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Authors

Kan, WH
Chen, D
Papp, JK
[et al.](#)

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Unravelling Solid-State Redox Chemistry in $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ Single-Crystal Cathode Material

Wang Hay Kan,[†] Dongchang Chen,[†] Joseph K. Papp,[‡] Alpesh Khushalchand Shukla,[†] Ashfia Huq,[§] Craig M. Brown,^{||} Bryan McCloskey,^{†,‡} and Guoying Chen^{*,†,§}

[†]Energy Storage and Distributed Resources Division, Lawrence Berkeley National Laboratory, Berkeley, California 94720, United States

[‡]Department of Chemical and Biomolecular Engineering, University of California, Berkeley, California 94720, United States

[§]Neutron Scattering Science Division, Oak Ridge National Laboratory, Oak Ridge, Tennessee 37831, United States

^{||}Center for Neutron Research, National Institute of Standards and Technology 100 Bureau Drive Stop 6102, Building 235 Room A116, Gaithersburg, Maryland 20899-6102, United States

Supporting Information

ABSTRACT: Recent reports on high capacities delivered by Li-excess transition-metal oxide cathodes have triggered intense interest in utilizing reversible oxygen redox for high-energy battery applications. To control oxygen electrochemical activities, fundamental understanding of redox chemistry is essential yet has so far proven challenging. In the present study, micrometer-sized $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ single crystals were synthesized for the first time and used as a platform to understand the charge compensation mechanism during Li extraction and insertion. We explicitly demonstrate that the oxidation of O^{2-} to O^{n-} ($0 < n < 2$) and O_2 loss from the lattice dominates at 4.5 and 4.7 V, respectively. While both processes occur in the first cycle, only the redox of $\text{O}^{2-}/\text{O}^{n-}$ participates in the following cycles. The lattice anion redox process triggers irreversible changes in Mn redox, which likely causes the voltage and capacity fade observed on this oxide. Two drastically different redox activity regions, a single-phase behavior involving only $\text{Mn}^{3+/4+}$ and a two-phase behavior involving $\text{O}^{2-}/\text{O}^{n-}$ ($0 \leq n < 2$), were found in $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ ($0 < x < 1.3$). Morphological damage with particle cracking and fracturing was broadly observed when O redox is active, revealing additional challenges in utilizing O redox for high-energy cathode development.

1. INTRODUCTION

Lithium-ion batteries (LIBs) have become increasingly important for energy storage in portable electronics, electric vehicles, and stationary application in power grids. Since the discovery of early generations of cathode materials (e.g., LiCoO_2 and LiMn_2O_4) in the 1980s, only a few compounds with other compositions or crystal structures have been reported in the last three decades.^{1–3} The conventional wisdom suggests that in O3-type layered oxides, the 3d orbitals of early transition metals (TMs), are partially overlapped with the 2p orbitals of oxygen.⁴ As such, only about half of the 3d electrons are available to participate in the redox reaction, which largely limits practical capacity (for example, ~160 mAh/g for LiCoO_2) despite their much higher theoretical capacities. To develop high-energy batteries, cathodes with higher specific capacity (>200 mAh/g) and operating voltage (>4.3 V vs Li^+/Li) are needed.^{1–3}

Recently, approaches to enable high-energy cathodes by utilizing redox reactions of both TM cations and oxygen anions have triggered intense interest.^{5–7} One of the most studied examples is the lithium and manganese-rich (LMR) layered

oxides with a general formula of $\text{Li}_{1+x}\text{Mn}_{1-x-y-z}\text{Ni}_y\text{Co}_z\text{O}_2$.^{8–10} Our recent work showed that, contrary to the common notion of a nanocomposite structure, the oxide has a single monoclinic phase (C2/m) with a large number of domains corresponding to different variants.¹¹ To involve the O 2p electrons in the following electrochemical reactions, the material typically undergoes an initial activation process signaled by a unique charging voltage profile that is much different from those of the subsequent cycles. Recent studies by Luo et al. suggested the formation of O^- holes in the intermediates, as evidenced by the progressive growth of a new peak on the O K-edge X-ray absorption spectroscopy (XAS) along with the use of a number of other characterization techniques, including isotopically labeled differential electrochemical mass spectroscopy (DEMS), X-ray absorption near edge structure (XANES), and resonant inelastic X-ray scattering (RIXS).⁹ However, this remains controversial as experimental evidence is difficult to

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obtain, largely due to their high reactivity toward carbonate-based electrolyte and/or self-combine to release carbon dioxide and oxygen gases, respectively. These processes also produce significant amount of oxide-ion vacancies in the unit cell, further destabilizing the crystal structure.¹² The activation process also triggers unfavorable phase transformations, possibly involving layered $\rightarrow$ spinel $\rightarrow$ rock-salt, as the cycling proceeds, leading to continuous voltage and capacity fades in the cathode.¹² Various optimization approaches such as doping,^{13,14} surface coating/engineering, and morphology control^{15–18} have been explored as means to mitigate the degradation, but all were met with limited success.

An alternative approach to address the phase instability is to confine O redox process within a rock-salt structure. This strategy was rarely used as it is generally believed that compounds with rock-salt crystal structure lack lithium-ion conduction pathways and are therefore electrochemically inactive. In 2003, Shigemura et al. first reported that the Li/Ti/FeO₂ system can deliver a capacity of ca. 200 mAh/g at a slow rate of 10 mAh/g.^{19,20} Investigation on lithium-ion conduction pathways in rock-salt Li_{1.211}Mo_{0.467}Cr_{0.3}O₂ was then reported by Lee et al. in 2014.²¹ Using density functional theory (DFT) calculation, they revealed the percolation of active diffusion channels in disordered Li-excess materials. In 2015, Yabuuchi et al. reported a new rock-salt Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ (LNMO) cathode with an impressive discharge capacity of ca. 300 mAh/g.²² With the use of soft/hard X-ray absorption spectroscopy, the authors proposed the involvement of both Mn^{3+/4+} and O^{2–}/O[–] redox reactions during charge. However, the detailed charge compensation mechanism and role of O, particularly the contribution from oxidation of O^{2–} to O^{*n*–} (0 < *n* < 2) and O loss from the lattice (in the form of O₂ gas, *n* = 0), remain unclear. In this study, well-formed Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ single crystals were synthesized by a molten-salt method for advanced diagnostic studies. The use of single crystals allows us to perform additional single-particle-based studies for systematic comparison of the changes in crystal structure, chemical state, and microstructure as a function of lithium content in the sample. The results provide us important insights on O activities in the oxide sublattice and how to control/tune them through engineering strategies such as chemical composition and surface morphology control.

2. EXPERIMENTAL SECTION

2.1. Synthesis. In a typical crystal synthesis procedure, stoichiometric amounts of Li₂CO₃, Nb₂O₅, and Mn₂O₃ precursors (Sigma-Aldrich, >99%) were milled together with ethanol solvent at 200 rpm for 12 h in a planetary ball mill (RETSCH PM100) using a zirconia jar with zirconia balls. About 10–15% extra Li₂CO₃ was used to compensate Li loss that often occurs during high temperature heating. The dried powder was further mixed with KCl flux in a molar ratio of 2.5–5 between the flux and TM precursors (defined as R ratio hereafter). The obtained mixture was then heated at 950 °C for 12 h in an Ar atmosphere. A ramp rate of 4 °C/min was used for both heating and cooling. After the reaction, KCl was dissolved in deionized water, and the final product was obtained after filtration and thorough washing with water. Chemical delithiation of Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ was achieved by reacting the pristine powder with various amounts of 0.1 M nitronium tetrafluoroborate (NO₂BF₄) in acetonitrile solution in an argon filled glovebox (O₂ < 1 ppm and H₂O < 1 ppm). The reaction was carried out at room temperature for 1–4 days. The resulting reaction mixtures were filtered, thoroughly washed with acetonitrile, and then dried overnight in a vacuum oven.

2.2. Characterization. Chemical composition of the samples was determined by an inductively coupled plasma optical emission

spectrometer (ICP-OES 720 Series). Scanning electron microscopy (SEM) and energy dispersive X-ray spectroscopy (EDS) images were collected on a JEOL JSM-7500F field emission microscope at a 10 kV accelerating voltage. Prior to the analysis, the powder samples were sputtered with a thin top layer of Au to reduce the charging effect. Samples for transmission electron microscopy (TEM) experiments were prepared by drop casting a sonicated solution of the crystals in anhydrous ethanol onto a carbon coated TEM grid or by sectioning using focused ion beam. Selected area electron diffraction (SAED) was performed in TEM mode at 300 kV on the sectioned sample, while high angle annular dark field (HAADF) imaging was performed on dispersed particles in scanning transmission electron microscopy (STEM) mode at 120 kV, both using an FEI Titan electron microscope.

Phase purity was first analyzed by using laboratory X-ray diffraction (XRD) collected on Bruker D2 powder X-ray diffractometer (Cu K α , 40 kV, 30 mA). Synchrotron X-ray diffraction patterns were collected at ambient temperature at 11-BM at the Advanced Photon Source (APS) in Argonne National Laboratory, which operates with a monochromatic X-ray of λ = 0.414 Å. The scans were collected between 0.5° and 50° (2 θ) at a step size of 0.0001°. The samples were first packed into Kapton capillary tubes before exposing to X-ray for 1 h. Time-of-flight (TOF) neutron diffraction data were collected at the POWGEN diffractometer at the Spallation Neutron Source, Oak Ridge National Laboratory. Samples were measured inside the vanadium sample cans, and a single bank wave with center wavelengths of 1.333 Å (2 h data collection) was used. Monochromatic (1.2 Å) neutron diffraction data were obtained using the high resolution powder diffractometer at BT1 at National Institute of Standards and Technology (NIST) Center for Neutron Research. The refinement of the diffraction data was carried out using GSAS/EXPGUI package.²³

Mn K-edges hard X-ray absorption spectroscopy data were collected in transmission mode using a (220) monochromator at SSRL beamline 2-2 and 2-3. The crystals were sandwiched between two Kapton films for data collection. Higher harmonics in the X-ray beam were reduced by detuning the Si (220) monochromator by 50% at the Mn edge. Energy calibration was accomplished by using the first inflection point at 6539 eV in the spectra of Mn metal foil reference. X-ray absorption near edge structure spectra were analyzed by Sam's Interface for XAS Package or SIXPACK software, with the photoelectron energy origin *E*₀ determined by the first inflection point of the absorption edge jump. Soft X-ray absorption spectroscopy measurements were carried out at beamlines 8-2 and 10-1 at SSRL. A thin layer of the crystals was spread onto a conductive carbon tape which was attached to an aluminum sample holder inside an Ar filled glovebox (O₂ < 1 ppm, H₂O < 1 ppm). Measurements were carried out on the 31-pole wiggler beamline 10-1 at SSRL with a ring current of 350 mA, a 1000 l mm^{–1} spherical grating monochromator with 20 mm entrance and exit slits, a 0.2 eV energy resolution, and a 1 mm² beam spot. Data were collected at room temperature under ultrahigh vacuum (10^{–9} Torr) in a single load using the total electron yield and fluorescence yield mode detectors.

2.3. Electrochemistry. To prepare the composite electrodes, the active material was first ball-milled with acetylene carbon black (Denka, 20 wt %) to reduce the particle size and improve its electronic conductivity. The mixture was then mixed with acetylene carbon black and a polyvinylidene fluoride (PVdF) binder (Kynar 2801) in a 8:1:1 weight ratio in an NMP solvent. The slurry was spread onto aluminum foil and dried overnight at 120 °C under vacuum. Cathode disks with an area of 1.6 cm² and a typical electrode loading of 2–3 mg cm^{–2} were cut from the electrode sheets and assembled into 2032-type coin cells in an argon-filled glovebox. Lithium foil (Alfa-Aesar) was used as counter and reference electrodes, Celgard 2400 polypropylene membrane as separators, and 1 M LiPF₆ in 1:1 (v/v) ethylene carbonate (EC):diethylene carbonate (DEC) (Novolyte Technologies Inc.) as electrolyte. The cells were galvanostatically cycled between 1.5 and 4.8 V using a VMP3 multichannel potentiostat/galvanostat controlled by EC-Lab v10.12 software (BioLogic Science Instruments). Cyclic voltammetry tests were carried out using the same coin cell configuration with Li foil as counter and reference electrodes. The

199 cells were scanned between 1.5 and 4.8 V with a constant scan rate of
200 5 mV/min. All electrochemical tests were carried out at room
201 temperature.

202 Differential electrochemical mass spectroscopy measurements were
203 performed in a customized Swagelok-type cell consisting of a lithium
204 anode, a liquid electrolyte (1.0 M LiPF₆ in EC/DEC 1:1 v/v), and the
205 crystal composite as the working positive electrode. The cell was
206 periodically pulsed with Ar gas at regular intervals to sweep the
207 evolved gases from the headspace to the mass spectrometer for
208 analysis. The experimental setup is described in detail elsewhere.²⁴
209 Electrochemical measurements were carried out with a constant
210 current rate of 25 mA/g between 1.5 and 4.8 V with a rest period of 25
211 min right after the cells reached the upper or lower cutoff voltages. All
212 data collection was controlled by a BioLogic SP-300 potentiostat.

3. RESULTS AND DISCUSSION

213 **3.1. Synthesis and Characterization of**
214 **Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ Single Crystals.** Synthesis of polycrystal-
215 line Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ cathode material was recently carried out
216 by Yabuuchi et al.²² The process involved ball-milling
217 stoichiometric amount of Li₂CO₃, Mn₂O₃, and Nb₂O₅, pressing
218 the resulting mixture into a pellet, and then heating the pellet in
219 Ar atmosphere at 900 °C for 24 h. Synthesis of
220 Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ single crystals, however, has not been
221 reported in the literature. Here, we adopted a molten-salt
222 method and systematically varied the synthesis conditions to
223 obtain phase-pure and discrete Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ single crystals
224 with uniform morphology and size distribution for the first
225 time. The following parameters were carefully optimized in our
226 synthesis: choice of transition-metal precursor salts and flux
227 salt, the molar ratio between the flux and TM precursors (R
228 ratio), choice and amount of lithium precursor, heating
229 temperature and time, and reaction atmosphere. It was found
230 that the best reaction time and temperature were 12 h and 950
231 °C, respectively. Li₃NbO₄ was often obtained as an impurity
232 when the reaction time was too short or the heating
233 temperature was below 950 °C. The loss of lithium at elevated
234 temperature was effectively compensated by using additional
235 10–15% of the lithium precursor. Various molten salts,
236 including LiCl, NaCl, KCl, CsCl, KOH, and Li₂SO₄, were
237 used as the flux, but only KCl (mp = 770 °C) led to the
238 formation of phase-pure samples. The optimized R ratio was
239 between 2.5 and 5. In addition, the presence of trace O₂ in the
240 synthesis atmosphere can lead to the formation of Li₂MnO₃
241 and Li₃NbO₄ impurities.

242 The nominal composition of the as-synthesized oxide was
243 analyzed by an inductively coupled plasma optical emission
244 spectrometer, which confirmed the chemical formula of
245 Li_{1.3}Nb_{0.3}Mn_{0.4}O₂. As shown in the scanning electron
246 microscopy image in Figure 1a and high angle annular dark
247 field scanning transmission electron microscopy image in
248 Figure 1b, the particles adopted a large spherical shape with an
249 average size of ca. 5–8 μm and absence of any grain
250 boundaries. This suggests that all the facets have similar
251 thermodynamic stability and grow at a similar rate under
252 current synthesis conditions. Contrast reversal toward the
253 center of the particles observed on HAADF STEM images
254 indicates that the pristine particles are too thick for detailed
255 analytical electron microscopy. Focused ion beam (FIB) milling
256 was then used to reduce the thickness and a SAED pattern in
257 [110] zone axis was taken over an area covering almost entire
258 single particle (Figure 1c). The pattern matches well with the
259 one simulated using *Fm* $\bar{3}$ *m* structure with a lattice parameter of
260 *a* = 4.2 Å (Figure 1d), confirming the single crystal nature of

Figure 1. Morphology and structure of Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ crystals: (a) secondary electron SEM image, (b) HAADF STEM image, (c) SAED pattern taken in [110] zone axis, and (d) simulated electron diffraction pattern corresponding to panel c.

the particle. It is worth noting that the SAED pattern from this
particle also shows diffuse scattering effects that are character-
istic of materials with short-range ordering.²⁵ At this point, it is
unclear whether this short-range order occurs broadly in the
entire sample or if it has any effect on oxide properties and
behavior. Both of them require further investigation with
complementary analytical techniques, and the results will be
reported in a future publication.

Bulk phase purity and crystal structure of the synthesized
samples were evaluated by using both synchrotron X-ray and
neutron diffraction studies. Figure 2 shows the Rietveld
refinement of the diffraction patterns, while Table 1 lists the
refined structural parameter. In both sets of diffraction patterns,
the high peak symmetry observed in a wide angular range
indicates a single phase with high crystallinity. Rietveld
refinement confirms a rock-salt crystal structure with a lattice
parameter of 4.1954(1) Å and cell volume of 73.685(1) Å³,
which is in good agreement with the values reported by
Yabuuchi et al. on polycrystalline samples.²² All cations (Li, Nb,
and Mn) were randomly located in the 4a sites, and all M–O
bonds had the same distance of 2.0965(1) Å, confirming the
disordered nature of the oxide crystals. The refined occupancies
agree well with the chemical formula of Li_{1.3}Nb_{0.3}Mn_{0.4}O₂.

3.2. Electrochemical Studies. The electrochemical
performance of Li_{1.3}Nb_{0.3}Mn_{0.4}O₂ crystals was evaluated at
room temperature in a half-cell configuration. Due to high
resistance and poor utilization of large particles, only limited
capacities were obtained on composite electrodes made with
the as-synthesized micrometer-sized crystals. In a modified
procedure, the crystal size was reduced and electronic
conductivity improved by ball-milling the oxide crystals
together with a carbon black additive in a
Li_{1.3}Nb_{0.3}Mn_{0.4}O₂:carbon weight ratio of 80:20. A SEM image
of the mixture is shown in Figure S1. The mixture was then
added into the standard carbon and PVdF slurry in an *N*-
methyl-2-pyrrolidone (NMP) solvent and cast onto the Al

Figure 2. Rietveld refinement of (a) XRD and (b) neutron diffraction patterns collected on pristine $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ single crystals (PWEGEN). Values in parentheses indicate standard deviation.

Table 1. Structural Parameters of Pristine $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ Crystals Determined from Joint Rietveld Refinements of Synchrotron X-ray and TOF Neutron Data^a

atom	position	Wyck. site	occu.	$d_{(\text{M}-\text{O})}$ (Å)	bond valence	$100(U_{\text{iso}})$ (Å ²)
Li	0,0,0	4a	0.65	2.0965(1)	1.092	1.13(1)
Mn	0,0,0	4a	0.2	2.0965(1)	2.43	1.13(1)
Nb	0,0,0	4a	0.15	2.0965(1)	3.66	1.13(1)
O	0,0,0.5	4b	1.0	2.9650(1)		1.72(1)

^a $Fm\bar{3}m$, a (Å) = 4.1954(1), V (Å³) = 73.845(1), wR_p = 9.22%, δ = 0.0(0).

active material loading of ca. 2–3 mg/cm² were then cut out and used as cathodes in coin cells, which were cycled between 1.5 and 4.8 V vs Li^+/Li at a constant current density of 10 mA/g. Figure 3a shows the voltage profiles of the first four cycles. Similar to what was observed in the LMR oxide cathodes, two distinct regions, a sloping region and a plateau region, are present during the first charge. In the sloping region, the voltage increased steadily from the open circuit voltage (OCV) of 3.0 to 4.3 V, which delivered a specific capacity of ~115 mAh/g. Whereas in the plateau region, the voltage increased only slightly from 4.3 to 4.5 V before it rapidly increased to the cut off voltage of 4.8 V. A specific capacity of ~215 mAh/g was obtained above 4.3 V on the first cycle, leading to a total charge capacity of 330 mAh/g. An overall sloping discharge profile was observed, which delivered a total discharge capacity of ~290

current collector. The final weight ratio of active material:carbon additive:PVdF binder in the thus-prepared electrodes was 56:34:10. Electrode disks with a size of 1.6 cm² and an

Figure 3. (a and b) Voltage profiles of $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ half-cell cycling. (c) Specific capacity and average voltage as a function of cycle number. (d) Cyclic voltammogram of the first 9 scans at a scan rate of 5 mV/min. Solid symbols and open symbols in panel c represent capacity and voltage, respectively.

315 mAh/g at the cutoff voltage of 1.5 V. The Coulombic efficiency
 316 for the first cycle is therefore $\sim 88\%$. Two guidelines, one
 317 showing the theoretical capacity originated from the $\text{Mn}^{3+}/$
 318 Mn^{4+} redox couple at 118 mAh/g and the other showing the
 319 theoretical capacity based on Li content at 383 mAh/g are also
 320 shown in Figure 3a. It is clear that both charge and discharge
 321 capacities (330 and 290 mAh/g, respectively) are significantly
 322 larger than the theoretical capacity of the $\text{Mn}^{3+}/\text{Mn}^{4+}$ redox
 323 (118 mAh/g). The charge capacity obtained in the sloping
 324 region is consistent with the capacity from the $\text{Mn}^{3+}/\text{Mn}^{4+}$
 325 redox couple, suggesting that $\text{Mn}^{3+}/\text{Mn}^{4+}$ may be the sole
 326 redox process occurring in this region below 4.3 V.

327 In the second cycle, there is an overall decrease in the
 328 charging voltage of the sloping region, which led to an
 329 increased capacity of ~ 180 mAh/g upon reaching 4.3 V. The
 330 plateau region became somewhat sloping, but voltage increase
 331 remained slow. With cycling, the degree of sloping gradually
 332 increased, and the capacity obtained from this plateau region
 333 continued to decrease. The same trend was also observed on
 334 the discharge capacity, which was reduced to 160 mAh/g after
 335 27 cycles, a loss of nearly 45% (Figures 3b and c). These results
 336 obtained from room-temperature cycling are comparable to
 337 what was obtained at 60 °C by Yabuuchi et al. Although more
 338 carbon was used in our electrode, which likely contributed to
 339 the improved performance, the results may also indicate that
 340 the crystal samples have a kinetic advantage compared to the
 341 polycrystalline samples synthesized using the solid-state
 342 method. Further analysis showed that the oxide experienced
 343 not only capacity fade but also voltage fade. As shown in Figure
 344 3c, the average discharge voltage, obtained by dividing the total
 345 cell energy (E) by the cell capacity (Q) at 1.5 V, continues to
 346 decrease along with cycling. The extent of decay appears to be
 347 sensitive to the discharge cutoff voltage, and much faster decay
 348 was observed when the cutoff voltage was reduced from 1.5 to
 349 1.0 V.

350 Further insights on the redox activities of $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$
 351 crystals were obtained from cyclic voltammetry performed
 352 between 1.5 and 4.8 V at a scan rate of 5 mV/min. Figure 3d
 353 shows the results obtained from the first nine cycles. A broad
 354 peak centered at 3.5 V and a split peak between 4.5 and 4.7 V
 355 were observed during the first oxidation, corresponding to the
 356 sloping and plateau regions on the charging voltage profile,
 357 respectively. In the following scans, the oxidation peak at 3.5 V
 358 gradually moves toward high voltage, and its intensity grew
 359 while that of the peak at 4.5 V decreased, along with the
 360 disappearance of the split peak at 4.7 V after the first cycle. As
 361 the peak around 4.5–4.7 V is often attributed to the oxidation
 362 of O^{2-} to O^{n-} ($0 \leq n < 2$), the disappearance of the split peak
 363 and the continuous decrease in peak intensity suggest the
 364 complex and irreversible nature of the processes. During the
 365 first reduction, only a single peak was observed around 3.2 V,
 366 which decreased its intensity along with the appearance and
 367 gradual growth of a new peak around 2.2 V in the following
 368 scans. The pair of peaks centered at 3.5 V during oxidation and
 369 3.2 V during reduction is likely associated with the $\text{Mn}^{3+}/\text{Mn}^{4+}$
 370 redox couple. The growth of the reduction peak at 2.2 V at the
 371 expense of the peak at 3.2 V suggests the gradual trans-
 372 formation of the Mn species and the increasing contribution
 373 from the lower voltage process along with cycling. This
 374 transformation was not observed when the upper cutoff voltage
 375 was limited to 4.0 V, before the onset of the oxygen oxidation
 376 peak. The results suggest that the oxidation of O^{2-} to O^{n-} at
 377 high voltage may be responsible for the irreversible conversion

of the Mn^{3+} redox species, which ultimately led to both capacity
 and voltage fade in this oxide cathode.

Operando differential electrochemical mass spectroscopy was
 used to further examine the oxidation process of O^{2-} to O^{n-} (0
 $\leq n < 2$) occurring between 4.5 and 4.7 V. Figure 4 shows the

Figure 4. (a) Voltage profile and (b) O_2 and CO_2 gas evolution analyzed by DEMS of a cathode containing 2.66 mg $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ active material.

results obtained during the first 2 cycles at a constant current
 density of 25 mA/g. The voltage profiles are similar to those
 obtained from the coin cell tests, validating the excellent quality
 of the in situ electrochemical cell. During the first cycle, both
 CO_2 and O_2 evolution were detected. CO_2 evolution began
 around 3.8 V during the charge (the first blue guideline in the
 figure) and completed around the mid-discharge cycle (the
 second blue guideline). There was an increase in intensity
 around the midplateau after the onset of O_2 evolution, and the
 peak CO_2 evolution was reached at the end of charge at 4.8 V.
 Oxygen gas was not detected until the midplateau region at
 about 4.5 V (the first red guideline in the figure), which also
 peaked at the upper cutoff voltage of 4.8 V. Oxygen evolution
 completes at the beginning of first discharge after the resting
 step (the second red guideline). The cumulative CO_2 and O_2
 evolved from the first cycle were 1.53 and 0.11 μmol ,
 respectively, from a cathode that contained 2.66 mg of active
 materials. Oxygen evolution, therefore, is much reduced
 compared to what was reported on the lithium and
 manganese-rich layered oxide system by Luo et al.^{9,10} In the
 second cycle, the amount of CO_2 was reduced from 1.53 to
 0.46 μmol , while negligible O_2 evolution was detected.

Several sources may contribute to the evolution of CO_2 gas.
 Li_2CO_3 , a precursor from synthesis or a byproduct formed
 during air exposure of Li-TM oxide samples, can decompose
 during charging and release CO_2 gas. Recent studies confirmed
 that in LMR oxide cathodes,²⁶ all CO_2 gas evolved during the
 first charge in the typical voltage region, results from residual
 Li_2CO_3 in the samples. Given the similar outgassing behavior of
 $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ cathode to LMR, we postulate that residual
 Li_2CO_3 likely accounts for most, if not all, CO_2 evolution
 observed in Figure 4. Nevertheless, carbonate-based electrolytes
 are usually unstable at voltages above 4.3 V, especially in the
 presence of high valence transition metals (such as Mn^{4+}) that
 may catalyze the process and further lower the decomposition
 voltage, and a small percentage of CO_2 evolution at high
 potentials may originate from this degradation process.
 Carbonate solvent decomposition due to the attack of oxygen
 species has also been reported in the context of Li– O_2
 batteries.^{27–29} This represents another possible mechanism

Figure 5. (a) Synchrotron X-ray diffraction and (b) monochromatic neutron diffraction patterns collected on chemically delithiated $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ crystals. For neutron measurement, Li1.3 and Li0 samples were measured at SNS (POWGEN) using TOF, while Li1.1, Li0.92, Li0.7, Li0.49, and Li0.22 samples were measured at NIST (BT-1) using a single wavelength of $\lambda = 1.19$ Å. Note that the Li contents were determined by ICP measurements where an error range of up to ± 0.06 is possible.

sample was examined by both techniques. Figure S2 and Table 2 show the results obtained from Rietveld refinement of these XRD patterns. The structural transformation can be separated into two regions: single phase (phase 1) behavior when $x > 0.9$ (region I) and a two-phase (phase 2 and 3) behavior when $0 < x < 0.9$ (region II). The unit cell volume and phase fraction as a function of Li content were constructed from the refinement and shown in Figures 6a and b, respectively. The Vegard's law is only followed in region I, where the total volume change is ca. 0.2%. This is consistent with the participation of simple cation redox from Mn^{3+} to Mn^{4+} along with the extraction of 0.4 Li^+ from the $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ structure. Complex mixed oxygen activities are likely involved in region II, where phases 2 and 3 with the same rock-salt crystal structure but different lattice dimension evolve along with the extraction of Li. Smaller sized phase 3 reaches the maximum fraction when x is ~ 0.5 , whereas phase 2 becomes the main phase with further Li removal. Combining with the results from the electrochemical studies, we propose that lattice O^{2-} oxidation to O^{n-} ($0 < n < 2$) mainly occurs in phase 3, whereas loss of O from the lattice mainly occurs in phase 2. This was further supported by results from the joint X-ray and neutron refinements (Figure S3), which revealed that ~ 2 and 7.8 mol % of oxygen ion vacancies exist in the delithiated $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_{2-\delta}$ with $x = 0.2$ and 0 , respectively. Compared to the oxygen ion vacancies in the electrochemically charged samples, the much higher concentration is likely due to the fact that chemical delithiation was

for lattice oxygen loss in addition to oxygen gas evolution. In the simplest case where we safely assume only O atoms in O_2 gas originate from $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ cathode material, the total lattice O loss from a cathode containing 2.66 mg of active materials was $0.11 \mu\text{mol}$, which corresponds to 0.38 mol % of oxygen ion vacancy in the delithiated cathode and a capacity of 4.48 mAh/g. Although some of the first charge capacity may also come from Li_2CO_3 oxidizes to CO_2 via a $2 \text{e}^-/\text{CO}_2$ process and side reactions from electrolyte decomposition, it is evident that the redox process of O^{2-} to O^{n-} ($0 < n < 2$) species accounts for the majority of the first charge capacity above 4.3 V (215 mAh/g) and the first discharge capacity (290 mAh/g) in $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ cathode.

The DEMS results also reveal that capacity contribution from O extraction from the oxide lattice occurs above ~ 4.5 V. This mechanism is active during the first charge which disappears in the following cycles. This is consistent with the observation on the CV studies, where the oxidation peak centered at 4.7 V was observed only during the first scan. The combined results suggest that the peak at 4.7 V is likely related to O^{2-} oxidation to O_2 gas. The peak centering at 4.5 V, therefore, is attributed to the oxidation O^{2-} anion to O^{n-} ($0 < n < 2$) species. While both O_2 gas evolution and lattice oxygen oxidation occur in the first charge, the following cycles involve only the redox of O^{2-} to O^{n-} ($0 < n < 2$) species, signaled by the single oxidation peak at 4.5 V. The continuous decrease in peak intensity along with scanning indicates that the lattice anion redox process is highly irreversible. After merely nine cycles, the O redox process no longer participates.

3.3. Structural, Chemical, and Morphological Evolution during the First Delithiation. To gain further insights on structural and chemical changes during the first charge, we resorted to chemical delithiation, which allows for the preparation of samples in the large quantity needed for a number of analytical techniques. This approach also allows us to monitor morphological evolution as large single crystals can be used directly without the prior milling process, a necessary step to electrochemically charge and discharge the sample. Pristine $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ crystals were mixed with various amount of nitronium tetrafluoroborate oxidant in acetonitrile to prepare a series of $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ ($x = 1.21, 1.1, 1.06, 1.03, 0.92, 0.87, 0.81, 0.76, 0.64, 0.56, 0.49, 0.40, 0.32, 0.22, 0.13, 0.01$, and 0) samples at different states of charge, which were subsequently examined by synchrotron X-ray and neutron diffraction, hard and soft X-ray absorption spectroscopy, and SEM. While the Li content was determined by ICP measurements, the Mn/Nb ratio in each sample was verified to be consistent with the calculated value of 1.33 by using energy dispersive X-ray spectroscopy. Figure 5a shows the powder XRD patterns collected at 11-BM at the Advanced Photon Source. As discussed earlier, the pristine $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ is phase pure with a rock-salt crystal structure ($Fm\bar{3}m$; $a = 4.1954(1)$ Å). The 4a sites are occupied by Li, Nb, and Mn cations, whereas the 4b sites are occupied only by O anions. Oxygen vacancy was not detected in the pristine sample. With Li extraction, a single rock-salt phase remained when the lithium content is above 0.9 ($x > 0.9$). Further removing Li from the structure led to peak broadening and peak shifting toward to high angle, suggesting an overall reduction in lattice dimension. Similar results were also observed in neutron patterns collected on several samples in the series, as shown in Figure 5b. Joint X-ray and neutron Rietveld refinement was performed in cases where the same

Table 2. Summary of Structural Parameters in $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_{2-\delta}$ ($0 \leq x \leq 1.3$) Determined from Rietveld Refinement of Synchrotron X-ray Diffraction and Neutron Data^a

x	1.3	1.21	1.1	0.92	0.87	0.49	0.22	0.13	0.01	0
V_1 (Å ³)	73.845(1)	73.839(1)	73.689(1)	73.544(1)						
V_2 (Å ³)					73.84(1)	73.295(1)	70.838(1)	68.93(1)	68.89(1)	
V_3 (Å ³)					72.877(2)	72.494(1)	68.80(2)	67.07(3)	66.69(1)	68.34(1)
n_1	1.0	1.0	1.0	1.0						
n_2					0.57(1)	0.24(1)	0.93(1)	0.84(1)	0.93(1)	
n_3					0.43(1)	0.76(1)	0.07(1)	0.16(1)	0.07(1)	1.0
d_1 (Å)	2.0965(1)	2.0976(1)	2.0962(1)	2.0948(1)						
d_2 (Å)					2.0976(5)	2.0736(7)	2.0690(2)	2.0500(4)	2.0495(3)	
d_3 (Å)					2.0885(4)	2.0667(4)	2.0541(3)	2.0315(3)	2.0276(7)	2.0444(1)
$100(U_{\text{iso}1})$ (Å ²)	1.13(1)	1.22(1)	1.44(1)	1.47(1)						
	1.72(1)	1.90(1)	2.18(1)	2.28(2)						
$100(U_{\text{iso}2})$ (Å ²)					1.55(3)	2.39(1)	4.11(1)	4.75(6)	3.03(4)	
					2.53(5)	3.15(1)	4.85(1)	6.08(9)	3.59(5)	
$100(U_{\text{iso}3})$ (Å ²)					2.92(4)	2.39(1)	4.11(1)	4.75(6)	3.03(4)	5.46(6)
					2.75(6)	3.15(1)	4.85(1)	6.08(9)	3.59(5)	6.15(8)
δ	0(0)		0(0)			0(0)	0.02(1)			0.08(1)
wR_p (%)	9.22	6.78	6.85	7.40	7.25	8.91	6.71	8.70	6.91	8.16

^a V_1 , V_2 , V_3 , n_1 , n_2 , n_3 , d_1 , d_2 , d_3 , $U_{\text{iso}1}$, $U_{\text{iso}2}$, and $U_{\text{iso}3}$ represent the unit cell volume, phase fraction, bond distance of Li/Mn/Nb–O, and thermal factors for Li/Nb/Mn and O in phase 1, 2, and 3, respectively. δ represents oxygen anion vacancy in each phase, while wR_p represent the residual factor of the refinement.

performed in an open system that shifts the equilibrium toward more gas evolution. We also like to point out that protons were detected on the chemically delithiated oxides based on the prompt-gamma activation analysis (PGAA). This is likely molecular in nature as delithiated samples generally have higher surface area (as shown in the SEM images in Figure 9) and tend to absorb water during brief air exposure. However, it is possible that the detected protons may originate from cation exchange between Li^+ and H^+ , as previously reported on the classic layered oxides.³⁰ Further analysis is needed to fully understand the source of protons in these samples.

The kinetic properties of delithiation were evaluated by the relationship between the molar ratio of oxidant/oxide used and the residual Li content in the sample (Figure 6c), as determined by ICP. While a linear relationship is apparent in region I, significant deviation is seen in region II, further confirming the complex nature of mixed anion activities at lower Li contents.

Hard X-ray absorption spectroscopy correlates the changes in the absorption edge energy to the variations in the average oxidation state of the absorbing atoms being analyzed in the bulk material.^{31,32} The technique also probes the local and electronic structures near the absorbing atoms and has the ability to reveal element-specific structural changes in the samples. Due to the energy limit of the XAS beamline at SSRL, Nb spectra were not collected in this study. A previous report from Yabuuchi et al. indicated that Nb remains at 5+ during the entire charge/discharge process.²² The normalized spectra of Mn *K*-edge and XANES spectra of the chemically delithiated $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ series are shown in Figures 7a and b, respectively. The spectrum of pristine $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ is consistent with that of Mn^{3+} collected on the standard reference. Substantial blue-shift of the edge energy was observed as the lithium content was reduced from 1.3 to 0.9, consistent with the oxidation of Mn^{3+} to Mn^{4+} . Negligible changes were observed in samples with the Li content below 0.9. The edge position, defined by Photoelectron Energy Origin (E_0), is a commonly used parameter for extracting edge energy level in XANES spectra.³³ Figure 7c compares the edge position

as a function of lithium content in $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ samples. It is clear that Mn *K*-edge energy shifts toward higher value in a near linear fashion when $x > 0.9$ (region I) whereas it remains nearly constant when $0 < x < 0.9$ (region II). This is consistent with the results from X-ray and neutron diffraction studies, further confirming that Mn is only redox active in region I.

Further changes were observed in the intensity of Mn pre-edge absorption peaks arising from the dipole forbidden $1s \rightarrow 3d$ electronic transitions, which are typically weak in the transition metals. In the presence of 3d and 4p orbital hybridization, often resulting from structural distortion in local symmetry or noncentrosymmetric environment between the metal and oxygen coordination, the transitions are made partially allowed, and the peaks become much stronger.^{34–36} As shown in Figure 7d, the intensity of the pre-edge absorption peaks is similar in region I, which becomes noticeably stronger in region II. This suggests that although Mn redox activities are not involved in region II, the nature of Mn–O bond is altered in these samples, providing further evidence for oxygen redox activities in this region as the processes are likely behind the changes in local symmetry and environment of metal–ligand coordination. Similar to the edge shift observed in Figures 7a and b, the absorption energy of the pre-edge peaks also experienced blue shift in region I while maintaining nearly constant in region II, corroborating with the observation of Mn redox activity in region I only.

Oxygen redox activities during delithiation of $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ were investigated by soft XAS that directly probes the chemical changes of the element as a function of penetration depth. The depth profiling from the surface to bulk were achieved by using two different detectors in total electron yield (TEY) and fluorescence yield (FY) modes, which have a typical probing depth of 5 and 50 nm, respectively.³⁷ Figures 8a and b show the evolution of TEY and FY O *K*-edge XAS spectra as a function of Li content in $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$. In both cases, the spectra can be separated into the pre-edge region and postedge region at the black dashed line (533 eV) shown in the figures. Features in the postedge region are often attributed to

Figure 6. (a and b) Cell volume and phase fraction as a function of x in delithiated $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ samples and (c) relationship between the molar ratio of oxidant/oxide and residue Li content in chemically delithiated $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ samples.

To better understand the oxygen redox activities as a function of Li content, quantifications of peak features in both TEY and FY O K-edge XAS spectra were performed, and the results are shown in Figures 8c and d. The detailed peak fitting method is also shown in Figure S4. For both TEY and FY spectra, the ratio between the pre-edge peaks and post-edge peaks continuously increase with lithium extraction, as shown in Figure 8c. This confirms electron extraction from O2p and TM 3d hybridization and agrees with the previous report by Yabuuchi et al.²² It is, however, unclear on the contribution from TM and O due to the hybridization nature between them. The quantification on the p -band intensity, on the other hand, provides an opportunity to directly probe oxygen redox activities. As shown in Figure 8d, the initial p -band intensity is fairly low in the pristine $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ and remains nearly constant below Li content of ~ 0.9 , suggesting negligible interactions between oxygen anions or oxygen redox activities. As x decreases to below 0.9, there is an increase in the p -band intensity, and the trend continues to full delithiation ($x = 0$). The results suggest a continuous increase in oxygen valence states in this region and the participation of oxygen redox for charge compensation. Combined with the results from the Mn K edge XAS spectra which shows that Mn is active when $x > 0.9$ and inactive when $x < 0.9$, the quantification of O K-edge XAS spectra enables the understanding of complete charge compensation mechanism during delithiation of $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ crystals. Note that the p -band intensity in the $x < 0.9$ region is generally higher in the FY mode than that in the TEY mode, consistent with a difference in oxygen activities, likely enhanced lattice O loss on the surface.

The evolution of particle morphology during the first delithiation is revealed by the SEM images collected on the series of delithiated crystal samples (Figure 9). Some surface roughing was observed but the particles remain intact when $x > 0.9$. With further lithium extraction, large cracks begin to appear, which tend to propagate across the entire particle, leading to the eventual fracturing of the crystals. The number of cracks on each crystal increases along with the decrease in x but appears to reach the maximum at a Li content of 0.5–0.6. This coincides with the occurrence of maximum fraction of phase 3, as shown in Figure 6b. The change in morphology is also consistent with the fact that Mn is redox active when $x > 0.9$ while O is redox active when $x < 0.9$. The participation of oxygen redox appears to cause significant morphological damage. Our results also suggest that lattice oxidation of O^{2-} to O^{n-} ($0 < n < 2$) is more damaging than O_2 gas evolution, although it is possible that particles become more stable when x is less than 0.5–0.6 as mechanical stress is released by certain cracks and fractures at that point. Further evaluation on oxygen oxidation and mechanical damage in Li-excess oxides is warranted, but we emphasize that particle cracking and fracturing create fresh surfaces which can contribute to increased O loss from the lattice and side reactions with the electrolyte. Successful surface stabilization strategies such as elemental segregation and coating treatment therefore need to take consideration in particle morphology evolution during cycling.

4. CONCLUSION

Disordered, Li-excess transition-metal oxide cathodes are capable of delivering high capacities and are promising next-generation LIB cathodes. In this study, uniform and phase-pure $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ single crystals were synthesized by a molten-

$\text{O}1s \rightarrow \text{O}2p\text{-TM}4s4p$ hybridization, while those in the pre-edge region are attributed to $\text{O}1s \rightarrow \text{O}2p\text{-TM}3d$ hybridization.^{38,39} Due to the crystal field effect and octahedral coordination of oxygen, the pre-edge peaks are often represented by multiplets between 528 and 533 eV. As the O2p and TM3d band electrons are of most interest to redox activities, we focus on the analysis in the pre-edge region. It is clear that as lithium is extracted from $\text{Li}_{1.3}\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$, both TEY and FY spectra exhibit a significant increase in the pre-edge peak intensity. This increase is often used as a qualitative measure for the generated holes left by electron extraction from either oxygen or TM at the O2p-TM3d energy levels.^{39–41} A new shoulder peak at ~ 529 eV also appeared and gradually grew along with delithiation, suggesting the involvement additional O-TM hybridization (referred to as p -band) during the process. Due to similarity between the p -band and the pre-edge absorption peak often observed in peroxides and superperoxides,^{22,40,42,43} we speculate this new O-TM hybridization arises from the interaction between oxygen anions as a result of oxygen redox activities. Note that similar results were previously reported on the LMR cathode materials.^{22,40}

Figure 7. (a and b) Mn K-edge hard XAS and XANES spectra, (c) XANES energy edge (E_0) as a function of Li content in $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ samples, and (d) Mn K-edge pre-edge spectra.

Figure 8. (a and b) O K-edge soft XAS spectra obtained in TEY and FY modes, (c) intensity ratio between the pre-edge and postedge absorption peaks as a function of lithium content, and (d) intensity of the p -band as a function of lithium content. In panels a and b, black dashed line divides the pre- and postedge regions, and the blue dashed arrow follows the evolution of the p -band intensity.

salt method for the first time and used as a diagnostic tool to understand the charge compensation mechanism during Li extraction and insertion. Electrochemical evaluation suggests structural irreversibility after the first cycle, leading to capacity and voltage fade of the cathode. Both CO_2 and O_2 gas evolution was found during the first cycle, with the latter only accounting for $\sim 1\%$ of the charge capacity, in addition to the contribution of $\sim 35\%$ capacity from the $\text{Mn}^{3+/4+}$ redox couple. A large fraction of capacity therefore originates from the redox process of $\text{O}^{2-}/\text{O}^{n-}$ ($0 < n < 2$). Joint refinement on

synchrotron X-ray and neutron diffraction patterns of $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ series showed that the oxide undergoes a single-phase reaction involving $\text{Mn}^{3+/4+}$ redox reaction when $0.9 < x < 1.3$ and a two-phase reaction involving mixed $\text{O}^{2-}/\text{O}^{n-}$ ($0 \leq n < 2$) reactions when $0 < x < 0.9$. For the fully delithiated sample, ~ 7.8 mol % of oxide-ion vacancies were found, resulting from O loss from the lattice. Cracks and fractures initiated and intensified with deep lithium extraction, which serves as a means to release the mechanical stress

Figure 9. SEM images of chemically delithiated $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ crystals: $x =$ (a) 1.3, (b) 1.1, (c) 0.87, (d) 0.76, (e) 0.56, and (f) 0.13.

endured by the large crystals during the oxygen redox processes.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.chemmater.7b05036.

SEM image of carbon mixed $\text{LiNb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ crystals after ball milling, Rietveld refinement of synchrotron X-ray diffraction patterns collected on chemically delithiated $\text{Li}_x\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ crystals, Rietveld refinement of synchrotron X-ray and neutron diffraction patterns collected on chemically delithiated $\text{Li}_0\text{Nb}_{0.3}\text{Mn}_{0.4}\text{O}_2$ crystals, and process used to quantify O K -edge XAS data (PDF)

AUTHOR INFORMATION

Corresponding Author

*E-mail: gchen@lbl.gov.

ORCID

Wang Hay Kan: 0000-0002-1663-2999

Joseph K. Papp: 0000-0002-7982-6096

Bryan McCloskey: 0000-0001-6599-2336

Guoying Chen: 0000-0002-3218-2609

Notes

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REFERENCES

- Goodenough, J. B.; Manthiram, A. A perspective on electrical energy storage. *MRS Commun.* **2014**, *4*, 135–142.
- Li, W.; Song, B.; Manthiram, A. High-voltage positive electrode materials for lithium-ion batteries. *Chem. Soc. Rev.* **2017**, *46*, 3006–3059.
- Manthiram, A.; Knight, J. C.; Myung, S.-T.; Oh, S.-M.; Sun, Y.-K. Nickel-Rich and Lithium-Rich Layered Oxide Cathodes: Progress and Perspectives. *Adv. Energy Mater.* **2016**, *6*, 1501010.
- Aydinli, M. K.; Kohan, A. F.; Ceder, G.; Cho, K.; Joannopoulos, J. Ab initio study of lithium intercalation in metal oxides and metal dichalcogenides. *Phys. Rev. B: Condens. Matter Mater. Phys.* **1997**, *56*, 1354–1365.
- McCalla, E.; Abakumov, A. M.; Saubanere, M.; Foix, D.; Berg, E. J.; Rousse, G.; Doublet, M.-L.; Gonbeau, D.; Novak, P.; Van Tendeloo, G.; Dominko, R.; Tarascon, J.-M. Visualization of O-O peroxo-like dimers in high-capacity layered oxides for Li-ion batteries. *Science (Washington, DC, U. S.)* **2015**, *350*, 1516–1521.
- Pearce, P. E.; Perez, A. J.; Rousse, G.; Saubanere, M.; Batuk, D.; Foix, D.; McCalla, E.; Abakumov, A. M.; Van Tendeloo, G.; Doublet, M.-L.; Tarascon, J.-M. Evidence for anionic redox activity in a tridimensional-ordered Li-rich positive electrode $\beta\text{-Li}_2\text{IrO}_3$. *Nat. Mater.* **2017**, *16*, 580–586.
- Sathiya, M.; Rousse, G.; Ramesha, K.; Laisa, C. P.; Vezin, H.; Sougrati, M. T.; Doublet, M. L.; Foix, D.; Gonbeau, D.; Walker, W.; Prakash, A. S.; Ben Hassine, M.; Dupont, L.; Tarascon, J. M. Reversible anionic redox chemistry in high-capacity layered-oxide electrodes. *Nat. Mater.* **2013**, *12*, 827–835.
- Armstrong, A. R.; Holzapfel, M.; Novak, P.; Johnson, C. S.; Kang, S.-H.; Thackeray, M. M.; Bruce, P. G. Demonstrating Oxygen Loss and Associated Structural Reorganization in the Lithium Battery Cathode $\text{Li}[\text{Ni}_{0.2}\text{Li}_{0.2}\text{Mn}_{0.6}]\text{O}_2$. *J. Am. Chem. Soc.* **2006**, *128*, 8694–8698.
- Luo, K.; Roberts, M. R.; Guerrini, N.; Tapia-Ruiz, N.; Hao, R.; Massel, F.; Pickup, D. M.; Ramos, S.; Liu, Y.-S.; Guo, J.; Chadwick, A. V.; Duda, L. C.; Bruce, P. G. Anion Redox Chemistry in the Cobalt

- 771 Free 3d Transition Metal Oxide Intercalation Electrode Li-
772 $[\text{Li}_{0.2}\text{Ni}_{0.2}\text{Mn}_{0.6}]\text{O}_2$. *J. Am. Chem. Soc.* **2016**, *138*, 11211–11218.
- 773 (10) Luo, K.; Roberts, M. R.; Hao, R.; Guerrini, N.; Pickup, D. M.;
774 Liu, Y.-S.; Edstrom, K.; Guo, J.; Chadwick, A. V.; Duda, L. C.; Bruce,
775 P. G. Charge-compensation in 3d-transition-metal-oxide intercalation
776 cathodes through the generation of localized electron holes on oxygen.
777 *Nat. Chem.* **2016**, *8*, 684–691.
- 778 (11) Shukla, A. K.; Ramasse, Q. M.; Ophus, C.; Duncan, H.; Hage,
779 F.; Chen, G. Unravelling structural ambiguities in lithium- and
780 manganese-rich transition metal oxides. *Nat. Commun.* **2015**, *6*, 8711.
- 781 (12) Mohanty, D.; Li, J.; Abraham, D. P.; Huq, A.; Payzant, E. A.;
782 Wood, D. L.; Daniel, C. Unraveling the Voltage-Fade Mechanism in
783 High-Energy-Density Lithium-Ion Batteries: Origin of the Tetrahedral
784 Cations for Spinel Conversion. *Chem. Mater.* **2014**, *26*, 6272–6280.
- 785 (13) Wang, D.; Huang, Y.; Huo, Z.; Chen, L. Synthesize and
786 electrochemical characterization of Mg-doped Li-rich layered Li-
787 $[\text{Li}_{0.2}\text{Ni}_{0.2}\text{Mn}_{0.6}]\text{O}_2$ cathode material. *Electrochim. Acta* **2013**, *107*,
788 461–466.
- 789 (14) Song, B.; Zhou, C.; Wang, H.; Liu, H.; Liu, Z.; Lai, M. O.; Lu, L.
790 Advances in sustain stable voltage of Cr-doped Li-rich layered
791 cathodes for lithium ion batteries. *J. Electrochem. Soc.* **2014**, *161*,
792 A1723–A1730.
- 793 (15) Li, G. R.; Feng, X.; Ding, Y.; Ye, S. H.; Gao, X. P. AlF₃-coated
794 $\text{Li}(\text{Li}_{0.17}\text{Ni}_{0.25}\text{Mn}_{0.58})\text{O}_2$ as cathode material for Li-ion batteries.
795 *Electrochim. Acta* **2012**, *78*, 308–315.
- 796 (16) Shi, S. J.; Tu, J. P.; Tang, Y. Y.; Liu, X. Y.; Zhang, Y. Q.; Wang,
797 X. L.; Gu, C. D. Enhanced cycling stability of Li-
798 $[\text{Li}_{0.2}\text{Mn}_{0.54}\text{Ni}_{0.13}\text{Co}_{0.13}]\text{O}_2$ by surface modification of MgO with
799 melting impregnation method. *Electrochim. Acta* **2013**, *88*, 671–679.
- 800 (17) Zhang, X.; Belharouak, I.; Li, L.; Lei, Y.; Elam, J. W.; Nie, A.;
801 Chen, X.; Yassar, R. S.; Axelbaum, R. L. Structural and Electrochemical
802 Study of Al_2O_3 and TiO_2 Coated $\text{Li}_{1.2}\text{Ni}_{0.13}\text{Mn}_{0.54}\text{Co}_{0.13}\text{O}_2$ Cathode
803 Material Using ALD. *Adv. Energy Mater.* **2013**, *3*, 1299–1307.
- 804 (18) Kim, K. J.; Jo, Y. N.; Lee, W. J.; Subburaj, T.; Prasanna, K.; Lee,
805 C. W. Effects of inorganic salts on the morphological, structural, and
806 electrochemical properties of prepared nickel-rich $\text{Li}[\text{Ni}_{0.6}\text{Co}_{0.2}\text{Mn}_{0.2}]\text{O}_2$.
807 *J. Power Sources* **2014**, *268*, 349–355.
- 808 (19) Shigemura, H.; Tabuchi, M.; Sakaebe, H.; Kobayashi, H.;
809 Kageyama, H. Lithium Extraction and Insertion Behavior of
810 Nanocrystalline $\text{Li}_2\text{TiO}_3\text{-LiFeO}_2$ Solid Solution with Cubic Rock Salt
811 Structure. *J. Electrochem. Soc.* **2003**, *150*, A638–A644.
- 812 (20) Tabuchi, M.; Nakashima, A.; Shigemura, H.; Ado, K.;
813 Kobayashi, H.; Sakaebe, H.; Tatsumi, K.; Kageyama, H.; Nakamura,
814 T.; Kanno, R. Fine $\text{Li}_{(4-x)/3}\text{Ti}_{(2-2x)/3}\text{Fe}_x\text{O}_2$ ($0.18 \leq x \leq 0.67$) powder
815 with cubic rock-salt structure as a positive electrode material for
816 rechargeable lithium batteries. *J. Mater. Chem.* **2003**, *13*, 1747–1757.
- 817 (21) Lee, J.; Urban, A.; Li, X.; Su, D.; Hautier, G.; Ceder, G.
818 Unlocking the Potential of Cation-Disordered Oxides for Rechargeable
819 Lithium Batteries. *Science (Washington, DC, U. S.)* **2014**, *343*, 519–
820 522.
- 821 (22) Yabuuchi, N.; Takeuchi, M.; Nakayama, M.; Shiiba, H.; Ogawa,
822 M.; Nakayama, K.; Ohta, T.; Endo, D.; Ozaki, T.; Inamasu, T.; Sato,
823 K.; Komaba, S. High-capacity electrode materials for rechargeable
824 lithium batteries: Li_3NbO_4 -based system with cation-disordered
825 rocksalt structure. *Proc. Natl. Acad. Sci. U. S. A.* **2015**, *112*, 7650–7655.
- 826 (23) Toby, B. H. EXPGUI, a graphical user interface for GSAS. *J.*
827 *Appl. Crystallogr.* **2001**, *34*, 210–213.
- 828 (24) Xu, J.; Renfrew, S.; Marcus, M. A.; Sun, M.; McCloskey, B. D.;
829 Tong, W. Investigating $\text{Li}_2\text{NiO}_2\text{-Li}_2\text{CuO}_2$ Solid Solutions as High-
830 Capacity Cathode Materials for Li-Ion Batteries. *J. Phys. Chem. C* **2017**,
831 *121*, 11100–11107.
- 832 (25) Edington, J. W. *Practical Electron Microscopy in Materials Science*;
833 Van Nostrand Reinhold Co., 1976.
- 834 (26) Renfrew, S.; McCloskey, B. D. Residual lithium carbonate
835 predominantly accounts for first cycle CO_2 and CO outgassing of Li-
836 stoichiometric and Li-rich layered transition metal oxides. *J. Am. Chem.*
837 *Soc.* **2017**, *139*, 17853–17860.
- 838 (27) Freunberger, S. A.; Chen, Y.; Peng, Z.; Griffin, J. M.; Hardwick,
839 L. J.; Barde, F.; Novak, P.; Bruce, P. G. Reactions in the Rechargeable
Lithium- O_2 Battery with Alkyl Carbonate Electrolytes. *J. Am. Chem.*
840 *Soc.* **2011**, *133*, 8040–8047.
- (28) Aurbach, D.; Daroux, M.; Faguy, P.; Yeager, E. The
842 electrochemistry of noble metal electrodes in aprotic organic solvents
843 containing lithium salts. *J. Electroanal. Chem. Interfacial Electrochem.*
844 **1991**, *297*, 225–44.
- (29) Aurbach, D.; Gofer, Y.; Langzam, J. The correlation between
846 surface chemistry, surface morphology, and cycling efficiency of
847 lithium electrodes in a few polar aprotic systems. *J. Electrochem. Soc.*
848 **1989**, *136*, 3198.
- (30) Choi, J.; Alvarez, E.; Arunkumar, T. A.; Manthiram, A. Proton
850 insertion into oxide cathodes during chemical delithiation. *Electrochem.*
851 *Solid-State Lett.* **2006**, *9*, A241–A244.
- (31) Terada, Y.; Yasaka, K.; Nishikawa, F.; Konishi, T.; Yoshio, M.;
853 Nakai, I. In Situ XAFS Analysis of $\text{Li}(\text{Mn}, \text{M})_2\text{O}_4$ ($\text{M} = \text{Cr}, \text{Co}, \text{Ni}$) 5 V
854 Cathode Materials for Lithium-Ion Secondary Batteries. *J. Solid State*
855 *Chem.* **2001**, *156*, 286–291.
- (32) Nam, K.-W.; Bak, S.-M.; Hu, E.; Yu, X.; Zhou, Y.; Wang, X.; Wu,
857 L.; Zhu, Y.; Chung, K.-Y.; Yang, X.-Q. Combining In Situ Synchrotron
858 X-Ray Diffraction and Absorption Techniques with Transmission
859 Electron Microscopy to Study the Origin of Thermal Instability in
860 Overcharged Cathode Materials for Lithium-Ion Batteries. *Adv. Funct.*
861 *Mater.* **2013**, *23*, 1047–1063.
- (33) Hu, E.; Bak, S.-M.; Liu, J.; Yu, X.; Zhou, Y.; Ehrlich, S. N.; Yang,
863 X.-Q.; Nam, K.-W. Oxygen-Release-Related Thermal Stability and
864 Decomposition Pathways of $\text{Li}_x\text{Ni}_{0.5}\text{Mn}_{1.5}\text{O}_4$ Cathode Materials. *Chem.*
865 *Mater.* **2014**, *26*, 1108–1118.
- (34) Kim, M. G.; Shin, H. J.; Kim, J.-H.; Park, S.-H.; Sun, Y.-K. XAS
867 Investigation of Inhomogeneous Metal-Oxygen Bond Covalency in
868 Bulk and Surface for Charge Compensation in Li-Ion Battery Cathode
869 $\text{Li}[\text{Ni}_{1/3}\text{Co}_{1/3}\text{Mn}_{1/3}]\text{O}_2$ Material. *J. Electrochem. Soc.* **2005**, *152*,
870 A1320–A1328.
- (35) Kim, M. G.; Yo, C. H. X-ray Absorption Spectroscopic Study of
872 Chemically and Electrochemically Li Ion Extracted $\text{Li}_y\text{Co}_{0.85}\text{Al}_{0.15}\text{O}_2$
873 Compounds. *J. Phys. Chem. B* **1999**, *103*, 6457–6465.
- (36) Ignatov, A. Y.; Ali, N.; Khalid, S. Mn K-edge XANES study of
875 the $\text{La}_{1-x}\text{Ca}_x\text{MnO}_3$ colossal magnetoresistive manganites. *Phys. Rev. B*
876 *Condens. Matter Mater. Phys.* **2001**, *64*, 014413/1–014413/16.
- (37) Stöhr, J. *NEXAFS Spectroscopy*; Springer, 1992.
- (38) Yoon, W.-S.; Balasubramanian, M.; Chung, K. Y.; Yang, X.-Q.;
879 McBreen, J.; Grey, C. P.; Fischer, D. A. Investigation of the Charge
880 Compensation Mechanism on the Electrochemically Li-Ion Deinter-
881 calated $\text{Li}_{1-x}\text{Co}_{1/3}\text{Ni}_{1/3}\text{Mn}_{1/3}\text{O}_2$ Electrode System by Combination
882 of Soft and Hard X-ray Absorption Spectroscopy. *J. Am. Chem. Soc.*
883 **2005**, *127*, 17479–17487.
- (39) de Groot, F. M. F.; Grioni, M.; Fuggle, J. C.; Ghijsen, J.;
885 Sawatzky, G. A.; Petersen, H. Oxygen 1s x-ray-absorption edges of
886 transition-metal oxides. *Phys. Rev. B: Condens. Matter Mater. Phys.*
887 **1989**, *40*, 5715–5723.
- (40) Oishi, M.; Yamanaka, K.; Watanabe, I.; Shimoda, K.; Matsunaga,
889 T.; Arai, H.; Ukyo, Y.; Uchimoto, Y.; Ogumi, Z.; Ohta, T. Direct
890 observation of reversible oxygen anion redox reaction in Li-rich
891 manganese oxide, Li_2MnO_3 , studied by soft X-ray absorption
892 spectroscopy. *J. Mater. Chem. A* **2016**, *4*, 9293–9302.
- (41) Ma, C.; Alvarado, J.; Xu, J.; Clément, R. J.; Kodur, M.; Tong,
894 W.; Grey, C. P.; Meng, Y. S. Exploring Oxygen Activity in the High
895 Energy P2-Type $\text{Na}_{0.78}\text{Ni}_{0.23}\text{Mn}_{0.69}\text{O}_2$ Cathode Material for Na-Ion
896 Batteries. *J. Am. Chem. Soc.* **2017**, *139*, 4835–4845.
- (42) Kang, J. S.; Kim, D. H.; Hwang, J. H.; Baik, J.; Shin, H. J.; Kim,
898 M.; Jeong, Y. H.; Min, B. I. Soft x-ray absorption and photoemission
899 spectroscopy study of superoxide KO_2 . *Phys. Rev. B: Condens. Matter*
900 *Mater. Phys.* **2010**, *82*, 193102.
- (43) Yilmaz, E.; Yogi, C.; Yamanaka, K.; Ohta, T.; Byon, H. R.
902 Promoting Formation of Noncrystalline Li_2O_2 in the Li- O_2 Battery
903 with RuO_2 Nanoparticles. *Nano Lett.* **2013**, *13*, 4679–4684.