



# High-performance carbon-coated mesoporous $\text{LiMn}_2\text{O}_4$ cathode materials synthesized from a novel hydrated layered-spinel lithium manganate composite†

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Carbon-coated mesoporous spinel  $\text{LiMn}_2\text{O}_4$  has been synthesized from a novel hydrated layered-spinel lithium manganate composite through a facile hydrothermal process and subsequent thermal treatment. Benefiting from the carbon coating and mesoporous structure, the  $\text{LiMn}_2\text{O}_4$  material exhibits superior high-rate capability and long-life cycling stability, delivering the initial discharge capacity of 117.8 mA h g<sup>-1</sup> at 30C with over 90% capacity retention after 1500 cycles. Moreover, the innovative employment of hydrated layered Li-deficient and spinel Li-rich intermediates might provide greater inspiration for other high-performance electrode materials with multiple layer architectures and optimized phase compositions.

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## Introduction

Nowadays, lithium-ion batteries (LIBs) have extensive applications in electrochemical energy storage and conversion including hybrid electric vehicles (HEVs), electric vehicles (EVs), portable electronic devices and energy storage systems (ESSs). However, at the present stage, lithium-ion batteries are still facing challenges to meet the increasing market demands for higher energy and power densities, longer cycling life, better security and lower cost.<sup>1–4</sup> Since cathode materials can severely restrict the electrochemical properties, breakthroughs in the cathodes become more essential and urgent.<sup>1,5–7</sup> Among all the appealing cathode materials, spinel  $\text{LiMn}_2\text{O}_4$  has attracted much attention by virtue of its low cost, environmental friendliness, high operating plateaus, acceptable energy density and excellent safety.<sup>8–10</sup> Nevertheless, the electrochemical properties of  $\text{LiMn}_2\text{O}_4$  seem far from satisfactory: (1) poor high-rate capability hampered by the low electronic and ionic conductivities; (2) capacity deterioration caused by the severe manganese dissolution and side reactions.<sup>9,11,12</sup>

To improve the high-rate capability and cycling stability of  $\text{LiMn}_2\text{O}_4$ , designs of carbon coating and mesoporous

structures have been extensively investigated.<sup>13–19</sup> Carbon coating can increase the electronic conductivities by forming a continuous electron transport route along the materials' surface, while mesoporous structures can increase the ionic conductivities by reducing the pathway of  $\text{Li}^+$  diffusion and providing more reaction sites, which contribute to the rate capability.<sup>13–17</sup> Meanwhile, upon lithium insertion/extraction, carbon coating can protect the materials from volume changes and side reactions, while the mesoporous structures can accommodate the structural strain, thus contributing to the cycling stability.<sup>13,16,18,19</sup>

As for the synthesis of  $\text{LiMn}_2\text{O}_4$ , various polymorphs of manganese oxides have been explored such as  $\alpha\text{-MnO}_2$ ,<sup>20,21</sup>  $\beta\text{-MnO}_2$ ,<sup>22,23</sup>  $\gamma\text{-MnO}_2$ ,<sup>21,24</sup>  $\delta\text{-MnO}_2$ ,<sup>25–27</sup> and  $\text{R-MnO}_2$ .<sup>28</sup> Among them, birnessite-type lithium manganese oxide ( $\delta$ -type or Li-birnessite) possesses a hydrated layered structure containing Li ions and crystal water between the layers,<sup>29–31</sup> contributing to the exotic properties in many fields.<sup>32–34</sup> Moreover, the layered phase is prone to generate the lamellar nanostructure<sup>32,33</sup> which can reduce the  $\text{Li}^+$  diffusion pathway in the thickness direction and further improve the rate performance. Unfortunately, though having been reported as precursors or immediate products for years, the Li-birnessite troubles with impurities such as Li-deficient  $\text{Li}_x\text{MnO}_2$  or  $\text{Mn}_2\text{O}_3$  and lamellar aggregation during the transformation to spinel  $\text{LiMn}_2\text{O}_4$ ,<sup>25,31,35</sup> resulting in inferior electrochemical performances. Thus, how to utilize the Li-birnessite with novel structure to synthesize high-performance  $\text{LiMn}_2\text{O}_4$  is worth further study.

In this work, we have employed a novel hydrated layered-spinel lithium manganese oxide composite to synthesize the carbon-coated mesoporous spinel  $\text{LiMn}_2\text{O}_4$  via a facile

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hydrothermal process and subsequent thermal treatment. Based on the gradual transformation from layered Li-birnessite phase to spinel  $\text{LiMn}_2\text{O}_4$  phase, the resultant mesoporous structure and carbon coating can effectively improve the rate capability and cycling stability. The outstanding electrochemical performance thus obtained, is of interest as a cathode material for lithium-ion batteries.

## Results and discussion

As illustrated in Fig. 1, the formation mechanism of the carbon-coated mesoporous  $\text{LiMn}_2\text{O}_4$  involves two steps: the hydrothermal reaction between manganese oxides (denoted as MO in Fig. 1a) and lithium hydroxide solution at  $160^\circ\text{C}$  (Step I) and the thermal treatment under vacuum at  $600^\circ\text{C}$  (Step II). The absorbed organic group  $\text{CH}_3\text{COO}^-$  (shown in Fig. 1a and b) is originated from the reduction of ethanol in the synthesis process of the initial MO. Through Step I, the obtained hydrothermal products (Fig. 1b) consist of two phases: layered Li-birnessite and spinel LMO. The blooming flower-like nanosheets at different levels were interlaced, contributing to the mesoporous structure. Then through Step II, carbon-coated mesoporous  $\text{LiMn}_2\text{O}_4$  (Fig. 1c) was obtained where the residual Li-birnessite fully transformed into the spinel LMO accompanied by the surface carbonization (see more experimental details in the ESI†).

The hydrothermal process of Step I is so critical through the whole synthesis process that deserves further investigation and analysis. X-ray diffraction (XRD) patterns of the prepared MO and hydrated hydrothermal products are shown in Fig. 2a. The weak peaks of MO indicate the disordered nature of the birnessite-type manganese oxides. While after the hydrothermal treatment, the layered monoclinic Li-birnessite ( $\text{Li}_{4-}\text{Mn}_{14}\text{O}_{27}\cdot x\text{H}_2\text{O}$ , JCPDS 50-0009) coexists with the spinel lithium manganese oxide (LMO). The layered Li-birnessite phase contains crystal water and Li-ions between layers with a basal spacing of  $\sim 0.7\text{ nm}$  (presented in Fig. 2b), which corresponds to the (001) diffraction peak at around  $12.8^\circ$ . Typical (111) diffraction at  $2\theta = 18.9^\circ$  confirms the appearance of Li-rich spinel phase  $\text{Li}_{1.27}\text{Mn}_{1.73}\text{O}_4$  (JCPDS 51-1582) which is a common hydrothermal product consistent with previous reports.<sup>36</sup> Fig. 2c shows the crystal structure of the spinel lithium manganese oxide. The two-phase mixture (denoted as hydrated L-S) consisting of both Li-deficient layered phase (compared with  $\text{Li} : \text{Mn} = 1 : 2$  (atom%) for stoichiometric  $\text{LiMn}_2\text{O}_4$ ) and Li-rich spinel phase can be significant for the subsequent changes. XRD patterns of hydrothermal products at intermediate time are depicted in Fig. S1a and c.† After 1 h

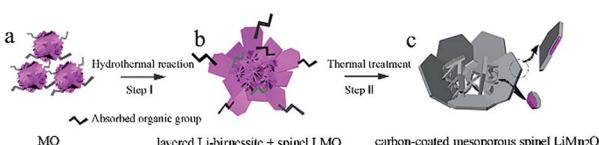


Fig. 1 A schematic illustrating the formation mechanism of the carbon-coated mesoporous spinel  $\text{LiMn}_2\text{O}_4$ .

reaction, only single layered Li-birnessite phase exists and when the reaction time increased, the spinel lithium manganese oxide (LMO) comes out and coexists with the layered Li-birnessite. The above results assist to understand the phase transformation process which are based on the 'hydrothermal-soft chemical reaction'.<sup>25,26,37-39</sup>  $\text{Li}^+$  inserts into the interlayer spaces of Li-birnessite (soft chemical reaction); the layered structure collapses along with the squeezing of water and changes into the spinel structure (hydrothermal reaction).

Furthermore, scanning electron microscopy (SEM) images in Fig. 2d, e, S1b and d† reveal the corresponding morphology evolution. The initial MO are bud-like nanoclusters of 200–300 nm assembled by interconnected thin nanosheets (Fig. 2d) and transmission electron microscopy (TEM) images show that these nanosheets are only a few nanometers in thickness (Fig. S2a and b†). Then after hydrothermal treatment, inhomogeneous nanostructures appear which consist of both pedal-like bigger nanosheets and bud-like smaller nanoclusters (Fig. 2e). Additionally, as can be seen from Fig. S2c and d,† the bigger nanosheets are of several hundred nanometers and the nanoclusters are assembled by small nanosheets (tens of nanometers). Fig. S1b† reveals that the morphology of nanosheets and nanoclusters originated from the layered Li-birnessite phase which is more liable to generate the laminar morphology. The obtained nanosheets with different sizes and the whole morphology evolution can be interpreted as Ostwald ripening, nucleation–dissolution–recrystallization and self-assembly processes in consistent with previous reports.<sup>25,39-42</sup> In particular, the MO nanodomains separated from the edge of original nanosheets and new  $\text{MnO}_2$  nuclei adsorbed with the  $\text{CH}_3\text{COO}^-$  group formed. Due to the coexistence of multi-phases in the hydrothermal system as described before and energy difference, more than one nucleus of the crystal units may be contained with different nucleation, orientation and growth rates,<sup>40,42</sup> which can explain the inhomogeneous outward migration of crystallites. Moreover, the relatively higher concentration of  $\text{Li}^+$  facilitates dissolution of manganese oxide<sup>39,42</sup> and thus favors the generation of those bud-like smaller nanoclusters according to the dissolution–recrystallization mechanism. Ultimately, the blooming flower-like nanosheets at different levels are self-assembled, the interconnection between which can result in the mesoporous structure. The mesoporous structure is further confirmed by the nitrogen adsorption–desorption isotherms with a type-IV curve and hysteresis loop (Fig. 2f). The pore size distribution curve calculated by using the density functional theory (DFT) shows a multimodal feature with peaks ranging from 3–30 nm. More importantly, X-ray photoelectron spectroscopy (XPS) investigation of C1s in Fig. 2g reveals peak locations at 284.6 eV (C–C), 285.8 eV (C–O) and 288.6 eV (C=O),<sup>15,43,44</sup> confirming organic groups terminated on the surface which guarantees the subsequent carbon coating.

Then subsequent thermal treatment under vacuum through Step II leads to the ultimate pure phase spinel  $\text{LiMn}_2\text{O}_4$  transforming from the intermediate of hydrated L-S. XRD characterization (Fig. 3a) of the heat-treated product reveals that all the reflections can be identified as spinel  $\text{LiMn}_2\text{O}_4$  (JCPDS



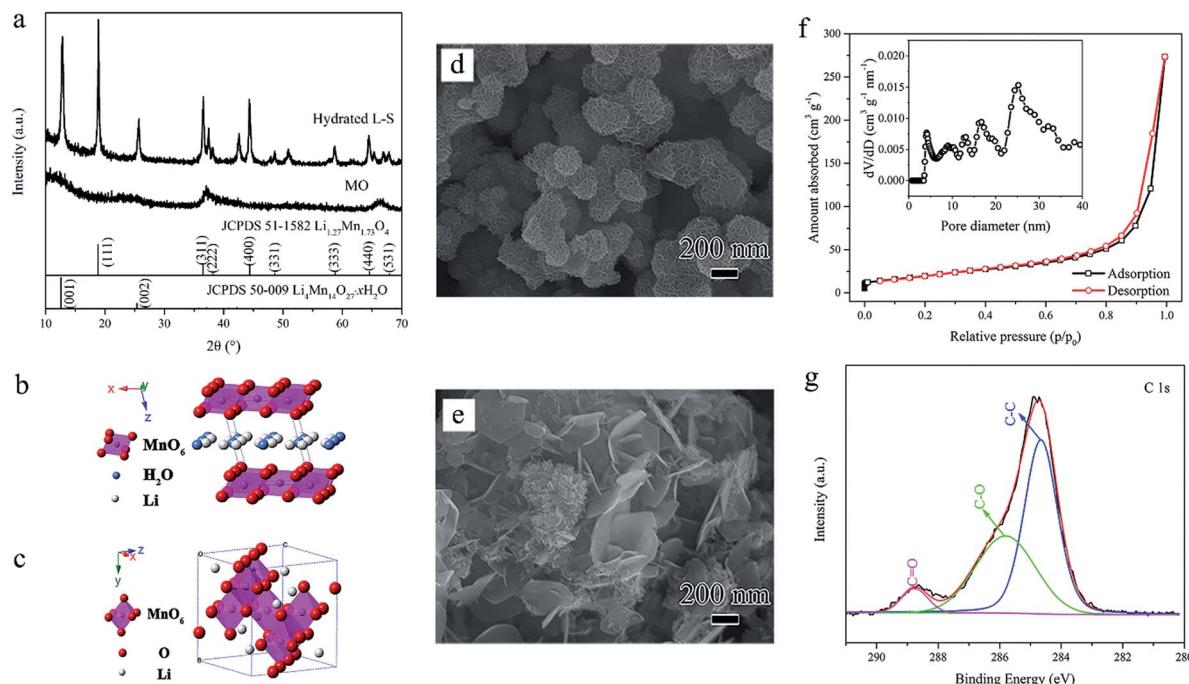


Fig. 2 (a) XRD patterns of MO and hydrated L-S. (b and c) Crystal structure of the layered hydrated Li-birnessite (b) and spinel LMO (c). (d and e) SEM images of MO and hydrated L-S. (f) N<sub>2</sub> adsorption–desorption isotherms and the corresponding pore size distribution of hydrated L-S. (g) XPS C1s spectrum of the initial manganese oxide (MO).

35-0782). The Li : Mn atomic ratio of 0.505 tested by inductively coupled plasma optical emission spectrometry (ICP-OES) method confirms the nearly stoichiometric composition. To further verify the disappearance of the Li-birnessite phase, thermogravimetric-differential thermal analysis (TG-DTA) was performed (Fig. S3a†). An obvious endothermic peak with remarkable weight loss of ~3.0% is observed at ~150 °C, which can be ascribed to dehydration of physically absorbed water and partial interlayer water.<sup>37,41,45</sup> XRD pattern of the sample annealed at 150 °C in Fig. S3b† also confirms the collapse of the

layered Li-birnessite. The weight loss over 150 °C can be assigned to the further dehydration of OH groups, release of oxygen molecules and structural rearrangement.<sup>37,41,45</sup> Moreover, when increasing the annealing temperature, the increasing intensity and slight left shift of (111) diffraction for the spinel structure can be observed (Fig. S3b†), implying the composition change of spinel phase. So far, the transformation from the two-phase mixture consisting of layered Li-deficient phase and spinel Li-rich phase to the single stoichiometric spinel LiMn<sub>2</sub>O<sub>4</sub> phase has been successfully completed.

Fig. 3b shows the SEM image of the as-prepared LiMn<sub>2</sub>O<sub>4</sub>. It can be observed that the heat-treated product inherits the former multiform lamellar structure but with coarsened nanoplates (200–500 nm) and distributed small nanoplatelets. The thickness of these nanoplates is only tens of nanometers and this is supposed to shorten the Li<sup>+</sup> diffusion pathways. Noteworthily, the smaller nanoplatelets spread around the bigger nanoplates, which can prevent the mutual aggregate and provide more interspaces. Nitrogen adsorption–desorption isotherms with the hysteresis loop shown in Fig. 3c confirm the mesoporous nature of the LiMn<sub>2</sub>O<sub>4</sub> product. Relatively smaller pore size is mainly due to the structural shrinkage during the annealing process compared with that of the hydrothermal product. Such mesoporous structure is beneficial to Li<sup>+</sup> storage by providing more convenient contact between the electrolyte and electrode materials.<sup>13,14</sup>

The crystal structure and surface feature of the LiMn<sub>2</sub>O<sub>4</sub> product were further studied by using high-resolution transmission electron microscopy (HRTEM) method combined with energy dispersive spectroscopy (EDS) point mapping analysis.

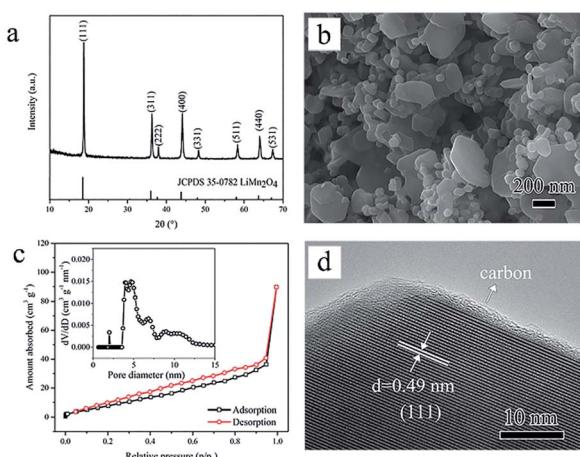


Fig. 3 Characterizations of the as-synthesized carbon-coated mesoporous spinel LiMn<sub>2</sub>O<sub>4</sub>. (a) XRD pattern, (b) SEM image, (c) N<sub>2</sub> adsorption–desorption isotherms together with the corresponding pore size distribution and (d) HRTEM image.



HRTEM image in Fig. 3d exhibits a clear interplanar distance of  $\sim 0.49$  nm, corresponding to the (111) plane of cubic spinel  $\text{LiMn}_2\text{O}_4$ . In addition, originated from surface carbonization, an apparent thin carbon layer wrapped on the surface can be distinguished. EDS point mapping results in Fig. S4 $\dagger$  indicates the higher relative concentration of C at the edge than that in the middle zone. The carbon content is 0.27 wt% measured by an element analyzer. The amorphous carbon layer can form a 3D conductive network and prevent carbon-coated  $\text{LiMn}_2\text{O}_4$  from excessive Mn dissolution during the charge/discharge cycling.<sup>16-19</sup>

Based on the above observations, the carbon-coated mesoporous spinel  $\text{LiMn}_2\text{O}_4$  (denoted as LMO-CM), as the final state, can be thus reasonably confirmed. When exploring the electrochemical performances as the cathode material for LIBs, we also introduced two control groups for comparison: the no carbon-coated spinel  $\text{LiMn}_2\text{O}_4$  (denoted as LMO-NC) which was annealed in air with the other experimental variables the same as those of LMO-CM and the commercial spinel lithium manganese oxide (denoted as LMO-COM). Their typical XRD patterns and SEM images are shown in Fig. S5. $\dagger$  Fig. 4a shows the first galvanostatic discharge–charge profiles between 3.2–

4.3 V vs.  $\text{Li}^+/\text{Li}$  at 0.2C ( $1\text{C} = 140 \text{ mA g}^{-1}$ ). All the three samples have two voltage plateaus at around 4.0 V and 4.1 V, which can be attributed to the  $\text{Mn}^{3+}/\text{Mn}^{4+}$  redox couple associated with the two-stage mechanism of  $\text{Li}^+$  insertion and extraction. The corresponding electrochemical conversion reactions can be expressed as follows:  $\text{LiMn}_2\text{O}_4 \rightarrow \text{Li}_{0.5}\text{Mn}_2\text{O}_4 + 0.5\text{Li}^+ + 0.5\text{e}^-$  (at low plateau);  $\text{Li}_{0.5}\text{Mn}_2\text{O}_4 \rightarrow 2\lambda\text{-MnO}_2 + 0.5\text{Li}^+ + 0.5\text{e}^-$  (at high plateau).<sup>10,46,47</sup> In contrast, LMO-CM has much flatter voltage plateaus, suggesting its better crystallinity property and reversibility; LMO-CM has the highest reversible discharge specific capacity of  $129.7 \text{ mA h g}^{-1}$  compared with  $116.1 \text{ mA h g}^{-1}$  and  $100.7 \text{ mA h g}^{-1}$  for LMO-NC and LMO-COM. These results are in accordance with the most distinguished cathodic/anodic peaks as well as the largest integral area surrounded by the relevant cyclic voltammetry (CV) plot (Fig. S6 $\dagger$ ).

Notably, LMO-CM exhibits superior rate capability as shown in Fig. 4b. It delivers high reversible capacities of  $129.4 \text{ mA h g}^{-1}$ ,  $121.7 \text{ mA h g}^{-1}$ ,  $115.4 \text{ mA h g}^{-1}$ ,  $114 \text{ mA h g}^{-1}$  and  $110.8 \text{ mA h g}^{-1}$  when cycled at 0.5C, 1C, 4C, 10C, 30C for each 10 cycles, respectively. Also, the capacity can be well recovered when cycled back to 1C. While LMO-NC and LMO-COM only remain  $91.0 \text{ mA h g}^{-1}$  and  $47.8 \text{ mA h g}^{-1}$  at 30C, much lower

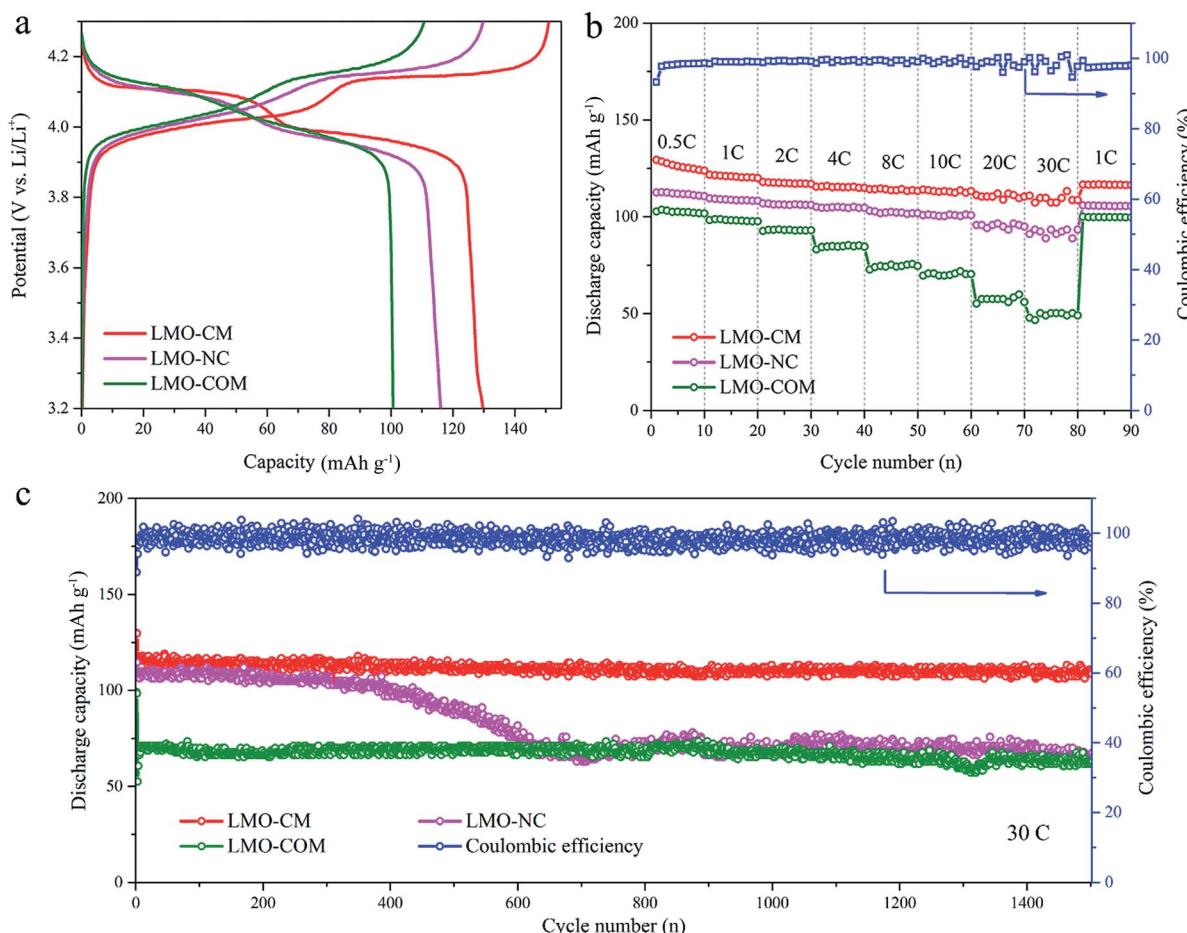


Fig. 4 Comparison of electrochemical performances for carbon-coated mesoporous LMO-CM, no carbon-coated LMO-NC and commercial LMO-COM. (a) The first galvanostatic discharge–charge profiles at 0.2C. (b) Discharge capacity with cycle number at different rates ranging from 0.5C to 30C. (c) Long-term cycling performance at 30C for all the three samples. Note that the given coulombic efficiency in (b) and (c) is only for LMO-CM.



than that of LMO-CM, besides, their corresponding capacity retention is  $\sim 83\%$  and  $\sim 49\%$  (*versus* capacity at 1C), inferior to  $\sim 91\%$  for LMO-CM. Rate performance of the as-synthesized LMO-CM was also compared with that of other typical high-rate LMO/C materials reported previously as displayed in Fig. S7,<sup>†</sup> showing the superior rate capability of our work especially at rates higher than 20C.

To investigate cycle performance at high rates, the three samples were cycled at 30C for 1500 cycles as presented in Fig. 4c. When charged and discharged at the same rate of 30C, the initial discharge capacity of LMO-CM is 117.8 mA h g<sup>-1</sup> (nearly 80% of the theoretical specific capacity of 148 mA h g<sup>-1</sup>), higher than that of LMO-NC (108.5 mA h g<sup>-1</sup>) and LMO-COM (52.5 mA h g<sup>-1</sup>). What's more, LMO-CM can retain  $\sim 92\%$  capacity (108.5 mA h g<sup>-1</sup>) after 1500 cycles with the coulombic efficiency retaining nearly 100%, whereas LMO-NC has a drastic capacity decay after about 500 cycles and ultimately retains  $\sim 61\%$  capacity (66.5 mA h g<sup>-1</sup>). As given in Fig. S8,<sup>†</sup> remarkably, at 60C (charged within 1 min actually), over 90% stable capacity retention is maintained with an initial capacity of 102.7 mA h g<sup>-1</sup>. Even cycled at 80C ( $\sim 34$  s) and 120C ( $\sim 28$  s), LMO-CM can still yield a discharge capacity as high as 102.5 mA h g<sup>-1</sup> and 88.6 mA h g<sup>-1</sup>, showing the unprecedented high-rate and long-life cycling stability. Though mainly confined to the laboratory level, the current results are still encouraging.

As schematically illustrated in Fig. 5, the outstanding electrochemical performance of the as-synthesized carbon-coated mesoporous LiMn<sub>2</sub>O<sub>4</sub> can be ascribed to three main factors. Firstly, the thin carbon layer decorated on the surface can not only reduce the interface resistance by forming a three dimensional electro-conductive network, but also improve the structural integrity by suppressing volume change and side reactions during cycling operations.<sup>16–18,48,49</sup> Electrochemical impedance spectroscopy (EIS) analysis in Fig. S9<sup>†</sup> suggests a smaller charge-transfer resistance of LMO-CM. Secondly, the novel mesoporous structure constructed by bigger nanoplates and distributed smaller nanoplates can ensure intimate contact between the electrolyte and electrode materials, thus providing more sites for Li<sup>+</sup> storage.<sup>13,14,50,51</sup> Finally, the thickness of all existing nanoplates is only tens of nanometers and such small dimension can support to shorten the pathways for Li<sup>+</sup>

diffusion.<sup>52–54</sup> Therefore, all the above factors enable the obtained LiMn<sub>2</sub>O<sub>4</sub> material with superior rate capability and remarkable cycling stability, which provides a perspective on the potential applications in high-power electric vehicles and smart grids.

## Conclusions

In summary, by employing the hydrated layered Li-deficient and spinel Li-rich composite, we have successfully synthesized a carbon-coated mesoporous LiMn<sub>2</sub>O<sub>4</sub> *via* a facile hydrothermal process and subsequent thermal treatment. Remarkably, the as-prepared LiMn<sub>2</sub>O<sub>4</sub> exhibits superior high-rate capability and long-life cycling stability, delivering the initial discharge capacity of 117.8 mA h g<sup>-1</sup> at 30C with over 90% capacity retention after 1500 cycles. Additionally, this work paves a fresh route to obtain pure and high-performance LiMn<sub>2</sub>O<sub>4</sub> with mesoporous structure by utilizing layered birnessite-type manganese oxides. The innovative employment of hydrated layered Li-deficient and spinel Li-rich intermediate product might provide inspiration to high-performance LiMn<sub>2</sub>O<sub>4</sub> synthesized from other methods or other high-performance electrode materials with multiple layer architecture and optimized phase composition.

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## Notes and references

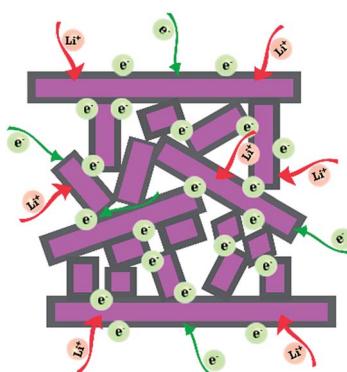


Fig. 5 Schematic illustration of electron conduction and Li<sup>+</sup> storage in LMO-CM.

- 1 R. Chen, T. Zhao, X. Zhang, L. Li and F. Wu, *Nanoscale Horiz.*, 2016, **1**, 423–444.
- 2 N.-S. Choi, Z. Chen, S. A. Freunberger, X. Ji, Y.-K. Sun, K. Amine, G. Yushin, L. F. Nazar, J. Cho and P. G. Bruce, *Angew. Chem., Int. Ed.*, 2012, **51**, 9994–10024.
- 3 V. Etacheri, R. Marom, R. Elazari, G. Salitra and D. Aurbach, *Energy Environ. Sci.*, 2011, **4**, 3243.
- 4 J. B. Goodenough and Y. Kim, *Chem. Mater.*, 2010, **22**, 587–603.
- 5 S. Myung, K. Amine and Y. Sun, *J. Power Sources*, 2015, **283**, 219–236.
- 6 R. Pitchai, V. Thavasi, S. G. Mhaisalkar and S. Ramakrishna, *J. Mater. Chem.*, 2011, **21**, 11040.
- 7 B. Xu, D. Qian, Z. Wang and Y. S. Meng, *Mater. Sci. Eng., R*, 2012, **73**, 51–65.
- 8 F. Mao, W. Guo and J. Ma, *RSC Adv.*, 2015, **5**, 105248.
- 9 S. Lee, Y. Cho, H. Song, K. T. Lee and J. Cho, *Angew. Chem., Int. Ed.*, 2012, **51**, 8748–8752.
- 10 K. Zhang, X. Han, Z. Hu, X. Zhang, Z. Tao and J. Chen, *Chem. Soc. Rev.*, 2015, **44**, 699–728.
- 11 G. Xu, Z. Liu, C. Zhang, G. Cui and L. Chen, *J. Mater. Chem. A*, 2015, **3**, 4092.
- 12 O. K. Park, Y. Cho, S. Lee, H.-C. Yoo, H.-K. Song and J. Cho, *Energy Environ. Sci.*, 2011, **4**, 1621–1633.



13 H. Lin, J. Hu, H. Rong, Y. Zhang, S. Mai, L. Xing, M. Xu, X. Li and W. Li, *J. Mater. Chem. A*, 2014, **2**, 9272.

14 J. Luo, Y. Wang, H. Xiong and Y. Xia, *Chem. Mater.*, 2007, **19**, 4791–4795.

15 W. Sun, H. Liu, Y. Liu, G. Bai, W. Liu, S. Guo and X. Zhao, *Nanoscale*, 2015, **7**, 13173–13180.

16 H. Zhang, Z. Li, S. Yu, Q. Xiao, G. Lei and Y. Ding, *J. Power Sources*, 2016, **301**, 376–385.

17 H. Li and H. Zhou, *Chem. Commun.*, 2012, **48**, 1201–1217.

18 A. R. Han, T. W. Kim, D. H. Park, S.-J. Hwang and J.-H. Choy, *J. Phys. Chem. C*, 2007, **111**, 11347–11352.

19 T. Yi, Y. Zhu, X. Zhu, J. Shu, C. Yue and A. Zhou, *Ionics*, 2009, **15**, 779–784.

20 H.-W. Lee, P. Muralidharan, R. Ruffo, C. M. Mari, Y. Cui and D. K. Kim, *Nano Lett.*, 2010, **10**, 3852–3856.

21 J. Luo, H. Xiong and Y. Xia, *J. Phys. Chem. C*, 2008, **112**, 12051–12057.

22 Y. Ding, J. Xie, G. Cao, T. Zhu, H. Yu and X. Zhao, *Adv. Funct. Mater.*, 2011, **21**, 348–355.

23 J. Lu, X. Fan, C. Zhou, Z. Liu, F. Zheng, K. S. Lee and L. Lu, *J. Electrochem. Soc.*, 2016, **163**, 197–202.

24 L. He, S. Zhang, X. Wei, Z. Du, G. Liu and Y. Xing, *J. Power Sources*, 2012, **220**, 228–235.

25 F. v. Bülow, H. Zhang and D. E. Morse, *Adv. Energy Mater.*, 2012, **2**, 309–315.

26 X. Lv, S. Chen, C. Chen, L. Liu, F. Liu and G. Qiu, *Solid State Sci.*, 2014, **31**, 16–23.

27 M. Tang, A. Yuan and J. Xu, *Electrochim. Acta*, 2015, **166**, 244–252.

28 B. Zou, X. Ma, Z. Tang, C. Ding, Z. Wen and C. Chen, *J. Power Sources*, 2014, **268**, 491–497.

29 R. M. McKenzie, *Mineral. Mag.*, 1971, **38**, 493–502.

30 V. V. Vol'Khin, O. A. Pogodina and G. V. Leont'eva, *Russ. J. Gen. Chem.*, 2002, **72**, 173–177.

31 X. Yang, W. Tang, Z. Liu, Y. Makita and K. Ooi, *J. Mater. Chem.*, 2002, **12**, 489–495.

32 L. Dang, C. Wei, H. Ma, Q. Lu and F. Gao, *Nanoscale*, 2015, **7**, 8101.

33 M. Qin, H. Zhao, W. Yang, Y. Zhou and F. Li, *RSC Adv.*, 2016, **6**, 23905.

34 A. K. Thapa, B. Pandit, R. Thapa, T. Luitel, H. S. Paudel, G. Sumanasekera, M. K. Sunkara, N. Gunawardhana, T. Ishihara and M. Yoshio, *Electrochim. Acta*, 2014, **116**, 188–193.

35 T. L. Christiansen, E. D. Bøjesen, M. Søndergaard, S. Birgisson, J. Becker and B. B. Iversen, *CrystEngComm*, 2016, **18**, 1996–2004.

36 B. J. Liddle, S. M. Collins and B. M. Bartlett, *Energy Environ. Sci.*, 2010, **3**, 1339.

37 L. Liu, Q. Feng, K. Yanagisawa, G. Bignall and T. Hashida, *J. Mater. Sci.*, 2002, **37**, 1315–1320.

38 Q. Feng, K. Yanagisawa and N. Yamasaki, *Chem. Commun.*, 1996, **14**, 1607–1608.

39 X. Huang, Q. Zhang, J. Gan, H. Chang and Y. Yang, *J. Electrochem. Soc.*, 2011, **158**, 139–145.

40 K. A. M. Ahmed and K. Huang, *Solid State Sci.*, 2014, **30**, 11–16.

41 X. Zhang, P. Yu, H. Zhang, D. Zhang, X. Sun and Y. Ma, *Electrochim. Acta*, 2013, **89**, 523–529.

42 Z. Zhang, G. Hu, Y. Cao, J. Duan, K. Du and Z. Peng, *RSC Adv.*, 2015, **5**, 81461–81467.

43 J. W. Lee, A. S. Hall, J.-D. Kim and T. E. Mallouk, *Chem. Mater.*, 2012, **24**, 1158–1164.

44 H. Luo, Y. Sun, F. Zhang, D. Zhang, Y. Yang and J. Zhang, *Electrochim. Acta*, 2014, **148**, 228–236.

45 Q. Feng, K. Yanagisawa and N. Yamasaki, *J. Porous Mater.*, 1998, **5**, 153–161.

46 T. Ohzuku, M. Kitagawa and T. Hirai, *J. Electrochem. Soc.*, 1990, **137**, 769–775.

47 K. K. A. G. W. Liu, *J. Electrochem. Soc.*, 1998, **145**, 459–465.

48 J. Wang, J. Yang, Y. Tang, R. Li, G. Liang, T.-K. Sham and X. Sun, *J. Mater. Chem. A*, 2013, **1**, 1579.

49 S. Xin, Y. Guo and L. Wan, *Acc. Chem. Res.*, 2012, **45**, 1759–1769.

50 J. Wang, W. Liu, S. Liu, J. Chen, H. Wang and S. Zhao, *Electrochim. Acta*, 2016, **188**, 645–652.

51 C. Zhu, G. Saito and T. Akiyama, *J. Mater. Chem. A*, 2013, **1**, 7077.

52 Y. Chen, K. Xie, Y. Pan and C. Zheng, *J. Power Sources*, 2011, **196**, 6493–6497.

53 M. Okubo, Y. Mizuno, H. Yamada, J. Kim, E. Hosono, H. Zhou, T. Kudo and I. Honma, *ACS Nano*, 2010, **4**, 741–752.

54 K. M. Shaju and P. G. Bruce, *Chem. Mater.*, 2008, **20**, 5557–5562.

