


Cite this: *RSC Adv.*, 2024, 14, 26516

# First-principles calculations to investigate the impact of fluorine doping on electrochemical properties of Li-rich $\text{Li}_2\text{MnO}_3$ layered cathode materials

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Li-rich layered oxides are promising candidates for high-capacity Li-ion battery cathode materials. In this study, we employ first-principles calculations to investigate the effect of F doping on Li-rich  $\text{Li}_2\text{MnO}_3$  layered cathode materials. Our findings reveal that both  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  exhibit significant volume changes (greater than 10%) during deep delithiation, which could hinder the cycling of more Li ions from these two materials. For  $\text{Li}_2\text{MnO}_3$ , it is observed that oxygen ions lose electrons to compensate for charge during the delithiation process, leading to a relatively high voltage plateau. After F doping, oxidation occurs in both the cationic (Mn) and anionic (O) components, resulting in a lower voltage plateau at the beginning of the charge, which can be attributed to the oxidation of  $\text{Mn}^{3+}$  to  $\text{Mn}^{4+}$ . Additionally, F doping can somewhat suppress the release of oxygen in  $\text{Li}_2\text{MnO}_3$ , improving the stability of anionic oxidation. However, the increase of the activation barriers for Li diffusion can be observed after F doping, due to stronger electrostatic interactions between  $\text{F}^-$  and  $\text{Li}^+$ , which adversely affects the cycling kinetics of  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ . This study enhances our understanding of the impact of F doping in  $\text{Li}_2\text{MnO}_3$  based on theoretical calculations.

Received 8th July 2024  
Accepted 12th August 2024  
DOI: 10.1039/d4ra04925j  
rsc.li/rsc-advances

## 1. Introduction

Li-ion batteries (LIBs) are extensively used in portable devices for their long cycle life, safety and high energy density.<sup>1</sup> However, entering new areas, like electric vehicles market, demands LIBs with higher energy density.<sup>2–4</sup> Cathode materials, a key component of LIBs, are seen as a significant bottleneck in developing high-energy-density LIBs.<sup>5</sup> Current transitional cathodes, such as  $\text{LiCoO}_2$  and  $\text{LiFePO}_4$ , are approaching their theoretical limits. Consequently, the need for new cathode materials with higher energy density has become more urgent than ever. Recently, Li-rich oxides have emerged as a promising

solution to achieve this goal.<sup>6,7</sup> Thanks to cumulative cationic and anionic redox reactions, these materials can achieve capacities exceeding  $250 \text{ mA h g}^{-1}$ .<sup>8–10</sup> Nevertheless, these materials have drawbacks such as significant voltage decay,<sup>10,11</sup> voltage hysteresis,<sup>12,13</sup> poor rate capability,<sup>14</sup> and capacity loss<sup>15</sup> during cycling, due to the structural transformation of host materials during the oxygen redox.<sup>10,12,13</sup> Therefore, numerous measures have been implemented to improve their structural stability.<sup>15–17</sup>

Elemental doping is regarded as a viable strategy to enhance the electrochemical properties of Li-rich oxides. For instance, Wang *et al.* showed that F-doping  $\text{Li}[\text{Li}_{0.133}\text{Mn}_{0.467}\text{Ni}_{0.2}\text{Co}_{0.2}]\text{O}_2$  suppresses voltage decay and enhances capacity retention compared to its undoped counterpart.<sup>18</sup> Additionally, Vanaphuti *et al.* demonstrated that co-doping Na and F into Li-rich Mn-base oxides improves structural stability and mitigates oxygen loss.<sup>19</sup> Mao *et al.* also proposed an effective, straightforward, and scalable co-doping strategy with trace amounts of Fe and F to enhance the rate capability and cycling performance of high-Ni Li-rich layered oxides.<sup>20</sup> These results show that F is a significant anion in the elemental doping of Li-rich materials. However, most studies focus on experimental analysis to highlight the key role of F doping in performance enhancement. There is a lack of exploration into the underlying mechanisms

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of F doping's effect on the electrochemical properties of Li-rich materials through theoretical calculations. In this study, using first-principles calculations, we investigate the impact of F doping on the electrochemical performance of Li-rich  $\text{Li}_2\text{MnO}_3$ , including voltage profile, redox reaction, structural stability, and ionic conductivity. Our results deepen the understanding of F doping's role in Li-rich materials.

## 2. Computational methods

The present calculations are carried out by using the projector-augmented wave (PAW)<sup>21</sup> representations within density functional theory (DFT) as implemented in the Vienna *Ab initio* Simulation Package (VASP).<sup>22,23</sup> The exchange and correlation energy is treated within the spin-polarized generalized gradient approximation (GGA) and parameterized by Perdew–Burke–Ernzerhof formula (PBE).<sup>24</sup> The effects due to the localization of the d electrons of the transition metal ions were taken into account with the GGA + *U* approach of Dudarev *et al.*<sup>25</sup> An effective single parameter *U*–*J* of 5.0 eV is used for Mn, which has been proved to be a good approximation in Mn-based compounds.<sup>26,27</sup> Wave functions are expanded in plane waves up to a kinetic energy cut-off of 550 eV. Brillouin-zone integrations are approximated by using special *k*-point sampling of Monkhorst–Pack scheme<sup>28</sup> with a *k*-point mesh resolution of  $2\pi \times 0.03 \text{ \AA}^{-1}$ . Lattice vectors together with the atomic coordinates were fully relaxed until the force on each atom is less than  $0.01 \text{ eV \AA}^{-1}$ .

The relative stability of  $\text{Li}_x\text{Host}$  polymorphs at each Li composition *x* was evaluated by

$$\Delta E_f = E(\text{Li}_x\text{Host}) - \left[ \frac{x}{2} E(\text{Li}_2\text{Host}) + \left( 1 - \frac{x}{2} \right) E(\text{Host}) \right] \quad (1)$$

where *E* is the calculated total energy for a given structure.

The average voltage (*V*) versus  $\text{Li/Li}^+$  was calculated as

$$V = \frac{E_{\text{tot}}(\text{Li}_{x_2}\text{Host}) - E_{\text{tot}}(\text{Li}_{x_1}\text{Host}) - (x_2 - x_1)E_{\text{tot}}[\text{Li}]}{(x_2 - x_1)e} \quad (2)$$

where  $x_2$  and  $x_1$  are the Li composition before and after the lithium extraction from the host structure, respectively.

The reaction energy associated with the formation of oxygen vacancy in  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  was calculated by

$$\Delta H = \frac{E_{\text{tot}}(\text{Li}_x\text{MnO}_{3-y}) + \left( \frac{y}{2} \right) E(\text{O}_2) - E_{\text{tot}}(\text{Li}_x\text{MnO}_3)}{y/2} \quad (3)$$

and

$$\Delta H = \frac{E_{\text{tot}}(\text{Li}_x\text{MnO}_{2.75-y}\text{F}_{0.25}) + \left( \frac{y}{2} \right) E(\text{O}_2) - E_{\text{tot}}(\text{Li}_x\text{MnO}_{2.75}\text{F}_{0.25})}{y/2} \quad (4)$$

The total energy of a single oxygen molecule ( $E(\text{O}_2)$ ) was calculated in a  $20 \times 20 \times 20 \text{ \AA}$  periodic box, and the correction proposed by Ceder *et al.*<sup>29</sup> was then added to  $E(\text{O}_2)$ .

The activation barriers for the Li migration in  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  were calculated using the climbing nudged

elastic band (NEB) method<sup>30</sup> in a  $2 \times 1 \times 2$  supercell containing 16 formula units. For the NEB calculations, the standard GGA functional was employed, and the lattice constants of a given structure were fixed as their equilibrium values, with all the internal degrees of freedom fully relaxed.

## 3. Results and discussion

### 3.1 Effect of F-doping on the voltage profile and volume change of $\text{Li}_2\text{MnO}_3$

The crystal structure of  $\text{Li}_2\text{MnO}_3$  considered in this study has a monoclinic symmetry with *C2/m* space group and includes 12 O atoms at 4i and 8j sites in the unit cell. We replace one of the O atoms with F (substituting O at 8j site is more stable) to construct F-doped  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  structure as shown in Fig. 1. The increase in volume following F doping is shown in Table 1, which indicates a 2.38% volume expansion. We also examined the impact of F-doping on the voltage profile of  $\text{Li}_2\text{MnO}_3$ . To calculate the theoretical voltage, the formation energies of delithiated structures were first investigated based on the eqn (1). For  $\text{Li}_x\text{MnO}_3$ , all possible Li distributions are considered, as shown in Fig. 2(a). The lowest energy  $\text{Li}_x\text{MnO}_3$  compounds were then used to replace O with F to construct F-doped  $\text{Li}_x\text{MnO}_{2.75}\text{F}_{0.25}$ , and their formation energies were calculated, as shown in Fig. 2(b). The delithiated structures that lie on the convex hull indicated stable intermediate phases. For  $\text{Li}_2\text{MnO}_3$ , two stable intermediate phases,  $\text{Li}_{0.5}\text{MnO}_3$  and  $\text{Li}_{0.125}\text{MnO}_3$ , are found, which is consistent with previous work.<sup>31</sup> For the F-doped  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ , the phases of  $\text{Li}_{1.75}\text{MnO}_{2.75}\text{F}_{0.25}$ ,  $\text{Li}_{0.5}\text{MnO}_{2.75}\text{F}_{0.25}$  and  $\text{Li}_{0.125}\text{MnO}_{2.75}\text{F}_{0.25}$  lie on the convex hull. Our

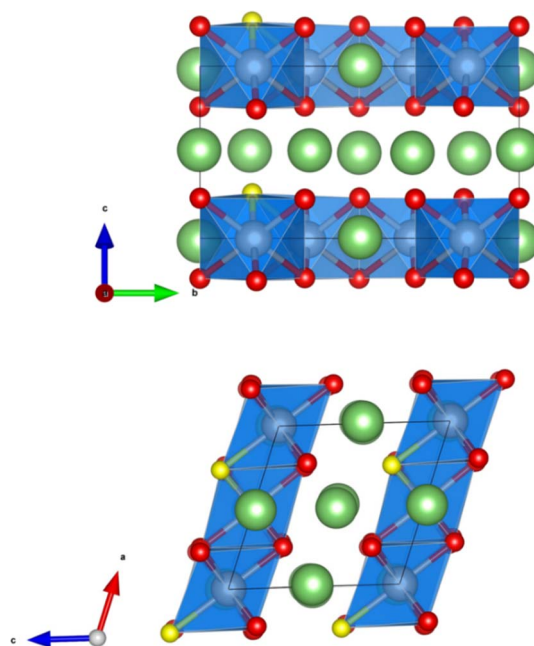


Fig. 1 The crystal structure of  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ . The Li and O atoms are denoted by green and red spheres, respectively. The substituted O atom by F is shown as a yellow sphere. The  $\text{MnO}_6$  octahedra are highlighted in blue.



**Table 1** The lattice parameters of  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ 

	$a, b, c$ (Å)	$\beta$ (°)	$V$ (Å <sup>3</sup> per f.u.)
$\text{Li}_2\text{MnO}_3$	5.01, 8.66, 5.08	109.48	51.99
$\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$	5.04, 8.80, 5.09	109.12	53.23

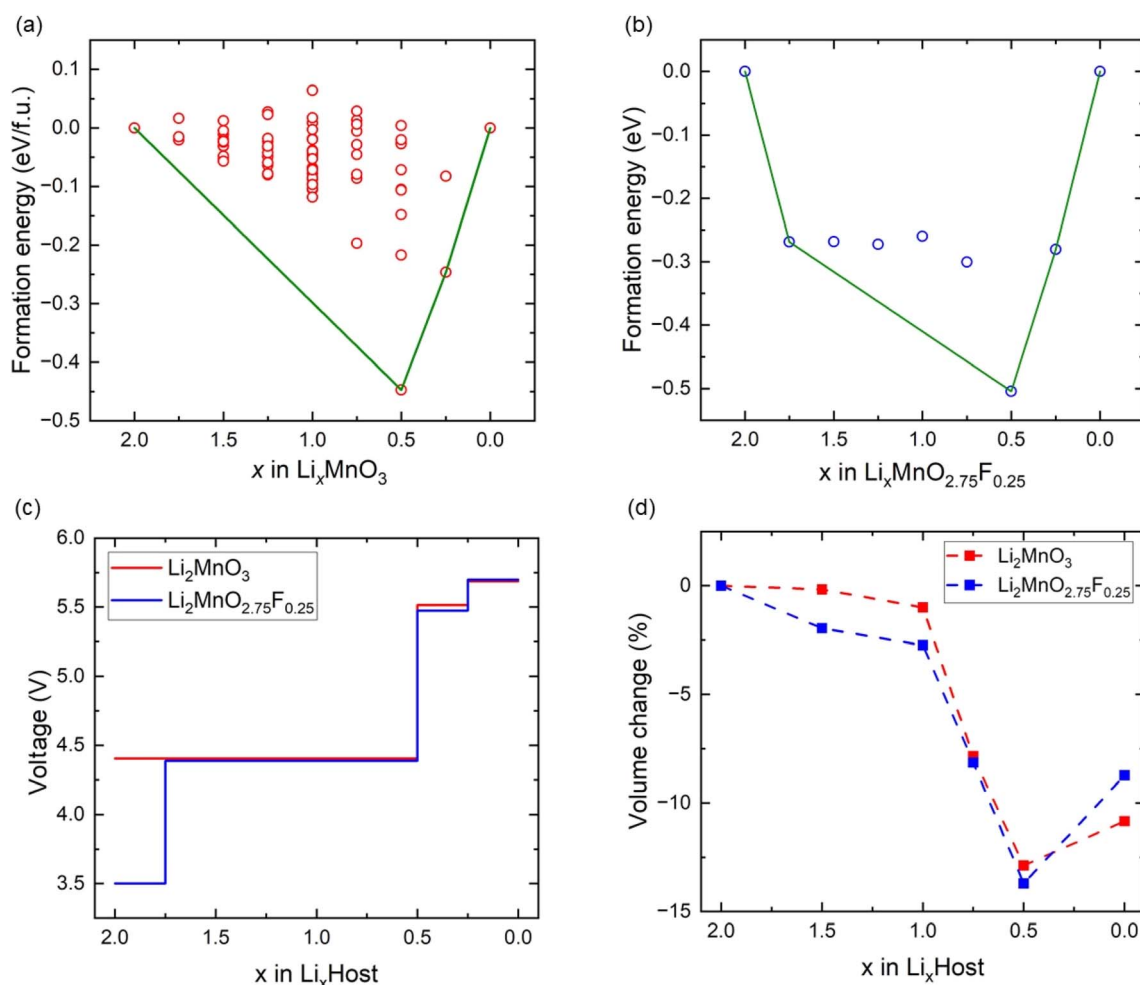
calculated results show that these three stable delithiated  $\text{Li}_x\text{MnO}_{2.75}\text{F}_{0.25}$  compounds all have F substituting the O 8j sites. Using these stable phases, we calculated the theoretical average voltages for  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  using eqn (2), as shown in Fig. 2(c). For  $\text{Li}_2\text{MnO}_3$ , a long voltage plateau at around 4.41 V is observed, which aligns well with experimental results.<sup>32</sup> When more Li ions were extracted ( $0.5 > x > 0$ ), two higher voltage plateaus at 5.52 and 5.69 V appear. After F-doping, a short plateau ( $2.0 > x > 1.75$ ) is observed, corresponding to a lower voltage of 3.50 V. As more Li ions were extracted, a long plateau can be seen at 4.39 V, followed by two higher plateaus between  $x = 0.5$  and  $x = 0$ . We will discuss the

changes in oxidation potential between  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  during delithiation later in this paper.

We also investigated the volume changes of  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  during delithiation, as shown in Fig. 2(d). For the first Li extraction, the volume changes of  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  are  $-1.1\%$  and  $-2.7\%$ , respectively. With further Li extraction ( $\text{Li}_{0.5}\text{MnO}_3$  and  $\text{Li}_{0.5}\text{MnO}_{2.75}\text{F}_{0.25}$ ), the calculated volume changes are  $-12.8\%$  and  $-13.7\%$ . Such substantial volume change could hinder the cycling of more Li ions from  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ , which suggests their structural instability upon deep delithiation.

### 3.2 Effect of F-doping on oxidation of $\text{Li}_2\text{MnO}_3$

Fig. 3(a) shows the calculated projected density of states (PDOS) for  $\text{Li}_x\text{MnO}_3$ , indicating that  $\text{Li}_2\text{MnO}_3$  is a semiconductor with a band gap of  $\sim 1.32$  eV. The valence band maximum is dominated by O 2p non-bonding states, attributed to a Li–O–Li configuration in the  $\text{Li}_2\text{MnO}_3$  structure.<sup>8</sup> As Li ions are extracted, the Fermi level shifts towards lower energy, leading to the gradual extraction of electrons from non-bonding O 2p states. Therefore, the observed voltage plateaus of 4.41, 5.52 and 5.69 V



**Fig. 2** The formation energies (a)  $\text{Li}_2\text{MnO}_3$  and (b)  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ . (c) The voltage profiles of  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ . (d) The volume change of  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ .



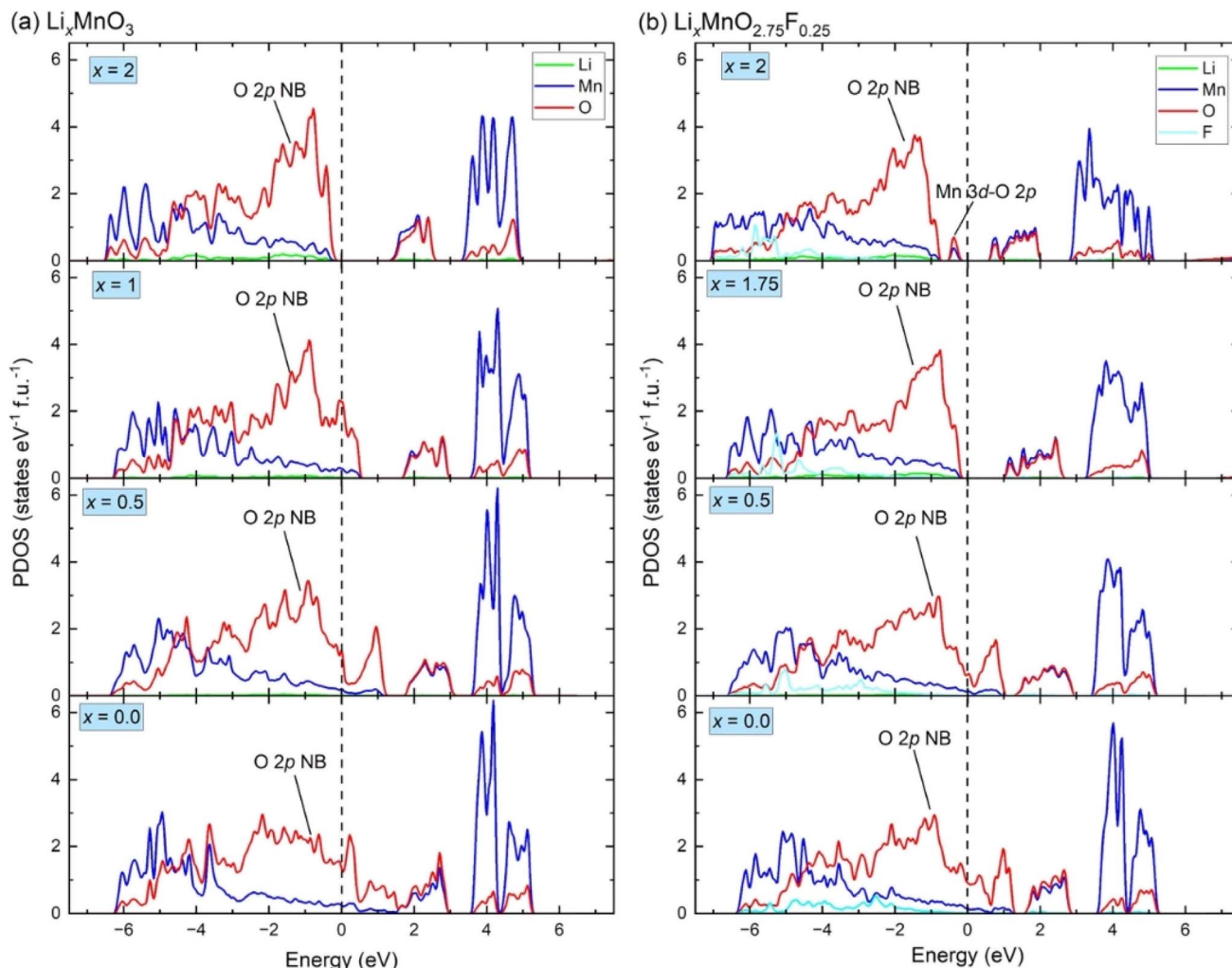


Fig. 3 The density of states for (a)  $\text{Li}_x\text{MnO}_3$  and (b)  $\text{Li}_x\text{MnO}_{2.75}\text{F}_{0.25}$ . The non-bonding O 2p states is denoted as "O 2p NB", the hybridized Mn 3d–O 2p states is denoted as "Mn 3d–O 2p".

for  $\text{Li}_2\text{MnO}_3$  (Fig. 2(c)) all correspond to oxygen oxidation. These results are consistent with the calculated magnetic moments for  $\text{Li}_2\text{MnO}_3$  (Fig. 4(a) and (b)). In  $\text{Li}_2\text{MnO}_3$  (4 Mn atoms and 12 O atoms per unit cell), all four Mn atoms display a magnetic moment of approximately 3.0, indicative of a valence state of +4 ( $3d^3$ , high-spin state  $e_g^1 t_{2g}^2$ ) as depicted in Fig. 4(c). Upon Li ion extraction, the magnetic moment of Mn remains almost unchanged, suggesting that Mn does not participate in oxidation. As illustrated in Fig. 4(b), all 12 O atoms in  $\text{Li}_2\text{MnO}_3$  exhibit a magnetic moment of approximately 0, indicating a pristine valence state of  $-2$  for O. During the delithiation process, the magnetic moment of O increases, indicating oxygen oxidation.

In F-doped  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ , the valence band maximum is dominated by hybridized Mn 3d–O 2p states, as shown in Fig. 3(b). During delithiation from  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  to  $\text{Li}_{1.75}\text{MnO}_{2.75}\text{F}_{0.25}$ , the hybridized Mn 3d–O 2p states shift to higher energy (above Fermi level), indicating that Mn atoms lose electrons to maintain charge balance. This observation aligns with the magnetic moment calculations depicted in Fig. 4(d). In

$\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ , one of the four Mn atoms exhibits a magnetic moment of about 4.0, indicative of a valence state of +3 ( $3d^4$ , high-spin state  $e_g^2 t_{2g}^2$ ) as shown in Fig. 4(c). At  $x = 1.75$ , oxidation from  $\text{Mn}^{3+}$  to  $\text{Mn}^{4+}$  occurs. Consequently, a lower voltage plateau of 3.50 V is observed between  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  and  $\text{Li}_{1.75}\text{MnO}_{2.75}\text{F}_{0.25}$ , as illustrated in Fig. 2(c). With further delithiation ( $x < 1.75$ ), non-bonding O 2p states begin to be extracted, as seen in Fig. 3(b) and 4(e), indicating electron loss from oxygen ions. The oxygen oxidation process occurring in  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  exhibits parallels to that observed in  $\text{Li}_2\text{MnO}_3$ , resulting in similar voltage plateaus at  $x < 1.75$  (Fig. 2(c)). The magnetic moment of F remains close to 0 in  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$  upon delithiation, showing that F does not participate in oxidation, as depicted in Fig. 4(f).

We also investigated the effect of F doping on the stability of oxygen redox in  $\text{Li}_x\text{MnO}_3$ , particularly concerning the formation of O vacancy ( $\Delta E$ ). As shown in Fig. 5, the  $\Delta E$  values decrease monotonically with the reduction in Li concentration, suggesting a downward trend in stability of  $\text{Li}_x\text{MnO}_3$  as  $x$  decrease. The  $\Delta E$  values become negative at  $x = 1.5$ , indicating that the





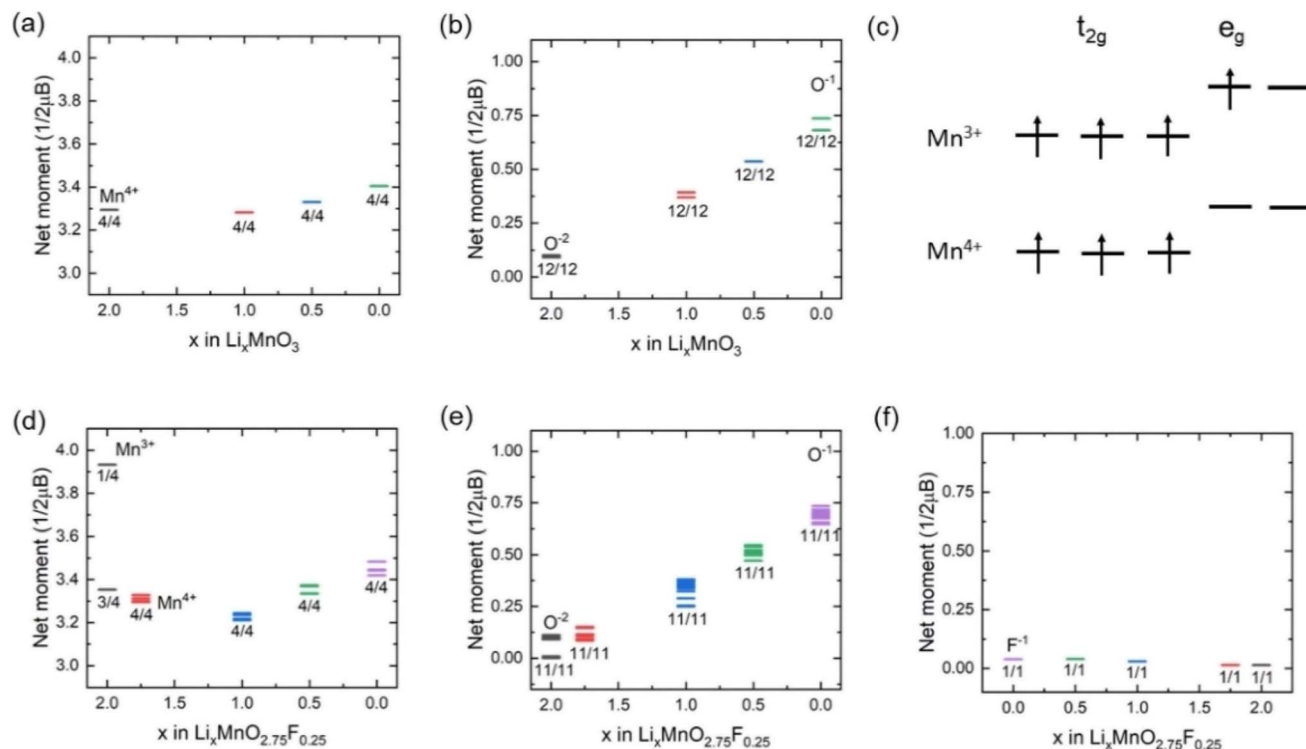


Fig. 4 Magnetic moment variation of (a) Mn and (b) O in Li<sub>2</sub>MnO<sub>3</sub>. (c) Schematic diagrams of Mn<sup>3+</sup> and Mn<sup>4+</sup>. Corresponding magnetic moment variation of (d) Mn, (e) O and (f) F in Li<sub>2</sub>MnO<sub>2.75</sub>F<sub>0.25</sub>.

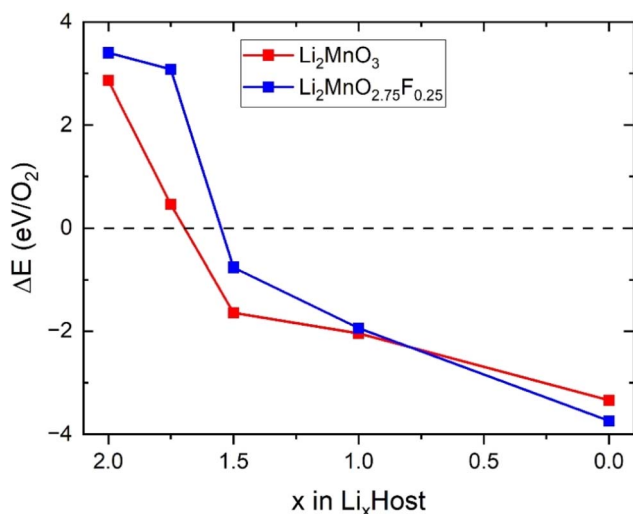


Fig. 5 Calculated formation energy for the reaction of O<sub>2</sub> in Li<sub>2</sub>MnO<sub>3</sub> and Li<sub>2</sub>MnO<sub>2.75</sub>F<sub>0.25</sub> during delithiation.

oxygen release reaction occurs spontaneously in delithiated Li<sub>1.5</sub>MnO<sub>3</sub>, which is agreed well with previous calculations.<sup>33</sup> Compared to Li<sub>2</sub>MnO<sub>3</sub>, smaller values of ΔE can be observed in Li<sub>2</sub>MnO<sub>2.75</sub>F<sub>0.25</sub> at x < 1.0, suggesting that F doping can somewhat suppress oxygen release. This phenomenon can be attributed to the F doping inducing a minor fraction of Mn to undergo oxidation, which in turn attenuate the oxidation of oxygen and enhances its stability.

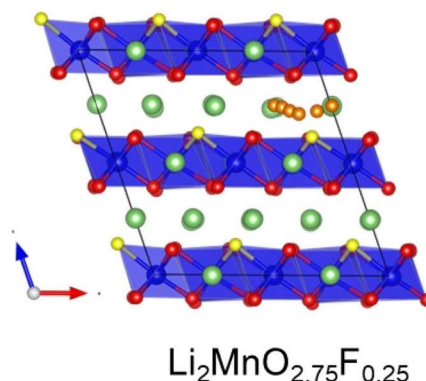
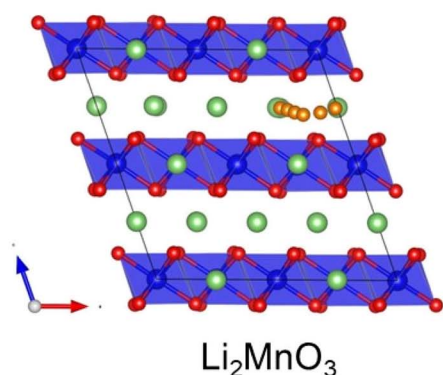
Since cation migration is recognized as a leading cause of voltage decay and hysteresis,<sup>10–13</sup> we investigated the effect of F-doping on Mn migration during deep charge. First, we constructed a 2 × 1 × 2 supercell with 16 formula units and moved one Mn ion to the Li layer in the deeply delithiated state Li<sub>0.5</sub>MnO<sub>3</sub>. After structural optimization, we found that O–O dimers form in this Li<sub>0.5</sub>MnO<sub>3</sub> with Mn migration, resulting in lower energy compared to pristine Li<sub>0.5</sub>MnO<sub>3</sub> (ΔE = 67 meV per atom). This suggests that the deep oxygen oxidation leads to O–O dimer formation and Mn migration, consistent with previous calculations.<sup>34</sup> When considering F doping, Mn migration and O–O dimer formation also occur, with the energy of the migrated Mn structure being lower (~70 meV per atom) than that of pristine Li<sub>0.5</sub>MnO<sub>2.75</sub>F<sub>0.25</sub>. These calculated results indicate that F-doping cannot suppress Mn migration at deep charge.

### 3.3 Effect of F-doping on ionic diffusion

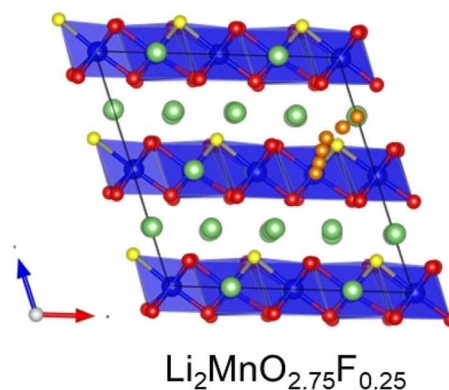
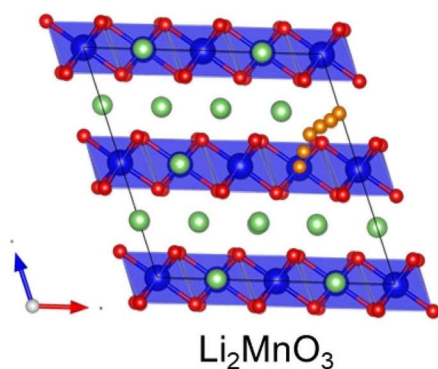
Climbing NEB calculations were used to investigate Li diffusion in Li<sub>2</sub>MnO<sub>3</sub> and Li<sub>2</sub>MnO<sub>2.75</sub>F<sub>0.25</sub>, assessing the impact of F-doping on diffusion kinetics. Previous calculations identified two types of Li<sup>+</sup> migration between the neighboring Li and LiMn<sub>2</sub> layers, and three pathways within Li layers.<sup>33</sup> We selected two of these pathways to study the effect of F-doping on ionic diffusion: path 1 involves Li<sup>+</sup> migration from the 4h site to the 2c site in the Li planes (Fig. 6(a)), and path 2 involves Li<sup>+</sup> migration from the 4h site to the 2b site from Li to LiMn<sub>2</sub> layers (Fig. 6(b)). The calculated activation barriers (E<sub>a</sub>) along these



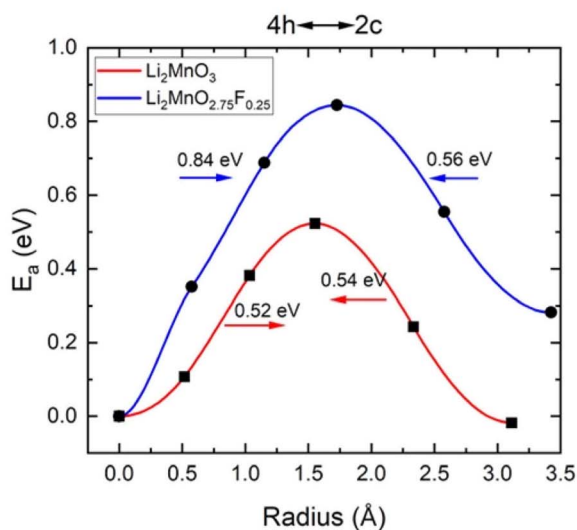
## (a) path1 4h-2c



## (b) path2 4h-2b



## (c)



## (d)

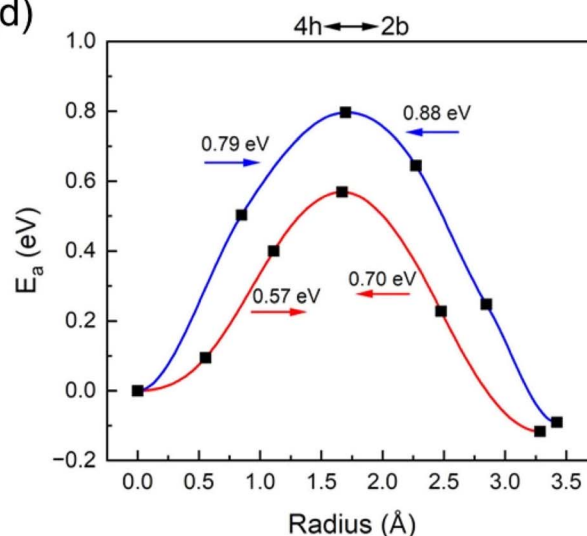


Fig. 6 Calculated kinetic properties of  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ . Schematic diagrams for (a) path 1 between 4h and 2c and (b) path 2 between 4h and 2b for  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ . Calculated activation barrier along (c) path 1 and (d) path 2. The Li and O atoms are denoted by green and red spheres, respectively. The yellow sphere is the substituted O by F atoms.  $\text{MnO}_6$  octahedron is colored in blue. Li diffusion paths are represented by the small orange spheres.

paths are shown in Fig. 6(c) and (d). For  $\text{Li}_2\text{MnO}_3$ , we observed that the energy barriers for Li diffusion between the 4h site and 2c site are 0.52 and 0.54 eV, aligning well with previous calculations (0.54 and 0.61 eV).<sup>33</sup> For Li hopping between the 4h site

and 2b site, the barriers are 0.57 and 0.70 eV, also consistent with Xiao *et al.*'s calculations.<sup>33</sup> However, after F-doping, the barriers of both paths change significantly. Here, F-doping is modeled by substituting the O site near the Li hopping trajectory



**Table 2** Calculated activation barriers ( $E_a$ ) and estimated diffusion coefficients ( $D_{300\text{ K}}$ ) for  $\text{Li}_2\text{MnO}_3$  and  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$

Structure	$E_a$ (eV)	$D_{300\text{ K}}$ ( $\text{cm}^2\text{ s}^{-1}$ )
$\text{Li}_2\text{MnO}_3$	0.52/0.54/0.57/0.70	$10^{-11}/10^{-11}/10^{-12}/10^{-14}$
$\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$	0.84/0.56/0.79/0.88	$10^{-16}/10^{-12}/10^{-15}/10^{-17}$

(Fig. 6(a) and (b)). Although the model is not exhaustive, it provides insights into the effect of F-doping on Li diffusion. As shown in Fig. 6(c) and (d), the path lengths of two pathways are increased after F doping, and  $E_a$  increases in both paths after F-doping due to the stronger interaction between  $\text{F}^-$  and  $\text{Li}^+$ . The diffusion coefficients of these pathways can be estimated according to  $D = d^2\nu \exp(-E_a/k_bT)$  ( $\nu = 10^{13}\text{ THz}$ ,  $T = 300\text{ K}$ , and  $d$  is the hopping distance)<sup>35</sup> as shown in Table 2. Consequently, F-doping leads to a deterioration in the kinetics of  $\text{Li}_2\text{MnO}_3$ .

## 4. Conclusions

Using the first-principles calculations, we conduct a comprehensive study on the impact of F-doping on the electrochemical performance of  $\text{Li}_2\text{MnO}_3$ . Our results indicate that both  $\text{Li}_2\text{MnO}_3$  and its F-doped variant,  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ , undergo significant volume changes (over 10%) during deep delithiation, which could impede more Li ions cycling of them. Analysis of the electronic structure and magnetic moments in the delithiated states reveals that charge compensation in  $\text{Li}_2\text{MnO}_3$  primarily involves the oxidation of  $\text{O}^{2-}$  anions. After F doping, both cationic (Mn) and anionic (O) oxidation are observed in  $\text{Li}_2\text{MnO}_{2.75}\text{F}_{0.25}$ , contributing to a lower voltage plateau at the beginning of charge due to the oxidation of  $\text{Mn}^{3+}$  to  $\text{Mn}^{4+}$ , which can somewhat mitigate oxygen release. Additionally, F doping appears to impair Li diffusion kinetics in comparison to pristine  $\text{Li}_2\text{MnO}_3$ , owing to enhanced interactions between  $\text{F}^-$  and  $\text{Li}^+$ . Our findings offer a deeper insight into the effect of F-doping on the electrochemical properties of  $\text{Li}_2\text{MnO}_3$  and provide strategic guidance for future optimization of this high-capacity cathode material.

## Data availability

We confirm that the data supporting the finding of this study are available within the main article.

## Author contributions

Xiang-Ming Zeng: conceptualization, investigation, formal analysis, writing – original draft, writing – review & editing; Jing Liu: methodology, formal analysis; Jiang-Bin Su: formal analysis, investigation; Fa-Hui Wang, conceptualization, investigation, formal analysis; Yan-Bing Li: methodology; Chang-Jun Zhang: conceptualization; Ming Liu: methodology; Run-Sheng Wu: formal analysis, investigation; Jun-Ping Hu: methodology; Feng Zheng: methodology; investigation, writing – review & editing; supervision.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The work at Xinyu University was supported by the Foundation of Jiangxi Province Educational Committee, China (Grant No. GJJ202317) and the Ministry of Education Collaborative Education Project for Industry-University Cooperation (Grant No. 231103117091418). The work at Jimei University was supported by the Natural Science Foundation of Xiamen, China (Grant No. 3502Z202372015), the Fujian Provincial Education Department, China (Grant No. JZ230025) and the Research Foundation of Jimei University (Grant No. ZQ2023013).

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