant value of $1 \times 10^{-12} \text{ cm}^2 \text{ s}^{-1}$ at a cell potential plateau of 1.7 V *vs.* Li⁺/Li (Fig. 8). The diffusion coefficient of a single EMIM cation in NiPS₃ measured in cells with a Pt counter electrode shows at the beginning of intercalation a minimum of $D_{\text{EMIM}} = 1.8 \times 10^{-16} \text{ cm}^2 \text{ s}^{-1}$, and increases again up to $1.7 \times 10^{-13} \text{ cm}^2 \text{ s}^{-1}$. From this value, the coefficient decreases slowly with increasing intercalation. In direct comparison to the average (EMIM,Li)

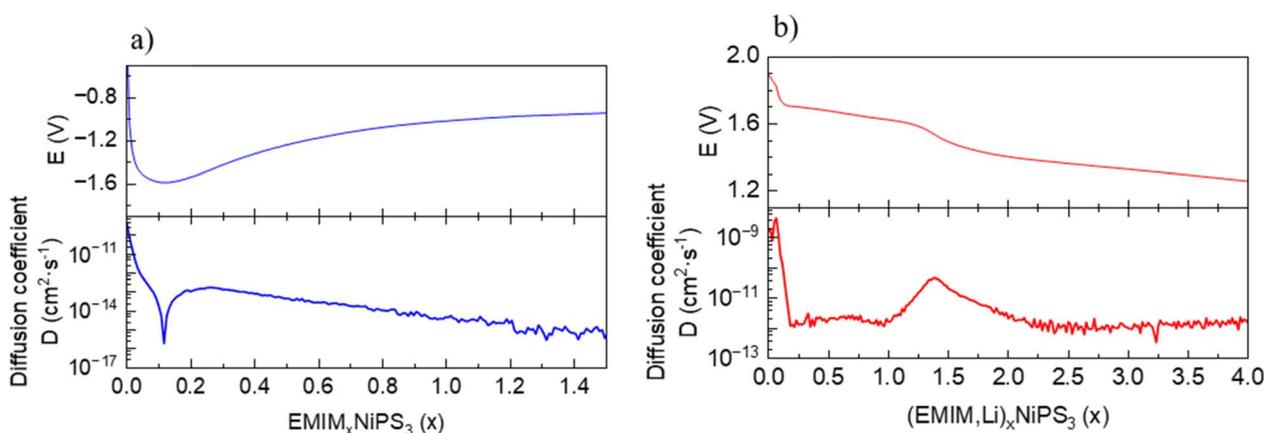


Fig. 8 Equilibrium potential and diffusion coefficient dependence on the state of charge for: (a) intercalation of EMIM in NiPS₃. (b) Co-intercalation of Li and EMIM in NiPS₃.



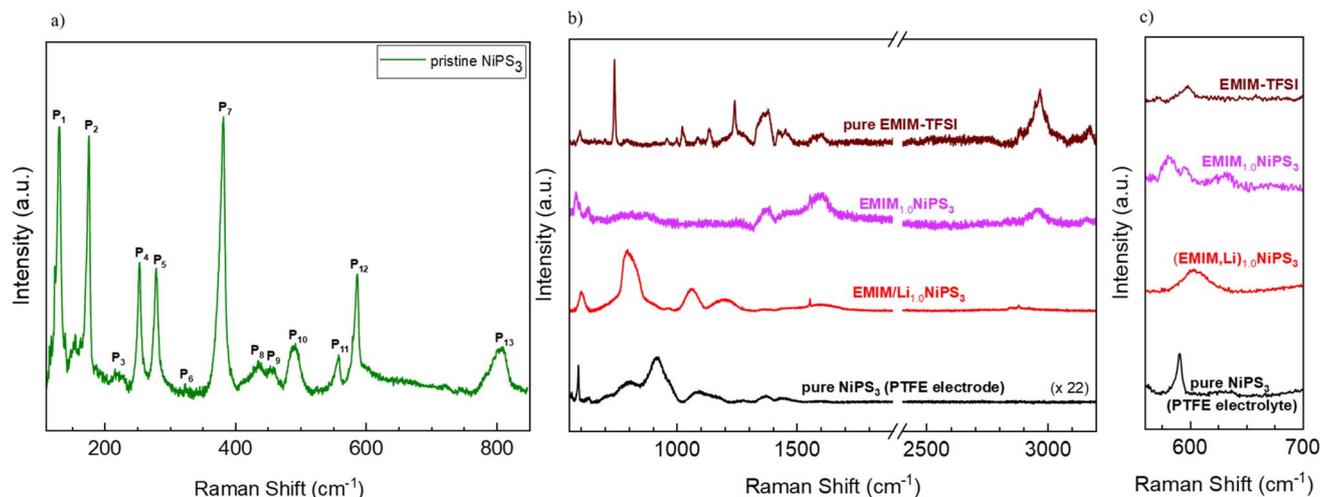


Fig. 9 (a) Raman spectrum of pristine NiPS₃ in a low-frequency range, using excitation at 514 nm. (b) Raman spectra in the high-frequency range of EMIM-TFSI, EMIM-intercalated NiPS₃, (EMIM,Li) co-intercalated NiPS₃, and pristine NiPS₃ combined with PTFE, using a 405 nm laser. (c) Zoomed-in view of the middle graph. All spectra were obtained using a 2400 grating. In (b) and (c), data are plotted with offset for better visualization.

diffusion in (EMIM,Li)_xNiPS₃, the diffusion coefficient of the single EMIM cation in (EMIM)_xNiPS₃ is always some orders of magnitude lower, since the smaller Li cation has a higher mobility.

3.7 Confirmation of cationic intercalation with Raman spectroscopy

Raman spectroscopy was also used to investigate the intercalation of EMIM moieties in the van der Waals interlayers of NiPS₃. For pristine NiPS₃, 514 nm excitation was used for the Raman measurement due to the lower cut-off filter available with this setup configuration, whereas 405 nm excitation was applied for EMIM-TFSI, pristine NiPS₃ (combined with PTFE), and intercalated samples considering an expected enhanced cross-section of the EMIM unit at this wavelength.⁵⁵ The characteristic peaks of pristine NiPS₃ can be seen in the low frequency range (Fig. 9a), matching the values reported in the literature.⁵⁶ Based on its crystal structure, NiPS₃ can have 15 active Raman modes (monoclinic, C_{2v}).⁵⁶ Due to the layered character of the structure with a small interlayer vibronic interaction (*D*_{3d} symmetry),⁵⁷ some vibrations become degenerate, *i.e.* symmetric and antisymmetric vibrations appear at the same frequency, reducing the number of peaks to 8 fundamental and some higher order modes as presented in Table S1.† Generally, the peaks P₁ and P₂ can be attributed to translational motions of Ni²⁺ ions whereas at higher frequencies, P₃, P₄, P₅, and P₇ are mostly due to the vibrations from the (P₂S₆)⁴⁻ units.⁵⁸

Characteristic Raman modes of EMIM-TFSI are very pronounced in the high-frequency region (Fig. 9b). The broad peaks between 1350 and 3200 cm⁻¹ belong mostly to vibrations of the EMIM-cation ring and the ethyl- and methyl-groups connected to the ring.⁵⁹ Two very intense and narrow signals at 760 cm⁻¹ and 1250 cm⁻¹ originate from vibrations in the TFSI-anion.⁶⁰ The effects of EMIM ion intercalation on the Raman spectrum of NiPS₃ are threefold: (i) the normal modes of the NiPS₃ host material

vanished (for example, P₁₃ at ~800 cm⁻¹), and some frequency shift occurred (P₁₂ at ~580 cm⁻¹), (ii) some bands in the spectrum of EMIM_{1.0}NiPS₃ can be assigned to EMIM-associated vibrational modes, for example at 1380 cm⁻¹, 1480 cm⁻¹, and 2970 cm⁻¹, and (iii) two new signals at 635 cm⁻¹ and 1600 cm⁻¹ arose (Fig. 9c). Although the signal at 1600 cm⁻¹ is already present in the EMIM-TFSI spectrum, a significant increase in the intensity could imply additional contribution from some processes. Note that the increased intensity in this region also has a contribution from the conductive carbon additive. Nevertheless, the absence of the very intense signals from TFSI⁻ in EMIM_{1.0}NiPS₃ confirms the incorporation of only EMIM ions into NiPS₃.

It is known from the literature that at sufficiently low potentials, EMIM cations can be electrochemically reduced on a Ag-electrode.⁶¹ Two additional signals at 1330 cm⁻¹ and 1607 cm⁻¹ were registered amongst others in the Raman spectra of reduction products under 638 nm laser excitation, which were ascribed to free radicals, carbenes, and different dimers.⁶¹ A reductive decomposition of EMIM cations might occur in contact with NiPS₃ as well. We assume that the EMIM cations are first intercalated into the van der Waals interlayers, as also evidenced by the layer expansion determined *via* XRD. Then, during the reduction process, dimers are possibly formed. On the other hand, reduction could also occur on the NiPS₃-surface followed by desorption or release of generated neutral molecules. However, other spectroscopic techniques such as proton nuclear magnetic resonance spectroscopy (H-NMR), infrared spectroscopy (IR) and electron paramagnetic resonance (EPR) spectroscopy have to be employed for further detailed investigations. Interestingly, (EMIM,Li) co-insertion results in a quite different Raman response, with a strong shift of signals of the NiPS₃ host (Fig. 9b). Unfortunately, in the literature, Raman studies of Li-intercalated NiPS₃ are limited to the low-frequency region,⁵³ pointing to a need for deeper Raman studies in a future work.



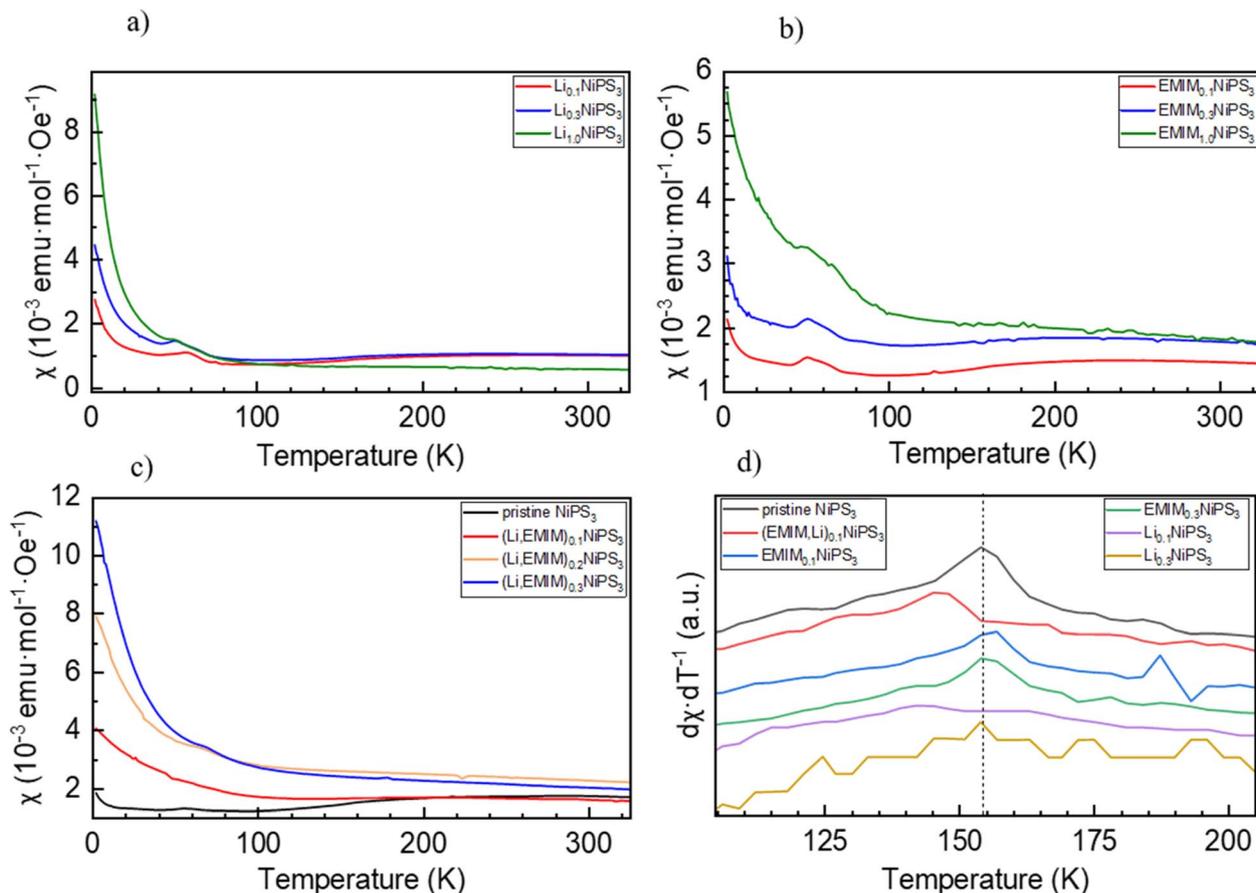


Fig. 10 Susceptibility (zero-field cooling) versus temperature plot for intercalated NiPS₃ powder samples. (a) Exclusively lithium ion intercalation, (b) exclusively EMIM intercalation, (c) Li and EMIM cation co-intercalation, and (d) first derivative of the temperature dependent susceptibility curves. The maximum corresponds to the Néel temperature of the compound.

3.8 Magnetization studies

As is known, NiPS₃ represents a quasi-two-dimensional anti-ferromagnet with a Néel temperature of 155 K.⁶² Above this temperature, there is a broad maximum in magnetization reflecting strong short-range magnetic interactions up to around 300 K.⁴⁰ It was also reported that intercalated Li_xNiPS₃ compounds exhibit a paramagnetic region above 300 K but decompose below 350 K.⁶³ Changes in magnetic properties of NiPS₃ after intercalation of Li and EMIM cations were evaluated using a SQUID magnetometer. Fig. 10 depicts the temperature dependence of susceptibility for the pristine compound and samples at different intercalation levels. The susceptibility curve of the initial NiPS₃ shows a broad maximum at 280 K, in accordance with the literature.^{35,57} Observations of a Curie-Weiss paramagnetic behaviour up to the upper temperature range of 330 K was not possible. A small peak at 55 K, observed in all measurements, came from oxygen traces in the system, which becomes solid at that point.⁶⁴ At temperatures below 10 K, there is another significant increase in magnetization, arising from defects in crystalline NiPS₃.

For all intercalated samples, the susceptibility increases in comparison to that of pristine NiPS₃, especially at low temperatures, while the broad maximum at 280 K vanishes. The Néel

temperature of intercalated materials was determined as a maximum of the first derivative of the susceptibility versus temperature plot. As one can see from Fig. 10d, all materials show nearly the same transition temperature of 154 K. Therefore, we conclude that the susceptibility curves emerge from a combination of the antiferromagnetic pristine phase and a paramagnetic-like intercalated phase. By comparing the magnetic behaviour of the three different intercalation systems (Li⁺, EMIM ion and co-intercalation), it is notable that co-intercalation leads to the highest susceptibilities, both in low and high temperature ranges. Applying the modified Curie-Weiss law (1) with a temperature-independent χ_0 contribution from core diamagnetism and Van Vleck paramagnetism in the temperature range of 300–330 K gives values of the Curie constant C of 0.07 emu K mol⁻¹ Oe⁻¹ for Li_{1.0}NiPS₃, 0.10 for EMIM_{1.0}NiPS₃ and 0.13 for (EMIM,Li)_{0.3}NiPS₃.

$$\chi = \frac{C}{T - \Theta} + \chi_0 \quad (1)$$

These values correspond to nearly similar paramagnetic moments, calculated as $\mu_{\text{exp}} = \sqrt{8 \cdot C}$, of 0.75 μ_B for Li_{1.0}NiPS₃, 0.93 μ_B for EMIM_{1.0}NiPS₃ and 1.0 μ_B for (EMIM,Li)_{0.3}NiPS₃,



which are significantly lower than the spin-only theoretical paramagnetic moment of $2.83 \mu_B$ for Ni^{2+} with a d^8 electron configuration ($S = 1$). This can indicate that there are still very strong magnetic interactions between Ni cations in this narrow temperature range of 300–330 K, which decrease the calculated paramagnetic moment. The more paramagnetic-like magnetization curve for intercalated materials points to decreased interaction between $NiPS_3$ -layers due to the inter-layer expansion. From these results, it is not possible to conclude whether Ni is reduced during intercalation. Formation of Ni^+ or a mixture of $0.5Ni^{2+}/0.5Ni^0$ would reduce the total paramagnetic moment of the system to $1.73 \mu_B$ or $2.0 \mu_B$, respectively, which is still much higher than the experimental value.

3.9 Single crystal X-ray diffraction (SCXRD) and TEM characterization of intercalated $NiPS_3$

A combination of electron diffraction, TEM, and X-ray diffraction was employed in order to analyse the local structure of $(EMIM,Li)_xNiPS_3$ (Fig. 11). 3D electron diffraction taken from $(EMIM,Li)_xNiPS_3$ powder exhibits characteristic in-plane d -spacings, $d(200) = 2.88 \text{ \AA}$ and $d(020) = 1.69 \text{ \AA}$. We can correlate the d -spacing values with the values obtained for the parent $NiPS_3$ and, hence, assign the (hkl) values. However, in the direction orthogonal to the sample plane, only lines with continuous intensity distribution are seen instead of spots due to a thin film diffraction effect. A d -spacing for the longest axis of about 10 \AA was taken from powder XRD data, in agreement with the above-discussed statement.

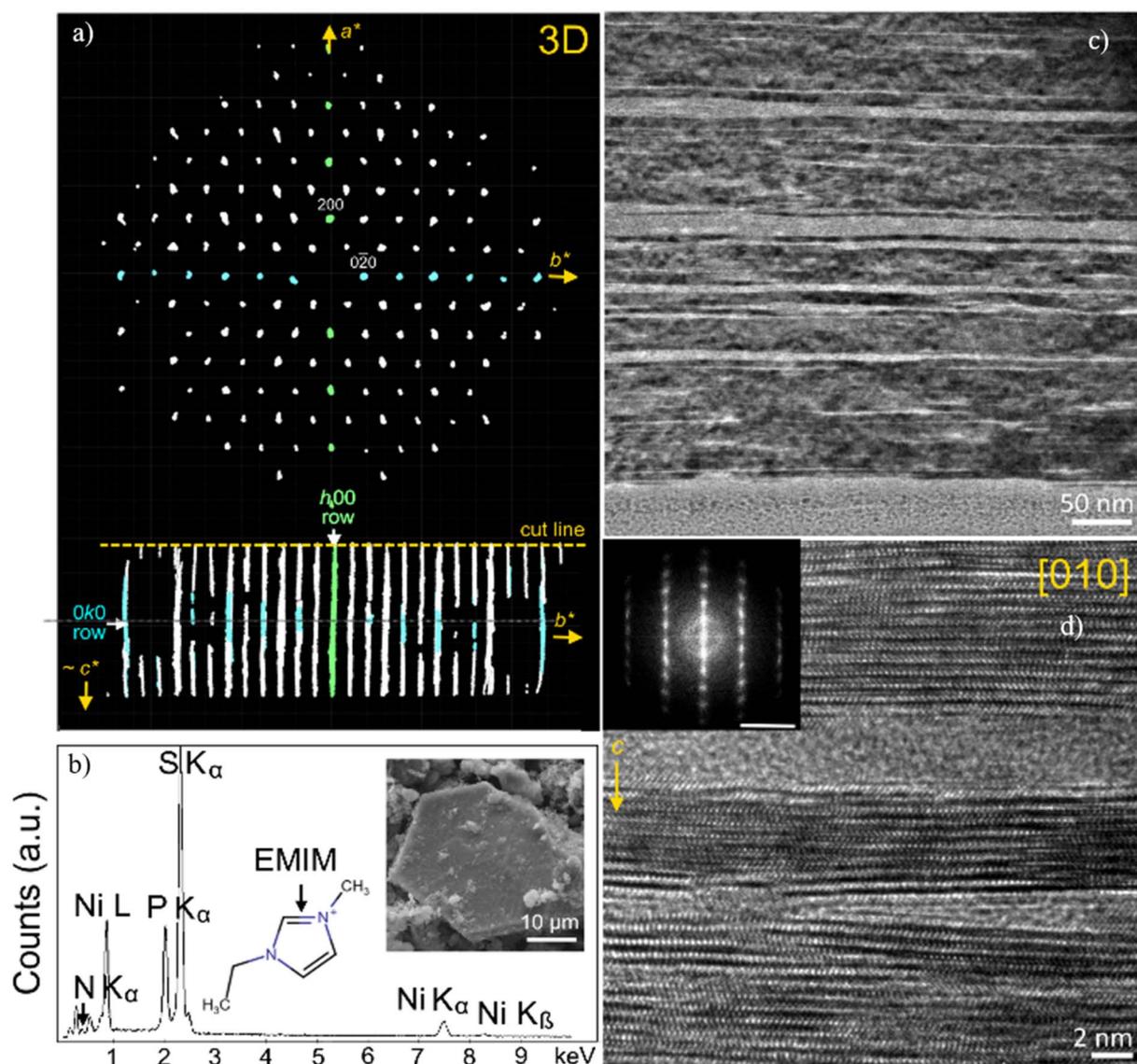


Fig. 11 (a) 3D ED pattern of $(EMIM)_xNiPS_3$ visualized by REDp software.⁶⁵ Two spots in the basal plane are indexed as (200) and (020) according to the corresponding diffraction pattern of the parent $NiPS_3$. In the c -axis direction, only lines with a continuous intensity distribution are seen due to a thin film diffraction effect. (b) SEM EDX spectra of $(EMIM)_xNiPS_3$. The inset shows the crystal used for data collection. (c and d) HRTEM images of the specimen after EMIM intercalation with regions of crystalline $NiPS_3$ and amorphous regions. The inset shows the FFT of the corresponding HRTEM image. The scale bar is 5 nm^{-1} .



SCXRD reflections collected from a $(\text{EMIM,Li})_x\text{NiPS}_3$ crystal can be indexed either in a monoclinic space group $C2/m$ with the cell parameters $a = 5.867(3) \text{ \AA}$, $b = 3.391(2) \text{ \AA}$, $c = 10.075(6) \text{ \AA}$, $\beta = 101.36(2)^\circ$, and $V = 196.5(2) \text{ \AA}^3$, or in a higher symmetry trigonal space group $R\bar{3}$ with the cell parameters $a = b = 3.394(2) \text{ \AA}$, $c = 29.72(3) \text{ \AA}$, $\beta = 101.36(2)^\circ$, and $V = 296.6(5) \text{ \AA}^3$. The latter is consistent with electron diffraction and powder XRD data and reflects a known behavior of intercalates when, due to a massive stacking disorder in the out-of-plane dimension, the space group symmetry for the average structure downgrades into a layer group symmetry. A trigonal lattice was used for further full matrix least-squares refinement, and the final cycle was based on 604 reflections measured ($4.11^\circ \leq 2\theta \leq 56.178^\circ$); of them 168 were unique ($R_{\text{int}} = 0.0445$, $R_{\text{sigma}} = 0.0463$) and converged to final $R_1 = 0.1321$ ($I > 2\sigma(I)$) and $wR_2 = 0.3223$ (all data). The largest peak/hole in the difference Fourier map ($F_o - F_c$) was 1.77 and $-1.45/e \text{ \AA}^{-3}$ respectively. Thus, the EMIM-NiPS₃ intercalate preserves the hexagonal lattice of Ni atoms bonded with six S atoms to form edge-sharing NiS₆ octahedra. The S atoms are connected to two P atoms located above and below the Ni plane, forming a $(\text{P}_2\text{S}_6)^{4-}$ anion with a pyramidal structure. While the monoclinic symmetry of the parent NiPS₃ structure exhibits the ABC stacking of the atomic layers along the *c*-axis, the intercalate structure reveals a significant degree of stacking disorder. The guest contribution to the structure factors, estimated by back-Fourier transformation of the electron density found in the solvent-accessible region (in the interlayer space) of a phase-optimized difference electron-density map,⁶⁶ is about 26 e⁻/unit cell, or approximately 1/3 of the EMIM cation per unit cell. It is worth noting that for this estimation, the EMIM cation was assumed to be the only guest molecule in the interlayer space. This finding supports the results of ICP-OES, revealing the $\text{EMIM}_{0.41}\text{Li}_{0.59}\text{NiPS}_3$ composition after cation intercalation in the cell with the Li anode and EMIM-TFSI electrolyte.

Cross sections of a larger $(\text{EMIM,Li})_x\text{NiPS}_3$ crystal and pristine NiPS₃ were prepared and analyzed by TEM on a nanometer scale. The TEM images highlight significant morphological changes after electrochemical intercalation with regions of crystalline NiPS₃ and amorphous regions (see Fig. 11c). The nanoscale inhomogeneity is seen at the nm-level as increased distances between atomic layers, especially in the vicinity of amorphous regions and numerous stacking defects (Fig. 11d). The characteristic fast Fourier transform of the HRTEM image in Fig. 11d corresponds to ED of the $\{010\}$ crystal plane in the NiPS₃ crystal structure. The calculated *d*-spacings varies between 6.31 Å, corresponding to *d*(001) of pristine NiPS₃ (ICSD entry 259148), 6.35 Å and 11.2 Å. Very minor changes in the *d*-spacings from 6.31 Å to 6.35 Å may indicate that the crystalline regions are not pristine NiPS₃, but that they additionally host some Li as a co-intercalate, in line with the literature.⁴⁰ Regions with a large distance of 11.2 Å between layers, caused by intercalation of EMIM molecules, border on some amorphous regions. This can reflect a weaker stability of $(\text{EMIM})_x\text{NiPS}_3$ domains upon electron beam irradiation, or a partial crystallinity loss associated with EMIM insertion.

4. Discussion

Intercalation of organic species into layered inorganic structures represents an interesting approach for creating ordered organic-inorganic nanocomposites with novel properties, which are distinct from the parent compounds. Upon cation insertion into layered MPS₃ with transition metal cations M, three possibilities for electronic transfer are usually discussed in the literature: (i) to a discrete atomic level, for example a 3d level of the transition metal, (ii) to a molecular level of a discrete polyatomic $(\text{P}_2\text{S}_6)^{4-}$ unit within the host structure, or (iii) to a part of a host conduction band.⁶⁷ The last case is common for layered transition metal chalcogenides such as FeS₂.⁶⁷

As we realized for NiPS₃, electrochemical insertion of Li and Na occurs *via* different charge compensation mechanisms, resulting most probably from the difference in ionic radii of alkali cations and different thermodynamic stabilities of the products. Upon insertion of Li cations, the electron density on the $(\text{P}_2\text{S}_6)^{4-}$ unit increases. It is somewhat surprising, since NiPS₃ is a charge-transfer insulator, and the first unoccupied density of states at ~ 1.3 eV above Fermi energy represents a narrow Ni 3d band hybridized with S 3p, with a dominant Ni character.⁵² Probably, intercalated Li cations changed the crystal structure and also exerted an electrostatic potential. In contrast, insertion of larger Na cations increases the electron density on Ni. Intercalation of much bigger EMIM cations leads, however, to very little changes in electronic states of the elements. The redox nature of EMIM intercalation is confirmed by the electrochemical process. However, direct contact of NiPS₃ and EMIM-TFSI in a beaker within an Ar-filled glove box for 24 h does not result in $(\text{EMIM})_x\text{NiPS}_3$ formation. Currently, two different reaction mechanisms of intercalation of big organic molecules into NiPS₃ are discussed in the literature. First, an increase in the electron density on Ni, P and S was observed after intercalation of cobaltocene,⁶⁸ tetrabutylammonium⁶⁹ or tetraheptylammonium cations.⁷⁰ The increase in the electron density on S, being in the formal oxidation state -2 in the pristine compound, is reasonable, since a partial charge transfer from S to Ni with a d⁹L dominant character was calculated for NiPS₃.⁵² The second reaction mechanism is displacive in nature, including creation of metal cation vacancies and intercalation of an appropriate number of organic cations for charge balancing without any change in the oxidation state of elements.¹⁸ A replacement mechanism in the case of EMIM insertion can be omitted, according to the chemical analysis results, which show the same Ni:P:S ratio in pristine and intercalated compounds. Reduction of Ni, P and S in NiPS₃ can be excluded as well, as confirmed by us with numerous spectroscopic studies. Therefore, reduction of EMIM cations upon electrochemical intercalation into NiPS₃ must be the only possible redox mechanism. As is known, imidazolium cations with a hydrogen atom in the 2-position can be considered as a protonated N-heterocyclic carbene, which can easily be deprotonated. A de-protonating agent can be the anion of the ionic liquid (for example, the acetate anion $[\text{OAc}]^-$),⁷¹ chemical reduction (with alkali metal hydrides NaH or KH),⁷² or



electrochemical reduction.⁷³ The stability of such imidazolium-derived carbenes depends on steric and electronic effects,⁷⁴ including electronic stabilization *via* a π -donation into the carbene out-of-plane p-orbital by the electron-rich π -system (N=C=C-N) and the σ -electronegativity effect of nitrogen atoms.

Some formally uncharged carbenes can be isolated as products of a redox reaction, such as, for example, 1,3,4,5-tetramethylimidazol-2-ylidene from 1,3,4,5-tetramethylimidazolium chloride, with a cationic composition resembling 1-ethyl-3-methylimidazolium (EMIM).⁷⁵ Electrochemical decomposition of 1-butyl-3-methylimidazolium tetrafluoroborate ionic liquid led to the formation of 1-butyl-3-methylimidazolium radicals that react with each other in a radical-radical coupling reaction, and in a disproportionation reaction.⁷⁶ Formation of C=C double bond dimers through dimerization of carbenes, and C-C single bond dimers through the reaction of single-electron radicals was considered in their work as possible products of the cathodic process of EMIM-TFSI on Ag electrodes.⁶¹ Similarities in the Raman spectra of (EMIM)_xNiPS₃ in our work, and the reaction product spectra in the work of Lu *et al.*⁶¹ point to the possibility of the EMIM cations here undergoing a similar reduction mechanism as presented there. Their reported onset potential for electrochemical EMIM reduction is also very close to the reaction plateau in the cell with a Li-anode during (EMIM,Li) co-intercalation that we present here. Chemical analysis of (EMIM)₁NiPS₃ yielded a smaller amount of inserted EMIM than that from galvanostatic measurements. Electrochemical reduction processes in EMIM can be the reason for this difference.

It is fascinating that EMIM reduction occurs between NiPS₃ layers and does not take place at the same potential on the surface of a Pt-electrode in symmetric Pt-Pt cells. A co-intercalation of Li and EMIM cations shows a reduction of the (P₂S₆)⁴⁻ unit, similar to pure Li-intercalation.

SQUID measurements of (EMIM)_xNiPS₃ and (EMIM,Li)_x-NiPS₃ materials show an increase in the absolute magnetization values with an increased amount of intercalated species. In both cases, the antiferromagnetic ordering vanished, reflecting the importance of a Ni-Ni interlayer interaction and its destruction upon insertion.

5. Conclusion

We investigated electrochemical cationic intercalation of Li, Na and EMIM into the 2D NiPS₃ structure. Three different mechanisms could be recognized based on comprehensive *operando* XRD, XAS (Ni K-edge) and *ex situ* XAS (P K-edge and S K-edge) as well as elemental analysis studies. Intercalation of small Li cations up to $x(\text{Li}) = 0.5$ in Li_xNiPS₃ occurs into free octahedral spaces between layers without any structural changes, according to the synchrotron XRD measurements. *Operando* XAS investigations did not show any changes in the Ni K-edge spectra, implying inertness of Ni upon Li insertion. In contrast, noticeable changes in the electronic structure of S and P were detected, revealing the role of the (P₂S₆)⁴⁻ unit in the redox process. Intercalation of more than 0.5 Li per NiPS₃ formula unit results in the formation of Li₄P₂S₆ as an

intermediate, and Li₂S as the final product of lithiation. Intercalation of Na into NiPS₃ is accompanied by a Ni redox process visible in changes in the Ni K-edge spectra. Insertion of the much larger organic EMIM cations leads to a significant expansion in interlayer distance but, surprisingly, does not result in any changes in electronic states of Ni, S, and P. The underlying process here is associated with the reduction of EMIM cations through their de-protonation, and a further possible dimerization. It is understood that the EMIM cations first enter between layers of the 2D material followed by the redox reaction of the organic cations. Li- and EMIM-insertion leads to vanishing of the initial antiferromagnetic ordering of NiPS₃ and to strengthening of its paramagnetic behavior.

A detailed understanding of the structural changes and element-specific redox and electronic modifications occurring during intercalation of alkali and larger organic cation species shall be of significance in the development of hybrid systems based on 2D materials towards desired applications in optoelectronics and catalysis.

Conflicts of interest

There are no conflicts to declare.

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