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Direct observation of alternating octahedral and prismatic sodium layers in O3-type transition metal oxides

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ABSTRACT

The oxygen stacking of O3-type layered sodium transition metal oxides (O3-NaTMO₂) changes dynamically upon topotactic Na extraction and reinsertion. While the phase transition from octahedral to prismatic Na coordination that occurs at intermediate desodiation by TM slab gliding is well understood, the structural evolution at high desodiation, crucial to achieve high reversible capacity, remains mostly uncharted. In this work, the phase transitions of O3-type layered NaTMO₂ at high voltage are investigated by combining experimental and computational approaches. We directly observe an OP2-type phase that consists of alternating octahedral and prismatic Na layers by in situ X-ray diffraction and high-resolution scanning transmission electron microscopy. The origin of this peculiar phase is explained by atomic interactions involving Jahn-Teller active Fe⁴⁺ and distortion tolerant Ti⁴⁺ that stabilize the local Na environment. We also rationalize the path-dependent desodiation and resodiation pathways in this material through the different kinetics of the prismatic and octahedral layers, presenting a comprehensive picture about the structural stability of the layered materials upon Na intercalation.

INTRODUCTION

Layered Na transition metal oxides with stoichiometry NaTMO₂ (TM: transition metal combination) show promising properties as high-performance cathode materials for low-cost Na-ion batteries.[1,2] In the discharged state, these materials are often isostructural with their lithium analogues: octahedrally-coordinated Na or Li ions sit between layers of edge-sharing TMO₆ octahedra with a repeated stacking unit of three (e.g., the O3 structure).[3] When charged, however, very different coordination preferences for Na⁺ and Li⁺ make their layered TM-oxides behave radically different.[4] As the preferred coordinations of Li (octahedral and tetrahedral) can both be achieved in the ABCABC anion rocksalt packing of the O3 structure, oxygen stacking changes upon delithiation of LiTMO₂ compounds are rare.[4] Rather, they tend to suffer from TM-ion migration driven by the tendency of Li to occupy tetrahedral sites.[5,6] In contrast, the octahedral and prismatic coordinations, which are preferred for Na, cannot be achieved in the same oxygen framework and require the gliding of oxygen layers to transform into each other. This gliding leads to a much more complex set of phase transformations for NaTMO₂ than that for the Li compounds.[7–10] While most of these transformations appear reversible upon Na extraction and
insertion, the hysteresis, inhomogeneity, and stresses associated with the more complex phase transitions in layered Na cathodes can lead to long term stability issues. Understanding these phases is particularly important as Na cathodes need to be charged to a relatively higher level of alkali extraction than Li compounds to make them competitive in light of their lower cell voltage.

For large-scale energy storage applications, Na-ion batteries utilizing Fe as a redox center are a promising low-cost alternative for Li-ion which mostly relies on Co and Ni. However, fully reversible use of Fe redox is difficult. Deintercalation of a large amount of Na can destabilize the layered structure, leading to irreversible changes especially in materials with large Fe content (e.g., NaFeO$_2$). The nature of the highly desodiated phase regarding how it forms and how it creates irreversibility is unclear. X-ray diffraction (XRD) has indicated the presence of stacking faults and pair-distribution function (PDF) analysis has been used to argue for TM migration. Yabuuchi et al. proposed a structural model for highly desodiated P2-type Na$_{0.67}$Fe$_{0.5}$Mn$_{0.5}$O$_2$. In their model, Na occupies a prismatic site in between two TMO$_6$ slabs with a repeating period of two (ABBA stacking in comparison to ABCABC in O3). They found that at 4.2 V the P2 stacking undergoes a phase transition into an unknown phase by comparing the simulated and observed XRD patterns. They suggested that the stacking of this unknown phase contains alternating octahedral and prismatic Na layers and named it “OP4” (ABBCABC oxygen stacking). Very recently, Somerville et al. investigated the structural evolution of P2-type layered Na$_{2/3}$Ni$_{1/6}$Mn$_{1/2}$Fe$_{1/3}$O$_2$ (33% Fe in the TM layer), and observed that in the high-voltage phase, prismatic and octahedral Na coexist.

In this work, the unusual phase observed at the very high state of charge in layered Na compounds is investigated. We provide a comprehensive perspective on the phase transitions in O3-NaTi$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$, a compound which shows high energy density and good cycle life. By combining in situ XRD, high resolution-scanning transmission electron microscopy (HR-STEM), X-ray absorption spectroscopy (XAS), and first-principles modeling, we visualize and rationalize the structural change in this O3-type transition metal oxide at high voltage. Our finding is that the strong Jahn-Teller distortion effect in Fe$^{4+}$ creates a Na-O-Fe$^{4+}$ interaction which can be better accommodated in a structure in which octahedral and prismatic Na stackings alternate, stabilizing an OP2-type (ABCAABCA stacking) structure. This result provides insight into the origin of the alternating O-P Na layer formation. We also relate our findings to other work on layered Na oxides that contain a certain concentration of Jahn-Teller ions, where the intergrowth
of different stackings is also observed.\textsuperscript{11–13,15} We propose a mechanism that may promote structural stability at high state of charge, potentially leading to better understanding of high voltage stability of Na-ion batteries.

**METHODS**

**Synthesis:** To obtain the target composition, 20\% excess Na$_2$O (80\%, Alfa Aesar) was combined with stoichiometric amounts of TiO$_2$ (99\%, Sigma Aldrich), Fe$_2$O$_3$ (99\%, Sigma Aldrich), Co$_3$O$_4$ (99\%, Alfa Aesar) and NiO (99\%, Alfa Aesar). The precursors were mixed with a SPEX 8000M Mixer/Mill in an argon-filled glovebox for 1 hour. The resulting mixture was pressed into a disc-shaped pellet, fired at 800$^\circ$C for 10 hours under flowing oxygen, and quenched to room temperature. The sample was transferred into the glovebox immediately after quenching to minimize air-exposure.

**Structural Characterization:** To analyze the structure of as-synthesized materials, Time-of-flight (TOF) powder neutron diffraction patterns were collected at room temperature on the Nanoscale Ordered Materials Diffractometer (NOMAD) in the Spallation Neutron Source at Oak Ridge National Laboratory and analyzed by Rietveld refinement using a TOPAS 4.2 software package. The pair-distribution function (PDF) data was fitted using a PDFFGUI software.\textsuperscript{18} The samples were sealed in a quartz tube in an Ar-filled glovebox.

**X-ray Absorption Spectroscopy (XAS):** XAS was carried out at the MRCAT 10-ID beamline in the Advanced Photon Source (APS), Argonne National Laboratory. K-edges of Ti, Fe, Co, and Ni spectra were recorded in transmission mode at room temperature with a SiO$_2$ monochromator. For XAS sample preparation, NaTi$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$, carbon black (Timcal, Super P) and polytetrafluoroethylene (PTFE, DuPont) were mixed and rolled into a film and sealed with a Kapton tape. The X-ray absorption near edge structure (XANES) and extended X-ray absorption fine structure (EXAFS) spectra were analyzed using Athena and Artemis software, respectively.

**Battery test:** The cathode film was made of 80 wt\% NaTi$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$, 15 wt\% carbon black, and 5 wt\% PTFE with a loading density of $\approx$3 mg cm$^{-2}$. A 2032-coin cell was assembled with the cathode film, Na metal (Sigma-Aldrich) as an anode, ethylene carbonate (EC, BASF)–diethyl carbonate (DEC, BASF) solution (1:1 by volume) with 1 M NaPF$_6$ (98\%, Sigma
Aldrich) as an electrolyte, and glass fiber (Whatmann GF/F) as a separator. The cells were tested at room temperature using a Maccor 4000 battery cycler.

**In situ X-ray Diffraction (XRD):** To probe the structural evolution of NaTi$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$ during electrochemical cycling, *in situ* XRD was carried out on a Bruker D8 Advance diffractometer in a reflection mode (6.5° < 2θ < 30°, Mo Kα). By using our custom-made *in situ* cell with a beryllium-Kapton window, the cathode was cycled at 10 mA g$^{-1}$ on a Solartron 1287 potentiostat, and XRD patterns were collected every 20 minutes, equivalent to a resolution of ~1.4% Na per formula unit.

**Transmission Electron Microscopy (TEM):** TEM samples were obtained at three different states of charge (pristine, charged at 3.3 V and 4.2 V) for NaTi$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$ and prepared by a drop casting method on a carbon grid. High-resolution scanning TEM (HR-STEM) images were acquired by aberration corrected TEM, JEM-ARM200CF in Condensed Matter Physics Department at Brookhaven National Laboratory and TEAM0.5 in National Center for Electron Microscope at Lawrence Berkeley National Laboratory, operated at 200 kV at a convergence angle of ~30 mrad using an annular dark-field imaging detector.

**Computation:** Density functional theory (DFT) calculations were performed with the projector augmented-wave approach, as implemented in a Vienna *ab initio* simulation package (VASP). The generalized gradient approximation (GGA) with the Perdew-Burke-Erzenhof (PBE) functional was used for the exchange-correlation with a Hubbard $U$ correction. We used $U$ = 4.3 eV for Fe and 6.0 eV for Ni. The plane-wave energy cutoff was 520 eV and reciprocal space was sampled with 25 k-points per Å$^{-1}$. All structures were converged to 10$^{-6}$ eV in total energy and 0.02 eV Å$^{-1}$ in forces. GGA+$U$ energies were used to construct the convex energy hull. The Strongly Constrained and Appropriately Normed (SCAN) functional (no $U$ correction) was used to calculate the structural parameters of the ground state structures. A plane-wave cutoff of 520 eV, total energy convergence of 10$^{-6}$ eV, and convergence of 0.02 eV Å$^{-1}$ on forces were used in SCAN calculations.

Supercells of O3, P3, and OP2 stacking were created starting from a two-formula unit cell of Na$_2$FeNiO$_4$. The TM ordering in the unit cells of Na$_{1-x}$Fe$_{0.5}$Ni$_{0.5}$O$_2$ was kept fixed while the Na-vacancy orderings for three compositions ($x = 0.25$, 0.50, and 0.75) were enumerated across different supercell shapes (sizes): 1×3×1 (3), 2×2×1 (4), 2×3×1 (6), 1×1×6 (6), 2×4×1 (8), 3×3×1 (9), 2×2×3 (12), 1×4×4 (16), 4×4×1 (16), 2×4×2 (16), 3×3×2 (18), 3×6×1 (18). For
each supercell shape and Na content, we selected the lowest 10 Na-vacancy orderings ranked by Ewald energy as calculated in the pymatgen library[26] and calculated their total energy in DFT. Phase stability as a function of sodiation level was determined by calculating the convex energy hull with respect to the fully desodiated O1-Fe\(_{0.5}\)Ni\(_{0.5}\)O\(_2\) structure and the fully sodiated O3-NaFe\(_{0.5}\)Ni\(_{0.5}\)O\(_2\) structure.

Regardless of the stacking in the initial structure used as input for the DFT calculation, we labeled the relaxed structures based on the local NaO\(_6\) environments using the Simple Chemical Environment Strategy in a LightStructureEnvironments pymatgen package.[26] This algorithm identifies whether a NaO\(_6\) local environment is prismatic or octahedral. If the structure contains only octahedral environments for Na, it is labeled O3; if it only contains prismatic environments, it is labeled P3; if there is an alternating mixture of octahedral and prismatic environments, the structure is labeled OP2. In total, there are 562 relaxed structures (181 P3 structures, 252 O3 structures, 129 OP2 structures).

To investigate details of the subtle structural changes imposed by Jahn-Teller active Fe\(^{4+}\) on the NaO\(_6\) environment, two different-sized supercells were created for each O3, P3, and OP2 stacking sequence: Small (O3-Na\(_{36}\)TM\(_{144}\)O\(_{288}\) with 9 Na layers, P3-Na\(_{36}\)TM\(_{144}\)O\(_{288}\) with 9 Na layers, OP2-Na\(_{50}\)TM\(_{200}\)O\(_{400}\) with 8 Na layers) and Large (O3-Na\(_{180}\)TM\(_{720}\)O\(_{1440}\) with 45 Na layers, P3-Na\(_{180}\)TM\(_{720}\)O\(_{1440}\) with 45 Na layers, OP2-Na\(_{250}\)TM\(_{1000}\)O\(_{2000}\) with 40 Na layers). In both cases, we randomly generated 200 structures, assign oxidation states to be Na\(^{+}\), Fe\(^{4+}\), TM\(^{3.67+}\), O\(^{2-}\), and rank them by Ewald energy using the pymatgen algorithms.[26] The lowest energy 100 structures are used for the simulation that will be described in the Discussion.
RESULTS

Figure 1. (a) Neutron diffraction pattern and (b) neutron pair-distribution function of as-synthesized NaTi$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$ (NaTFCN) with the right panel giving the detail for the low $r$ region. (c) The voltage-capacity profile of the NaTFCN cathode with respect to the Na metal anode for the first three cycles.
Neutron scattering was used to identify the stacking of the structure, lattice parameters, and site occupancies for the synthesized material. Figure 1a shows a neutron Bragg diffraction pattern obtained from as-prepared NaTi_{0.25}Fe_{0.25}Co_{0.25}Ni_{0.25}O_{2} (NaTFCN). The structure fits well to the \( R\bar{3}m \) symmetry with the ABCABC O3-stacking model. From Rietveld refinement results we obtain \( a = 2.9893(4) \) Å and \( c = 16.0155(0) \) Å, in good agreement with the literature. The occupancies for Na in the 3a site and for each TM in the 3b site are 1 and 0.25, respectively. Ordering between transition metals can be identified by neutron diffraction as the neutron scattering power depends on the coherent scattering length of an atom, not on its atomic number. As Ti, Fe, Co, and Ni have coherent scattering lengths of –3.44, 2.49, 9.45, and 10.3 fm, respectively, TM ordering in the 3b site, if it exists, between Ti, Fe, and either Co or Ni, can therefore be identified by Rietveld refinement. We examined the neutron diffraction pattern in the low-Q region (the Figure 1a inset), where the TM ordering peaks would appear, and found no sign of any superstructure. The refined pattern of a structural model with random transition metal distribution agrees well with the observed one. Thus, we find that in-plane TM ordering is unlikely. Tables S1 in supplementary information summarizes the refinement results.

Figure 1b shows the neutron pair distribution function (PDF) which contains information about the local structure. The PDF result for large \( r \) (3.3–30 Å) could be fitted well using the same \( R\bar{3}m \) model used for Rietveld refinement in Figure 1a (also shown in Figure S1), confirming that the bulk material has the O3-type stacking. Without changing the fitting parameters, we extended the fitting range to the small \( r \) region (1–3.3 Å) and found that the PDF data deviate from the calculated profile at ~2 Å (the green-shaded region). This deviation is mostly due to the asymmetric shape of the first peak which accounts for the TM-O bond length. The result implies that TM-O bond lengths have a significant deviation from their average value in NaTFCN. The local structural information according to the PDF analysis can be found in Table S2.

Figure 1c shows the first three galvanostatic cycles of the NaTFCN cathode at C/20 (1C = 242 mA g\(^{-1}\)) at room temperature. The first charge and discharge capacities are 204 and 165 mAh g\(^{-1}\), respectively. In subsequent cycles, the reversible capacity stabilizes at ~160 mAh g\(^{-1}\). The first cycle capacity loss can be due to SEI formation, electrolyte decomposition, and/or the oxygen evolution at 4.2 V.
Figure 2. An intensity contour plot for the in situ XRD peaks of Na$_{1-x}$TFCN in 2–4.2 V.
To better understand the Na (de)intercalation behavior of NaTFCN, we monitor the structure evolution as a function of the Na content upon charge and discharge in 2–4.2 V using *in situ* XRD. The result is shown in **Figure 2**. At $x = 0$ in $\text{Na}_{1-x}\text{Ti}_{0.25}\text{Fe}_{0.25}\text{Co}_{0.25}\text{Ni}_{0.25}\text{O}_2$ (Na$_{1-x}$TFCN), the XRD pattern shows the O3-type layered structure, characterized by the (003)$_\text{O3}$ and (104)$_\text{O3}$ peaks at 7.6° and 18.8°, respectively. Upon desodiation, the (003)$_\text{O3}$ peak shifts toward a lower angle in $0 < x < 0.25$, indicating $c$-lattice parameter expansion due to the increasing Na interlayer spacing. In the same region, the (104)$_\text{O3}$ peak shifts toward a higher angle as a result of decreasing in-plane TM-TM bond length, reflecting the oxidation of transition metals. As desodiation proceeds to $0.25 < x < 0.33$, the (003)$_\text{O3}$ peak position appears to change discontinuously, and the (104)$_\text{O3}$ peak vanishes. A new set of peaks appears near 7.3° and 20.3°, which belong to the P3-type structure indexed as (003)$_\text{P3}$ and (015)$_\text{P3}$, respectively. P3 has prismatic Na sites created by the octahedral TMO$_6$ slabs above and below, which repeat with a period of three (ABBCCA stacking). The continuous peak shifting for $0.33 < x < 0.69$ indicates that topotactic Na extraction occurs by a single-phase mechanism: the Na slab spacing gradually expands until the cell voltage reaches 4 V ($x = 0.69$).

Charging Na$_{1-x}$TFCN into the region $0.69 < x < 0.82$ requires a potential higher than 4 V and leads to a peculiar structural change. The (003)$_\text{P3}$ peak at 7.2° abruptly shifts toward higher 20, implying that the Na slab spacing collapses. New peaks also appear at 16.9° and 20.1°. *In situ* XRD could not fully reveal the structure due to the limited number of diffraction peaks, substantial peak broadening, as well as peak overlapping with the Al current collector of the cell. It should be emphasized that the structure at the end of charge can be fitted to neither the P3 nor the O3 structure model.
**Figure 3.** HR-STEM images obtained from Na$_{1-x}$TFCN in three different states of charge: pristine with $x = 0$ (a, b), charged to 3.3 V with $x = 0.35$ (c, d), and charged to 4.2 V with $x = 0.78$ (e, f). Simulated HR-STEM images using the multislice method with frozen phonon approximation are insets (white outline) in images b, d, and f. (scale bar: 2 nm in a, c, e; 1 nm in b, d, f).
To identify the high voltage structure in atomic resolution, we employed high-angle annular dark-field (HAADF) STEM, in which the contrast is sensitive to the atomic number (Z) of periodic atoms in the sample (Z⁻¹.⁷). Three samples were prepared by electrochemical desodiation: pristine NaTFCN, Na₀.₆₅TFCN (charged to 3.3 V), and Na₀.₂₂TFCN (to 4.2 V). HR-STEM images for all samples were taken consistently in the same [100] direction.

In a pristine NaTFCN particle (Figure 3a), HR-STEM displays arrays of bright spots representing TM constituents (i.e., Ti, Fe, Co, and Ni) and dimmer spots corresponding to Na, visualizing the typical layered structure. Figure 3b shows the atomic arrangement of NaTFCN at a higher magnification. The interlayer distance, measured by the distance between transition metals in distinct layers, is 5.2 Å. The O3 unit cell with the R̅3m symmetry overlaid onto the HR-STEM image matches well with the atomic arrays of Na and TM in NaTFCN. The simulated HR-STEM O3 result (inset in Figure 3b, white box) also reproduces the observed STEM image well.

Figures 3c and d show HR-STEM images with low and high magnification obtained from desodiated Na₀.₆₅TFCN. The arrays of bright spots correspond to the TM contrast. The P3 model structure with the R3m symmetry (i.e., no inversion center) is projected onto the observed HR-STEM image in Figure 3d. The atomic positions of the TM layer in the P3 unit cell and the simulated HR-STEM image of the P3 structure (framed inset in Figure 3d) match well with the bright spots in the observed structure. The TM-TM interlayer distance is 5.7 Å. It should be noted that for the P3 structure the Na contrast in HR-STEM (both observed and simulated) is much less pronounced than that for O3. This weak Na contrast can be attributed to the disordering of Na over two distinct prismatic sites available in the P3 structure. From the electron beam direction [100], the two prismatic Na positions do not coincide but overlap only slightly with one another (Figure S2), resulting in a loss of intensity by a process known as the de-channeling effect. This effect of disordering is consistent with the simulation shown in the inset in Figure 3d which also lacks intensity from the Na layer. Weak Na contrast is in fact one of the key features that differentiates P3 from O3 stacking and has been observed in other reports.

To explain the HR-STEM results obtained from Na₀.₂₂TFCN at 4.2 V charge (Figures 3e-f), we first describe O3 and P3 using the conventional nomenclature by which consecutive anion layers are labeled by their “A”, “B”, or “C” position. O3 and P3 are then described by:

i) O3: A₀-C_TM-B₀ (A_Na) C₀-B_TM-A₀ (C_Na) B₀-A_TM-C₀ (B_Na),

ii) P3: A₀-C_TM-B₀ (C_Na/A_Na) B₀-A_TM-C₀ (A_Na/B_Na) C₀-B_TM-A₀ (B_Na/C_Na),
where subscripts of O, TM, and Na indicate the position of the oxygen, transition metal, and Na layer. As Na in the P3 structure can occupy two different sites (face-sharing with the top/facesharing with the bottom of a transition metal octahedron), we have labeled both. Most crucially, the bolded letters emphasize the transition metal layers which are most visible in HR-STEM. By examining their order, the reversed stacking sequence of the transition metal layers becomes evident: O3 has $C_{TM}B_{TM}A_{TM}$ stacking while P3 has $A_{TM}B_{TM}C_{TM}$ stacking.

These reversed TM stacking sequences in the O3 and P3 structures can be seen in HR-STEM. In Figures 3b and d, we indicate TMO$_6$ octahedra (the blue rhombuses) to represent their orientations. As all the rhombuses for the O3 and P3 phases in HR-STEM are aligned in the same orientation, the TM stackings can be directly compared by looking at the ($\bar{1}02$) plane that connects Na and TM in the O3 structure (the solid line in Figure 3b) and also in the P3 structure (the dashed line in Figure 3d). The ($\bar{1}02$) orientation in the O3 structure is flipped compared with that in the P3 structure, depicting that the TM stacking sequence is reversed.

HR-STEM images with low and high magnifications from highly desodiated Na$_{0.22}$TFCN (4.2 V) are shown in Figures 3e and f. The intensity from transition metals is clearly observable and shows that the structure remains layered at 4.2 V, although there is some rippling that is not seen in the O3 and P3 HR-STEM images. Unlike in the HR-STEM images in Figures 3a-d, we observe that the bright and dark contrasts from the Na layers alternate. In addition, no single line connecting transition metals can extend more than two slabs, indicating that the stacking sequence represents neither O3 or P3 (or O1).

Rather, we infer from the HR-STEM that two different types of stacking sequences alternate in the sample at 4.2 V. In Figure 3f, we note two different interlayer distances. TM-TM interlayer distances around bright (octahedral) and dark (prismatic) Na layers are 4.7 Å and 5.1 Å, respectively. The appearance of Na contrast in the bright layer indicates the stacking is octahedral, and the solid line tracing the transition metal and Na columns corresponds to the ($\bar{1}02$) plane of the O3 structure. In the dark Na layer the stacking is displaced from $(C_{TM}B_{TM}A_{TM})$, where it would be if O3 continued, to $(C_{TM}B_{TM}'C_{TM}')$. This section is represented by the dashed line corresponding to the ($\bar{1}02$) plane of the P3 structure (where noted by P). As previously explained, there is no Na contrast in this layer. Therefore, the dark Na layers are assigned to be prismatic stacking. The simulated HR-STEM image of Na$_{0.22}$Ti$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$ with alternating
octahedral and prismatic stacking (inset, white box in Figure 3f) reproduces Figure 3f well, further corroborating these results.

As illustrated, the direction of the (102) plane can be used to characterize the relative displacement of the transition metal slabs with respect to each other. In Figure 3f, the (102) planes change direction consistent with changes in the contrast of Na layers in O and P. Indeed, this reversal of the (102) plane indicates that TM-O bonding directions, exhibited by the blue rhombuses alternate between the directions presented for pure O3 and P3 (Figure 3b and 3d). This analysis of the HR-STEM data indicates that the high voltage structure is formed by a regular gliding of TM-O slab. Note that while the (102) plane is used to describe the features of structures, other planes could have been chosen. The dark/bright contrast variations characteristic of alternating P and O environments for Na appears striped can be unambiguously observed in a lower magnification in Figure S3. On occasion two subsequent prismatic layers can also be observed (Figure S4). When two prismatic stacking sequences are adjacent, the interlayer distance is larger at around ∼5.3 Å. Because the Na0.22TFCN high voltage structure exhibits alternating octahedral and prismatic stacking we refer to the structure as OP2, as suggested by Yabuuchi et al.12

Now that the high voltage structure is identified, the evolution of the OP2 upon resodiation can be reexamined in Figure 2. Na insertion at 4.2 V causes the (002)OP2 peak at 7.6° to shift to lower angles for Na content 0.82 > x > 0.71 in Na1-xTFCN, indicating that the c-lattice parameter increases. Concurrently, the (101)OP2 and (103)OP2 peaks at 16.9° and 20.1° slightly change to the lower angles. The down-shift of the (002)OP2 peak continues for 0.71 > x > 0.5 to lower angle but stagnates around 7.4° at x = ∼0.5. As sodiation continues, the (002)OP2 peak then shifts gradually to higher angle for 0.5 < x < 0.42. Over this whole range 0.71 > x > 0.42, the (101)OP2 and (103)OP2 peaks continuously shift to 16.6° and 19.7°, respectively. It is surprising that upon discharge the Na1-xTFCN phase does not show the characteristic set of peaks for O3 or P3 in the range 0.71 > x > 0.42 as P3 mostly appears in this composition range during charge. Rather, the in situ XRD patterns confirms the continuous presence of OP2 with the (002)OP2, (101)OP2, and (103)OP2 all substantially shifting from their position in the highly charged state at x = 0.82. In the region 0.42 > x > 0.37 strong (003)O3 and (104)O3 peaks at 7.5° and 19° indicating that the O3 phase reappears via a two-phase reaction. Further sodiation leads the (003)O3 peak to shift to 7.6° and the (104)O3 peak to 18.8° at the end of discharge (x = 0.15). To summarize, Na1-xTFCN undergoes an O3-P3-
OP2 phase transition upon charge between 2.5–4.2 V, but upon discharge the OP2 structure bypasses the P3 phase and transforms directly to the O3 phase.

**Figure 4.** *Ex situ* XAS for the quaternary transition metals upon charging. XANES for (a) Ti, (b) Fe, (c) Co, and (d) Ni and EXAFS for (e) Ti, (f) Fe, (g) Co, and (h) Ni. Samples 0, 1, 2, 3, 4, 5, and 6 are prepared by electrochemical desodiation from NaTFCN, in which the capacities are 0, 35, 71, 106, 141, 176, and 209 mAh g⁻¹, respectively.
Using XAS we investigate the redox reactions in the quaternary transition metals during the first charge and discharge and their relation to the local environment changes around different transition metals. Twelve *ex situ* XAS samples were prepared electrochemically (1 pristine, 6 different states of charge, and 5 different states of discharge) at conditions shown on the voltage profile in Figure S5. XAS upon TM oxidation is shown in Figure 4. XAS upon TM reduction can be found in Figure S6 but is essentially the same (in reverse) as in charge.

Figures 4a-d show X-ray absorption near edge structure (XANES) of Ti, Fe, Co, and Ni in NaTFCN at different states of charge. For pristine NaTFCN (sample 0), the oxidation states of each transition metal were previously determined as $\text{Ti}^{4+}$, $\text{Fe}^{3+}$, $\text{Co}^{3+}$, and $\text{Ni}^{2+}$.[16,17] In Figure 4a, the Ti K-edge during charging is invariant, indicating that Ti is electrochemically inert between 2.5 and 4.2 V. However, the Ti pre-edge changes noticeably at the end of charge at 4.2 V (sample 6), suggesting that TiO$_6$ octahedra undergo substantial distortion. The Fe K-edge in Figure 4b does not shift in the range 2.5–3.25 V (samples 0-3), but Fe oxidation is observed at high voltage between 3.25 and 4.2 V, (samples 4-6). Its contribution to the overall capacity is small as the edge shifts only slightly to higher energy.[12] We also observe a change in the pre-edge of Fe at high state of charge, suggesting its environment becomes distorted. The Co K-edge is shown in Figure 4c and shifts at low voltage (samples 0-3) but not in the high voltage region (samples 4-6), indicating that Co oxidation compensates for desodiation at low state of charge. Figure 4d shows a substantial shift in the Ni K-edge, suggesting oxidation occurring over the whole voltage range (2–4.2 V). In summary, $\text{Ni}^{2+}$ and $\text{Co}^{3+}$ oxidize first at low voltage, followed by $\text{Ni}^{3+}$ and some $\text{Fe}^{3+}$ oxidation at high voltage. On discharge, the redox couples are reduced in the opposite order as on charging (Figure S6).

Figures 4e-h show the extended X-ray absorption fine structure (EXAFS) spectra of Ti, Fe, Co, and Ni for the same states of charge as used in Figures 4a-d. (Results for discharge can be found in Figure S6.) EXAFS gives element-specific information for the TM-O and TM-TM local environment. The EXAFS in Figure 4e indicate that Ti-O bond length does not change in samples 0 to 5 (2.5 – 3.9 V), further supporting that Ti is not involved in the electrochemical reaction. The Ti-TM bond length becomes shorter upon desodiation due to the oxidation of other TMs. At the end of charging (sample 6) both the Ti-O and Ti-TM bond lengths decrease abruptly while their EXAFS signals broaden and intensities decrease, indicating a substantial distortion of the O$_6$ point symmetry of the TiO$_6$ octahedron. Figure S6e shows that Ti recovers its octahedral symmetry upon
discharge. The EXAFS spectra for Fe in Figure 4f show that the Fe-O bond length is invariant at low desodiation level (samples 0-3, 2.5–3.25 V) but shortens upon higher charge, concomitant with a decrease of EXAFS intensity. At the end of charging (4.2 V), the Fe-O and Fe-TM bond lengths abruptly decrease and the related EXAFS intensity weakens. These changes are consistent with oxidation to Fe$^{4+}$. The strong Jahn-Teller activity of Fe$^{4+}$ lowers the symmetry around the ion leading to a reduction of peak intensity.$^{[38]}$ The peak and intensity changes for Fe are reversible as resodiation removes the local distortion around Fe and recovers the octahedral symmetry (Figure S6f). Consistent with the XANES data, the EXAFS spectra in Figure 4g indicate that the Co-O and Co-TM bond lengths only decrease at low voltage (samples 0-3, 2.5–3.25 V), indicating Co oxidation. Unlike the cases of Ti and Fe, the EXAFS intensities for Co-O and Co-TM remain unchanged consistent with Co$^{3+}$ and Co$^{4+}$ preferring an almost ideal octahedral environment around them.$^{[39,40]}$ The EXAFS data for Ni, shown in Figure 4h, is consistent with its oxidation to Ni$^{2+}$ and Ni$^{4+}$ in charging. For samples 0-2, charged between 2.5 and 3 V, the EXAFS intensity and Ni-O bond length decrease due to the Jahn-Teller distortion of Ni$^{3+}$. Upon further desodiation (samples 3-5, 3.25–3.9 V), this intensity increases, consistent with recovery of octahedral symmetry around Ni$^{4+}$. At the end of desodiation, the EXAFS intensity slightly decreases again, likely due to the interaction with other TM such as Ti and Fe, which is commonly observed in layered oxide cathodes.$^{[28,41]}$ In summary, the TiO$_6$ and FeO$_6$ octahedra are distorted at high state of charge. The Co-O environment is invariant throughout desodiation. The Ni-O environment undergoes distortion at intermediate desodiation, recovers its octahedral symmetry, and then distorts again at high state of charge.
Figure 5. Formation energies of O3, P3, OP2, and O1 structures as a function of the Na content in \( \text{Na}_{1-x}\text{Fe}_{0.5}\text{Ni}_{0.5}\text{O}_2 \) (convex-hull construction) and the ground state orderings at \( x = 0, 0.25, 0.5, 0.75, \) and 1.

To understand better why OP2 forms we use first-principles computations to investigate a model system containing only \( \text{Fe}^{3+/4+} \) and \( \text{Ni}^{3+/4+} \) ions as these are the active ions in the highly charged state. We calculate the energy of the different structures that can be obtained by slab-gliding from the O3 structure (O3, P3, OP2, and O1). For each of these, several hundred Na-vacancy orderings at different Na compositions are evaluated as described in the Methods. **Figure 5** shows formation energies as a function of the Na content in \( \text{Na}_{1-x}\text{Fe}_{0.5}\text{Ni}_{0.5}\text{O}_2 \) (\( x = 0, 0.25, 0.5, 0.75, \) and 1). Thermodynamically-stable structures, based on the convex-hull construction\(^{[19,21]}\) are boxed in red, indicating that; at \( x = 0 \) and 0.25 the ground state structure is O3 (circle), at \( x = 0.5 \) the structure is P3 (plus), and at \( x = 0.75 \) the structure is OP2 (cross). The ground state of the fully desodiated endpoint, at \( x = 1 \), is the O1 phase (triangle). These 0 K ground states, illustrated in Figure 5 and described in Table 1, are also maintained at 300 K (see **Figure S7**) when the free energy includes a simple ideal-solution-like approximation for the configurational entropy.
Table 1. Description of Na occupancies and lattice parameters in the lowest energy phases calculated by SCAN

<table>
<thead>
<tr>
<th>x</th>
<th>Stable Structure</th>
<th>Na Layer</th>
<th>Na Environment</th>
<th>Na Content</th>
<th>TM-TM interlayer distance (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>O3</td>
<td>All</td>
<td>Octahedral</td>
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<td>5.44</td>
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<tr>
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<tr>
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<td></td>
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<tr>
<td>0.75</td>
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<tr>
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<tr>
<td></td>
<td></td>
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<td></td>
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<tr>
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<td>O1</td>
<td>All</td>
<td>N/A</td>
<td>0</td>
<td>4.52</td>
</tr>
</tbody>
</table>

Table 1 summarizes the Na coordination environment and average TM-TM interlayer distance in O3, P3, and OP2 ground state structures on the hull from Figure 5. Note that the TM-TM interlayer distance refers to the vertical distance from one TM layer to another. The effective Na content is listed when there are multiple layers within the unit cell. The ground state structures are geometrically optimized using SCAN functional which has been shown to reproduce known experimental lattice parameters more accurately than the PBE functional.\cite{20} Recent work observed that the SCAN functional includes medium-range intermediate van der Waals forces without compromising known thermodynamic stability rules and so in this case does not systematically over-stabilize features such as larger c-lattice parameters.\cite{25}

Consistent with the experimental result, a transition occurs from a stable O3-type structure a large Na concentration to the P3 structure around \( x \approx 0.5 \). The lowest energy calculated P3 structure at \( x = 0.5 \) contains 0.375 Na in one layer and 0.625 in the next. The OP2 ground state structure at \( x = 0.75 \) in Na\(_{1-x}\)Fe\(_{0.5}\)Ni\(_{0.5}\)O\(_2\) has an even distribution of Na in the layers. The O1-type structure identified by computation is similar to what is found in some fully desodiated layered TM\(_2\)O\(_2\).\cite{42}

According to the results in Table 1, the average slab spacing in O3 increases from 5.44 Å to 5.50 Å upon desodiation to \( x = 0.25 \), further increases to 5.88 Å when the P3 phase becomes

stable at \( x = 0.5 \) in \( \text{Na}_{1-x}\text{Fe}_{0.5}\text{Ni}_{0.5}\text{O}_2 \), and decrease to 5.78 Å in the OP2 phase at \( x = 0.75 \). These slab height changes are qualitatively consistent with the experimental results shown in Figures 2 and 3. Using the calculated structures, we generated XRD patterns for the O3, P3, and OP2 ground states and compare them with \textit{ex situ} XRD patterns with the corresponding Na content, as shown in Figure S8. The simulation captures the observed lattice parameter changes during charging well.

**DISCUSSION**

In this paper we provided evidence for the appearance of the OP2 phase at the top of charge in NaTFCN and directly visualized the structure using HR-STEM. This structure is highly unusual as it periodically repeats alternating O and P stacking leading to a unit cell length along the \( c \)-axis of \( \sim 16.2 \) Å. The fact that a periodic structure of O and P blocks is stable rather than a phase separation between distinct O and P phases indicates that some physical effect provides a positive interaction between the O and P layers at this composition. The EXAFS data suggest that it is associated with strong distortions around the Ni, Ti, and Fe site. Other factors correlating with its stability can be gleaned from the chemistries for which OP2 occurs such as \( \text{Na}_{1-x}\text{Mn}_{0.5}\text{Fe}_{0.5} \) and \( \text{Na}_{1-x}\text{Mn}_{0.4}\text{Fe}_{0.2}\text{Ni}_{0.4} \) for large \( x \). But not all layered Na compounds display an OP2 structure at high state of charge. For example, in \( \text{NaMn}_{0.25}\text{Fe}_{0.25}\text{Co}_{0.25}\text{Ni}_{0.25}\text{O}_2 \) the O3-P3-O3 phase transition is observed rather than O3-P3-OP2. To rationalize these different observations, we analyze how the Na-O-TM interactions affect the OP2 phase stability at Na = 0.25. In Figure S9 and Description S9 we show arguments why these are the most likely interactions responsible for OP2’s stability.

The concentration at which OP2 forms coincides with the formation of Fe\(^{4+}\), as demonstrated by \textit{in situ} XRD (Figure 2) and \textit{ex situ} XAS (Figure 4). The \( e_g \)-level degeneracy of high-spin Fe\(^{4+}\) in FeO\(_6\) octahedra causes a strong Jahn-Teller (JT) distortion, buckling the layered structure.\[^{44}\] We demonstrate here that this JT activity is crucial for forming OP2. Note that the P3 phase, rather than OP2, is observed above 4 V in various Fe-free Ni-containing NaTMO\(_2\),\[^{13,22,45}\] suggesting that OP2 stabilization at high state of charge by Ni\(^{3+}\) is less likely. This is consistent with the fact that the Ni\(^{3+}\) JT distortion is considerably weaker than for Mn\(^{3+}\) and Fe\(^{4+}\) and the fact that much of Ni\(^{3+}\) is oxidized to the non-JT-active Ni\(^{4+}\).
Figure 6. (a) Normed histogram of Na-O bond lengths in seven OP2 structures which are within 50 meV/O₂ above the sodiation hull. Red indicates the Na-O bond is connected to long axis of the Jahn-Teller-active Fe⁴⁺_JT while grey indicates an alternate connection (e.g. short axis of the Jahn-Teller-active Fe⁴⁺ ion, or any bond with Ni³⁺ or Ni⁴⁺). (b) Schematic of an Fe⁴⁺O₆ octahedron in OP2. The Fe⁴⁺O₆ octahedron (brown outline) next to non-Jahn-Teller-activated TMO₆ octahedron (red outline) is shown to interact with prismatic and octahedral Na. The green three-atom cluster indicates the favorable Na-O-Fe⁴⁺ triplet interaction.

A Jahn-Teller (JT) ion, in essence, enables a transition metal octahedron to distort while lowering the electronic energy of the metal. For example, the Q3 JT mode by which one octahedral axis becomes longer and the other two become shorter can be oriented along any of the three octahedral axes,[⁴⁶] enabling a degree of freedom whereby any of the faces of the oxygen octahedron can be displaced at no energy cost in order to satisfy the interaction of those oxygen ions with other ions. Connectivity of a Na ion to the long axis of the JT ion Fe⁴⁺, via the
intermediate O anion, thus also affords the Na an additional degree of freedom in an otherwise highly constrained environment. This is because the size of the triangular base of the prism or octahedron is controlled by the size of the transition metal, leaving only the slab spacing adjustable for Na to optimize its six-fold coordination.

We thus consider the influence of the long axis of the oxygen octahedron around the JT Fe\(^{4+}\) ion on Na-O bonds in all OP2 structures (n = 7) which are within 50 meV/O\(_2\) above the sodiation hull. **Figure 6a** shows the normed distribution of Na-O bond lengths categorized by the connectivity of its bound O anion, which can be to either the long axis of the JT ion (indicated by Fe\(^{4+}\)\(_{JT}\), in red) or not (in grey). The histogram shows how the Fe\(^{4+}\) JT distortion accommodates shorter Na-O bond lengths compared to those formed by a non JT-ion. This effect is also observed in all other competitive structures less than 50 meV/O\(_2\) above the hull (**Figure S10**). We reason that the shorter Na-O bond made possible by Fe\(^{4+}\)\(_{JT}\) improves the effective coordination of Na. (See **Description S10** for discussion on the ideal Na-O bond length in NaO\(_6\) six-fold environments.) As a result, this optimization of Na coordination leads to a preference for Na-O-Fe\(^{4+}\)\(_{JT}\) triplets in which the long-axis of the Fe\(^{4+}\) JT bond is connected to Na. **Figure 6b** illustrates one such Na-O-Fe\(^{4+}\)\(_{JT}\) triplet in green, showing how the long axis of the JT ion extends its O anion into the Na octahedral (prismatic) slab, thereby reducing strain on the edge-sharing octahedral (prismatic) site below (above). In summary, we reason that a Jahn-Teller ion, such as Fe\(^{4+}\) is effectively a way to distort the underlying TM-octahedron at no energy cost and enables the system to lower its energy by improving the effective coordination for Na.

If the Na-O-Fe\(^{4+}\)\(_{JT}\) triplet is the favored Na-O-TM triplet, then we hypothesize that OP2 accommodates the greatest number of such triplets. **Figure 7a** illustrates this interaction in an example OP2 structure. An oxygen atom that is part of the octahedral coordination around the Na atom is connected to a Jahn-Teller-activated Fe\(^{4+}\) arm. This connectivity thus forms the Na-O-Fe\(^{4+}\)\(_{JT}\) triplet as circled in bold black lines.

To test our hypothesis, we carry out a simulation to investigate which of the three stacking sequences accommodates the highest probability of Na-O-Fe\(^{4+}\)\(_{JT}\) triplets. The description of the cell sizes used and structure generation algorithms are given in the Methods. We assume each structure contains 25% randomly distributed Fe\(^{4+}\), in which one pair of axial bonds in each FeO\(_6\) is designated as the long Jahn-Teller Fe\(^{4+}\)-O bond. Na ions are also randomly distributed. If the activated Fe\(^{4+}\)\(_{JT}\) is connected to Na, it is referenced in the simulation as a stabilizing interaction.
and assigned a unitless energy of $-1$. Note that 1) one O is always connected to three transition metals, so as long as one of these connections is Na-O-Fe$^{4+}_{JT}$ we consider the Na-O bond to be a Na-O-Fe$^{4+}_{JT}$ triplet. The maximum number of Na-O-Fe$^{4+}_{JT}$ connections per Na atom is six. 2) The overall Na content of the test structures is constrained to 0.25 but no requirement that each layer contains the same concentration is imposed. Finally, the total number of Na-O-Fe$^{4+}_{JT}$ triplets are normalized per formula unit (Na$_{0.25}$M$_{0.5}$Fe$_{0.5}$O$_2$), yielding the normalized unitless simulation energy shown in Figures 7b and 7c for small and large cells, respectively.

**Figure 7.** (a) Na-O-Fe$^{4+}_{JT}$ triplet marked by black lines in the OP2 structure (a single Na atom in a P3 layer is shown for clarity). Jahn-Teller modes across different Fe$^{4+}$ (indicated by orange lines) are not assumed to be correlated. (b) The cumulative probability of having Na-O-Fe$^{4+}_{JT}$ triplets in a small cell and (c) in a large cell.
Since the cumulative probability illustrates which structure type has the greatest number of stabilizing Na-O-Fe\(^{4+}\)JT triplets, it also indicates which stacking has the greatest number of lower-energy arrangements of the ions. We observe that regardless of cell size, the OP2 structure has the greatest probability of low energy structures precisely due to its greatest number of Na-O-Fe\(^{4+}\)JT triplets. In summary, by enumerating structures with fully randomized Na and TM occupancy, with no assumption about Na-Na interactions or correlated Fe Jahn-Teller modes, we find evidence that the OP2-type stacking indeed affords the greatest number of connections between Na and Jahn-Teller-activated Fe\(^{4+}\).

It should be pointed out that even though NaMn\(_{0.25}\)Fe\(_{0.25}\)Co\(_{0.25}\)Ni\(_{0.25}\)O\(_2\) contains Fe\(^{4+}\), it goes through an O3-P3-O3 phase transition, without an OP2 phase appearing.\(^{[43]}\) The difference between this composition and ours suggests that Ti\(^{4+}\) may also play a role in relieving the strain needed to adjust the Na coordination. This is not surprising since Ti\(^{4+}\) is a d\(^0\) element which has been shown to accommodate highly distorted octahedral sites because it lacks the valence electrons that induce preference for certain bond geometries.\(^{[47,48]}\) XANES results shown in Figure 4 indicate indeed that octahedral TiO\(_6\) is strongly distorted upon Fe oxidation. The electronic structure of Ti may therefore facilitate the displacement of oxygen ions to improve the Na coordination.

We also remark on the relatively low Fe concentration in NaTFCN, which is notably below the Fe doping threshold of 30% as proposed by Li et al. to suppress Fe migration.\(^{[38]}\) They argued that locally-rich Fe clusters lower the energy barrier for Fe migration into tetrahedral sites. Consistent with that threshold, we do not observe any sign of Fe migration in desodiated NaTFCN by STEM analysis. Therefore, we emphasize that using Fe is a sensible approach for designing NaTMO\(_2\) with reversible phase transformations and greater cyclability. Solutions that rely on an even higher Fe concentration would be highly valuable.

P3 and OP2 are stable at the intermediate (\(x = 0.5\)) and low (\(x = 0.75\)) Na content, respectively in Na\(_{1-x}\)TFCN, leading to the O3-P3-OP2 phase transition upon desodiation. However, the observation that the P3 phase does not appear upon resodiation, as shown in Figure 2, implies that the formation of OP2 triggers different phase transformation paths upon discharge from those on charge. We speculate that the difference in desodiation and resodiation pathways arises from the Na intercalation kinetics in O and P layers. Na ions diffuse faster through face-sharing prismatic sites than through edge-sharing octahedral sites.\(^{[49]}\) One piece of evidence may be found in the different behavior of the \(c\)-lattice parameter in charge and in discharge.
**Figure 7. In situ XRD for (00l) peak evolution as a function of the Na content**

**Figure 7** replots the (00l) peaks from Figure 2. The c-lattice parameter of Na$_{1-x}$TFCN at $x = -0.7$ in discharge is much smaller than at $x = -0.7$ in charge (red circles). This suggests that the discharging structure remains “collapsed” as the OP2 phase and does not form P3. By carefully inspecting *in situ* XRD upon discharge, we found that there are three different stages for the (00l) peak shift. The peak shifts to lower angle from 7.62° to 7.51° in 0.71 < $x$ < 0.82 and from 7.51° to 7.4° in 0.5 < $x$ < 0.71, but to higher angle from 7.4° to 7.45° in 0.42 < $x$ < 0.5, suggesting three different modes of c-lattice parameter variation.

While equal Na occupancy is predicted in the O and P layers at equilibrium, sodiation into the P layer may occur kinetically faster than sodiation into the O layer. As a result, the Na slab spacing of the O and P layers in the OP2 phase may change at different rates. During the early resodiation stage, for 0.82 > $x$ > 0.71, while the OP2 structure is maintained, the P slab spacing is seen to increase faster than the O slab spacing. As sodiation continues, the slab spacing of the P layer which contains more Na than that of the O layer will start to decrease due to the reduced interlayer Coulombic repulsion, whereas the slab spacing for the O layer increases due to its lower
population of Na. This may lead to the stagnated c-lattice parameter change that is observed for $0.71 > x > 0.5$. For $0.5 > x > 0.42$, the c-lattice parameter gradually decreases as more Na in the O layers can now screen the Coulombic repulsion. When $x = 0.42$ is reached in discharge, the c-lattice parameter of the discharged phase is still smaller than that of the charged P3 phase at $x = 0.42$ (blue squares). It is possible that the OP2 structure starts to develop faults as P layers turn into O layers when they have reached a high Na content. This defected structure finally turns into the O3 phase through a two-phase reaction between $0.42 > x > 0.37$. We note that in spite of the path-dependent Na (de)intercalation, the O3 phase is reasonably recovered at Na$_{0.85}$TFCN after the first discharge.[38]

In summary, the difference in desodiation and resodiation pathway is consistent with the more rapid Na intercalation in the P layers of the OP2 phase. We consider that this coordination-dependent Na mobility causes the P layers to transform to O before the O layers absorb enough Na to transform to P, kinetically suppressing the formation of the P3 phase upon discharge. While the voltage vs. capacity profile of NaTFCN upon charge and discharge is asymmetric, the O3 phase is recovered in the discharged structure and can sustain reversible Na extraction and insertion in subsequent cycles.

CONCLUSION

We directly observe and rationalize the formation of a new phase called OP2, consisting of alternating octahedral-prismatic oxygen stacking, which is formed at a high state of charge in the O3-NaTi$_{0.25}$Fe$_{0.25}$Co$_{0.25}$Ni$_{0.25}$O$_2$ cathode. The redox process is highly reversible after the first cycle and generates a specific capacity of ~160 mAh/g when charged to 4.2 V. We suggest that the OP2 phase is thermodynamically stable due to the local optimization of oxygen coordination around Na enabled by Jahn-Teller-active Fe$^{4+}$ and distortion-tolerant Ti$^{4+}$. As layered Na transition metal oxide cathodes utilizing a large amount of Fe are known to face stability challenges, it is remarkable that a specific concentration of Fe, ensuring oxidation to Fe$^{4+}$, remains redox-active while undergoing complex phase transitions. Therefore, we hope to motivate future studies in layered Na-cathodes that incorporate some fraction of Fe as a crucial mechanical stabilization ingredient, enabling stability at higher charge and achieving greater intercalation in Na-ion batteries.
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CONFLICT OF INTEREST
The authors declare no conflict of interest.

REFERENCES


