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Direct synthesis and characterization of mixedvalent $Li_{0.5-\delta}CoPO_4$, a Li-deficient derivative of the *Cmcm* polymorph of LiCoPO₄⁺

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While the majority of research activities on LiCoPO₄ is focussed on the thermodynamically stable olivinetype Pnma polymorph, the metastable Pna21 and Cmcm modifications have recently attracted considerable attention due to their interesting material properties. In this study, we present the first Li-deficient structural derivative of the Cmcm modification with the nominal composition $Li_{0.5-\delta}CoPO_4$. As opposed to the substoichiometric olivine (Pnma) phases $Li_x CoPO_4$ (x = 0; 2/3), which are exclusively accessible by electrochemical or chemical Li extraction techniques, this is also the first time that a direct soft-chemical synthesis route towards a Li_xCoPO₄-type material is accomplished. X-ray and neutron diffraction studies indicate that Cmcm-type Li_{0.5- δ}CoPO₄ shows vacancies on both the Li and Co sites, whereas X-ray absorption spectra demonstrate that the structure features heterovalent Co ions (+2/+3) to compensate for the Li deficit. Magnetic measurements reveal a long-range antiferromagnetic order below 10.5 K. A thorough investigation of the thermal stability using thermogravimetric analysis, differential scanning calorimetry, and temperature-dependent in situ X-ray powder diffraction demonstrates that $Li_{0.5-\delta}CoPO_4$ is metastable and exhibits a complex, multi-step thermal decomposition mechanism. In the first step at 394 °C, it decomposes to α -Co₂P₂O₇ (P2₁/c) and LiCoPO₄ (Cmcm) upon O₂ release. The LiCoPO₄ (Cmcm) intermediate is then irreversibly transformed to olivine-type LiCoPO₄ (Pnma) at 686 °C. The material properties of $Li_{0.5-\delta}CoPO_4$ are further compared to the fully lithiated, isostructural LiCoPO₄ (Cmcm) phase, for which an improved structure solution as well as Co L2.3-edge X-ray absorption spectra are reported for the first time.

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Introduction

In the last two decades, thermodynamically stable, olivine-type (space group: Pnma) LiCoPO₄ polymorphs have been extensively studied as a high-voltage cathode material for lithium-ion batteries (operating voltage: \sim 4.8 V vs. Li/Li⁺; theoretical capacity: 167 mA h g^{-1}).¹⁻⁴ The three-dimensional network structure features $[CoO_6]$ octahedra, $[PO_4]$ tetrahedra, and Li⁺ ions in octahedral voids.5 The majority of research activities have been focused on optimizing the electrochemical performance of the material.^{4,6,7} However, despite intensive efforts, the nature of the intermediate phase Li_xCoPO₄, which occurs upon the two-step Li insertion-extraction reaction of LiCoPO₄,⁸ is still under investigation and debated in the literature. Earlier

details from PXRD data, (3) Rietveld refinement details from neutron powder diffraction data, (4) additional illustrations of the crystal structure, (5) electrochemical characterization, (6) SEM and EDS analysis, (7) full IR spectra, (8) additional magnetic data, (9) Rietveld refinement details of the PXRD pattern of the sample obtained from the TGA/DSC measurement of Li_{0.5-6}CoPO₄ under air, (10) thermal stability under Ar (TGA/DSC, Rietveld refinement), (11) Rietveld fits and crystallographic data of the in situ PXRD patterns (30-700 °C; air), (12) additional in situ PXRD patterns (800 °C, 900 °C, and 25 °C; air). The cif files containing the crystallographic data of Li_{0.5-0}CoPO₄ and Li_{1-y}CoPO₄ can be obtained from FIZ Karlsruhe, 76344 Eggenstein-Leopoldshafen, Germany (fax: +49 7247 808 666; e-mail: crysdata@fiz-karlsruhe.de), on quoting the CSD deposition numbers 432850 and 432851. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7ra04043a

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[†] Electronic supplementary information (ESI) available: (1) Comparison of the PXRD patterns of $Li_{0.5-\delta}CoPO_4$ and $Li_{1-\gamma}CoPO_4$, (2) Rietveld refinement

investigations suggested compositions of Li_{0.7}CoPO₄ (ref. 8) and $Li_{0.6}CoPO_4$,⁹ respectively, whereas a later report¹⁰ stated a Li_x - $CoPO_4$ (x = 0.20-0.45) composition. Recently, the lithiation state of the intermediate was determined to be Li_{2/3}CoPO₄ by two independent studies.^{11,12} Since the completely delithiated phase CoPO₄ is unstable and undergoes amorphization when exposed to air or moisture,^{8,9} the application of inert gas atmospheres and/or in situ techniques is crucial. According to Bramnik et al.,13 both lithium-poor, Co³⁺-containing phases are intrinsically instable and exhibit a low thermal stability. Charged LiCoPO₄ electrodes were shown to decompose rapidly at temperatures below 200 °C, leading to gas evolution and the crystallization of $LiCoPO_4$ (only in the case of Li_xCoPO_4) and $Co_2P_2O_7$. In contrast, Theil et al.¹⁴ claimed that Li_xCoPO_4 is thermally stable up to 550 °C and that the thermal instability of charged LiCoPO₄ electrodes can be solely ascribed to the instability of the CoPO₄ phase. To the best of our knowledge, these Li-deficient phases are exclusively accessible by electrochemical or chemical Li extraction from LiCoPO₄ (Pnma),^{9,13} and a direct synthesis route has not been reported to date. Interestingly, in contrast to previous reports,15,16 we recently demonstrated17 that also the fully lithiated olivine-type LiCoPO4 does not exhibit unlimited thermal stability since it transforms to the (at room temperature) metastable Pna21-LiCoPO4 phase around 900 °C.

The less common, metastable LiCoPO₄ modifications, which crystallize in the space groups Pna2₁ (ref. 16, 18 and 19) and Cmcm, (ref. 16, 20 and 21) have recently attracted attention because of their interesting material properties and potential applicability as cathode materials for Li-ion batteries. The Pna21 modification exhibits a network of [PO₄] and [CoO₄] tetrahedra and Li⁺ ions on tetrahedral sites.¹⁸ To date, the polymorph has only been accessible by microwave-assisted synthesis techniques.^{16,18,19} Pna2₁-type LiCoPO₄ shows the highest redox potential of \sim 5.0 V vs. Li/Li⁺ compared to the other two LiCoPO₄ polymorphs.^{16,18} A single redox peak was observed upon cycling, indicating that the compound is delithiated in one step. However, the electrochemical performance was found to be poor (maximum capacity: 33 mA h g⁻¹).^{16,18,19} Magnetic measurements indicated a paramagnetic Curie-Weiss-like behavior at high temperatures, and a long-range antiferromagnetic order below $T_{\rm N} = 11$ K.¹⁸ Recently, a structure redetermination suggested that the material is non-stoichiometric and shows Li-Co anti-site defects, which provide an explanation for this poor performance.19 A thorough investigation of the thermal stability revealed that LiCoPO₄ (Pna2₁) converts to the olivine LiCoPO₄ (Pnma) modification at 527 °C.19 Interestingly, the Pna21 structure re-emerges as a stable high-temperature phase above 800 °C.19

The LiCoPO₄ (*Cmcm*) polymorph was first reported by Amador *et al.*²⁰ using a high-pressure, high-temperature synthesis route (6 GPa, 900 °C). Alternative pathways using lowtemperature procedures such as microwave-assisted solvothermal¹⁶ and polyol²¹ synthesis have been demonstrated recently. The structure is built from $[CoO_6]$ and $[PO_4]$ units, with Li⁺ ions occupying tetrahedral sites. LiCoPO₄ (*Cmcm*) shows a single redox peak at ~4.3 V vs. Li/Li⁺ upon cycling,¹⁶ which corresponds to the lowest redox potential of all the LiCoPO₄ polymorphs. A discharge capacity of only 6 mA h g⁻¹ has been reported, which was associated with the poor conductivity of the material.¹⁶ The magnetic characterization suggested a longrange antiferromagnetic order below $T_{\rm N} = 11$ K at low fields (10 kOe) and the presence of a metamagnetic transition.²¹ Investigations on the thermal stability showed that the structure is metastable and transforms to olivine-type LiCoPO₄ (*Pnma*) at 575 °C, which then transforms to the *Pna2*₁ modification at 675 °C. The thermodynamically stable *Pnma*-LiCoPO₄ phase was obtained after cooling.²¹

Based on our previous work on the three LiCoPO₄ polymorphs,^{17,19,21} we herein present the first Li-deficient structural derivative of the Cmcm modification with the nominal composition $\text{Li}_{0.5-\delta}$ CoPO₄. To the best of our knowledge, this is the first time that a sub-stoichiometric Li_xCoPO₄ phase has been synthesized directly (bottom-up) by a soft-chemical polyol approach as opposed to electrochemical or electrochemical Li extraction (top-down) techniques described in the literature.9,13 The structure, morphology, oxidation state as well as electrochemical and magnetic properties of the novel Cmcm-type phase $Li_{0.5-\delta}CoPO_4$ are investigated. Moreover, the thermal properties are studied using thermogravimetry, differential scanning calorimetry, and temperature-dependent in situ X-ray powder diffraction. The results are discussed in context of the 'fully lithiated' LiCoPO4 (Cmcm) phase, for which an improved structure solution (revealing a sub-stoichiometry reflected by the revised empirical formula $Li_{1-\gamma}CoPO_4$) as well as X-ray absorption spectra are presented for the first time.

Experimental

Synthesis of $Li_{0.5-\delta}CoPO_4$ (*Cmcm*) and $Li_{1-\gamma}CoPO_4$ (*Cmcm*)

Cmcm-type LiCoPO₄ samples with varied Li contents were obtained from a polyol process as described in our previous work²¹ using LiCH₃COO (ChemPur, 99+%), (Co(CH₃COO)₂·4H₂O, Merck, 99.99%), and H₃PO₄ (Merck, 85 wt%) as precursors, and tetraethylene glycol (TTEG, Merck, $\geq 99.0\%$) as solvent. For the synthesis of $\text{Li}_{1-\gamma}\text{CoPO}_4$ (*Cmcm*), a Li : Co : P molar ratio of 3 : 1 : 10 was used, whereas $Li_{0.5-\delta}CoPO_4$ (*Cmcm*) was obtained from a modified process using a ratio of 1 : 1 : 10. First, H₃PO₄ was added dropwise to a solution of cobalt acetate in 125 mL TTEG. Then, a second solution containing lithium acetate in 75 mL TTEG was added. The resulting mixture was refluxed at 185 °C for 14 h in a round-bottom flask. After cooling, the precipitate was recovered by centrifugation (1500 rpm, 20 min, three times) and washed with ethanol (VWR AnalaR NORMA-PUR, 99.95%). The light pink powder (cf. graphical abstract) was collected by filtration, washed with acetone (99%), and dried in air at 100 °C for 14 h. Note that in contrast to the delithiated Pnma structures Li_xCoPO₄ and in particular CoPO₄, which are sensitive to air and moisture,8,9 the Cmcm-derivative Li0.5-6-CoPO₄ is stable under air for at least several months.

X-ray powder diffraction (PXRD) and Rietveld refinement details

Room-temperature PXRD data of the ground powders sealed in borosilicate glass capillaries (0.5 mm, Hilgenberg) were

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collected on a Stoe STADI P diffractometer (Mo $K_{\alpha 1}$ radiation, $\lambda = 0.70930$ Å; Ge(111) monochromator; Dectris MYTHEN DCS 1K silicon solid-state detector) in a 2θ range of $3-60^{\circ}$ (PSD step: 0.015° ; time per step: 30 s, three ranges, total measurement time: 12 h). The diffraction patterns were calibrated using an external silicon standard. The Jana2006 software²² was used for the structure refinement by the Rietveld method, using the recently reported structure solution of LiCoPO₄ (*Cmcm*; ICSD no. 432186)²¹ as a starting model. Details on the Rietveld refinement strategy and parameters used can be found in our previous work.²¹

Neutron powder diffraction (NPD) experiments

Neutron powder diffraction data were collected using the diffractometer D2B at the Institut Laue-Langevin (ILL, Grenoble, France), working at a calibrated wavelength of 1.5942 Å in a 2θ range of 5–160°. The data were recorded at 296 K with a collection time of 4 h per pattern. The data were analyzed by the Rietveld method with the FULLPROF program.²³ The line shape of the diffraction peaks was generated by a pseudo-Voigt function. The instrumental contribution to the peak broadening was determined using an instrument resolution function built from the refinement of a Na₂Ca₃Al₂F₁₄ standard, while the wavelength was refined using a Si standard.

Elemental analysis

Analysis of the Li, Co, and P contents was carried out by atomic absorption spectroscopy (AAS, Varian AA280FS sequential device) and photometry (Shimadzu UV-160 photometer). A Hekatech Euro EA CHNSO combustion analyzer was used to determine the C, H, N, and S amounts.

Soft X-ray absorption spectroscopy (soft XAS)

Co $L_{2,3}$ -edge soft XAS spectra were collected at beamline 8-2 of Stanford Synchrotron Radiation Lightsource (SSRL), operating the spherical grating monochromator (SGM, ruling: 1100 mm⁻¹) with 40 \times 40 μ m slits (resolution: ~0.3 eV), as described in our previous work.17 The XAS spectra presented in this report are derived from the total electron yield (TEY), measured via the drain current (probing depth: 2-5 nm). We also recorded Auger electron yield (AEY) and total fluorescence yield (FY) spectra via a cylindrical mirror analyzer and a silicon diode (AXUV100). These modes, which probe ~ 2 nm and 50-100 nm deep, respectively, ensured that the best quality TEY spectra was bulk representative and that the contribution from surface contamination was small. All spectra were normalized to the incoming flux and the energy scale was calibrated to match that of ref. 24, followed by a background subtraction to a line, and a final area normalization for comparison. In order to produce the most stable and reproducible fits, the fitting was limited to the region just around the L_3 edge (774–784 eV), in which the normalization was further restricted through another line subtraction and area normalization.

Fourier-transform infrared (FTIR) spectroscopy

IR spectra were collected on a Varian 670 FTIR spectrometer equipped with a PIKE GladiATR diamond ATR stage. The measurements were performed using 132 scans in a wave-number range of 400–4000 cm⁻¹.

Magnetic measurements

DC magnetization measurements were performed using a Quantum Design MPMS XL7 SQUID magnetometer in the temperature range from 2 K to 300 K with a magnetic field of up to 7 T. The powder samples were placed inside gelatin capsules. The temperature-dependent magnetic moments were recorded under an applied magnetic field of 1 kOe after cooling the samples under a magnetic field of 7 T (field-cooled, FC) and 0 T (zero field-cooled, ZFC). Magnetic hysteresis measurements were carried out with magnetic field strengths between -7 T and 7 T at 2 K, 11 K, and 300 K, respectively.

Thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC)

The stability of the material (specimen weight: ~10 mg) upon heating was assessed using a simultaneous Mettler Toledo TGA/ DSC 1 STAR device, which did not allow to monitor the cooling cycle due to the setup. The measurement was run in a temperature range of 30–900 °C (heating rate: 10 °C min⁻¹) in an argon stream (10 mL min⁻¹), and additionally synthetic air (10 mL min⁻¹) in order to ensure comparability of the data with the *in situ* PXRD experiment.

Temperature-dependent *in situ* X-ray powder diffraction (PXRD)

Temperature-controlled PXRD was performed *in situ* under air using a PANalytical X'Pert Pro diffractometer (Bragg–Brentano geometry; Cu K_{α} radiation; XCelerator detector) equipped with an Anton Parr HTK-1200 hot stage and a TCU 1000N temperature controller. The material was placed inside a corundum flat plate sample holder and heated up to 900 °C in steps of 100 °C (heating rate: 5 °C min⁻¹), with each temperature being held for 5 min before collecting the data. The scans were recorded in a 2θ range of 15–70° (step: 0.022°; time per step: 209.5 s; total experiment time: 145.5 h).

Results and discussion

Rietveld refinement of X-ray and neutron powder diffraction data

Fig. 1a shows the Rietveld fit of the X-ray powder diffraction pattern of the as-prepared title compound $\text{Li}_{0.5-\delta}\text{CoPO}_4$. The pattern of a reproduced sample of *Cmcm*-type LiCoPO₄ (Fig. 1b) is in good agreement with our previous work.²¹ In both cases, all the reflections can be indexed in the orthorhombic space group *Cmcm* and no additional reflections are observed, indicating that both materials are phase pure. While the pattern of $\text{Li}_{0.5-\delta}\text{CoPO}_4$ appears to be roughly similar to the one of LiCoPO₄, suggesting that the crystal structure of the Li-deficient





Fig. 1 Rietveld fits of the X-ray powder diffraction data (transmission geometry, Mo $K_{\alpha 1}$ radiation) of (a) Li_{0.5- δ}CoPO₄ (*Cmcm*; refined composition: Li_{0.39(2)}Co_{0.96(1)}PO₄, $\delta = 0.11(2)$), and (b) Li_{1- γ}CoPO₄ (*Cmcm*, refined composition: Li_{0.94(2)}Co_{0.96(1)}PO₄, $\gamma = 0.06(2)$).

compound is strongly correlated with the one of the fully lithiated material, some shifts in the peak positions can be recognized. The most significant feature of the $\text{Li}_{0.5-\delta}\text{CoPO}_4$ pattern is the narrowing of the (200) and (112) reflections at 15.0° and 15.9° 2 θ . Furthermore, a completely different peak pattern can be observed in the 2 θ region of 25.5–27.5° (for a detailed view, please refer to Fig. S1, ESI†).

To gain further insights into the structural differences causing the peak shifts, a Rietveld refinement was performed, using the previously published structure solution of *Cmcm*-type LiCoPO₄ (ICSD no. 143186)²¹ as a starting model. Since the elemental analysis indicated an approximate 50% deficit in Li for Li_{0.5- δ}CoPO₄ compared to *Cmcm*-LiCoPO₄ (*cf.* Table 2) within standard deviations, the structures were at first refined with fixed Li site occupancy factors of 50% and 100%, respectively, resulting in good reliability factors (Table S1, ESI†). Taking into account that the empirical formulas derived from elemental analysis indicated a deficit in both Li and Co for the two materials (empirical formulas: Li_{0.45(5)}Co_{0.93(3)}P_{1.00(2)}O₄ and Li_{0.93(5)}Co_{0.91(3)}P_{1.00(2)}O₄; *cf.* Table 2), we tentatively refined the occupancy factors of the Li and Co sites after having applied an

absorption correction.25 In both cases, the free refinement resulted in statistically significant values for the occupancies (39(2)% Li and 96.4(5)% Co for Li_{0.5- δ}CoPO₄; $\delta = 0.11(2)$ and 94(2)% Li and 95.5(5)% Co for LiCoPO₄, cf. Table S2, ESI[†]), indicating that both structures feature vacancies in the cationic substructures and are non-stoichiometric. In both cases, the reliability factors were significantly improved over the previous structure models with fixed occupancies (cf. Tables 1 and S1, ESI[†]). To simplify the sum formulas of both compounds while still reflecting the off-stoichiometry from the idealized formulas $Li_{0.5}COPO_4$ and $LiCoPO_4$ (within three standard deviations), the compounds are referred to as Li_{0.5-d}CoPO₄ for the Li-deficient phase, and $\text{Li}_{1-\gamma}\text{CoPO}_4$ (with $\gamma = 0.06(2)$) for *Cmcm*-type LiCoPO₄ in this work. It is worth noting that on basis of these refinements, there was no indication for the occurrence of antisite defects, which are profound for materials synthesized at low temperatures (as observed e.g. in Pna21-type LiCoPO4).19 Furthermore, in contrast to Li_{0.94(2)}Co_{0.96(1)}PO₄, the composition of the Li-deficient Cmcm derivative Li_{0.39(2)}Co_{0.96(1)}PO₄ would not be charge-balanced assuming that Co is only present in the oxidation state +2. We therefore assume that the deficit in positive electric charge caused by the lower Li⁺ content is compensated by Co³⁺ in the framework, which was confirmed by X-ray absorption spectroscopic studies discussed later.

The refined cell parameters (Table 1) indicate a significant contraction along the *a* axis and an expansion along the *c* axis for $\text{Li}_{0.5-\delta}\text{CoPO}_4$ compared to $\text{Li}_{1-\gamma}\text{CoPO}_4$, while *b* is not significantly changed, hence providing an explanation for the peak shifts observed in the PXRD patterns. Furthermore, the respective cell volumes ($V = 278.116(19) \text{ Å}^3 \text{ vs. } 276.28(2) \text{ Å}^3$, corresponding to an increase of 0.7%) reveal that the $\text{Li}_{0.5-\delta^-}$ CoPO₄ structure is less dense, which is consistent with the decrease in crystal densities. This is surprising since for the delithiated phases Li_xCoPO_4 and CoPO₄ derived from olivine-

Table 1 Crystallographic parameters of (a) Li_{0.5- δ}CoPO₄ in comparison with (b) Li_{1- γ}CoPO₄ (both *Cmcm*, *Z* = 4) as refined from X-ray powder diffraction data (*T* = 298 K)^{*a*}

| Sample | (a) Lio 5 SCoPO4 | (b) Li₁ "CoPO₄ |
|--------------------------------------|----------------------------------|------------------------------|
| Empirical formula | $Li_{0,39(2)}CO_{0,96(1)}PO_{4}$ | $Li_0 q_1(2)CO_0 q_6(1)PO_4$ |
| $M_{\rm r}$ (g mol ⁻¹) | 154.3 | 158.1 |
| Crystal system | Orthorhombic | Orthorhombic |
| Space group (no.) | <i>Cmcm</i> (63) | <i>Cmcm</i> (63) |
| Z | 4 | 4 |
| a (Å) | 5.3385(2) | 5.4432(3) |
| <i>b</i> (Å) | 8.1763(3) | 8.1695(4) |
| <i>c</i> (Å) | 6.3716(2) | 6.2128(3) |
| $V(Å^3)$ | 278.116(19) | 276.28(2) |
| F(000) | 297 | 302 |
| ρ (calcd) (g cm ⁻³) | 3.684(1) | 3.800(1) |
| R _p | 0.0257 | 0.0196 |
| <i>R</i> _{wp} | 0.0327 | 0.0255 |
| R _{exp} | 0.0267 | 0.0252 |
| R _F | 0.0143 | 0.0106 |
| R _B | 0.0254 | 0.0184 |
| χ^2 | 1.23 | 1.01 |
| Data/restraints/parameter | 3800/0/59 | 3835/0/57 |
| | | |

^{*a*} The estimated standard deviations were calculated by the Berar's procedure and are indicated in parentheses.

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type LiCoPO₄ (*Pnma*), a significant decrease in cell volume of up to ~7% (CoPO₄) was observed due to the smaller ionic radius of Co³⁺ compared to Co²⁺.^{8,9,11} The slight increase in cell volume might be explained by the fact that our Li-poor *Cmcm* material was produced from a kinetically controlled synthesis as opposed to *Pnma*-type CoPO₄, which was obtained by electrochemical Li extraction. As a result, the Li ions and voids are likely to be statistically distributed within the *Cmcm* structure. This would also be in line with investigations on olivine-type LiFePO₄, which revealed that materials synthesized at low temperature are prone to disorder, resulting in larger cell volumes than expected.²⁶ However, a thorough investigation of the Li⁺/vacancy distributions in the structures is beyond this work.

In order to further verify the Li contents and the structure model, we performed additional neutron powder diffraction (NPD) studies of both samples. Rietveld refinements of the NPD data were carried out using the structure parameters obtained from the PXRD experiment as a starting model. Then, the atomic coordinates, thermal parameters as well as occupation factors of Li and Co were refined (see Fig. S2 (ESI†) for the Rietveld fits). The refined lattice parameters (Table S4, ESI†) are in good agreement with the X-ray data. The refinement of the site occupancy factors revealed 37(4)% Li and 98(2)% Co for Li_{0.5- δ}CoPO₄ (δ = 0.13(4)), and 90(3)% Li and 95(6)% Co for Li_{1- γ}CoPO₄ (γ = 0.10(3)) (Table S5, ESI†). The compositions and structural parameters of Li_{0.37(4)}Co_{0.98(1)}PO₄ and Li_{0.90(2)}-Co_{0.95(6)}PO₄ are very close to the ones derived from the X-ray experiment and hence confirm the structure model.

Crystal structure

The structures of both Cmcm-type LiCoPO₄ materials feature rows of edge-sharing, distorted [CoO₆] octahedra which are running along the [001] direction as illustrated in Fig. 2a.^{20,27} The $[CoO_6]_{\infty}$ rows are cross-linked along [100] by alternating pairs of edge-sharing, asymmetric [PO₄] and [LiO₄] units (note that the tetrahedrally coordinated Li ions are shown in Fig. 2 instead of [LiO₄] tetrahedra for clarity). As a result, layers of the composition ($[CoO_6][LiO_4][PO_4]$)_{∞} are formed in the *ac* plane (Fig. 2b). In these layers, every $[CoO_6]$ octahedron shares two opposite O2–O2 edges with neighboring [CoO₆] units, and two apical O2 atoms with two different [PO₄] and [LiO₄] tetrahedra, which connect the $[CoO_6]_{\infty}$ strands (for details see Fig. S3, ESI†). The layers are in turn stacked in the sequence AB (indicated by black and grey boxes in Fig. 2a and c) along [010] with a displacement of a/2 via the apical O1 atoms of the [CoO₆] units so that a three-dimensional framework is formed. It has to be emphasized that despite the fact that Li channels seem to occur along [100] (Fig. 2a), the Li–Li distances are very large (d =5.3385(4) Å and 5.4432(5) Å, cf. Table S3b and d, ESI[†]), so that these sites are considered isolated. Hence, Li migration will require a very high activation energy along this path and the Li mobility is extremely low in both structures.20,27 The lack of suitable Li migration pathways is confirmed by electrochemical measurements (see Section 5 and Fig. S4, ESI[†]) that revealed a very poor performance for both Cmcm materials (discharge



Fig. 2 Polyhedral illustration of the crystal structures of *Cmcm*-type $\text{Li}_{0.5-\delta}\text{CoPO}_4$ and $\text{Li}_{1-\gamma}\text{CoPO}_4$ viewed along (a) [100], (b) [010] (showing one A layer as indicated by the black boxes in a, c), and (c) [001]. [CoO₆] octahedra are displayed in grey, [PO₄] tetrahedra in red, and tetrahedrally coordinated Li ions in blue (for clarity, no T_d representation was used for the [LiO₄] units). The structure is built from ([CoO₆][LiO₄][PO₄])_{∞} layers in the *ac* plane (b), which are stacked along *b* in the sequence AB (c). The layers consist of rows of edge-sharing [CoO₆] octahedra (a), which are connected by alternating pairs of [LiO₄] and [PO₄] units. The occupancies on the Li and Co sites are 39(2)% Li and 96.5(5)% Co for Li_{0.5- δ}CoPO₄, and 94(2)% Li and 95.5(5)% Co for Li_{1- γ}CoPO₄.

capacities of < 3 mA h g⁻¹), and is also in line with a previous report.¹⁶ Hence, due to the intrinsically low Li-ion conductivity, the Li_{0.5- δ}CoPO₄ and Li_{1- γ}CoPO₄ phases cannot be considered suitable for battery applications.

The structural differences between $Li_{1-\gamma}CoPO_4$ and its Lideficient analogue $Li_{0.5-\delta}CoPO_4$ can be derived from the cell parameters (Table 1) and the interatomic distances (Table S3b and d, ESI[†]). In general, the framework of $Li_{0.5-\delta}CoPO_4$ is contracted by ~ 0.10 Å in the *a* dimension and expanded by ~0.16 Å and ~0.01 Å along c and b, respectively, as also indicated by the distances between the Co centers in and between the layers. The average Co-O (2.122 Å in Li_{0 5-0}CoPO₄ vs. 2.123 Å in LiCoPO₄) and P-O (1.539 Å vs. 1.538 Å) distances in the [CoO₆] and [PO₄] units remain virtually unchanged, which is surprising since one would expect a decrease in the Co-O distances due to the occurrence of a definite amount of the smaller Co^{3+} ion compared to Co^{2+} in the structure. The individual bond lengths, however, reveal that both the $[CoO_6]$ and [PO₄] units show a higher degree of distortion. This is reflected by the fact that the Co-O2 and P-O2 distances (in the $([CoO_6])$ [LiO₄][PO₄])_∞ layers) are shortened, whereas the Co–O1 and P– O1 bonds (connecting the layers along b) are expanded, resulting in an increase of the distance between the A-B layers (cf. Co-P distances). On the other hand, the mean Li-O distances are increased by \sim 0.4 Å, leading to an expansion of the "channels" along [100]. This is consistent with the increase of the Li-Li distances by \sim 0.10 Å. However, it has to be noted that we cannot provide any information about the ordering of the Li⁺, Co²⁺, and Co³⁺ ions or the vacancies in the structure. Based on considerations on the charge distribution, it is likely that the vacancies in the $Li_{0.5-\delta}CoPO_4$ framework are located next to the Co³⁺ centers, which would be consistent with DFT (density functional theory) studies¹² on the Li⁺/vacancy distribution in olivine-type Li2/3CoPO4.

Elemental analysis

The results of the elemental analysis of $\text{Li}_{0.5-\delta}$ CoPO₄ (*Cmcm*) are compared to the values of *Cmcm*-type LiCoPO₄ from our recent report²¹ in Table 2. The CHNS analyses are similar and show small amounts of hydrogen and carbon in both materials, which arise from residual tetraethylene glycol (TTEG) solvent or the decomposition products of TTEG and the acetate precursors. Whereas the Co and P contents (in wt%) are comparable within three standard deviations, it is evident that the novel compound contains about half the amount of Li (2.0(2) wt%)

| Table 2 | Elemental analysis of (a) $Li_{0.5-\delta}CoPO_4$ (<i>Cmcm</i>) in comparison |
|----------|---|
| with (b) | $Li_{1-\gamma}CoPO_4$ (<i>Cmcm</i> , data reproduced from ref. 21) ^{<i>a,b</i>} |

| Sample | (a) $\operatorname{Li}_{0.5-\delta}\operatorname{CoPO}_4$ | (b) $Li_{1-\gamma}CoPO_4$ |
|---------------------------------|---|---|
| C (wt%) | 0.4(3) | 0.8(3) |
| H (wt%) | 0.5(3) | 0.4(3) |
| Li (wt%) | 2.0(2) | 4.1(2) |
| Co (wt%) | 35(1) | 34(1) |
| P (wt%) | 20.0(3) | 19.6(3) |
| $n(\mathrm{Li}): n(\mathrm{P})$ | 0.45(5) | 0.93(5) |
| n(Co): n(P) | 0.93(3) | 0.91(3) |
| Empirical formula | Li _{0.45(5)} Co _{0.93(3)} P _{1.00(2)} O ₄ | Li _{0.93(5)} Co _{0.91(3)} P _{1.00(2)} O ₄ |

^{*a*} The molar composition is calculated from the experimental values and normalized to the P content (standard deviations in parentheses). ^{*b*} The N and S contents were below the detection limit in both samples (= 0).

compared to the LiCoPO₄ (*Cmcm*) material (4.1(2) wt%). This is confirmed by the respective Li:Co:P molar ratios of 0.45(5):0.93(3):1.00(2)0.93(5):0.91(5):1.00(2). and EDS measurements further substantiate the Co : P ratios found by elemental analyses. (Images of the obtained crystal morphologies and semi-quantitative EDS analyses are provided in Fig. S5, ESI[†].) The empirical sum formulas derived from elemental analysis, Li_{0.45(5)}Co_{0.93(3)}P_{1.00(2)}O₄ and Li_{0.93(5)}-Co_{0.91(3)}P_{1.00(2)}O₄, indicate a deficit both in Li and Co for both compounds and are also consistent with the results of the Rietveld refinements (cf. Table 1) within standard deviations. Based on the results of the elemental analysis, δ in this case is 0.05(5) in $\text{Li}_{0.5-\delta}\text{CoPO}_4$ (whereas γ is 0.07(5) in $\text{Li}_{1-\gamma}\text{CoPO}_4$), which is slightly smaller than derived from X-ray and neutron diffraction. Due to the kinetically controlled synthesis procedure, however, it is possible that the composition will slightly vary from batch to batch. Summarizing all efforts from diffraction experiments and elemental analysis to determine the Li and Co contents of the present phases, the notations $Li_{0.5-\delta}CoPO_4$ and $Li_{1-\gamma}CoPO_4$ seem therefore to best reflect the off-stoichiometries found by the different techniques within three standard deviations.

As shown in our previous work²¹ on Cmcm-type LiCoPO₄, the synthesis method strongly affects the phase composition. Whereas it was observed that the composition (and also the morphology) can be slightly varied by changing the synthesis technique (solvothermal vs. polyol), the molar ratio of the Li, Co, and P precursors represents another, more effective approach towards compositional tuning. For the synthesis of $Li_{1-\gamma}CoPO_4$, the Li : Co : P molar ratio of the starting materials used was 3:1:10, whereas $Li_{0.5-\delta}CoPO_4$ was obtained from a Li : Co : P ratio of 1 : 1 : 10 (cf. experimental part). As a result, different amounts of Li are incorporated in the crystal structures of the products, although the molar amounts of the precursors and contents found in the obtained materials are not correlated linearly. Adjusting the molar ratio of the precursors might therefore provide a synthetic strategy towards other lithium cobalt or transition metal phosphate materials with modified Li contents. Given the fact that the delithiated Pnma phases Li2/3CoPO4 and CoPO4 are only accessible by chemical or electrochemical Li extraction from the fully lithiated olivinetype LiCoPO₄ (Pnma) material^{9,13} and also very instable, this bottom-up approach might also pave the way towards the direct and simple soft-chemical preparation of these Li-deficient intermediates. In that matter, the influence of the synthesis route on the chemical composition will have to be thoroughly examined in further experiments.

X-ray absorption spectroscopy

 $L_{2,3}$ -edge X-ray absorption spectroscopy is highly sensitive to the oxidation state as well as the spin state and chemical environment of transition metals.^{28,29} We have therefore applied Co $L_{2,3}$ -edge XAS to investigate the chemical state (valency and symmetry) of the cobalt ions in $\text{Li}_{0.5-\delta}\text{CoPO}_4$ (*Cmcm*) and $\text{Li}_{1-\gamma}\text{CoPO}_4$ (*Cmcm*). Fig. 3 displays the normalized TEY spectra over the L_3 -edge region from this study along with reference



Fig. 3 Normalized Co L_3 -edge XAS spectra in the TEY mode for (a) $Li_{0.5-\delta}CoPO_4$ (*Cmcm*, blue), (b) $Li_{1-\gamma}CoPO_4$ (*Cmcm*, red) along with reference spectra for (c) O_h high-spin Co^{2+} in CoO (adapted from ref. 24, black), and (d) low- and (e) high-spin Co^{3+} in EuCoO₃ and Sr₂-CoO₃Cl (from ref. 30, both black). (f) shows the difference spectrum resulting from a subtraction of 71% Co^{2+} from $Li_{0.5-\delta}CoPO_4$ (orange, $Li_{1-\gamma}CoPO_4$ subtraction), representing the trivalent Co^{3+} ion in the compound. The Co^{3+} association and the lower energy shoulder spectral weight are apparent (see text). All datasets have been aligned to match the common energy scale of ref. 24. The vertical dashed lines indicate the energies corresponding to $O_h Co^{2+}$ (776.4 eV) and $O_h Co^{3+}$ (779.4 eV).

spectra (adapted from (ref. 24 and 30)) for octahedral (O_h) highspin (HS) Co^{2+} as well as both low-spin (LS) and high-spin (HS) Co^{3+} . $\text{Li}_{1-\gamma}\text{CoPO}_4$ exhibits all characteristic features of octahedrally (O_h) coordinated Co^{2+} , including the low-energy peak at 776.4 eV that is uniquely ascribed to O_h Co^{2+} .^{24,29} Furthermore, overall sharp features can be observed, which indicate a low degree of covalency and energetic alignment with an average octahedral crystal field strength in the order of 1 eV.²⁹ In fact, a broadening of the CoO spectrum results in a near identical spectrum to $\text{Li}_{1-\gamma}\text{CoPO}_4$, indicating high purity Co^{2+} , with local disorder being the main difference.

Charge balance arguments suggest that $\text{Li}_{0.5-\delta}\text{CoPO}_4$ (*Cmcm*) bears cobalt ions in nominal oxidation states of both +2 and +3. Fitting of the spectrum using principal Co²⁺ and Co³⁺ components results in relative contributions of approximately (71 ± 3)% Co²⁺ and (29 ± 3)% Co³⁺ (as opposed to (97 ± 3)% Co²⁺ and (3 ± 3)% Co³⁺ for Li_{1-\gamma}CoPO₄), where the symmetry and spin state of the trivalent Co has some, but not dominating effects on the distribution. These values are in line with the Co²⁺ and Co³⁺ contents expected on basis of the nominal composition Li_{0.5- δ}CoPO₄ (~50% Co²⁺ and ~50% Co³⁺), but where the Co³⁺ contribution derived from XAS is lower. The discrepancy can partly be explained by a small but noticeable reduction at the surface, in line with the fact that Co^{3^+} is significantly less stable than Co^{2^+} . Moreover, the material was produced using a TTEG solvent, which also acts as a weak reducing agent^{31,32} and hence, might reduce the Co^{3^+} concentration on the particle surface (*cf.* probing depth of TEY: 2–5 nm). We note that the more bulk sensitive spectra (FY, not shown) indicated higher spectral weight towards higher energies (and thus more Co^{3^+} in the bulk), consistent with this hypothesis, but the spectra are not of high enough quality to be analyzed or discussed further.

In order to learn more about the symmetry and spin of the Co³⁺ sites, we have subtracted the fitted Co²⁺ contribution from the $Li_{0.5-\delta}COPO_4$ spectrum (Fig. 3f). We note that while the main intensity difference is centered around the energy associated with the main peak of Co³⁺ (779.4 V), there is significant intensity on the low-energy side of this peak that is not accounted for by the LS Co^{3+} . Comparison with Co^{3+} ref. 30 of different spin indicates that the trivalent Co ions are primarily high-spin, which can be rationalized based on the tetragonal distortions in the [CoO₆] octahedra (cf. Table S3, ESI[†]) and the analogous HS Co³⁺ *L*-edge spectral assignment upon axial elongations and equatorial contractions in various perovskites^{30,33,34}. The XAS thus indicates that the Li vacancies are indeed inducing distortions near the Co³⁺ sites that favor the HS Co³⁺ state. The HS Co³⁺ state is also consistent with a larger average Co-O distance (*i.e.*, larger ionic radius) than what would be expected from the (smaller) LS Co³⁺ that only occupies the t_{2g} orbitals and is associated with a stronger, more covalent Co-O interaction (cf. Table S3, ESI[†]). The presence of HS Co³⁺ is further consistent with the large magnetic moment (see later).



Fig. 4 Comparison of the FTIR spectra of (a) $Li_{0.5-\delta}CoPO_4$ (*Cmcm*, blue), and (b) $Li_{1-\gamma}CoPO_4$ (*Cmcm*, red, data reproduced from ref. 21). The omitted region of 1700–4000 cm⁻¹ (*cf.* Fig. S6, ESI†) does not show any absorption bands of water or other impurities.

Infrared spectroscopy

Fig. 4 compares the infrared spectra of $Li_{0.5-\delta}CoPO_4$ (*Cmcm*) and $Li_{1-\gamma}COPO_4$ (*Cmcm*; data reproduced from ref. 21) in the range of 400–1700 cm⁻¹. The omitted region from 1700 cm⁻¹ to 4000 cm⁻¹ is presented in Fig. S6 (ESI^{\dagger}) and does not show any absorption bands of water or other impurities. As expected, both spectra are dominated by the four fundamental vibrations of the pseudo-tetrahedral phosphate groups. Two regions can be distinguished, with the symmetric (v_1) and asymmetric (v_3) stretching vibrations of P-O being observed at higher frequencies, as well as the symmetric (v_2) and asymmetric (v_4) bending vibrations of the O-P-O group at lower frequencies. According to previous reports,^{16,21} the absorption peaks of *Cmcm*-type $Li_{1-\gamma}CoPO_4$ (Fig. 4b) can be assigned as follows: the ν_1 and ν_3 stretching modes are observed at 1070 cm⁻¹, 1016 cm⁻¹, and 939 cm⁻¹, whereas the ν_2 and ν_4 deformation vibrations appear in the region of $455-602 \text{ cm}^{-1}$. Despite the structural similarity of the compounds, the IR spectrum of $Li_{0.5-\delta}CoPO_4$ (*Cmcm*, Fig. 4a) shows some distinct differences, in particular a shift of the band frequencies to slightly lower energies and band broadening, and the appearance of additional weak bands at 1512 cm⁻¹ and 1138 cm⁻¹.

The changes observed for the Li-deficient phase $Li_{0.5-\delta}$ -CoPO₄ are in line with reports^{35,36} on olivine-type Li_x FePO₄ (0 \leq $x \leq 1$) which demonstrated that the absorption modes of the $[PO_4]^{3-}$ groups are extremely sensitive to the delithiation of LiFePO₄ and the associated oxidation of Fe^{2+} to Fe^{3+} . As discussed, the average P-O distances in $Li_{1-\gamma}CoPO_4$ (Cmcm) remain virtually unchanged when less Li is incorporated in the structure (cf. Table S3, ESI[†]). Hence, the energies of the absorption modes are not changed drastically, indicating similar local structures of the $[PO_4]^{3-}$ units. This is in good agreement with a report by Popović and co-workers,37 which suggested a linear correlation between the P-O bond lengths and stretching frequencies. The observed band splitting, on the other hand, is correlated with interactions between ions, in this case between the $[PO_4]^{3-}$ units and the adjacent Li⁺ and Co^{2+/3+} cations, *i.e.* the stronger the interaction, the larger the factor group splitting effects.³⁶ In fact, the P-Li and P-Co distances are reduced in $\text{Li}_{0.5-\delta}$ CoPO₄ compared to $\text{Li}_{1-\gamma}$ CoPO₄ (Table S3, ESI†), which leads to stronger interactions. Furthermore, it was shown^{38,39} that the factor group splitting of the ν_3 modes increases with the second ionization potential of the transition metal due to the formation of strong bonds with the oxygen atoms of the [PO₄] units, which causes a redistribution of electron density in the P-O bonds. This is reflected in the observation that the P-O1 bonds are expanded, and the P-O2 bonds shortened by ~0.02 Å each in $\text{Li}_{0.5-\delta}\text{CoPO}_4$ (cf. Table S3, ESI[†]). Due to the significantly higher ionization potential of Co^{3+} compared to Co^{2+} , the larger factor group splitting in the IR spectrum of $Li_{0.5-\delta}CoPO_4$ is therefore the result of the mixed valence state of the Co ions in the structure (Co^{2+}/Co^{3+}) as opposed to $Li_{1-\gamma}CoPO_4$ which contains Co^{2+} only. However, a thorough analysis of the spectra, including the assignment of the additional absorption band at 1512 cm⁻¹, would require a complete structural model, including the ordering of the Li⁺,

 Co^{2+} , and Co^{3+} ions in the framework as well as DFT calculations, which is beyond the scope of this work. It is likely that the occurrence of additional modes is the result of a lower local symmetry in $\text{Li}_{0.5-\delta}$ CoPO₄ due to a higher defect concentration.

Magnetic properties

Fig. 5 shows the magnetic susceptibility as function of temperature measured at a magnetic field of 0.1 T in fieldcooled (FC) condition as well as the magnetic hysteresis recorded at a temperature of 2 K of $Li_{0.5-\delta}CoPO_4$ in comparison with $Li_{1-\gamma}CoPO_4$ (both *Cmcm*). The temperature dependence of the magnetic susceptibilities indicates a long-range antiferromagnetic to paramagnetic transition at $T_{\rm N} = 10.5$ K for ${\rm Li}_{0.5-\delta^{-}}$ CoPO₄, and $T_{\rm N} = 12$ K for Li_{1- γ}CoPO₄, respectively. The low transition temperature T_N of 12 K of the Li_{1- γ}CoPO₄ (*Cmcm*) phase is in agreement with our previous study²¹ on Cmcm-type LiCoPO₄ and is comparable to the non-olivine, metastable LiCoPO₄ (*Pna*2₁) phase ($T_N = 11$ K),¹⁸ but much lower than the well investigated olivine-type LiCoPO₄ (*Pnma*, $T_{\rm N} = 21.6$ K).^{40,41} This finding demonstrates the close relation between structural and magnetic properties. However, the further reduction of the observed transition temperature T_N of the Li-deficient $\text{Li}_{0.5-\delta^-}$ CoPO₄ (Cmcm) compound might be related to the vacancies on the Co site, weakening the Co-O-Co super exchange interaction. Above the transition temperature, the magnetic susceptibility of both compounds follow the Curie-Weiss law (see Fig. S7, ESI[†]), $\chi = \chi_0 + (N_A \mu_{eff}^2)/(3k_b(T - \theta_C))$, where χ_0 is a temperature-independent contribution, μ_{eff} the effective magnetic moment, $N_{\rm A}$ the Avogadro number, $\theta_{\rm C}$ the Weiss temperature, and $k_{\rm b}$ the Boltzmann constant. Fitting the hightemperature magnetic susceptibility with the Curie-Weiss law yields a higher Weiss temperature of -21.8 K of Li_{0.5-d}CoPO₄ compared to the stoichiometric $\text{Li}_{1-\gamma}\text{CoPO}_4$ (-28.2 K; cf. inset of Fig. S7, ESI[†]), which reflects the difference in transition



Fig. 5 Magnetic susceptibility as function of temperature of $\text{Li}_{0.5-\delta}$ -CoPO₄ (*Cmcm*, blue data points) and $\text{Li}_{1-\gamma}\text{CoPO}_4$ (*Cmcm*, red) measured at a magnetic field of 0.1 T in field-cooled condition. The vertical dashed lines mark the respective transition temperatures of both phases (10.5 K and 12 K). The inset shows the corresponding magnetic hysteresis curves recorded at T = 2 K.

temperatures $T_{\rm N}$. Furthermore, similar effective magnetic moments of $\mu_{\rm eff} = (5.20 \pm 0.02) \,\mu_{\rm B}$ for ${\rm Li}_{0.5-\delta}{\rm CoPO}_4$ and $(5.08 \pm 0.02) \,\mu_{\rm B}$ for ${\rm Li}_{1-\gamma}{\rm CoPO}_4$ were found. Both values exceed the spin-only value of high-spin ${\rm Co}^{2+}$ and ${\rm Co}^{3+}$ (3.9 $\mu_{\rm B}$ and 4.8 $\mu_{\rm B}$, respectively), demonstrating a non-negligible orbital contribution.⁴² Taking into account the finite angular momentum, effective magnetic moments of 5.2 $\mu_{\rm B}$ and 5.5 $\mu_{\rm B}$ for high-spin ${\rm Co}^{2+}$ and ${\rm Co}^{3+}$ in the O_h symmetry (${}^{5}{\rm t}_{2g}$ and ${}^{4}{\rm t}_{1g}$ ground state) are expected in the weak spin–orbit coupling limit. Assuming the presence of only ${\rm Co}^{2+}$ in the ${\rm Li}_{1-\gamma}{\rm CoPO}_4$ compound, the observed values are slightly smaller than the expected ones. However, the larger effective magnetic moment $\mu_{\rm eff}$ of ${\rm Li}_{0.5-\delta^-}$ CoPO₄ clearly supports the findings of a mixed valence state of Co in the Li-deficient compound as discussed in the previous sections.

The inset of Fig. 5 reveals a more distinct difference of the magnetic properties of $Li_{1-\gamma}CoPO_4$ and the Li-deficient $Li_{0.5-\delta}CoPO_4$ compounds. LiCoPO₄ exhibits a magnetic double-hysteresis loop at 2 K, demonstrating an antiferromagnetic ground state at 0 T (for the hysteresis curves at 11 K and 300 K see Fig. S8, ESI†). Furthermore, the double-hysteresis loop indicates a spin-flip transition at a critical field of around ± 3 T, which is much lower than for Pnma-type LiCoPO4.43 Again, this can be attributed to the different structural properties. In contrast, an almost linear dependence of the magnetization as a function of the applied field is observed for Li_{0.5-ô}CoPO₄ below $T_{\rm N}$. No hysteresis with a finite remanence caused by a weak ferromagnetic phase due to the mixed-valence state of Co ions were found.⁴⁰ However, the difference of the magnetic susceptibility recorded under field-cooled (FC) and zero fieldcooled (ZFC) conditions suggests the formation of magnetic domains below $T \approx 5$ K. Below this temperature, the finite amount of Co³⁺ ions as well as the observed defects on the Co sites might cause competing magnetic interactions resulting in a complex antiferromagnetic state as, for instance, described by Jensen and co-workers.44

Thermal stability

The thermal stability of $Li_{0.5-\delta}CoPO_4$ (*Cmcm*) was investigated using TGA/DSC (Fig. 6) and temperature-dependent in situ X-ray powder diffraction, both performed under air (Fig. 7). The thermal behavior of LiCoPO₄ (Cmcm) has been discussed in detail in ref. 21. In the TGA curve (Fig. 6a), an overall mass loss of ~4.8 wt% is observed between 30 °C and 900 °C, which proceeds in several steps. The mass loss of \sim 1.3 wt% up to 360 °C is probably correlated with the decomposition of residues of the TTEG solvent and the acetate precursors (cf. CHNS analysis, Table 2). The DSC curve shows two pronounced, broad endothermic peaks at 394 °C and 686 °C, respectively, with the first signal being accompanied by a weight loss step of ~ 2.5 wt% and the second one of \sim 0.1 wt%. The corresponding X-ray powder diffraction pattern (Fig. 6b, refinement details see Tables S7-S11[†]) of the dark violet powder (cf. graphical abstract) obtained after the TGA/DSC measurement reveals that a mixture of 44.9(6) wt% olivine-type LiCoPO₄ (Pnma, ICSD no. 431999)⁴⁵ and 55.1(6) wt% of the low-temperature modification



Fig. 6 (a) DSC (black) and TGA (blue) curves of Li_{0.5- δ}CoPO₄ (*Cmcm*) measured in a temperature range of 30–900 °C (heating rate: 10 °C min⁻¹, atmosphere: synthetic air). Two endothermic DSC signals are observed at 394 °C (accompanied by a TGA weight loss step of ~2.5 wt%) and 686 °C, respectively. (b) Rietveld fit of the X-ray powder diffraction data (transmission geometry, Mo $K_{\alpha 1}$ radiation) of the dark violet post TGA/DSC-material demonstrating that mixture of olivine-type LiCoPO₄ (*Pnma*) and α -Co₂P₂O₇ (*P*2₁/*c*) was formed upon heating.

of cobalt pyrophosphate, α -Co₂P₂O₇ (*P*2₁/*c*, ICSD no. 280959)⁴⁶ was formed. Note that data obtained from a similar experiment using an argon atmosphere are in good agreement (Fig. S9 and S10, Tables S12–S16, ESI†).

In order to understand the signals observed in the TGA/DSC study and to identify possible intermediates of the heating process, we performed a temperature-controlled *in situ* PXRD experiment between room temperature (30 °C) and 900 °C with a step size of 100 °C (Fig. 7). The Rietveld fits of the individual PXRD patterns at each temperature up to 700 °C can be found in Fig. S11 (ESI†). The refined phase fractions and crystallographic details (atomic coordinates, thermal displacements parameters, bond lengths), reflecting the structural changes, are presented in Tables S17–S25.† (Note that the patterns at $T \ge 800$ °C, which are shown in Fig. S12 (ESI†), were of insufficient quality for a refinement because of the occurrence of strong reflections from the corundum sample holder.) Up to 300 °C, no change of



Fig. 7 *In situ* X-ray powder diffraction patterns (Bragg–Brentano geometry, Cu K_α radiation) of Li_{0.5-δ}CoPO₄ (*Cmcm*) measured between 30 °C and 700 °C under air (heating rate: 5 °C min⁻¹; for patterns at 800 °C, 900 °C, and after cooling see Fig. S12, ESI†). The phase undergoes several transitions upon heating. The theoretical patterns of the involved phases, which were calculated from room temperature data, are displayed in color. Between 300 °C and 400 °C, Li_{0.5-δ}CoPO₄ (*Cmcm*, this work, blue) decomposes to LiCoPO₄ (*Cmcm*, ICSD no. 432186,²¹ red) and α-Co₂P₂O₇ (*P*2₁/*c*, ICSD no. 280959,⁴⁶ green). Above 500 °C, an irreversible transition of LiCoPO₄ (*Cmcm*, blue) to olivine-type LiCoPO₄ (*Pnma*, ICSD no. 431999,⁴⁵ orange) occurs. The reflections marked with red asterisks (*****) in the pattern at 600 °C can be assigned to residues of LiCoPO₄ (*Cmcm*).

the diffraction patterns occurs, indicating that the $\text{Li}_{0.5-\delta}\text{CoPO}_4$ (*Cmcm*) phase is thermally stable up to that temperature. Between 300 °C and 400 °C, a mixture of the 'fully lithiated' $\text{Li}_{1-\gamma}\text{CoPO}_4$ (*Cmcm*) structure (referred to as LiCoPO₄ with $\gamma = 0$ in Fig. 7 since ICSD no. 432186 (ref. 21) was used for the theoretical pattern) and α -Co₂P₂O₇ (*P*2₁/*c*) is observed. Hence, the endothermic DSC signal at 395 °C can be explained by the decomposition of $Li_{0.5-\delta}CoPO_4$ (*Cmcm*) to these phases (the simultaneous mass loss will be explained later in the text). Between 500 °C and 600 °C, the *Cmcm*-type $\text{Li}_{1-\gamma}\text{CoPO}_4$ intermediate starts to convert to the thermodynamically more stable olivine *Pnma* structure, whereas the peaks of α -Co₂P₂O₇ remain unaltered. The fact that the low-temperature α -modification of $Co_2P_2O_7$ (P2₁/c) does not transform to the high-temperature β - $Co_2P_2O_7 (A2/m)^{47}$ phase, which would be expected at ~480 °C (ref. 48) for the pure material, suggests that the transformation is either kinetically hindered or very slow. Hence, the DSC signal at 688 °C corresponds to the transformation of the Cmcm to the Pnma LiCoPO₄ phase. The lower transition temperature found in the in situ PXRD study compared to the DSC data might be related to slightly different atmospheres (air vs. synthetic air) and heating rates (10 °C min⁻¹ νs . 5 °C min⁻¹) being used. The phase transformation is not completed until 700 °C because traces of $Li_{1-\gamma}COPO_4$ (*Cmcm*) can be identified in the diffraction pattern at 600 °C. Compared to the thermal stability of the pure, lithiated LiCoPO₄ (Cmcm) material (transformation to singlephase LiCoPO₄ (Pnma) at 575 °C),²¹ the phase transition temperature is significantly increased. No further phase changes are observed in the PXRD patterns up to 700 °C, yet the diffraction peaks are shifted to lower angles, indicating bigger lattice dimensions due to thermal expansion. Unfortunately, it cannot be derived from the patterns at $T \ge 800$ °C (see Fig. S12, ESI[†]) whether the *Pna2*₁-type LiCoPO₄ structure reappears as a high-temperature phase as reported for all three, fully lithiated LiCoPO₄ polymorphs (Pnma,¹⁷ Pna2₁,¹⁹ and Cmcm²¹). The pattern of the cooled sample (25 °C; Fig. S12, ESI†) is consistent with the ex situ PXRD experiment (Fig. 6b) and shows reflections of α -Co₂P₂O₇ and *Pnma*-type LiCoPO₄, indicating that both phase transitions are irreversible.

The results of the thorough investigation of the thermal stability of the Li-deficient compound Li_{0.5-d}CoPO₄ demonstrate that the phase exhibits a complex behavior upon heating which involves several phase transitions. Based on the combined approach using TGA, DSC (Fig. 6), and in situ PXRD studies (Fig. 7), a decomposition mechanism can be proposed according to Scheme 1. Note that since it is not possible to determine the composition of the decomposition products and intermediates (which are likely to be deficient in Li and Co as well), the mechanism is presented on basis of the nominal composition $\text{Li}_{0.5-\delta}\text{CoPO}_4$ with $\delta = 0$. In the first step (eqn (I)), LiCoPO₄ (*Cmcm*) and $Co_2P_2O_7$ (*P*2₁/*c*; both with Co oxidation state +2) are formed from four equivalents of the Co^{2+}/Co^{3+} mixed-valent starting material $Li_{0.5-\delta}CoPO_4$ (Cmcm). This step is based on a redox reaction, in which the two Co³⁺ equivalents from the four $\text{Li}_{0.5-\delta}\text{CoPO}_4$ ($\delta = 0$) units are reduced by O^{2-} ions (eqn (Ia)), which are released upon the pyrophosphate formation (*i.e.*, coupling of isolated $[PO_4]$ tetrahedra to $[P_2O_7]$ units via shared corners). The O²⁻ ions are in turn oxidized to form elemental oxygen (eqn (Ib)) that is released. The O_2 release from the phosphate groups corresponds to an approximate mass loss of ~2.5 wt% which in full agreement with the TGA mass loss step observed at 394 °C (cf. Fig. 6a). The estimated mass fractions of $\text{Li}_{1-\gamma}\text{CoPO}_4$ (*Cmcm*, with $\gamma = 0$) and $\text{Co}_2\text{P}_2\text{O}_7$ after the

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Scheme 1 Proposed two-step mechanism of the thermal decomposition of $\text{Li}_{0.5-\delta}\text{CoPO}_4$ (*Cmcm*, with $\delta = 0$) and theoretically expected weight fractions (in wt%) of the involved phases during the reaction as well as in the solid remainder after O₂ release. It can be inferred that the TGA weight loss (*cf.* Fig. 7) at 394 °C is due to the release of oxygen as a result of a redox process. The color code of the phases is related to Fig. 6 and 7.

oxygen release are ~52.4 wt% and ~47.6 wt%, respectively, and are very close to the refined values obtained from our Rietveld analysis (*cf.* Fig. 6, and Table S17, ESI†). The slight deviation of our refined values from the calculated ones can be explained by the fact that the nominal composition was used for this hypothesis and that the refinement of phase fractions on basis of the reflection intensities depends on the particle size and crystallinity. In the second step at 686 °C (eqn (II)), which was not accompanied by a significant mass loss in the TGA (Fig. 6a), LiCoPO₄ (*Cmcm*) is transformed to the thermodynamically more stable olivine-type LiCoPO₄ (*Pnma*) as observed in the PXRD study (Fig. 6b).

Interestingly, the decomposition leads to the crystallization of same phases (Pnma-type LiCoPO₄ and Co₂P₂O₇) under oxygen evolution as reported for the olivine-like Li-poor phase Li_r-CoPO₄.¹³ However, there are some significant mechanistic differences between $\operatorname{Li}_{0.5-\delta}\operatorname{CoPO}_4(Cmcm)$ and $\operatorname{Li}_x\operatorname{CoPO}_4(Pnma)$, which might be due to the fact that our studies are based on the pristine material whereas the studies on Li_xCoPO₄ were based on charged LiCoPO₄ electrodes. First of all, as reported by Bramnik et $al_{,1}^{,13}$ the decomposition of Li_xCoPO_4 cathodes occurs at much lower temperatures (<200 °C) suggesting that the Li-poor, Cmcm-type $\text{Li}_{0.5-\delta}\text{CoPO}_4$ phase (decomposition at 394 °C) is significantly more stable. (Note that compared to the results of Theil *et al.*,¹⁴ on the other hand, *Cmcm*-type $\text{Li}_{0.5-\delta}$ -CoPO₄ seems to be less stable than Li_xCoPO₄, which was found to be thermally stable at least up to 550 °C. This discrepancy might be explained by different particle sizes and carbon contents.). Furthermore, in contrast to our material, the Co₂P₂O₇ crystallization was found to be not proceeding simultaneously, but at higher temperature than the Li_xCoPO_4 (Pnma) decomposition, which was related to a possibly amorphous intermediate.13 In addition, the decomposition process was suggested to be promoted by carbon present in samples, which reacts with the released oxygen to form CO2 gas.13 The crucial

influence/destabilizing effect of carbon on the thermal stability was also confirmed for charged LiCoPO₄ electrodes containing CoPO₄.¹⁴ Based on this work, however, there is no indication that carbon affects the decomposition of the pristine $\text{Li}_{0.5-\delta^-}$ CoPO₄ (*Cmcm*) material since the carbon content of our material is not significant (0.4(3) wt%, *cf.* Table 2). To clarify whether this is also the case for the pristine, carbon-free Li_xCoPO₄ (x = 0, 0.7; space group *Pnma*) olivine phases (as opposed to the studies^{13,14} based on carbon-containing charged LiCoPO₄ electrodes), our direct synthetic approach might provide a pathway to get a deeper understanding of their intrinsic thermal stabilities as well.

Conclusions

In this study, a polyol synthesis pathway towards the first Lideficient structural derivative of the *Cmcm* polymorph of LiCoPO₄ with the nominal composition $\text{Li}_{0.5-\delta}\text{CoPO}_4$ and its material properties were presented. To the best of our knowledge, this is also the first time that a sub-stoichiometric Li_x-CoPO₄ phase was synthesized using a soft-chemical bottom-up approach as opposed to common chemical and electrochemical Li extraction techniques starting from LiCoPO₄-type materials.

Neutron and X-ray powder diffraction experiments as well as elemental analysis suggested that $Li_{0.5-\delta}CoPO_4$ (*Cmcm*) is nonstoichiometric and deficient both in Li and Co, which generates vacancies on both cation sub-lattices in the crystal structure. The occurrence of vacancies was also observed in the course of a structure redetermination of the 'fully lithiated' *Cmcm* phase, resulting in the revised formula $Li_{1-\gamma}CoPO_4$. Co $L_{2,3}$ -edge X-ray absorption spectroscopy indicated that, unlike Li1-7CoPO4 which exclusively contains octahedrally coordinated Co²⁺ ions, the Li-deficient structure bears both Co²⁺ and Co³⁺ ions to compensate for the Li deficit. Due to the reduced Li content and amount of electrochemically active Co²⁺ ions, the material exhibited a poor electrochemical performance. The thermal stability of $Li_{0.5-\delta}CoPO_4$ has been studied thoroughly using thermogravimetry, differential scanning calorimetry, and temperature-dependent in situ X-ray powder diffraction experiments. $Li_{0.5-\delta}CoPO_4$ (*Cmcm*) is metastable and shows a complex, two-step decomposition mechanism upon heating. At 394 °C, it decomposes to α -Co₂P₂O₇ (P2₁/c) and Li_{1- γ}CoPO₄ (Cmcm) in an endothermic reaction upon which oxygen is released as a result of a redox reaction. The $Li_{1-\gamma}CoPO_4$ (Cmcm) phase then irreversibly converts to the thermodynamically more stable LiCoPO₄ (*Pnma*) olivine structure at 686 °C.

To conclude, the present work paves the way towards the direct and simple soft-chemical preparation and investigation of Li-deficient structures derived from lithium transition-metal phosphates. Our methodology provides fundamental insights into the material properties, and hence to study Li-deficient intermediates that are probably involved in the charge/discharge steps of LiCoPO₄-type cathodes. It further helps to understand the complex structure chemistry of this class of cathode materials for Li-ion batteries. In that matter, future studies should focus on compositional tuning (*e.g.* by

modifying the amounts of the precursors in the synthesis) in order to identify other partially lithiated structural derivatives.

Author contributions

J. L. performed the material characterization using *in situ* and *ex situ* PXRD, SEM/EDS, IR spectroscopy, and electrochemical measurements, and analyzed the data (under supervision of T. N. and M. M. D.). C. A.-S. performed the synthesis. S. G. and D. N. performed and analyzed magnetic and XAS measurements, respectively. I. P. O. and C. A.-S. conducted neutron diffraction experiments. J. L. wrote the manuscript. All authors have given approval to the final version of the manuscript.

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Abbreviations

| AAS | Atomic absorption spectroscopy |
|-------|---|
| AEY | Auger electron yield |
| DFT | Density functional theory |
| DSC | Differential scanning calorimetry |
| EDS | Energy-dispersive X-ray spectroscopy |
| FC | Field-cooled |
| FTIR | Fourier-transform infrared |
| FY | Fluorescence yield |
| HS | High-spin |
| ICSD | Inorganic Crystal Structure Database |
| LS | Low-spin |
| NPD | Neutron powder diffraction |
| PXRD | Powder X-ray diffraction |
| SEM | Scanning electron microscope |
| SQUID | Superconducting quantum interference device |
| TEY | Total electron yield |
| TGA | Thermogravimetric analysis |
| TTEG | Tetraethylene glycol |
| XAS | X-ray absorption spectroscopy |
| ZFC | Zero field-cooled |
| | |

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