Discrete Li-occupation versus pseudo-continuous Na-occupation and their relationship with structural change behaviors in Fe₂(MoO₄)₃

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The key factors governing the single-phase or multi-phase structural change behaviors during the intercalation/deintercalation of guest ions have not been well studied and understood yet. Through systematic studies of orthorhombic Fe₂(MoO₄)₃ electrode, two distinct guest ion occupation paths, namely discrete one for Li and pseudo-continuous one for Na, as well as their relationship with single-phase and two-phase modes for Na⁺ and Li⁺, respectively during the intercalation/deintercalation process have been demonstrated. For the first time, the direct atomic-scale observation of biphasic domains (discrete occupation) in partially lithiated Fe₂(MoO₄)₃ and the one by one Na occupation (pseudo-continuous occupation) at 8d sites in partially sodiated Fe₂(MoO₄)₃ are obtained during the discharge processes of Li/Fe₂(MoO₄)₃ and Na/Fe₂(MoO₄)₃ cells respectively. Our combined experimental and theoretical studies bring the new insights for the research and development of intercalation compounds as electrode materials for secondary batteries.

Intercalation compounds as energy storage materials have been extensively studied for secondary batteries. All of these intercalation materials for the secondary batteries allow the guest ions to move in and out without significant damage of their host frameworks. The composition variations in intercalation compounds during the intercalation/deintercalation of guest ions are often accompanied by structural changes. Most of intercalation compounds (Supplementary Table 1) fall into the single-phase solid solution mode or the two-phase transformation mode or three-phase separation mode as a result of the composition variations in the certain concentration range of guest ions. For example, the layered LiₓCoO₂ (0.5 < x ≤ 0.75) deintercalates/intercalates Li⁺ via a single-phase process, while the layered NaₓCoO₂ (0.5 ≤ x ≤ 1) shows various single-phase or two-phase domains depending on Na⁺ concentration. The olivine-type LiFePO₄ exhibits a two-phase transformation reaction (LiFePO₄/FePO₄) by undergoing a phase interface propagation based on steady-state results, and some non-equilibrium single-phase solid solution processes as predicted by ab initio calculations and confirmed by in situ diffraction experiments. The various phase transformation mechanisms of Li⁺ ions in LiFePO₄/FePO₄ are revealed to be dependent on the rate. Understanding these structural change mechanisms during the intercalation/deintercalation process is very important for the development of high energy density and long cycle-life batteries. Here, through the systematic studies of an intercalation compound of Fe₂(MoO₄)₃, we report the single-phase structural change behavior for Na⁺ intercalation/deintercalation, but two-phase reaction mode for Li⁺ intercalation/deintercalation. The framework structure remains unchanged in the entire concentration range of Na⁺ or Li⁺. More interestingly, such single-phase and two-phase reactions are closely related to the guest ion occupation paths during intercalation/deintercalation, as clearly demonstrated by the aberration-corrected scanning transmission electron microscopy (STEM) results. These results provide new insights into the origin of structural changes in the guest-host material systems.
Fe₂(MoO₄)₃ is one of the most promising cathode materials for rechargeable lithium/sodium battery as an environment friendly energy storage material from the viewpoints of the inexpensive and non-toxic of iron. From X-ray diffraction studies, it is known that Fe₂(MoO₄)₃ has two types of crystal structures: low temperature monoclinic structure and high temperature orthorhombic structure. Although there have been several reports on the monoclinic Fe₂(MoO₄)₃ as the cathode materials for lithium (or sodium) battery, the Li⁺ (or Na⁺) intercalation/deintercalation mechanisms remain unclear or contradict with each other. For example, some of literatures indicated a two-phase reaction during the intercalation/deintercalation of both Na⁺ and Li⁺ into the monoclinic Fe₂(MoO₄)₃, whereas single-phase solid solution reaction of NaₓFe₂(MoO₄)₃ (0 < x < 2) was also observed. This may be due to the structural complex or thermodynamic unfavorableness of monoclinic Fe₂(MoO₄)₃. In this work, orthorhombic Fe₂(MoO₄)₃ was studied as the cathode material for lithium and sodium batteries. Its electrochemical properties and structural change behaviors during charge and discharge processes are investigated by synchrotron based X-ray diffraction (XRD), X-ray absorption spectroscopy (XAS), aberration-corrected scanning transmission electron microscopy (STEM) and first-principles thermodynamic calculations. The discrete Li occupation path and pseudo-continuous Na occupation path in Fe₂(MoO₄)₃ during intercalation/deintercalation process and their relationship with the two-phase and single-phase reactions are proposed.

**Results**

**Electrochemical characterization.** The intercalation/deintercalation behaviors of alkali (A = Li or Na) metal ions in the Fe₂(MoO₄)₃ were examined in Li and Na cells in Fig. 1. As shown in Fig. 1a and b, the initial discharge capacity of 89.5 mAh g⁻¹ can be obtained for both Li and Na cells at the current rate of C/20. This value corresponds to the intercalation number of 2.0 Li or Na per Fe₂(MoO₄)₃ unit. The discharge/charge curves in the lithium cell (Fig. 1a) show a flat plateau at about 3.0 V vs. Li⁺/Li during the discharging process and a flat plateau at about 3.02 V vs. Li⁺/Li during the charging process in a large range. By contrast, the discharge/charge curves of Na/Fe₂(MoO₄)₃ cell show a slope type in the voltage range of 2.5 to 2.7 V vs. Na⁺/Na in Fig. 1b. The capacity fades of Li/Fe₂(MoO₄)₃ and Na/Fe₂(MoO₄)₃ cells during the first 20 cycles are about 0.3% and 0.9% per cycle, respectively, indicating a better capacity retention of Li/Fe₂(MoO₄)₃ cell than that of Na/Fe₂(MoO₄)₃ cell. The discharge and charge curves of Li/Fe₂(MoO₄)₃ cell at a current density of C/5 shown in Supplementary Fig. 1 indicates a good cyclic performance up to 400 cycles with a capacity fading less than 0.02% per cycle. The shapes of one pairs of cathodic peak and anodic peak in the cyclic voltammogram (CV) curves of Li/Fe₂(MoO₄)₃ cell exhibit the feature of mirror-symmetry as shown in Fig. 1c. Such an appearance of the peak is related to the typical diffusion and reaction kinetics at around half-discharging and half-charging processes. For the CV curves of Na/Fe₂(MoO₄)₃ cell, two couples of reduction/oxidation peaks at around 2.65/2.73 V and 2.54/2.62 V (Fig. 1d) are in good agreement with the discharge/charge curves.
**Ex situ XRD patterns.** To investigate the structural evolutions of Fe₂(MoO₄)₃ during Li⁺ and Na⁺ intercalation/deintercalation, a series of synchrotron based XRD patterns were collected at different charge/discharge states. As shown in Fig. 2a for LiₓFe₂(MoO₄)₃ (x = 0.0, 0.5, 1.0, 1.5 and 2.0) during the first discharge-charge cycle in a Li/Fe₂(MoO₄)₃ cell. All diffraction peaks of the pristine and fully lithiated Fe₂(MoO₄)₃ (after discharging to 2.5 V) can be well indexed to Fe₂(MoO₄)₃ (JCPDS card No. 852278) and Li₂Fe₂(MoO₄)₃ (JCPDS card No. 841001) with the same orthorhombic structure, respectively. During the discharge process, the intensity of the major peaks (111), (211), (012), (021), (310), (212) and (231) representing Fe₂(MoO₄)₃ decreased gradually and finally disappeared when the 2.5 V discharge limit was reached. At the meantime, a new Li₂Fe₂(MoO₄)₃ phase was formed as observed through the appearing and growing intensity of a new set of (111), (211), (021), (310), (212) and (231) peaks at lower 2θ angles relative to those original ones. No peak shifts are observed for both Fe₂(MoO₄)₃ and Li₂Fe₂(MoO₄)₃ phases in the entire discharge process. During the recharge process, the peak evolution is exactly in the opposite way to the discharge process. After the initial cycle, the Fe₂(MoO₄)₃ phase can be fully recovered, indicating an excellent structural reversibility. The coexistence of both Fe₂(MoO₄)₃ and Li₂Fe₂(MoO₄)₃ phases with changing ratio, demonstrates a typical two-phase reaction during the discharge and charge process in the lithium cell.

Fig. 2b shows the XRD patterns of NaₓFe₂(MoO₄)₃ (x = 0.0, 0.5, 1.0, 1.5 and 2.0) during the first discharge-charge cycle in a Na/Fe₂(MoO₄)₃ cell. Interestingly, a completely different structural change behavior other than that in the lithium cell was observed. No new set of peaks, but only peak shifts were observed throughout the entire discharge/charge process. During the discharge process, major peaks (111), (211), (102), (021), (310), (212), (022), (130) and (231) all gradually moved toward lower 2θ angles with increasing x from 0.0 to 2.0 in NaₓFe₂(MoO₄)₃. During the recharge process, all of these peaks reversibly moved back to their original positions with decreasing x from 2.0 to 0.0. The reversible peak shifts are attributed to the continuous lattice expansion and contraction during the discharge and charge respectively. This result demonstrates a typical single-phase (solid-solution) reaction during the discharge and charge process in the sodium cell.

**DFT simulations.** It is quite interesting to note that the same Fe₂(MoO₄)₃ orthorhombic structure shows such different structural change behaviors during Li⁺ and Na⁺ intercalation/deintercalation process. The crystal structure and thermodynamics of the orthorhombic Fe₂(MoO₄)₃ during guest alkali ion insertion were further studied. As shown in Fig. 3a and b, the crystal structure of the orthorhombic Fe₂(MoO₄)₃ with a space group of Pbcn is composed of MoO₄ tetrahedra sharing all four corners with FeO₆ octahedra and...
FeO₆ octahedra sharing all six corners with MoO₄ tetrahedra. Such an open three dimensional framework structure is suitable for guest (A = Li and Na) ions accommodation and diffusion. After the alkali ion intercalation, the crystal structure of Li₂Fe₂(MoO₄)₃ is isostructural to Na₂Fe₂(MoO₄)₃ based on ab initio calculations. A₂Fe₂(MoO₄)₃ (A = Li, Na) have the same orthorhombic structure as Fe₂(MoO₄)₃. The eight A ions occupy the 8d tetrahedra interstitial sites in a unit cell. Their lattice parameters and unit cell volumes obtained from Le Bail fitting of the XRD patterns are listed in Supplementary Table 2. In order to understand the thermodynamic origin about the intercalation/deintercalation behaviors of A ions in the Fe₂(MoO₄)₃, the formation energy (Eᵣ) of AₓFe₂(MoO₄)₃ with respect to Fe₂(MoO₄)₃ and A₂Fe₂(MoO₄)₃ via:

\[ Eᵣ = E[AₓFe₂(MoO₄)₃] - \left[ \frac{(2-x)/2}E[Fe₂(MoO₄)₃]ight] - \frac{x/2}E[A₂Fe₂(MoO₄)₃] \]

where three energy terms represent the total energies of per AₓFe₂(MoO₄)₃, Fe₂(MoO₄)₃ and A₂Fe₂(MoO₄)₃ formula unit (f.u.), respectively. Here a completely random A/vacancy distribution at 8d site in AₓFe₂(MoO₄)₃ is given, the configurational entropy, \( S_{\text{con}} = k_B \left[ (x/2)\ln(x/2) + (1 - x/2)\ln(1 - x/2) \right] \), is included in the total

![Figure 3 | DFT simulations. Crystal structures of (a) Fe₂(MoO₄)₃ and (b) AₓFe₂(MoO₄)₃. (c) Formation energies of AₓFe₂(MoO₄)₃ at different x values (A = Li or Na). Optimized lattice parameters of (d) LiₓFe₂(MoO₄)₃ and (e) NaₓFe₂(MoO₄)₃ at different x values.](image-url)
energy of $A_xFe_2(MoO_4)_3$. To obtain the most energetically favorable configuration, all possible $A$ occupancies were considered. There are 248 possibilities in all, namely, $\sum_{i=0}^8 C_i^5$, where $C_i$ ($i = 0, 1, 2, 3, 4, 5, 6, 7, 8$) stands for possible arrangements of different numbers of $A$ in a unit cell ($Z = 4$). Only the calculated lowest formation energies of $A_xFe_2(MoO_4)_3$ at different $x(x = i/Z)$ values are shown in Fig. 3c. It can be found that the single-phase solid solution of $Li_xFe_2(MoO_4)_3$ is not thermodynamically favorable with the positive formation energies of $\sim 30 \text{ meV/\text{f.u.}}$, which implies the phase separation of $Fe_3(MoO_4)_3$ and $Li_xFe_2(MoO_4)_3$. Conversely, $NaFe_2(MoO_4)_3$ exhibits the negative formation energies ($-12 \sim -40 \text{ meV/\text{f.u.}}$), which is representative of the single-phase solid solution instead of the two-phase separation. Dependences of calculated lattice parameters $(a, b$ and $c)$ on $x$ in $A_xFe_2(MoO_4)_3$ are shown in Fig. 3d and e. Results indicate that the variation of the lattice parameters with $x$ in $Li_xFe_2(MoO_4)_3$ disobeys Vegard’s law, thus invalidating the solid solution reaction for the lithium ion intercalation process. Nonetheless, the Vegard’s law is true for the case of $Na_xFe_2(MoO_4)_3$, indicating the single phase solid solution reaction for the sodium ion intercalation process. This suggests that the first-principles thermodynamical calculations draw the identical conclusion as the ex situ XRD patterns in Figure 2.

Considering an interface between $Li_xFe_2(MoO_4)_3$ and $Fe_2(MoO_4)_3$, spherical aberration-corrected STEM imaging was employed to obtain a direct observation at the atomic resolution. Considering the relative less structural stability of lithium-containing compound based on our own experience and reported works$^{23}$, we decreased the probe current to about 20 pA and the pixel dwell time to 10 ms, to avoid the electron beam damage or phase transformation during STEM analysis. The schematic drawings for $Fe_2(MoO_4)_3$ and $Li_xFe_2(MoO_4)_3$ projected along the [001] direction were given in Fig. 4a. All annular-bright-field (ABF) images of partially lithiated $Fe_2(MoO_4)_3$ at the 1/4, 1/2 and 3/4 discharge states were examined and the same two different regions could be observed in these ABF images. The typical ABF image of partially lithiated $Fe_2(MoO_4)_3$ at the 1/2 discharge state is shown in Fig. 4b, in which one boundary is marked with a red dash line between regions 1 and 2. Unfortunately, lithium ions which are supposed to occupy both sides of the shoulder of ellipse cannot be obviously visualized because of the wide atomic number gap between Li and Mo. Therefore, line profiles (Fig. 4c) were acquired across two regions through the purple line in the ABF image (Fig. 4b) to confirm the lithium contrast with respect to oxygen. The corresponding purple lines in the $Fe_2(MoO_4)_3$ and $Li_xFe_2(MoO_4)_3$ unit structures are also shown in Fig. 4a. The Li and O positions are displayed as arrows in the line profile (Fig. 4c). The corresponding line profile shows two distinctly different periodic characteristics in these two regions.

STEM imaging. To further confirm the two-phases and single-phase during the lithiation and sodiation processes of $Fe_2(MoO_4)_3$, STEM imaging of partially lithiated $Fe_2(MoO_4)_3$ were carefully examined, but only one uniform region is found in all of these images. As shown in Fig. 5a, b and c, the repeat unit can be clearly visualized (shown in the green ellipses).
in these ABF images, which has the identical cage structure with Fe_2(MoO_4)_3. In the ABF images of sodiated Fe_2(MoO_4)_3 at the 1/4, 1/2 and 3/4 discharged states, four spots representing Mo 8d sites on the shoulder of ellipse do not exhibit the same blackness as that in the pristine Fe_2(MoO_4)_3 (Supplementary Fig. 3). Therefore, some intriguing contrasts of four black spots representing Mo 8d sites on the shoulder of ellipse between the pristine and partially sodiated Fe_2(MoO_4)_3 in ABF images could provide the information of Na occupancy in Fe_2(MoO_4)_3. Interestingly, one spot marked by yellow circle on the green ellipse at the 1/4 discharge states, in which other three dark spots were marked by white circles (Fig. 5a). According to the atomic occupancies of simulated ABF image of Na_0.5Fe_2(MoO_4)_3 based on the energetically most favorable configuration (Supplementary Fig. 5), one Na 8d site (shown as the red arrow in Fig. 5d) resides in the vicinity of one of four Mo 8d sites on the shoulder of ellipse in the construction of Na_0.5Fe_2(MoO_4)_3, resulting in that one of four spots representing Mo 8d sites at every repeated unit structure exhibits more blackness than any other three spots. The Na columns as red dash lines with the periodicity and homogeneity can be clearly observed in the ABF image (Fig. 5a). After the depth of discharge to 1/2, two spots representing Mo 8d sites (yellow circles) on both sides of the shoulder of ellipses with asymmetric distribution are found to be blacker than other two spots (white circles) as shown in Fig. 5b, which perfectly coincides with simulated ABF image of NaFe_2(MoO_4)_3 structure based on first principle calculations (Supplementary Fig. 5), in which two Na 8d sites (shown as the red arrow in Fig. 5e) reside in the vicinity of the two of four Mo 8d sites with Na ordering on the shoulder of ellipse. When the Na/Fe_2(MoO_4)_3 cell is discharged to 3/4 of the fully discharged state, the ABF image is different from one of any other ABF images for the pristine and partially sodiated Fe_2(MoO_4)_3 at the 1/4 and 1/2 discharged states. In four spots representing Mo 8d sites on the shoulder of ellipse, three spots (yellow circles) are blacker than other one spot (white circles) in the Fig. 5c. They represent Na-occupied or Na-unoccupied in Na_1.5Fe_2(MoO_4)_3 structure based on first principle calculations (Supplementary Fig. 5) and the simulated ABF image (Fig. 5f). Na columns as red dash lines with the periodicity and homogeneity are also observed in all region of these images, indicating the feature of single domain. In the other word, ABF images at various discharge states not only characterize the solid solution mode of Na” intercalation into Fe_2(MoO_4)_3, but also provide the preferred

**Figure 5** | The STEM images of partially sodiated Fe_2(MoO_4)_3. (a) at the 1/4, (b) at the 1/2 and (c) at the 3/4 discharged states viewed along [001] projection and their corresponding simulated ABF images (d), (e) and (f). Repeated unit structures are labeled by green ellipse. The Na occupied sites are marked by yellow circles and the unoccupied sites are marked by white circles in (a), (b) and (c).

**Figure 6** | Comparison between sodiation and lithiation process in Fe_2(MoO_4)_3. Schematic diagrams of “discrete occupation” and “pseudo-continuous occupation” during Li and Na ions intercalation into Fe_2(MoO_4)_3. Solid red circles and dash green ellipses stand for Li” (or Na”) and Fe_2(MoO_4)_3 frameworks, respectively.
occupancy patterns of Na⁺ in Fe₂(MoO₄)₃ during the electrochemical intercalation process.

Discussion

The high resolution atomic images provide a direct evidence on the different atomic occupancies of Li and Na in Fe₂(MoO₄)₃ from the repeating Fe₂(MoO₄)₃ unit of ellipse along the [001] projection in the partially lithiated and sodiated Fe₂(MoO₄)₃. The existence of biphasic domains and the lithium occupancy near four of the Mo 8d sites taking a discrete manor (all empty or all occupied) during the discharge process of Li₄Fe₂(MoO₄)₃ cell are confirmed. The new formed LiₓFe₂(MoO₄)₃ phase is revealed with four Li 8d sites occupying both sides of ellipse configuration in symmetric distribution apart from the original Fe₂(MoO₄)₃ phase. In contrast, the single-phase domains with various compositions were observed at the different discharge states of NaₓFe₂(MoO₄)₃ cell. Four Na 8d vacancies in the ellipse configuration are occupied one by one during the discharge process. According to the structural characteristics of Fe₂(MoO₄)₃, its unit of ellipse along the [001] projection can hold four A atoms distributed on both sides of ellipse configuration. The electrochemical intercalated Li ions capture all four sites simultaneously to form LiₓFe₂(MoO₄)₃ at the different discharge states. This feature can be depicted as a “discrete-occupation” path (shown in Fig. 6) that defines that the Li occupation for the four available 8a sites is either all empty or all occupied, in forming the LiₓFe₂(MoO₄)₃. The amount of LiₓFe₂(MoO₄)₃ phase increases with increasing Li content at the expense of Fe₂(MoO₄)₃ phase. In contrast, Na ion intercalation in Fe₂(MoO₄)₃ can be described as a “pseudo-continuous-occupation” path, where Na ions progressively occupy four of the 8d sites in the ellipse unit of Fe₂(MoO₄)₃ one after another. The holistic occupation of Na ions in Fe₂(MoO₄)₃ results in the formation of a series of pseudo continued NaₓFe₂(MoO₄)₃ solid solutions in which x value increases from 0.0 to 2.0. Apparently, the different occupation behaviors of Li and Na ions in Fe₂(MoO₄)₃ lead to two different structural change modes. Based on the ex situ XRD and electrochemical characterization results, the deintercalation of A ions from AₓFe₂(MoO₄)₃ is reversible during the charge process.

The single-phase and two-phase modes of Fe₂(MoO₄)₃ with the intercalation/deintercalation of Li and Na ions are revealed on atomic-scale. The orthorhombic Fe₂(MoO₄)₃ electrode is a quite interesting material for studying the key factors governing the solid-solution and two-phase reactions during the ion intercalation/deintercalation process. Based on the calculated lattice information in Supplementary Table 2, the unit-cell volume changes of Fe₂(MoO₄)₃ after the full lithiation and sodiation are ca. 2.96% and 4.72%, respectively. These values are significantly smaller than that of the LiFePO₄ (6.8%)26. In addition, the calculated diffusion constants of Li⁺ and Na⁺ are 3.45 × 10⁻⁶ and 4.94 × 10⁻⁷ cm²/s, respectively (Supplementary Fig. 6 and Supplementary Table 3). The smaller volume change and faster ion diffusion in the Li₂Fe₂(MoO₄)₃ cell contribute to its better cycle performance as compared with that in the Na₂Fe₂(MoO₄)₃ cell. Thus, Fe₂(MoO₄)₃ electrode is also a good example to bridge the understanding of the relationship between the electrochemical properties and the intercalation/deintercalation process.

In summary, the two-phase and single-phase mechanisms were revealed during the intercalation/deintercalation of Li and Na ions into Fe₂(MoO₄)₃. The “discrete occupation” and “pseudo-continuous” were proposed to describe the distinctly different occupation paths of Li and Na ions into Fe₂(MoO₄)₃. The first-principle thermodynamic calculations and direct atomic-scale observation by STEM provide further insight on the intercalation/deintercalation process. Most importantly, the discrete occupation path for Li and pseudo-continuous occupation path for Na, and their relationship with two-phase reaction for Li and single-phase reaction for Na, respectively during the intercalation/deintercalation process in Fe₂(MoO₄)₃ may very well be extended to the knowledge of other intercalation compounds. Noticeably, the present results were made at a relatively low rate of 1/20 C, which is close to the equilibrium state. The above mentioned intercalation/deintercalation reactions of Li and Na ions into Fe₂(MoO₄)₃ should be further investigated at the high rates which is far away from the equilibrium state in the future work. All in all, our experimental and theoretical studies could provide very valuable information for the research and development of intercalation compounds as electrode materials for secondary batteries.

Methods

Sample Preparation and Characterization. Orthorhombic Fe₂(MoO₄)₃ microcrystals were synthesized by a simple hydrothermal method25 using Fe(NO₃)₃·9H₂O and Na₂MoO₄·2H₂O as precursor. Scanning electron microscopy (SEM, Cambridge S-360) was employed to study the morphology and particle size of Fe₂(MoO₄)₃ (Supplementary Fig. 7). Powder X-ray diffraction (XRD) patterns (Supplementary Fig. 8) were collected at beamline X14A of the National Synchrotron Light Source (NSLS) at Brookhaven National Laboratory using a linear position sensitive silicon detector. The wavelength used was 0.774 Å. X-ray absorption spectroscopy (XAS) was performed at beamline X19A of the NSLS. Fe K-edge XAS was collected in transmission mode (Supplementary Fig. 9). The XAS data was processed using Athena22.

Electrochemical measurements. A slurry of 80 wt % Fe₂(MoO₄)₃, 10 wt % carbon black, and 10 wt % polyvinylidenefluoride (PVDF, Sigma-Aldrich) dispersed in N-methylpyrrolidone (NMP, Sigma-Aldrich) was prepared and cast on aluminum foil. The electrodes were dried at 20°C, and were punched to small circular pieces of diameter of 14 mm. Electrochemical cells were assembled into coin cells in an Ar-filled glovebox (M Braun, Germany). Sodium pieces and lithium pieces were used as a counter electrode for sodium and lithium batteries, respectively. The electrolytes consisted of 1 M NaPF₆ (Alfa-Aesar) and LiPF₆ (Alfa-Aesar) in a nonaquaous solution of ethylene carbonate (EC, Alfa-Aesar) and dimethyl carbonate (DMC, Alfa-Aesar) with a volume ratio of 1:1. Galvanostatic discharge-charge measurements were carried out at room temperature with a Land CT 2001A battery test system. The current densities and capacities of electrodes were calculated based on the weight of active materials. On the basis of two electrons transfer in the Fe₂(MoO₄)₃, 1C was calculated to correspond to about 9.10 mA g⁻¹.

DFT simulations. All the first-principle total energy calculations were performed using a plane-wave basis set and the projector-augmented wave (PAW) method22 as implemented in the Vienna ab initio simulation package (VASP)27. Generalized gradient approximation (GGA) in the parametrization of Perdew, Burke, and Ernzerhof (PBE)28 pseudopotential was used to describe the exchange–correlation potential and a Hubbard-type correction U was taken into account due to the strongly correlated nature of the Fe 3d electrons29. Referring to the DFT calculations on LiFePO₄ and FePO₄ materials, the effective U value was set to 4.3 eV27. A kinetic energy cutoff of 550 eV was used in all calculations. Geometry optimizations were performed by using a conjugate gradient minimization until all the forces acting on ions were less than 0.02 eV/Å per atom. (1 × 2 × 2) and (1 × 1 × 2) Monkhorst-Pack grids were used for the bulk and interface supercells. Activation energies for Na⁺ and Li⁺ ion diffusion in Fe₂(MoO₄)₃ were calculated using the nudged-elastic-band (NEB) method30 using seven images and two endpoint structures, which is a reliable method to search the minimum-energy path (MEP) when the initial and final states are known. An interpolated chain of configurations (images) between the initial and final positions are connected by springs and relaxed simultaneously to the minimum-energy path. The structure diagrams were drawn by VESTA22.

STEM imaging. STEM samples were made by sonication of the discharged cathode films in anhydrous dimethyl carbonate inside an argon-filled glove box, and sealed in airight bottles before immediately transfer into the STEM column. A JEM-ARM200F STEM operated at 200 kV and equipped with double aberration-correctors for both probe-forming and imaging lenses was used to perform high-angle annular-dark-field (HAADF) and ABF imaging. The attainable spatial resolution of the microscope is 78 pm at the incident semi-angle of 25 mrad. To observe Li directly using ABF collection geometry, the acceptance semi-angle in this study was fixed between 12 and 25 mrad. The STEM ABF and HAADF images were taken simultaneously at the optimal defocus value of the HAADF imaging condition, which was more defocused than the optimal ABF imaging condition on this instrument. Thus, the contrast in the ABF image is reversed with the bright area corresponding to the atomic positions11.

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Author contributions

Z.-W.F. and X.-Q.H. planned the study and supervised all aspects of the research. J.-L.Y. and Z.-W.F. wrote the manuscript. J.-L.Y. and X.-Q.H. tested the electrochemical performance. Y.-N.Z. and X.-Q.H. performed the computations were performed on the high performance computing platform of Shanghai University.

Additional information

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