Redox Chemistry and the Role of Trapped Molecular O₂ in Li-Rich Disordered Rocksalt Oxyfluoride Cathodes

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ABSTRACT: In the search for high energy density cathodes for next-generation lithium-ion batteries, the disordered rocksalt oxyfluorides are receiving significant attention due to their high capacity and lower voltage hysteresis compared with ordered Li-rich layered compounds. However, a deep understanding of these phenomena and their redox chemistry remains incomplete. Using the archetypal oxyfluoride, Li₂MnO₂F, we show that the oxygen redox process in such materials involves the formation of molecular O₂ trapped in the bulk structure of the charged cathode, which is reduced on discharge. The molecular O₂ is trapped rigidly within vacancy clusters and exhibits minimal mobility unlike free gaseous O₂, making it more characteristic of a solid-like environment. The Mn redox process occurs between octahedral Mn³⁺ and Mn⁴⁺ with no evidence of tetrahedral Mn⁵⁺ or Mn⁷⁺. We furthermore derive the relationship between local coordination environment and redox potential; this gives rise to the observed overlap in Mn and O redox couples and reveals that the onset potential of oxide ion oxidation is determined by the degree of ionicity around oxygen, which extends models based on linear Li–O–Li configurations. This study advances our fundamental understanding of redox mechanisms in disordered rocksalt oxyfluorides, highlighting their promise as high capacity cathodes.

INTRODUCTION

Advances in high energy density cathodes are crucial for the development of next-generation lithium-ion batteries for portable electronics and electric vehicles. Lithium-rich cathode materials are attracting considerable attention as they offer increased capacities by invoking redox chemistry on both the transition metal and oxide ions, but rather than on only the transition metal as found in traditional oxide-based intercalation compounds.

Recently, there has been growing interest in disordered Li-rich intercalation materials, especially disordered rocksalt structures, including early work on systems based on Li₁ₓNbO₃ and LiₓVO₂F [refs 27–29]. House et al. presented for the first time an all-manganese oxyfluoride, Li₁₀Mn₀.₇⁹OₓF₀.₉₉, with a disordered rocksalt structure, which exhibits a large capacity utilizing both Mn and O redox. This LiₓMnO₂F-based cathode has a discharge capacity of ~280 mA h g⁻¹ (corresponding to 960 W h kg⁻¹) after the initial charge, making it comparable to Li-rich layered oxides such as Li₁₀NiₓMn₀.₇⁹Co₀.₃₁O₃ and greater than conventional cathodes such as LiCoO₂ (170 mA h g⁻¹) and NMC-Li(Ni,Mn,Co)O₂ (200–220 mA h g⁻¹). Cathodes comprised of manganese (rather than cobalt or nickel) are also attractive due to its low cost, low toxicity, and high natural abundance.

Layered Li-rich cathode materials commonly undergo extensive structural rearrangement during the first charge/discharge cycle leading to a large voltage hysteresis and O₂ gas evolution at the surface; this involves a substantial loss of voltage and therefore energy density. In contrast, the disordered rocksalt LiₓMnO₂F does not exhibit such large first cycle voltage hysteresis (shown in Figure 1) and also shows minimal oxygen loss, which are major advantages of this system. These differences raise the important question: to what extent does the transition metal and oxygen redox chemistry in the ordered Li-rich layered compounds translate to disordered rocksalt systems?

In this work, combined operando X-ray absorption spectroscopy (XAS), high resolution resonant inelastic X-ray scattering (RIXS), and ab initio modeling techniques are used to elucidate and quantify the Mn and O redox chemistry as well as local structural changes upon delithiation. We show, for the first time, by experimental methods (O K-edge RIXS) and ab
initio modeling that Li removal from the rocksalt oxyfluoride is accompanied by the formation of molecular O₂ trapped inside the cathode particles. Ab initio molecular dynamics simulations show that the trapped O₂ exhibits substantially reduced freedom of mobility, making it more characteristic of a solid-like environment in line with recent solid state ¹⁷O NMR measurements for O₂ in the layered cathode Li₁.₂Ni₀.₁₃Mn₀.₅₄Co₀.₁₃O₂ [ref 26]. Previously the significance of the Li⁺−O−Li⁺ configurations in Li-rich oxides in pinning O₂p states at the top of the oxygen valence band and hence accessible for O-redox has been emphasized. Here, we show that the onset potential of oxygen oxidation varies with the number of coordinating Li⁺ ions and that, along with the strong modulation of Mn redox potential by its O/F anionic coordination environment, leads to overlap of the Mn and the O redox processes, i.e., transition metal and oxygen redox occur together in the disordered rocksalts.

RESULTS AND DISCUSSION

Disordered Rocksalt Structural Properties. Li₂MnO₂F was prepared by mechanochemical ball-milling, and the oxidation state of Mn was subsequently confirmed as +3.00(5) from iodometric titration. Li₂MnO₂F possesses a cubic rock-salt structure where each cation (Li⁺ or Mn³⁺) is octahedrally coordinated to six anions (O₂⁻ or F⁻) and vice versa (illustrated in Figure 2). To investigate the possibility of local ordering, Mn K-edge extended X-ray absorption fine structure (EXAFS) and neutron pair distribution function (PDF) analysis were performed on pristine Li₂MnO₂F (the experimental methods are detailed in the Supporting Information (SI), Section S1).

The EXAFS technique is an element-specific probe of local coordination environment, in this case around Mn, giving a plot of the nearest-neighbor (NN) atoms as a function of distance from the central atom. As shown in Figure 2a, the first two peaks in the EXAFS spectrum for Li₂MnO₂F are a close match in shape and relative position to that of MnO, which has a well-defined cubic rocksalt structure. These peaks correspond to the first and second NN shells of atoms, anions and cations, respectively. Normalizing each spectrum by the first peak area, since O₂⁻ and F⁻ are indistinguishable with EXAFS, permits direct comparison of the second peak area. Here, Mn is a much stronger scatterer of the photoelectron wave than Li, which is a very weak scatterer, so Mn dominates the second peak intensity. The difference in peak area between the two materials indicates a much lower amount of second NN Mn for Li₂MnO₂F than MnO, consistent with the presence of Li on the cation sites in the former. Measuring the peak intensity relative to a baseline of zero scattering from 12 Li and

Figure 1. Charge−discharge curves. Representative first cycle load curves for disordered rocksalt Li₂MnO₂F (blue) and layered Li₁₂Ni₀.₁₃Co₀.₁₃Mn₀.₅₄O₂ (red) at a current rate of 20 mA g⁻¹. (Second cycle data are presented in Supporting Information, Figure S1).

Figure 2. Structure of Li₂MnO₂F. (a) Mn K-edge EXAFS for pristine Li₂MnO₂F compared with a MnO reference, each with cubic rocksalt crystal structure. The spectra are normalized to the area under the first peak corresponding to the occupancy of the first nearest neighbor anion site which is the same for both. A good fit of the EXAFS data can also be obtained with a rocksalt model with 4 × 2nd nearest neighbor Mn. (b) Neutron PDF data fitted to structural models of distortion-free random cubic rocksalt and the Monte Carlo derived model for pristine Li₂MnO₂F. Li, Mn, O, and F atoms are indicated by green, purple, red, and gray spheres, respectively. There is very good agreement between the models and the PDF data showing Li₂MnO₂F exhibits a close to completely disordered rocksalt structure. The slight asymmetry of the first peak at around 1.9 Å may indicate some element-specific preference for shorter bond length. Refined cell parameters a = 4.117 Å and a = 12.152 Å, b = c = 8.336 Å, and Uiso values 0.027 and 0.014, respectively.
maximum scattering from 12 Mn (as in MnO), shows an average of 3.6 Mn as second NN for Li₂MnO₂F. This is in line with that expected from a completely random distribution of Li and Mn in a 2:1 ratio, i.e., four Mn. As further confirmation, a good fit was obtained of the EXAFS data using a rocksalt structure with six degenerate first NN O and four degenerate second NN Mn atoms, Figure 2a and Table S1.

PDF is another powerful tool for probing local structure, giving a superposed plot of all atom−atom pairs throughout the structure resolved as a function of increasing separation. Unlike EXAFS, neutron PDF can probe over much longer correlation lengths and is much more sensitive to Li, since neutrons are more strongly scattered by Li than X-rays. The fitted PDF data (Figure 2b), show that the local structure of Li₂MnO₂F can be well-described by a disordered rocksalt model indicating minimal short-range order. Together with the EXAFS data, the experimental evidence supports a close to completely disordered rocksalt structure for Li₂MnO₂F.

To obtain a computationally tractable structural model for Li₂MnO₂F capturing this disorder, a Monte Carlo random sampling approach was employed to generate a 3 × 2 × 2 unit cell which possessed a representative distribution of different sites (computational methods applied to battery cathode materials are well established11,53,54 and detailed in the SI, Section S1). The validity of this Monte Carlo-derived model was checked by fitting to the neutron PDF data. The quality of the fit was even better than the distortion-free one showing it is a closer match to the experimentally observed structure. Furthermore, the calculated mean lattice parameter, 4.146 Å, compare well with the experimental value (a = b = c = 4.118 Å) from X-ray diffraction studies.36 Our ab initio simulations confirm that the disordered rocksalt structure of Li₂MnO₂F does not exhibit the cooperative Jahn−Teller distortion usually associated with Mn³⁺ in ordered structures, which often leads to poor cycling. The full structural data set for the pristine Li₂MnO₂F computational model is given in the SI, section S2.

Charge-compensation on Lithium Ion Extraction. To investigate the redox processes occurring over the first cycle in Li₂MnO₂F, operando Mn K-edge XANES was performed. Operando experiments allow the intercalation reaction in the cathode to be followed under operating conditions, eliminating the effect of any relaxation phenomena. As shown in Figure 3, a continuous shift in the Mn K-edge is observed during charge and discharge in line with the expected oxidation and reduction from Mn³⁺ toward Mn⁵⁺. Near the top of charge, around x ≈ 1.2 in LiₓMnO₂F, the changes become less pronounced as oxygen oxidation starts to dominate the redox process; this corresponds to a slight inflection in the electrochemical load curve. Note that previous operando electrochemical mass spectrometry studies50 indicate that
there is negligible oxygen loss. Given that the edge continues to evolve and the voltage profile remains sloped throughout this region (Figure 1), there must be a significant degree of overlap between the Mn and O redox couples. Close analysis of the Mn K-edge pre-edge (Figure 3d), which is generally considered to be a better measure of oxidation state than the main edge, reveals a similar trend. The pre-edge shape (insets of Figure 3b,c) does not appear to change much, but there is evidence of a slight increase and decrease in intensity of the twin peaks. These peaks arise from the quadrupole-allowed transition from the Mn 1s to the Mn 3d states, which are subdivided by crystal field splitting, and are weak due to the centro-symmetry of octahedral coordination. The intensity gain can be attributed to a slight distortion of this centro-symmetry allowing mixing between the Mn 3d and 4p states. In contrast, pre-edge features for tetrahedral geometries tend to be of significantly larger intensity, often of comparable height to the main edge, reveals a similar trend.

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Interestingly, recent reports for the vanadium-based disordered rocksalt systems, Li$_{1.25}$Nb$_{0.25}$V$_{0.5}$O$_2$ and Li$_2$VO$_2$F, show XANES data indicating a strong increase in pre-edge intensity, characteristic of vanadium in a noncentrosymmetric coordination environment such as tetrahedral V$^{5+}$. Baur et al. recently confirmed the presence of tetrahedral vanadium by PDF in Li$_2$VO$_2$F, and Chang et al. report superoxide formation in this oxyfluoride from computational and EPR studies. We note that Lun et al. studied the Li−Mn−O−F chemical space to derive a capacity map of Li percolation and redox properties. In the context of Li-rich oxide structures, Hong et al. report that oxygen redox may be stabilized in the local coordination environments created through cation vacancies, and Gent et al. find that the defect formation energy landscape is a key factor controlling the electrochemical reversibility of high valent redox. Our previous simulation study on Li-rich layered Li$_2$MnO$_3$ suggests that delithiation leads to oxygen dimerization and eventually to the formation of molecular O$_2$. To further probe the changes in local environment around the Mn during the first cycle, the Mn K-edge EXAFS were
analyzed. As shown in Figure 3e,f, the first and second neighbor atoms to Mn do not change as a function of state of charge. This clearly shows Mn remains octahedrally coordinated throughout the charge/discharge cycle, with no evidence of tetrahedral Mn.

To complement our XANES and EXAFS work, we used DFT methods to examine local structures and to quantify the redox chemistry on lithium extraction from Li$_x$Mn$_2$O$_2$F. As in previous studies,9,11 we stress that high level hybrid functionals were employed as they are found to be important in reproducing accurately the electronic structure of oxygen states (further details in the SI, S1 Methods). Figure 4a illustrates the overall contribution of Mn vs O redox as a function of Li content in Li$_x$Mn$_2$O$_2$F derived from the ab initio calculations (Figure S2 shows the change in the average oxidation states of all the component elements of Li$_x$Mn$_2$O$_2$F as Li is removed).

The results clearly show significant overlap between Mn and O redox couples with O redox activity starting from about x ≈ 1.5, well before all of the Mn has been oxidized and in accord with the XANES results; this is attributed to the disordered structure with a range of local ion environments (which we return to below). This behavior contrasts with the redox activity found in layered Li-rich ordered oxides (further details in the SI, S1 Methods). Figure 4a illustrates the overall contribution of Mn vs O redox as a function of Li content in Li$_x$Mn$_2$O$_2$F derived from the ab initio calculations (Figure S2 shows the change in the average oxidation states of all the component elements of Li$_x$Mn$_2$O$_2$F as Li is removed).

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The projected density of states for Li$_x$Mn$_2$O$_2$F (x = 2.0, 1.5, 0.75) (Figure S) also show the strong hybridization between the Mn 3d and O 2p states, with the energy of the O-2p states raised in the charged systems, which promotes O redox activity.

Regarding charge compensation on Li extraction, Mn undergoes oxidation from Mn$^{3+}$ at x = 2.0 toward Mn$^{4+}$ at x = 0.75, with no evidence of any change from octahedral to tetrahedral coordination. Our DFT structural analysis also indicate that tetrahedral Mn$^{3+}$ is not formed during delithiation of Li$_x$Mn$_2$O$_2$F which agrees with the *operando* Mn K-edge XANES and EXAFS results. As expected, the oxidation states of Li and F do not change on delithiation. When considering the overall Li deintercalation process (x = 2.00 to 0.75 in Figure 4a), Mn redox activity accounts for approximately 75% of the total capacity, while O redox accounts for the remaining 25%.

**Role of Local Environment.** To gain an atomistic understanding of the overlapping nature of Mn and O redox, the local coordination environment of each atom was investigated as a function of lithium content (x). Figure 4b shows the oxidation potentials of Mn and O as a function of their coordination environment in Li$_x$Mn$_2$O$_2$F; the value of x for Li$_x$Mn$_2$O$_2$F in the plot is related to the experimentally measured voltage at that state of charge (from Figure 1). Figure 4c compares the calculated change in the average oxidation state of Mn atoms that are initially coordinated by three or more F atoms with those that are coordinated by fewer than three F atoms. Figure 4d compares the change in the average oxidation state of O atoms with their coordination environments.

At a given Li content, Figure 4c indicates that Mn atoms with low coordination to F (e.g., Mn(O$_4$F$_3$)) are more oxidized than those Mn atoms coordinated to three or more F atoms (e.g., Mn(O$_4$F$_4$)) and helps to explain why O redox is predicted before all the Mn are oxidized. In other words, the redox potential of the Mn$^{3+}$/4+ couple is raised by an increasing number of F in its coordination shell, shown in Figure 4b. These results suggest that the substitution of F by Li leads to greater Mn redox overlap with oxygen.

With respect to the O environments, Figure 4d shows greater oxidation for O atoms with five Li nearest-neighbors (O(Li$_5$Mn$_5$)) at the fully lithiated state than those with four or three Li ions. This trend is consistent with that found by Seo et al.,11 who showed that the anion redox chemistry in a variety of Li-intercalation oxide cathodes, including LiNiO$_2$ and Li$_x$Mn$_2$O$_3$, is dependent on the anion nearest-neighbor coordination environment. They focused on how the presence of linear Li−O−Li configurations promote labile oxygen electrons in the adjoining 2p orbital that effectively pin them at a set energy above the bonding electrons. With the extension of this model, our results show a more continuous variation in the O-redox potential dependent on the number of Li coordinated to a given O$^{2−}$ ion. We note that recent computational screening work on layered oxide cathodes$^{61}$ report trends in O-redox activity associated with the electrostatic (Madelung) energy at oxygen sites. In general, our findings indicate that a greater number of coordinating Li can promote O oxidation at a lower potential (as shown in Figure 4b). This highlights the more general role of ionicity of the coordination environment around O in tuning its oxidation potential.

In summary, the effect of the local coordination environment has a major effect on the redox potentials of both Mn and O leading to increased competition between the Mn and O redox couples across the voltage range. The results presented here indicate that the O−Li environments in Li$_x$Mn$_2$O$_2$F encourage O oxidation at lower potentials than the typically observed 4.6 V plateaus for layered Li-rich oxide materials. Meanwhile, the presence of Mn−F bonds increases the voltage of Mn oxidation, leading to increased overlap between Mn and O redox processes.
Trapped Molecular O₂ in the Bulk Structure. Detailed analysis of the ab initio simulated structures of Li₅MnO₂F on Li removal indicate that the possibility of dimerization of oxidized O species is heavily dependent upon the local environment. At high degrees of Li deintercalation (high states of charge) corresponding to the Li-deficient composition Li₀.₇₅MnO₂F, some of the oxygen ions will be undercoordinated (fewer than three cations). The calculations indicate that such undercoordination results in molecular O₂ formation in the bulk structure (shown in Figure 6a); for this delithiated composition Li₀.₇₅MnO₂F, when the cations are arranged in such a way that the cation vacancies are clustered together, O₂ molecules form during energy minimization to give a structure with a significantly lower total energy (>320 meV/formula unit) than other configurations. Hence, bulk O₂ formation is triggered by driving the oxyfluoride electrode to high degrees of Li deintercalation, such that the local coordination around oxidized O ions is reduced to below three with local Li vacancies, e.g., OMn□₄. For instances where the oxidized O species are bonded to three or more cations, dimerization does not occur and the O–O separation remains at 2.3–2.8 Å. The formation of molecular O₂ inside the particles is only possible due to the loss of lithium ions at high states of Li deintercalation to create vacancy clusters that accommodate the O₂ molecules. From local analysis of the simulated relaxed structure of Li₀.₇₅MnO₂F, the diameter of the vacancy cluster is approximately 6.5 Å. The undercoordination of oxygen results in O 2p orbitals that no longer interact with any cations, and are at a high energy (shown in the pDOS plots, Figure 5c), which are where the oxygen holes are concentrated. Such undercoordinated oxygen will be unstable in the lattice and leads to molecular O₂ formation; this accords with our previous suggestions that undercoordinated oxygen more easily form O–O species, and now shown explicitly in disordered rocksalt oxyfluorides.

Figure 6a illustrates the local environment of the O₂ molecule with a calculated O–O bond length of 1.23 Å, directly comparable to that of molecular O₂, 1.21 Å, and in accord with RIXS results (Figure 7a) discussed in detail below. Analysis of the calculated charge density of O₂ within this cavity (Figure S4) indicate the electron density is heavily localized on the O₂ molecule as expected from the strong, covalent O=O bonding, with only weak chemical interaction with its neighboring environment. Once O₂ is formed, some Mn octahedra around the vacancy cluster may experience greater distortion to accommodate the defect. However, the amount of O₂ that is expected to form is small (around 5% of all the oxide ions). Hence, the impact on the distortion to Mn octahedra overall is likely to be limited, consistent with the slight increase in the pre-edge intensity of the Mn K-edge on charge.

In addition, ab initio molecular dynamics simulations of the trapped molecular O₂ in the oxyfluoride structure were performed for the first time. We stress that our focus here was to probe the degree of oxygen mobility within the vacancy cluster, rather than long-range diffusion. The variation in O–O bond distance and Mn–O₂ separation over the simulation time was analyzed (Figure 6b); we also examined oxygen mobility in the cavity through the mean square displacements (MSDs),
which were directly compared with MSDs from simulations of free gaseous O2 molecules within a volume of the same size (Figure 6c).

Two key features emerge. First, the trapped O2 molecule has an O−O bond length that remains directly comparable to that of gaseous molecular O2 (1.21 Å) whereas the nearest-neighbor Mn−O2 distance (mean value of 2.33 Å) is always longer than the Mn−O bond (2.0 Å) in the solid lattice, again confirming weak O2 interactions with the host lattice. Even when the O2 molecule makes very close approach to its nearest-neighbor Mn atom, the O−O distance remains around 1.2 Å.

Second, the increase in the mean square displacement with time for free gaseous O2 (Figure 6c) clearly indicates significant molecular diffusion as expected, whereas this is not the case for the O2 in the solid particle. The results therefore indicate that while the O−O bond length might be similar between the two, the trapped molecular O2 is different from free, gaseous O2 in exhibiting substantially reduced freedom of mobility, making it more characteristic of a solid-like environment (Figure 6c); this result is in line with recent solid state 17O NMR measurements for O2 in Li12Ni0.13Mn0.54Co0.13O2.26 The rigid trapping of O2 within the host lattice is also included showing a peak spacing consistent with a peroxide O2−2 moiety. In contrast, the trapped O2 molecule within a similar environment (Figure 6c) is found from our ab initio simulations. The similarity of the peak spacing to that of gaseous molecular O2 suggests that there is minimal bonding interaction with the host lattice in agreement with the ab initio simulations, which would accord with the expectation for strong localization of electron density in the heavily hybridized O=O bond. However, as has been noted previously,44,45,46 the excitation energy at which these O-redox RIXS features appear is slightly higher relative to O2 in the gas phase by about 0.5 eV. This is consistent with the O2 molecules being trapped in a solid-like environment.

To probe experimentally the nature of oxidized oxygen in charged Li2MnO2F, O K-edge RIXS was performed at a higher resolution than previously achieved for this material.36 Previous RIXS data for charged electrodes revealed a more prominent elastic peak when exciting at ~531 eV and a new energy loss feature, attributed to the formation of localized electron holes on oxygen. The new high-resolution data in Figure 7a show that the broad elastic peak can in fact be resolved into a progression of sharp peaks, as also observed for the layered O-redox material Li1.2Ni0.13Mn0.54Co0.13O2.26 This peak progression arises from the molecular vibrations of an O=O diatomic with well-defined frequency matching that of molecular O2 (1550 cm−1) and clearly distinguishable from superoxide O2− and peroxide O22− which have vibrational frequencies of around 1100 and 790 cm−1, respectively.66 The peak spacing decreases linearly with increasing energy loss (Birge–Sponer plot Figure 7b) consistent with an anharmonically oscillating diatomic. A reference RIXS spectrum for Li2O2 is also included showing a peak spacing consistent with a peroxide O2−2 as expected, demonstrating the ability of RIXS to distinguish different O−O bond orders.

The other energy loss feature at 8 eV also belongs to molecular O2 and can be assigned to the filled π molecular orbitals. These results show that the localized electron hole states appearing at 531 eV reside on O2 molecules, which, since the RIXS experiment was performed under ultrahigh vacuum conditions, must be trapped within the bulk of the cathode as found from our ab initio simulations. The similarity of the peak spacing to that of gaseous molecular O2 suggests that there is minimal bonding interaction with the host lattice in agreement with the ab initio simulations, which would accord with the expectation for strong localization of electron density in the heavily hybridized O=O bond. However, as has been noted previously,44,45,46 the excitation energy at which these O-redox RIXS features appear is slightly higher relative to O2 in the gas phase by about 0.5 eV. This is consistent with the O2 molecules being trapped in a solid-like environment.

On discharge, the signal is no longer evident indicating that the O2 that formed in the fully charged samples is no longer present in the bulk material. The lack of O2 gas evolution at
the surface during discharge from differential electrochemical mass spectrometry measurements, which, coupled with the disappearance of the O₂ signal from the RIXS data, indicates that the trapped O₂ is reduced back to O²⁻. It is worth mentioning that O²⁻ ions and molecular O₂ are the most stable forms of oxygen.

To investigate the possibility of the RIXS features being beam-induced, we undertook low temperature measurements (20 K) to suppress sample heating by the beam. The data, Figure S5, show negligible difference between the spectra indicating no such effect. Taken together, the full outgassing of the electrode under UHV conditions and the reversible reduction of O₂ rule out O₂ being trapped anywhere other than in the particle bulk, where it can still be reversibly reincorporated back into the structure as O²⁻. Overall, the reversible O-redox process involves O²⁻ being oxidized to form bulk molecular O₂ on charge, followed by its reduction on discharge to reform O²⁻.

The observation of molecular O₂ in layered O-redox cathode materials and here, for the first time, in disordered rocksalts, suggests the two systems share the same O-redox mechanism, (i.e., 2O²⁻ ↔ O₂ + 4e⁻). However, Li₄MnO₂F does not exhibit the commonly observed O-redox charging plateau at 4.6 V vs Li⁺/Li, nor such large first cycle voltage hysteresis (Figure 1). Both phenomena have been recently linked with the irreversible loss of highly ordered honeycomb superstructures belonging to the layered cathodes. In Li₁₂Ni₉Se₅Co₀₁₃O₁₂, all oxide ions in the honeycomb lattice will be coordinated by at least two transition metal (TM) ions (O(Li₄TM₃)) which, during charge, are oxidized to O²⁻ at a high potential of 4.6 V. However, this honeycomb arrangement of O²⁻ is highly unstable. In-plane TM migration to form vacancy clusters occurs, causing some O to become coordinated by fewer than two TM ions which then dimerize to form stable O₂ molecules. On discharge, these vacancy clusters are repopulated by Li leading to O(Li₄TM) and O(Li₄) configurations which remain in the structure explaining the lack of further voltage plateaus. In contrast, Li₄MnO₂F is already intrinsically disordered in the pristine state, possessing a range of coordination environments including some O(Li₄Mn) and O(Li₄) regions. Therefore, O⁻ redox can occur without such severe structural rearrangement and hence less pronounced voltage hysteresis. After the first cycle, the load curves for both compounds (Figure S2) exhibit a similar degree of voltage hysteresis in line with the presence of preformed sites for O₂ formation in both materials after discharging of the TM ions within the TM layer of Li₁₂Co₀₁₃Ni₀₁₃Mn₀₂₅₄O₁₂.

Our observations on Li₄MnO₂F are unexpected in the context of previous work, which cannot be simply translated from ordered Li-rich layered compounds to disordered rocksalt oxyfluorides, and are important in future strategies to develop new high capacity cathodes.

### Conclusion

The oxygen redox mechanism in the disordered rocksalt cathode, Li₄MnO₂F, involves the formation of molecular O₂ trapped inside the bulk structure of the charged material, which is reversibly reduced to O²⁻ on discharge. Combined RIXS and ab initio simulation studies show that molecular O₂ is held within vacancy clusters in the structure. Bulk O₂ formation is triggered by driving the disordered rocksalt oxyfluoride structure to high degrees of Li deintercalation, such that the local coordination number decreases around oxidized O ions with local Li vacancies. The trapped molecular O₂ also exhibits minimal mobility unlike free gaseous O₂, making it more characteristic of a solid-like environment. This rigid trapping of O₂ within close proximity to cation centers also helps to rationalize how it could be reduced to O²⁻ with ease on discharge.

The Mn redox process occurs between 3+ and 4+, with no evidence of tetrahedral Mn³⁺ or Mn⁷⁺. We show that the significant overlap between the Mn and O redox couples is determined by the different local coordination environments in the disordered oxyfluoride structure: more ionic Li-rich O environments (e.g., O(Li₄Mn)) are oxidized at lower voltages than the typically observed 4.6 V plateaus for layered Li-rich oxides, whereas F-rich Mn coordination (e.g., Mn(F₃O₃)) increases the voltage for Mn oxidation, leading to the overlapping nature of the Mn and O redox processes.

Since Li₂MnO₂F already possesses an intrinsically disordered structure, it avoids the extensive structural rearrangement observed in layered honeycomb cathodes resulting in reduced voltage hysteresis on the first charge/discharge cycle. This work advances our understanding of fundamental redox mechanisms in Li-rich disordered rocksalts and highlights their promise as more structurally stable oxygen-redox cathodes.

### Associated Content

#### Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/jacs.0c10270.

Experimental and computational methods, EXAFS fitting, Buckingham potentials, DFT-computed structural data, change in oxidation states and magnetic moments, charge density of the O₂ and local environments, reduced voltage hysteresis on the first charge/discharge cycle. This work advances our understanding of fundamental redox mechanisms in Li-rich disordered rocksalts and highlights their promise as more structurally stable oxygen-redox cathodes.

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