



Cite this: *RSC Adv.*, 2018, 8, 26325

High-pressure synthesis and electrochemical properties of tetragonal LiMnO_2 †

Takeshi Uyama, ^{*,a} Kazuhiko Mukai ^a and Ikuya Yamada ^b

Tetragonal structured LiMnO_2 (t- LiMnO_2) samples were synthesized under pressures above 8 GPa and investigated as a positive electrode material for lithium-ion batteries. Rietveld analyses based on X-ray diffraction measurements indicated that t- LiMnO_2 belongs to a γ - LiFeO_2 -type crystal structure with the $I4_1/amd$ space group. The charge capacity during the initial cycle was 37 mA h g^{-1} at 25°C , but improved to 185 mA h g^{-1} at 40°C with an average voltage of 4.56 V vs. Li^+/Li . This demonstrated the superiority of t- LiMnO_2 over other lithium manganese oxides in terms of energy density. The X-ray diffraction measurements and Raman spectroscopy of cycled t- LiMnO_2 indicated an irreversible transformation from the γ - LiFeO_2 -type structure into a $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel structure by the displacement of 25% of the Mn ions to vacant octahedral sites through adjacent octahedral sites.

Received 1st May 2018
 Accepted 14th July 2018

DOI: 10.1039/c8ra03722a

rsc.li/rsc-advances

Introduction

Lithium manganese oxides (LMOs) have been extensively studied as a potential positive electrode material for lithium-ion batteries (LIBs), due to their low cost and environmental friendliness.^{1–6} Since such properties are crucial for large-scale applications for LIBs such as electric vehicles and grid energy storage systems,⁷ research efforts have been devoted to the development of LMOs with high energy densities (W_s). Table 1 summarizes the structure types, synthesis methods, and electrochemical properties of previously reported LMOs, and their crystal structures are shown in Fig. 1a–g. Of these LMOs, a cubic spinel-structured LiMn_2O_4 (Fig. 1a) is widely accepted as the most practical positive electrode material.^{1,2,8} LiMn_2O_4 exhibits a rechargeable capacity (Q_{recha}) of approximately 120 mA h g^{-1} with an average voltage (E_{ave}) of 4.1 V vs. Li^+/Li , and its W approaches the theoretical limit value due to the moderate theoretical capacity (Q_{theo}) of 148 mA h g^{-1} .

In terms of the Q_{theo} , LiMnO_2 and its derivatives are appealing, and their structural and electrochemical properties vary depending on their composition and method of synthesis. A monoclinic layered Li_2MnO_3 , also written as $\text{Li}[\text{Li}_{1/3}\text{Mn}_{2/3}]\text{O}_2$

(Fig. 1b), has a large Q_{theo} of 458 mA h g^{-1} . However, Li_2MnO_3 is essentially electrochemically inactive because of the difficulty in further oxidizing the Mn^{4+} species,⁹ unless a proton exchange¹⁰ and/or an oxygen loss¹¹ in its lattice proceed during the initial charge reaction. An orthorhombic LiMnO_2 (o- LiMnO_2)^{12–18} and a monoclinic LiMnO_2 (m- LiMnO_2)^{4,19} have a formal oxidation state of Mn^{3+} (Fig. 1c and d) and their charge capacities (Q_{cha}) reach a maximum of 230 mA h g^{-1} for o- LiMnO_2 and 270 mA h g^{-1} for m- LiMnO_2 . However, delithiated o- and m- LiMnO_2 can be irreversibly and spontaneously transformed into a $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel; thus, the Q_{recha} values for both of these compounds decrease in a manner similar to that seen in LiMn_2O_4 as the changes in the charge and discharge curves echo those of $\text{Li}_x\text{Mn}_2\text{O}_4$.^{12,15–19} In contrast, O2-type layered $\text{Li}_{2/3}[\text{Li}_{1/6}\text{Mn}_{5/6}]\text{O}_2$ (ref. 20) and $\text{Li}_p[\text{Li}_{1/4}\text{Mn}_{3/4}]\text{O}_2$ with $p < 1$ (ref. 21) (Fig. 1e and f) were reported to display good cycleability without transformation into the $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel. The oxygen stacking, which is ABACAB for $\text{Li}_{2/3}[\text{Li}_{1/6}\text{Mn}_{5/6}]\text{O}_2$ and $\text{Li}_p[\text{Li}_{1/4}\text{Mn}_{3/4}]\text{O}_2$ (unlike the ABCABC stacking seen in m- LiMnO_2), plays an important role in this. As a result, the Q_{recha} is 150 mA h g^{-1} for $\text{Li}_{2/3}[\text{Li}_{1/6}\text{Mn}_{5/6}]\text{O}_2$ and 200 mA h g^{-1} for $\text{Li}_p[\text{Li}_{1/4}\text{Mn}_{3/4}]\text{O}_2$. In the initial cycle, the Q_{cha} ranges between 20 and 50 mA h g^{-1} , and is smaller than the Q_{recha} due to lithium deficiency in these compounds. This triggers a decrease in the Q_{recha} whenever these compounds are used in full cells with negative electrode materials that do not contain residual lithium, e.g. graphite and silicon.

Sugiyama *et al.* synthesized a tetragonal LiMnO_2 (t- LiMnO_2) from o- LiMnO_2 by a high-pressure and high-temperature method, with pressures between 4 and 6 GPa and temperatures between 900 and 1200°C .²² Rietveld analyses based on X-ray diffraction (XRD) measurements showed that t- LiMnO_2 has a γ - LiFeO_2 -type structure with $I4_1/amd$ space group (Fig. 1g). In this structure, MnO_6 octahedra form a three-dimensional

^aToyota Central Research and Development Laboratories, Inc., Nagakute, Aichi, 480-1192, Japan. E-mail: e1599@mosk.tytlabs.co.jp

^bDepartment of Materials Science, Graduate School of Engineering, Osaka Prefecture University, 1-2 Gakuen, Sakai, Osaka 599-8590, Japan

† Electronic supplementary information (ESI) available: Cycle performances at 25°C of the o- LiMnO_2 , LMO (5 GPa), LMO (8 GPa) and LMO (12 GPa) samples, charge and discharge curves of the lithium cells with the LMO (5 GPa) and LMO (8 GPa) samples operated at 40°C , structural parameters of the LMO (5 GPa) and LMO (8 GPa) samples, Raman spectrum of the charged LMO (12 GPa) sample, Raman spectra of the cycled LMO (5 GPa) and LMO (8 GPa) samples, and Raman spectra of PTFE and AB. See DOI: 10.1039/c8ra03722a



Table 1 Crystal structures, synthesis methods, and electrochemical properties of various lithium manganese oxides

Compounds	Structure types (space group)	Synthesis methods	$Q_{\text{theo}}^a/\text{mA h g}^{-1}$	Electrochemical properties	Remarks	References
LiMn_2O_4	Cubic spinel ($Fd\bar{3}m$)	Solid-state reaction	148 ($n = 1$)	$Q_{\text{recha}} \sim 120 \text{ mA h g}^{-1}$, $E_{\text{ave}} \sim 4.1 \text{ V}$	Electrochemically inactive	1, 2 and 8
Li_2MnO_3	Monoclinic layer ($C2/m$)	Solid-state reaction	459 ($n = 2$)	$Q_{\text{recha}} \sim 0 \text{ mA h g}^{-1}$		9 and 10
$(\text{Li}[\text{Li}_{1/2}\text{Mn}_{2/3}\text{O}_2])$	$\beta\text{-NaMnO}_2$ ($Pmmn$)	Solid-state reaction	285 ($n = 1$)	$Q_{\text{cha}} = 180\text{--}230 \text{ mA h g}^{-1}$, $E_{\text{ave}} \sim 3.6 \text{ V}$ (first charge only)	Transformation into spinel phase on the charge reaction	12–18
m-LiMnO ₂	$\alpha\text{-NaMnO}_2$ ($C2/m$)	Ion exchange method	285 ($n = 1$)	$Q_{\text{cha}} = 270 \text{ mA h g}^{-1}$, $E_{\text{ave}} \sim 3.8 \text{ V}$ (first charge only)	Transformation into spinel phase on the charge reaction	4 and 19
$\text{Li}_{2/3}[\text{Li}_{1/6}\text{Mn}_{5/6}\text{O}_2]$	O2-type layer ($P3m1$)	Ion exchange method	193 ($n = 2/3$)	$Q_{\text{recha}} = 150 \text{ mA h g}^{-1}$, $E_{\text{ave}} \sim 3.2 \text{ V}$	Decrease of the Q_{recha} in a full cell	20
$\text{Li}_p[\text{Li}_{1-4p}\text{Mn}_{3/4}\text{O}_2]$ ($p < 1$)	O2-type layer ($P6_3mc$)	Ion exchange method	<327 ($n < 1$)	$Q_{\text{recha}} = 200 \text{ mA h g}^{-1}$, $E_{\text{ave}} \sim 3.2 \text{ V}$	Decrease of the Q_{recha} in a full cell	21
t-LiMnO ₂	$\gamma\text{-LiFeO}_2$ ($I4_1/amd$)	High-pressure and high-temperature method	285 ($n = 1$)	Unknown		This work

^a Theoretical capacity (Q_{theo}) is calculated assuming that a charge-transfer number is n .

framework by sharing their edges, resulting in a straight channel of Li^+ ions along the [111] direction. This configuration between MnO_6 octahedra and Li^+ ions is clearly different from the framework composed of two-dimensional MnO_6 layers in the LiMnO_2 -based compounds mentioned above (see Fig. 1b–f); rather, the crystal structure of t- LiMnO_2 is comparable to that of LiMn_2O_4 . Therefore, in addition to the high Q_{theo} (285 mA h g^{-1}), t- LiMnO_2 is also a desirable material for understanding the relationships between the structural and electrochemical properties in a group of LMOs.

Despite its unique crystallographic character, the electrochemical properties of t- LiMnO_2 remain unclear. This is likely due to the difficulty in conducting large-scale synthesis *via* the high-pressure method because a tiny sample container is generally used to generate high pressure ($>10 \text{ GPa}$).²³ However, recent technological developments, including the use of belt-type and multi-anvil-type equipment, enabled us to evaluate a variety of functional materials.^{24–27} In the current study, we synthesized t- LiMnO_2 under high pressures up to 12 GPa and reported its electrochemical performance as a positive electrode material for LIBs for the first time, in order to clarify the relationship between the structural and electrochemical properties of t- LiMnO_2 . The Q_{cha} of t- LiMnO_2 reached 185 mA h g^{-1} with an E_{ave} of 4.56 V upon increasing the operating temperature to $40 \text{ }^\circ\text{C}$, which is superior to other LMOs. This information will be helpful in designing advanced LMOs with high W in terms of their structure and composition.

Experimental

Sample preparations

t- LiMnO_2 was synthesized from o- LiMnO_2 by the high-pressure and high-temperature method using a Walker-type apparatus.²⁸ o- LiMnO_2 was first synthesized *via* solid-state reactions. Stoichiometric amounts of $\text{LiOH}\cdot\text{H}_2\text{O}$ (Wako Pure Chemical Industries Ltd.) and Mn_2O_3 (Wako Pure Chemical Industries Ltd.) were mixed in ethanol and the mixture was dried and pressed into a 5 mm-thick pellet with a diameter of 15 mm. The pellet was heated at $1000 \text{ }^\circ\text{C}$ for 12 h under argon gas flow at a heating rate of $200 \text{ }^\circ\text{C h}^{-1}$ and a cooling rate of $1 \text{ }^\circ\text{C min}^{-1}$. The obtained o- LiMnO_2 was crushed and re-pressed into a 5 mm-thick pellet with a diameter of 2.8 mm before being packed into a platinum capsule. The capsule was then placed in a BN insulation sleeve, which was placed in a cylindrical graphite heater. The assembled sample was placed in a (Mg, Co)O octahedral pressure medium with side lengths of 14 mm. The (Mg, Co)O octahedra were slowly compressed to 5, 8, or 12 GPa by eight tungsten carbide truncated 8 mm edges. The compressed samples were then heated at $1000 \text{ }^\circ\text{C}$ for 30 min and subsequently quenched to room temperature, and the pressure was slowly released until an ambient pressure was achieved. Hereafter, the o- LiMnO_2 samples treated at 5, 8, and 12 GPa are represented as LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa), respectively, to avoid a misunderstanding of the actual phase purity. A powder sample of LiMn_2O_4 was also synthesized by heating a mixture of $\text{LiOH}\cdot\text{H}_2\text{O}$ and MnO_2 (Kojundo Chemical Laboratory Co., Ltd.) at $1000 \text{ }^\circ\text{C}$ for 12 h under oxygen gas flow.



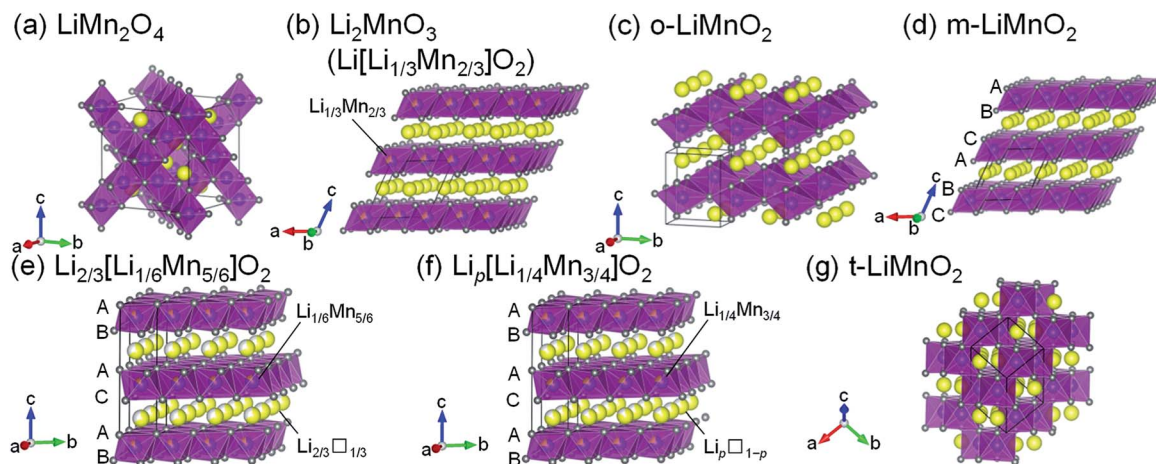


Fig. 1 Crystal structures of seven lithium manganese oxides: (a) LiMn_2O_4 , (b) Li_2MnO_3 ($\text{Li}[\text{Li}_{1/3}\text{Mn}_{2/3}]\text{O}_2$), (c) $o\text{-LiMnO}_2$, (d) $m\text{-LiMnO}_2$, (e) $\text{Li}_{2/3}[\text{Li}_{1/6}\text{Mn}_{5/6}]\text{O}_2$, (f) $\text{Li}_p[\text{Li}_{1/4}\text{Mn}_{3/4}]\text{O}_2$, and (g) $t\text{-LiMnO}_2$. Yellow, purple, and gray spheres indicate Li, Mn, and O atoms, respectively. Unit cells are shown by the solid black lines. The oxygen stacking manner is shown in Fig. 1d–f.

Characterization of the samples

The powder samples of $o\text{-LiMnO}_2$, LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa) were characterized by scanning electron microscopy (SEM), synchrotron XRD measurements, and Raman spectroscopy. The SEM images were recorded using an S-3600N (Hitachi High-Technologies, Company Ltd.) at an accelerating voltage of 15 kV. The XRD measurements were conducted at a BL5S2 beamline available at the Aichi Synchrotron Radiation Center. The samples were packed into borosilicate glass capillary tubes with a diameter of 0.3 mm (WJM-Glas Müller GmbH). The XRD patterns were collected using a two-dimensional detector, PILATUS 100K (Dectris Ltd., Baden-Daettwil), over a 2θ range between 5° and 95° . The incident X-ray wavelength (λ) was determined to be 0.799323(2) or 0.799670(2) Å from the XRD patterns of silicon powders (NIST 640d). Rietveld analyses and drawings of crystal structures were carried out using the RIETAN-FP²⁹ and VESTA softwares,³⁰ respectively. The Raman spectra were collected on an NRS-3300 (Jasco Co. Ltd.) using an excitation laser wavelength of 532 nm and a laser power of 0.1 mW. The duration of exposure was 600 s.

Electrochemical properties

The electrochemical reactivities of the $o\text{-LiMnO}_2$, LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa) samples were examined with sandwich-type two-electrode cells. The powdered sample, acetylene black (AB, Denka Co., Ltd.), and polytetrafluoroethylene (PTFE, Daikin Industries Ltd.) were combined in a ratio of 70 : 20 : 10 to give a viscoelastic mixture, which was then pressed onto an aluminum mesh current collector with a diameter of 16 mm. The mixture functioned as a working electrode. Lithium metal pressed onto 19 mm-wide stainless steel was used as a counter electrode. The electrolyte was made from a solution of 1 M LiPF_6 dissolved in ethylene carbonate (EC)/dimethyl carbonate (DMC) at an EC : DMC ratio of 3 : 7 v/v (Kishida Chemical Company Ltd.) and the separator was constructed from two sheets of porous polyethylene membrane

(Tonen-General Sekiyu K. K.). The cells were assembled in an argon-filled glove box and operated at a current density of 0.025 mA cm^{-2} . The voltage ranged from 1.8 to 4.8 V and the operating temperature was set at 25 or 40 °C.

After cycling test at 25 °C, crystal structures of the samples were investigated using synchrotron XRD and Raman measurements to clarify both macroscopic and microscopic structural changes during the cycling. The cells were charged and discharged twenty times at 25 °C. The cycled samples were then taken from the working electrodes, which had been thoroughly rinsed with a diethyl carbonate solution. Each of the samples was packed into a capillary tube with a diameter of 0.7 mm and put into a quartz glass cell (GL Sciences Inc.) for the XRD measurements and for the Raman spectroscopy, respectively. All the procedures were conducted in the argon-filled glove box, so as not to contact with the atmosphere. The laser power of the Raman spectroscopy was set at 1.0 mW in consideration of absorbance of the quartz cell. The Raman spectra of the charged LMO (12 GPa) sample, LiMn_2O_4 , PTFE, and AB were also measured.

Results and discussion

Morphological and structural characterization

Fig. 2a–d show SEM images for the $o\text{-LiMnO}_2$, LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa) samples, respectively. Particles of $o\text{-LiMnO}_2$ exhibit flake-like shapes with widths between 2–10 μm and thicknesses less than 2 μm . The particle shapes appear to round out and become uniform in size with increasing applied pressure, *i.e.*, rough shapes with a dispersive size of 1–10 μm at 5 GPa and smooth morphologies with a dominant size of $\sim 8 \mu\text{m}$ at 8 GPa and 12 GPa were observed. This morphological change is likely due to a phase transformation, as will be discussed below.

Fig. 3 shows the results of Rietveld analyses for the (a) $o\text{-LiMnO}_2$, (b) LMO (5 GPa), (c) LMO (8 GPa), and (d) LMO (12 GPa) samples. The crystal structure of the $o\text{-LiMnO}_2$ sample was



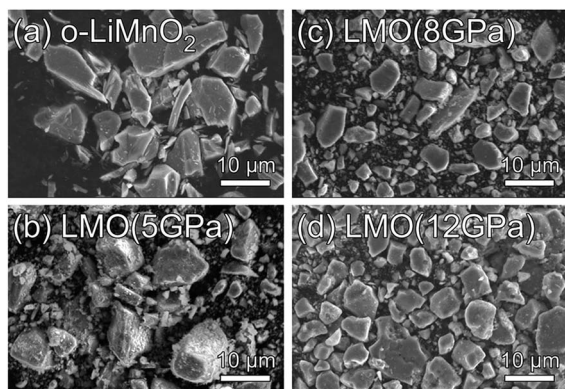


Fig. 2 SEM images for the (a) o-LiMnO₂, (b) LMO (5 GPa), (c) LMO (8 GPa), and (d) LMO (12 GPa) samples.

assigned as an orthorhombic β -NaMnO₂-type structure with *Pmmn* space group; that is, a zigzag layered structure in which Li and Mn atoms occupy each of the octahedral *2b* sites (Fig. 1c). However, the o-LiMnO₂ sample is not in a single phase but contained some impurity phases as shown by asterisk marks at around $2\theta = 10^\circ$ and 16° , and these were found to be hausmannite (Mn₃O₄ with *I4₁/amd* space group) and spinel LiMn₂O₄

by examining the possible crystal structures of LMOs and manganese oxides. Thus, we performed Rietveld analyses with three coexisting phases, in which a Mn under-stoichiometric Li_{1+ δ} Mn_{1- δ} O₂ is adopted to refine the crystal structure of o-LiMnO₂. Here, δ represents the Mn deficiency due to impurities. Moreover, we considered Li and Mn atoms to randomly occupy each of the *2b* sites, as a cation mixing of Li and transition metal ions occasionally occurs in lithium insertion materials.³¹ The structure parameters of the o-LiMnO₂ sample are listed in Table 2. The lattice parameters of o-LiMnO₂ were calculated to be $a_o = 2.80700(2)$ Å, $b_o = 4.57721(2)$ Å, and $c_o = 5.75210(3)$ Å. The actual composition was determined to be Li_{1.005}Mn_{0.995}O₂ ($\delta = 0.005$), where 1.5% of the Mn ions occupied Li (*2b*) sites. The weight fractions were 98.9 wt% for Li_{1.005}Mn_{0.995}O₂, 0.7 wt% for Mn₃O₄, and 0.4 wt% for LiMn₂O₄, giving a Li/Mn ratio of 0.999/1.000, which was consistent with the Li/Mn ratio of the starting material.

For the LMO (5 GPa) sample, the diffraction line at $2\theta = 12.8^\circ$ decreases, but the diffraction line at $2\theta = 12.3^\circ$ clearly generates, due to the formation of the t-LiMnO₂ phase (see the inset of Fig. 3b). As listed in Table S1,[†] the mass fractions of o-LiMnO₂ and t-LiMnO₂ were found to be 86.6 and 8.6 wt%, respectively. For the LMO (8 GPa) and LMO (12 GPa) samples,

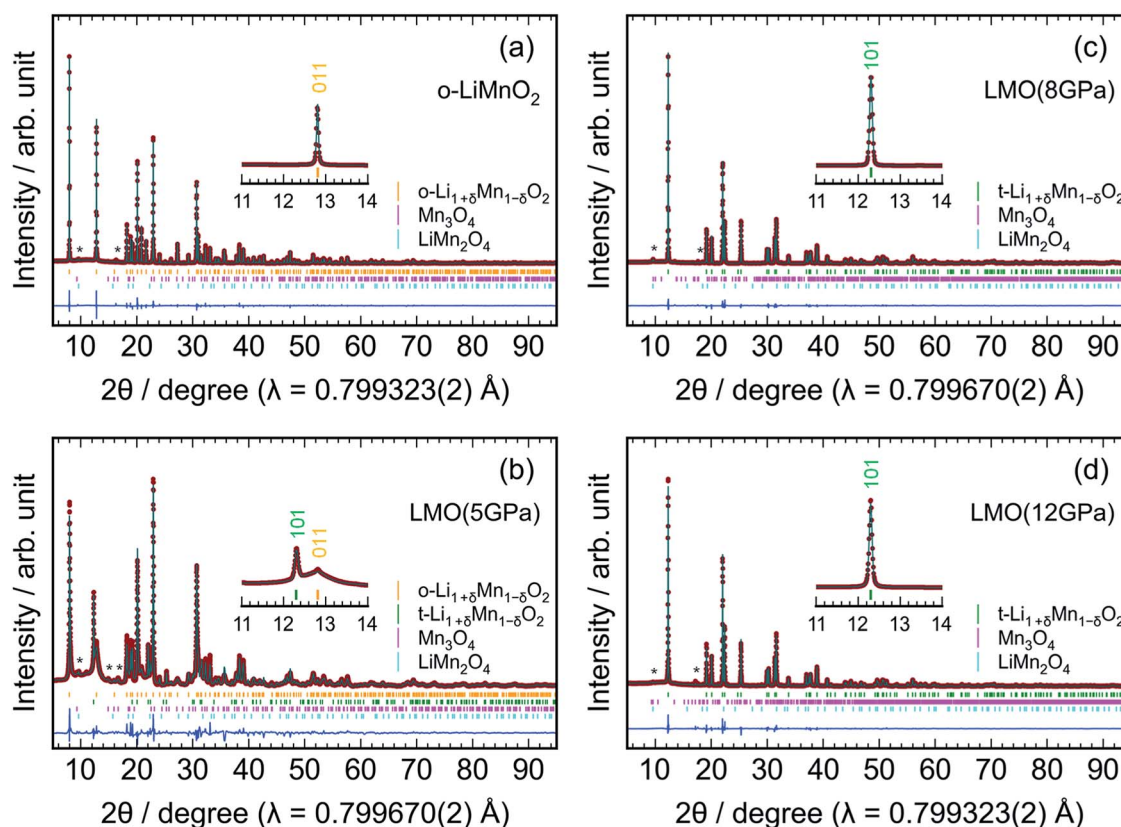


Fig. 3 Rietveld results for the (a) o-LiMnO₂, (b) LMO (5 GPa), (c) LMO (8 GPa), and (d) LMO (12 GPa) samples. Enlarged patterns between 11° and 14° are also shown in the insets to clarify a phase transformation. The orange, green, magenta, and cyan vertical lines indicate the Bragg positions for orthorhombic Li_{1+ δ} Mn_{1- δ} O₂ with *Pmmn* space group, tetragonal Li_{1+ δ} Mn_{1- δ} O₂ with *I4₁/amd*, Mn₃O₄ with *I4₁/amd* or *Pbcm* space group, and LiMn₂O₄ with *Fd $\bar{3}m$* space group, respectively. The diffraction lines from the Mn₃O₄ and LiMn₂O₄ impurities are denoted by asterisks (*). Red dots and solid green lines indicate observed and calculated profiles, respectively, whereas solid blue lines show the differences between the observed and the calculated profiles.



Table 2 Structure parameters of the o-LiMnO₂ sample determined by Rietveld analyses

Phase	Space group	Atom	Wyckoff position	g^a	x	y	z^a	$B^a/\text{\AA}^2$
o-LiMnO ₂	<i>Pmmn</i>	Li1	2 <i>b</i>	0.985(1)	1/4	3/4	0.1188(9)	1.32(8)
		Mn1	2 <i>b</i>	0.015(1)	1/4	3/4	0.1188(9)	1.32(8)
		Mn2	2 <i>b</i>	0.980(1)	1/4	3/4	0.6347(1)	0.55(1)
		Li2	2 <i>b</i>	0.020(1)	1/4	3/4	0.6347(1)	0.55(1)
		O1	2 <i>a</i>	1	1/4	1/4	0.1432(3)	0.58(2)
		O2	2 <i>a</i>	1	1/4	1/4	0.6004(3)	0.58(2)

Composition: Li_{1.005}Mn_{0.995}O₂ ($\delta = 0.005$), $a_o = 2.80700(2)$ Å, $b_o = 4.57721(2)$ Å, and $c_o = 5.75210(3)$ Å

Reliable factors: $R_{wp} = 5.658\%$, $R_p = 4.184\%$, and $S = 0.4501$

Mass fractions: 98.9 wt% for o-LiMnO₂, 0.7 wt% for Mn₃O₄, and 0.4 wt% for LiMn₂O₄

^a Constraints: $g(\text{Mn1}) = 1 - g(\text{Li1})$, $g(\text{Mn2}) = g(\text{Li1}) - \delta$, $g(\text{Li2}) = 1 + \delta - g(\text{Li1})$, $z(\text{Mn1}) = z(\text{Li1})$, $z(\text{Li2}) = z(\text{Mn2})$, $B(\text{Mn1}) = B(\text{Li1})$, $B(\text{Li2}) = B(\text{Mn2})$, and $B(\text{O2}) = B(\text{O1})$.

the diffraction line at $2\theta = 12.8^\circ$ disappears, and almost all diffraction lines can be assigned as the tetragonal γ -LiFeO₂-type structure with *I4₁/amd* space group (Fig. 3c and d). The hausmannite impurity also transformed into a CaMn₂O₄-type structure (*Pbcm*) with an applied pressure up to 8 GPa.³² Table 3 shows the structure parameters of the LMO (12 GPa) sample as determined by Rietveld analyses under identical assumptions as the o-LiMnO₂ sample. The tetragonal lattice parameters, a_t and c_t , were determined to be 4.18278(2) and 8.22922(6) Å, respectively. Note that the actual composition of t-LiMnO₂, Li_{1.012}Mn_{0.988}O₂ ($\delta = 0.012$), was almost identical to that of o-LiMnO₂. From the occupancy factor, g , it was shown that Mn ions occupy 0.8% of the Li (4*a*) sites. As shown in Table S3,[†] there were no significant differences in the structural parameters, including the a_t , c_t , composition, atomic coordination, and g between the LMO (8 GPa) and LMO (12 GPa) samples.

The change in the XRD patterns indicated that o-LiMnO₂ gradually transformed into t-LiMnO₂ at 5 GPa, with the phase transformation occurring as the pressure reached 8 GPa. Conversely, Sugiyama *et al.* reported that this phase transformation was completed at a pressure of 5 GPa in Mn over-stoichiometric Li_{0.93}Mn_{1.07}O₂.²² Therefore, the transformation from o-LiMnO₂ to t-LiMnO₂ is very sensitive to the Li/Mn ratio of o-LiMnO₂; Mn under-stoichiometric Li_{1.1}Mn_{0.9}O₂ ($\delta = 0.1$) actually transformed into a rock-salt structure rather than into t-LiMnO₂ at 5 GPa and 1000 °C.²²

Fig. 4a–d show Raman spectra of the o-LiMnO₂, LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa) samples, respectively. Factor group analyses³³ predicted twelve Raman bands of 4A_g +

4B_{2g} + 4B_{3g} for o-LiMnO₂ structure and eight Raman bands of A_{1g} + 3B_{1g} + 4E_g for t-LiMnO₂ structure. The Raman spectrum for the o-LiMnO₂ sample shows three major Raman bands at 411, 556, and 657 cm⁻¹ and three minor Raman bands at 191, 363, and 488 cm⁻¹. Although all of the Raman bands are not fully assigned, according to the previous reports,^{34–36} the major Raman bands are attributed to vibrations in the MnO₆ octahedra, that is, the O–Mn–O stretching modes at 657 cm⁻¹, the O–Mn–O bending modes at 556 cm⁻¹, and the Mn–O–Mn deformation modes at 441 cm⁻¹. The Raman spectrum of the LMO (5 GPa) sample is similar with that of the o-LiMnO₂ sample, however, the intensities of the Raman bands at ~411, 556, and 657 cm⁻¹ become weak compared with its original Raman spectrum. Furthermore, new and weak Raman bands are observed at 406, 510, and 617 cm⁻¹, and these Raman bands are clearly observed in the Raman spectra of the LMO (8 GPa) and LMO (12 GPa) samples. There are at least ten Raman bands in the Raman spectrum of the LMO (12 GPa) sample, although the theoretical calculation predicted eight Raman bands. This difference is probably due to the nonstoichiometry and/or impurity phases in the LMO (12 GPa) sample. Anyway Raman spectroscopy clarified that the transformation from o-LiMnO₂ to t-LiMnO₂ is achieved under pressures above 8 GPa, which is consistent with the results of the XRD measurements.

Electrochemical properties

Fig. 5a shows charge and discharge curves of the Li cell with the LiMnO₂ sample, operated at temperature of 25 °C. The cell voltage (E) increases rapidly from an open circuit voltage (~3.1

Table 3 Structure parameters of the LMO (12 GPa) sample determined by Rietveld analyses

Phase	Space group	Atom	Wyckoff position	g^a	x	y	z	$B^a/\text{\AA}^2$
t-LiMnO ₂	<i>I4₁/amd</i>	Li1	4 <i>a</i>	0.992(1)	0	3/4	1/8	1.31(9)
		Mn1	4 <i>a</i>	0.008(1)	0	3/4	1/8	1.31(9)
		Mn2	4 <i>b</i>	0.980(1)	0	1/4	3/8	0.51(1)
		Li2	4 <i>b</i>	0.020(1)	0	1/4	3/8	0.51(1)
		O1	8 <i>e</i>	1	0	1/4	0.1413(1)	1.2(2)

Composition: Li_{1.012}Mn_{0.988}O₂ ($\delta = 0.012$), $a_t = 4.18278(2)$ Å, and $c_t = 8.22922(6)$ Å

Reliable factors: $R_{wp} = 6.403\%$, $R_p = 4.544\%$, and $S = 0.4712$

Mass fractions: 97.8 wt% for t-LiMnO₂, 1.8 wt% for Mn₃O₄, and 0.4 wt% for LiMn₂O₄

^a Constraints: $g(\text{Mn1}) = 1 - g(\text{Li1})$, $g(\text{Mn2}) = g(\text{Li1}) - \delta$, $g(\text{Li2}) = 1 + \delta - g(\text{Li1})$, $B(\text{Mn1}) = B(\text{Li1})$, and $B(\text{Li2}) = B(\text{Mn2})$.



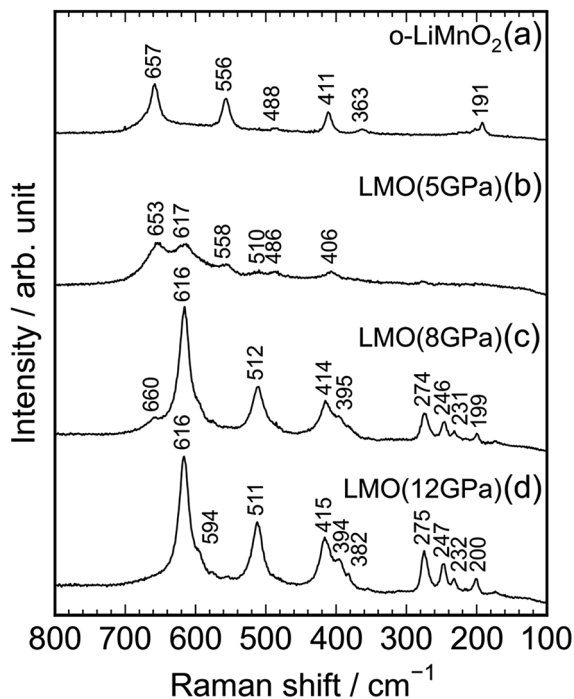


Fig. 4 Raman spectra of the (a) o-LiMnO₂, (b) LMO (5 GPa), (c) LMO (8 GPa), and (d) LMO (12 GPa) samples.

V) before plateauing at ~ 3.5 V when the Q_{cha} reaches 120 mA h g^{-1} . Then, the E climbs to 4.8 V with a gentle gradient during the initial charge. In the discharge curve, however, the E drops sharply to 3.5 V, and gradually decreases to ~ 3.0 V without any clear voltage plateaus. The Q_{cha} is 160 mA h g^{-1} , whereas discharge capacity (Q_{dis}) is only 70 mA h g^{-1} . New voltage plateaus appeared at ~ 3.0 and 4.0 V in subsequent charge and discharge curves. The differences in charge and discharge curves between 1st and subsequent cycles is caused by an irreversible structural transformation to the $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel during the initial cycle, as previously reported.^{12,18,37}

Fig. 5b–d display the charge and discharge curves of the lithium cells with the LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa) samples, respectively. The charge and discharge curves of these three samples are clearly different from those of o-LiMnO₂; *i.e.*, the E monotonically increases from 3.2 V to 4.8 V without any apparent voltage plateaus. The Q_{cha} of LMO (5 GPa) is 154 mA h g^{-1} , while the Q_{cha} values of LMO (8 GPa) and LMO (12 GPa) are 41 and 37 mA h g^{-1} , respectively, which are approximately 13% of the Q_{theo} . The initial discharge curves have a similar E profile with a broad center at ~ 3.0 V. The Q_{dis} values of the LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa) samples are 52, 21, and 22 mA h g^{-1} , respectively. Besides the decline in the E , as similarly noted for o-LiMnO₂, both the charge and discharge curves display multiple voltage plateaus with an increasing cycle number. This implied that t-LiMnO₂ irreversibly transformed into the $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel with the electrochemical cycling. The cycle performances of these four samples are given in Fig. S1.† The Q_{cha} and Q_{dis} values of o-LiMnO₂ [LMO (5 GPa)] increase with the cycle numbers until tenth cycles, and then maintain at ~ 150 (100) mA h g^{-1} . On the

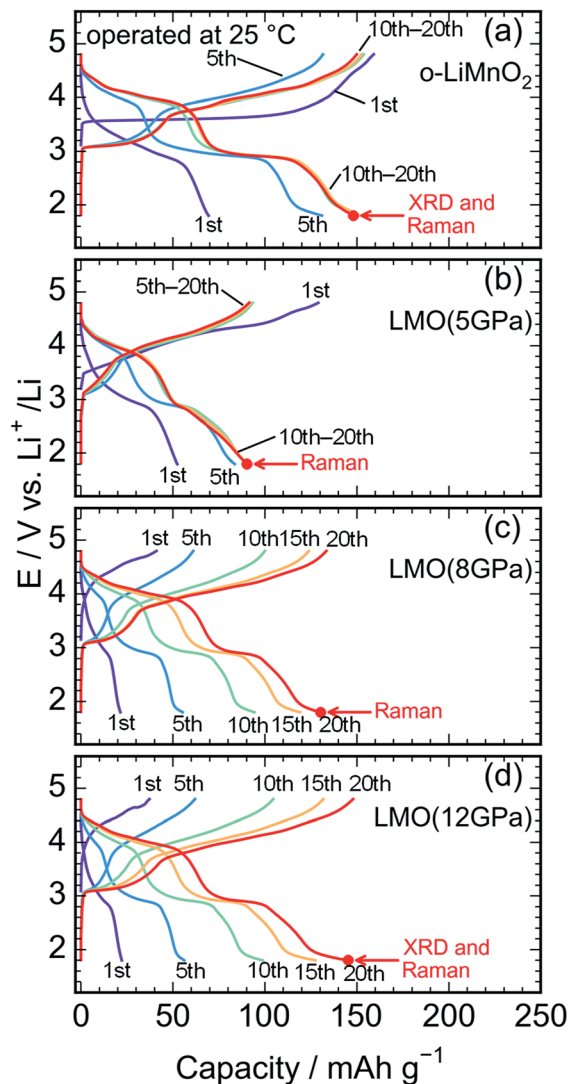


Fig. 5 Charge and discharge curves of the lithium cells with the (a) o-LiMnO₂, (b) LMO (5 GPa), (c) LMO (8 GPa), and (d) LMO (12 GPa) samples. The cells were operated at temperature of 25 °C. *Ex situ* XRD measurements and Raman spectroscopy were conducted at the discharged state, as indicated by the red arrows.

other hand, the Q_{cha} and Q_{dis} values of LMO (8 GPa) and LMO (12 GPa) increase with the cycle numbers up to twenty cycles. The Q_{cha} and Q_{dis} values of LMO (12 GPa) are similar with those of o-LiMnO₂.

Fig. 6a and b show charge and discharge curves of the lithium cells with the o-LiMnO₂ and LMO (12 GPa) samples, respectively, operated at temperature of 40 °C. The initial Q_{cha} of o-LiMnO₂ is approximately 200 mA h g^{-1} , which is larger than the value ($=160 \text{ mA h g}^{-1}$) obtained at 25 °C. This originates from the appearance of a moderate voltage plateau over 4.0 V with a Q_{cha} of $\sim 50 \text{ mA h g}^{-1}$, as reported for an o-LiMnO₂/Li cell at 55 °C by Cho.³⁸ Subsequent charge and discharge curves reveal the increases in Q_{cha} from 200 mA h g^{-1} to 225 mA h g^{-1} and in Q_{dis} from 155 mA h g^{-1} to 220 mA h g^{-1} . The irreversible transformation into the $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel is thought to be accelerated, compared to the measurements at 25 °C.



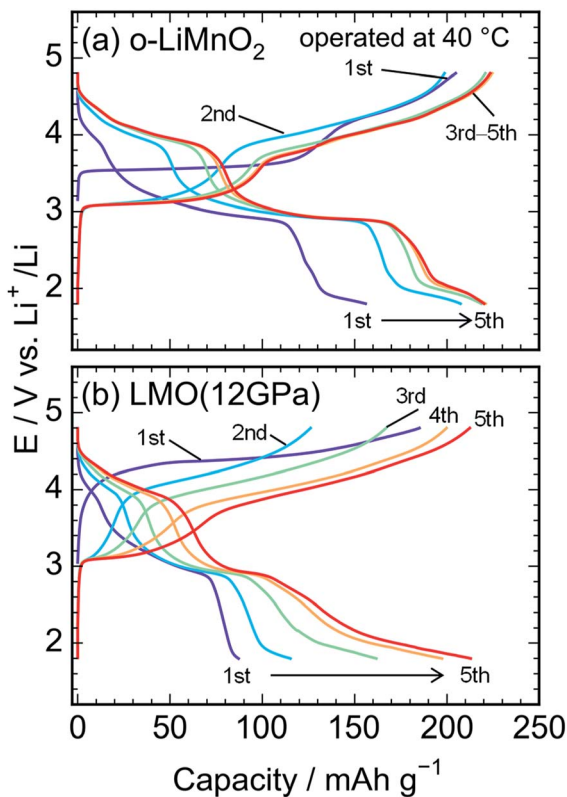


Fig. 6 Charge and discharge curves of the lithium cells with the (a) o-LiMnO₂ and (b) LMO (12 GPa) samples. The cells were operated at temperature of 40 °C.

The E of LMO (12 GPa) slowly increases from an open circuit voltage of approximately 3.0 V to ~ 4.4 V with a Q_{cha} of ~ 40 mA h g⁻¹ during the initial charge. Afterwards, the E remains flat until finally reaching the cut-off voltage of 4.8 V. In contrast to the situation observed at 25 °C, the Q_{cha} increases dramatically to 185 mA h g⁻¹, in comparison to the Q_{theo} of 65%. The redox reaction at approximately 4.4 V is not observed in LMO (5 GPa), whereas it is clearly seen in the LMO (8 GPa) (see Fig. S2a and b†). According to the first-principle calculations and numerical simulations reported by Ceder's group,^{39–42} the Li⁺ ion diffusion barrier for γ -LiFeO₂-type structured materials is higher than that for other positive electrode materials such as LiCoO₂ with an α -NaFeO₂-type structure and LiMn₂O₄ with a spinel-type structure. Moreover, the Mn ions that occupied 0.8% of the 4a lithium sites (Table 3) would block the lithium diffusion path at 25 °C, as reported for LiFePO₄, which has a one-dimensional lithium ion path along the b axis.⁴³ Therefore, by increasing the temperature up to 40 °C, Li⁺ ions could be kinetically removed from the lattice during the initial charge reaction. Moreover, the initial charge curve exhibited an E_{ave} of 4.56 V, indicating that the Mn³⁺/Mn⁴⁺ redox potential in t-LiMnO₂ was superior to that of o-LiMnO₂ ($E_{\text{ave}} = 3.86$ V). Since the W for positive electrode materials is calculated as a product of the Q_{recha} and E_{ave} , the W for t-LiMnO₂ during the initial charge was estimated to be 844 mW h g⁻¹ using a $Q_{\text{cha}} = 185$ mA h g⁻¹ and $E_{\text{ave}} = 4.56$ V. This value was much larger than the W for LiMn₂O₄ (~ 500 mW h g⁻¹) which is already

commercially available as the positive electrode material.⁸ A high E_{ave} is favorable for the potential application of t-LiMnO₂ in this field.

The initial discharge curve, by contrast, could not maintain this E_{ave} and exhibited a Q_{dis} of 87 mA h g⁻¹, resulting in a W value of 273 mW h g⁻¹. This was due to the type of polarization, which was similar to that seen in the results obtained at 25 °C. With an increasing cycle number, the Q_{cha} and Q_{dis} improved to approximately 210 mA h g⁻¹. Nevertheless, the E_{ave} was about 2.9 V at discharge and 4.0 V at charge over the course of more than two cycles. The W in the charge and discharge curves during the fifth cycle was 815 mW h g⁻¹ and 618 mW h g⁻¹, respectively.

Crystal structure change upon electrochemical cycling

Ex situ XRD measurements and Raman spectroscopy were conducted on the cycled electrodes to understand the phase transformation of t-LiMnO₂ during cycling. The XRD patterns of the cycled o-LiMnO₂ and LMO (12 GPa) samples are shown in Fig. 7a and b, respectively. The XRD pattern of o-LiMnO₂ is assigned as a mixture of the Li _{x} Mn₂O₄ spinel ($Fd\bar{3}m$) with $x < 1$, lithiated Li _{y} Mn₂O₄ tetragonal ($I4_1/amd$) phase with $y > 1$, and PTFE originated from the binder in the electrode. The lattice parameters, which were calculated using the least squares method with more than five non-overlapping diffraction lines, are found to be $a_c = 8.224(9)$ Å for Li _{x} Mn₂O₄, and $a_t = 5.666(6)$ Å and $c_t = 9.151(9)$ Å for Li _{y} Mn₂O₄. Since these lattice parameters correspond to those for Li_{0.98}Mn₂O₄ ($a_c = 8.230$ Å) and Li_{1.82}Mn₂O₄ ($a_t = 5.654$ Å and $c_t = 9.202$ Å),⁴⁴ the x and y values are estimated to be 0.98 and 1.82, respectively.

As seen in Fig. 7b, the XRD pattern of the cycled LMO (12 GPa) sample is assigned as a mixture of the Li _{x} Mn₂O₄ spinel, the Li _{y} Mn₂O₄ tetragonal, t-Li₂MnO₂, and PTFE. The evolution of the Li _{x} Mn₂O₄ spinel and Li _{y} Mn₂O₄ tetragonal phases confirms the transformation to the spinel structure from t-LiMnO₂ during cycling, as in the case for o-LiMnO₂. The lattice parameters were calculated to be $a_c = 8.252(4)$ Å for Li _{x} Mn₂O₄, $a_t = 5.657(4)$ Å and $c_t = 9.173(7)$ Å for Li _{y} Mn₂O₄, and $a_t = 4.1854(8)$ Å and $c_t = 8.230(2)$ Å for t-Li₂MnO₂. Thus, the x , y , and z values were estimated to be 0.98, 1.82, and ~ 1 , respectively.

As seen in Fig. S3,† the Raman spectrum of the charged LMO (12 GPa) sample is similar with that of the pristine LMO (12 GPa) sample, except for the Raman band at 654 cm⁻¹. This indicates that the local structure of LMO (12 GPa) is maintained during the initial charge reaction. However, Fig. 8 clarifies that the extended twenty cycle test converts the t-LiMnO₂ structure into the spinel structure. That is, there are only three broad Raman bands at ~ 650 , 620, and 490 cm⁻¹ in the cycled o-LiMnO₂ and LMO (12 GPa) samples, and these Raman spectra are similar with the Raman spectrum of the pristine LiMn₂O₄ sample. The Raman bands at ~ 650 , 620, and 490 cm⁻¹ are also observed in the cycled LMO (5 GPa) and LMO (8 GPa) samples (see Fig. S4†). It should be noted that the Raman spectra of the cycled (or charged) o-LiMnO₂, LMO (5 GPa), LMO (8 GPa), and LMO (12 GPa) samples contain contributions of PTFE and AB, which were used for preparing the working electrodes. As seen



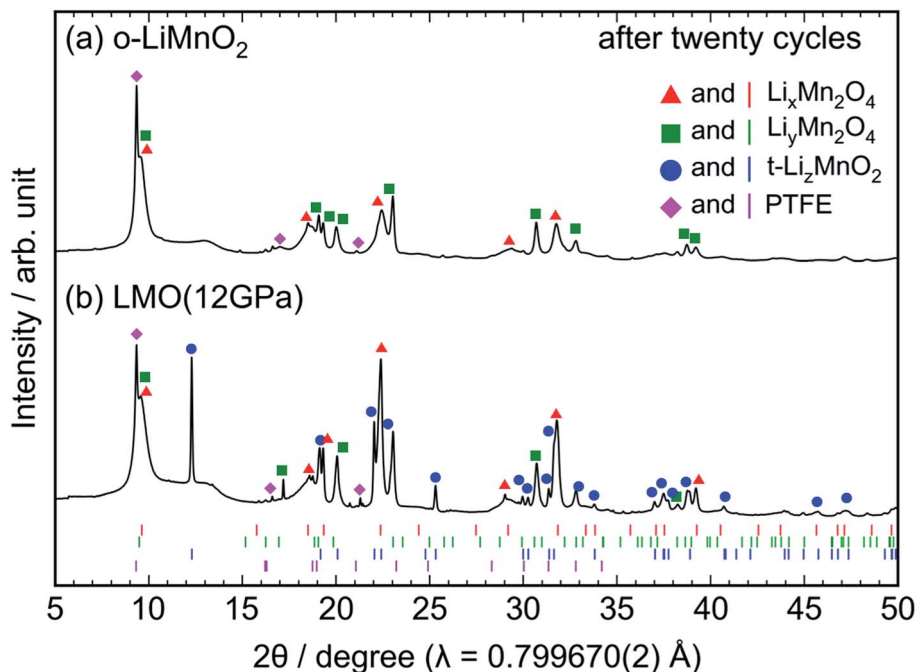


Fig. 7 *Ex situ* XRD patterns of the cycled (a) o-LiMnO₂ and (b) LMO (12 GPa) samples. The measurements were performed at the discharged states, as indicated by the red arrows in Fig. 5a and d. Diffraction lines represented by red triangles, green squares, blue circles, and magenta diamonds, originate from those from Li_xMn₂O₄ with the spinel structure, Li_yMn₂O₄ with tetragonal structure, t-Li_zMnO₂, and PTFE, respectively. The Bragg positions for each of these phases are also shown in the bottom using identically colored vertical lines.

in Fig. S5,[†] the Raman spectrum of PTFE shows three major Raman bands at 733, 385, and 290 cm⁻¹, while that of AB is featureless. The contributions of PTFE and AB are, hence, negligibly small to the Raman spectra of the cycled (or charged) LMO samples.

Fig. 9 shows possible mechanisms of the transformation into the spinel structure from the delithiated (a) o-LiMnO₂ and

(b) t-LiMnO₂. Oxide ions in o-LiMnO₂, t-LiMnO₂, and Li_xMn₂O₄ spinel are arranged in the same ABCABC stacking manner along the [012] direction for o-LiMnO₂, [112] for t-LiMnO₂, and [111] for Li_xMn₂O₄, although the orderings between Mn and vacancy (or Li) octahedral sites are different each other. According to Thackeray *et al.*,¹⁵ Li_xMn₂O₄ is formed from two repeating types of MnO₂ sheets, as shown in the left and right sides of Fig. 9. Thus, there are two ways for the transformation into the spinel from o-LiMnO₂ or t-LiMnO₂. In the case of o-LiMnO₂, the transformation is caused by the displacement of half of the Mn ions to adjacent vacant octahedral sites.^{15–17} By contrast, the transformation of t-LiMnO₂ is achieved by the displacement of a quarter of the Mn ions in the 4b sites into the vacant octahedral sites (4a) *via* adjacent octahedral sites without a rearrangement of the ABCABC oxygen packing (Fig. 9b). In this transformation, t-LiMnO₂ has 50% lower amounts of migrated Mn ions, resulting in approximately twice longer in routes of Mn ions, compared to the case for o-LiMnO₂.

The transformation mechanism of t-LiMnO₂ shows that a suppression of the Mn displacement at the charged state is essential for realizing electrochemical properties such as the Q_{rechar} , E_{ave} , and W in the initial charge state over several cycles. Although the blocking effect of the movement of Mn ions in o-LiMnO₂ has not been reported for the substitution of different cations with Li ions, a small amount of doping could be suitable for t-LiMnO₂ due to the small amount of migrated Mn ions and their long displacement distance. Recently, electrochemical properties of tetragonal structured Li_{0.35}MnO₂ (ref. 45) and Li_{0.59}MnO₂ (ref. 46) nanoparticles were reported; *i.e.*, the initial Q_{cha} and Q_{dis} values of the nanosized Li_{0.35}MnO₂ are 72.9 and

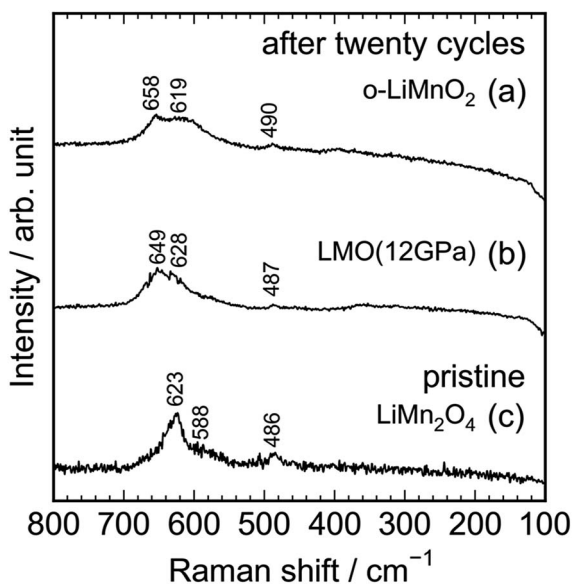


Fig. 8 *Ex situ* Raman spectra of the cycled (a) o-LiMnO₂ and (b) LMO (12 GPa) samples together with the Raman spectrum of the pristine (c) LiMn₂O₄. Raman spectra of (a) and (b) were taken at the discharged state, as indicated by the red arrows in Fig. 5a and d.



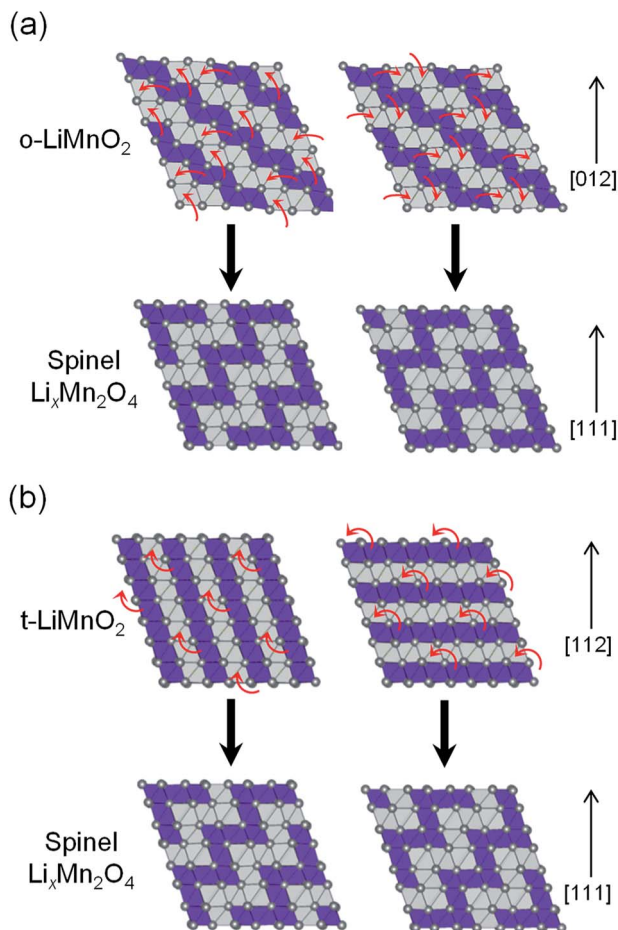


Fig. 9 Schematics of the transformations into $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel: (a) from o- LiMnO_2 and (b) from t- LiMnO_2 . The [012] in o- LiMnO_2 , [112] in t- LiMnO_2 , and [111] in $\text{Li}_x\text{Mn}_2\text{O}_4$ directions correspond to those for oxygen closed-packing array. Purple and gray polyhedra are MnO_6 and LiO_6 (or O_6 : is vacancy) octahedra, respectively. Red arrows represent the Mn migration paths to vacant sites.

178.8 mA h g^{-1} , respectively. The E profile is similar to the discharge curve of the $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel, although the crystal structure of $\text{Li}_{0.35}\text{MnO}_2$ is maintained even after twenty cycles.⁴⁵ Since the electrochemical properties and structural stabilities of t- LiMnO_2 differ from those of $\text{Li}_{0.35}\text{MnO}_2$ (ref. 45) (or $\text{Li}_{0.59}\text{MnO}_2$),⁴⁷ both Li content (x) and particle size also play important roles in stabilizing structure during cycling.

Conclusions

We first investigated the electrochemical performances of t- LiMnO_2 to obtain structural and electrochemical information regarding a series of LiMnO_2 compounds with potential for the development of LMOs with high W . The t- LiMnO_2 samples with γ - LiFeO_2 -type structure ($I4_1/amd$) were prepared from o- LiMnO_2 under pressures up to 8 GPa and at a temperature of 1000 °C, as evident by XRD measurements and Raman spectroscopy. Rietveld analyses indicated that the actual composition of t- LiMnO_2 is $\text{Li}_{1.012}\text{Mn}_{0.988}\text{O}_2$, in which Mn ions partially occupy the 4a lithium sites. The Q_{cha} of t- LiMnO_2 rose to 185 mA h g^{-1} at 40 °C

during the initial charge, even though it was only 37 mA h g^{-1} at 25 °C. Furthermore, the initial charge curve showed that the E_{ave} (4.56 V) with $\text{Mn}^{3+}/\text{Mn}^{4+}$ redox was the highest among LMOs such as o- LiMnO_2 and LiMn_2O_4 . However, the initial discharge curve did not maintain the E_{ave} , and exhibited a Q_{dis} of 87 mA h g^{-1} due to an irreversible phase transformation into the $\text{Li}_x\text{Mn}_2\text{O}_4$ spinel during the initial charge. Substitution of different cations and optimization of Li content and particle size will be necessary to maintain the electrochemical properties of t- LiMnO_2 throughout long cycles.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We appreciate Dr Yuichi Kato of TCRDL for help with Raman spectroscopy. The XRD measurements were conducted on the BL5S2 beamline of the Aichi Synchrotron Radiation Center, Aichi Science & Technology Foundation, Japan (Proposal No. 201702017 and No. 201706077).

Notes and references

- M. M. Thackeray, W. I. F. David, P. G. Bruce and J. B. Goodenough, *Mater. Res. Bull.*, 1983, **18**, 461.
- M. M. Thackeray, P. J. Johnson, L. A. de Picciotto, P. G. Bruce and J. B. Goodenough, *Mater. Res. Bull.*, 1984, **19**, 179.
- M. M. Thackeray, S. H. Kang, C. S. Johnson, J. T. Vaughey, R. Benedek and S. A. Hackney, *J. Mater. Chem.*, 2007, **17**, 3053.
- A. R. Armstrong and P. G. Bruce, *Nature*, 1996, **381**, 499.
- K. Yamaura, Q. Huang, L. Zhang, K. Takada, Y. Baba, T. Nagai, Y. Matsui, K. Kosuda and E. T. Muromachi, *J. Am. Chem. Soc.*, 2006, **128**, 9448.
- M. Freire, N. V. Kosova, C. Jordy, D. Chateigner, O. I. Lebedev, A. Maignan and V. Pralong, *Nat. Mater.*, 2016, **15**, 173.
- O. K. Park, Y. Cho, S. Lee, H.-C. Yoo, H.-K. Song and J. Cho, *Energy Environ. Sci.*, 2011, **4**, 1621.
- M. M. Thackeray, *Prog. Solid State Chem.*, 1997, **25**, 1.
- P. Kalyani, S. Chitra, T. Mohan and S. Gopukumar, *J. Power Sources*, 1999, **80**, 103.
- A. D. Robertson and P. G. Bruce, *Chem. Mater.*, 2003, **15**, 1984.
- D. Y. W. Yu, K. Yanagida, Y. Kato and H. Nakamura, *J. Electrochem. Soc.*, 2009, **156**, A417.
- I. Koetschou, M. N. Richard, J. R. Dahn, J. B. Soupart and J. C. Rousche, *J. Electrochem. Soc.*, 1995, **142**, 2906.
- C.-H. Lu and H.-C. Wang, *J. Eur. Ceram. Soc.*, 2004, **24**, 717.
- Y.-I. Jang, B. Huang, H. Wang, D. R. Sadoway and Y.-M. Chiang, *J. Electrochem. Soc.*, 1999, **146**, 3217.
- R. J. Gummow, D. C. Liles and M. M. Thackeray, *Mater. Res. Bull.*, 1993, **28**, 1249.
- H. Wang, Y.-I. Jang and Y.-M. Chiang, *Electrochem. Solid-State Lett.*, 1999, **2**, 490.



- 17 Y.-M. Chiang, H. Wang and Y.-I. Jang, *Chem. Mater.*, 2001, **13**, 53.
- 18 L. Croguennec, P. Deniard and R. Brec, *J. Electrochem. Soc.*, 1997, **144**, 3323.
- 19 P. G. Bruce, A. R. Armstrong and R. L. Gizendanner, *J. Mater. Chem.*, 1999, **9**, 193.
- 20 J. M. Paulsen, C. L. Thomas and J. R. Dahn, *J. Electrochem. Soc.*, 1999, **146**, 3560.
- 21 N. Yabuuchi, R. Hara, M. Kajiyama, K. Kubota, T. Ishigaki, A. Hoshikawa and S. Komaba, *Adv. Energy Mater.*, 2014, **4**, 1301453.
- 22 J. Sugiyama, T. Noritake, T. Hioki, T. Itoh, T. Hosomi and H. Yamauchi, *Mater. Sci. Eng.*, 2001, **B84**, 224.
- 23 W. B. Holzapfel and N. S. Isaacs, *High-Pressure Techniques in Chemistry and Physics*, Oxford University Press, London, 1997.
- 24 P. F. McMillan, *Nature*, 2002, **1**, 19.
- 25 I. Yamada, H. Fujii, A. Takamatsu, H. Ikeno, K. Wada, H. Tsukasaki, S. Kawaguchi, S. Mori and S. Yagi, *Adv. Mater.*, 2017, **29**, 1603004.
- 26 K. Mukai and I. Yamada, *J. Electrochem. Soc.*, 2017, **164**, A3590.
- 27 I. Yamada, K. Tsuchida, K. Ohgushi, N. Hayashi, J. Kim, N. Tsuji, R. Takahashi, M. Matsushita, N. Nishiyama, T. Inoue, T. Irifune, K. Kato, M. Takata and M. Takano, *Angew. Chem., Int. Ed.*, 2011, **50**, 6579.
- 28 D. Walker, M. A. Carpenter and C. M. Hitch, *Am. Mineral.*, 1990, **75**, 1020.
- 29 F. Izumi and K. Momma, *Solid State Phenom.*, 2007, **130**, 15.
- 30 K. Momma and F. Izumi, *J. Appl. Crystallogr.*, 2011, **44**, 1272.
- 31 Y. Makimura, T. Sasaki, T. Nonaka, Y. F. Nishimura, T. Uyama, C. Okuda, Y. Itou and Y. Takeuchi, *J. Mater. Chem. A*, 2016, **4**, 8350.
- 32 A. F. Reid and A. E. Ringwood, *Earth Planet. Sci. Lett.*, 1969, **6**, 205.
- 33 E. Kroumova, M. I. Aroyo, J. M. Perez-Mato, A. Kirov, C. Capillas, S. Ivantchev and H. Wondratschek, *Phase Transitions*, 2003, **79**, 155.
- 34 E. Kroumova, M. I. Aroyo, J. M. Perez-Mato, A. Kirov, C. Capillas, S. Ivantchev and H. Wondratschek, *Phase Transitions*, 2003, **79**, 155.
- 35 C. M. Julien and M. Massot, *Mater. Sci. Eng.*, 2003, **B100**, 69.
- 36 T.-J. Kim, D. Son, J. Cho and B. Park, *J. Power Sources*, 2006, **154**, 268.
- 37 L. Z. Zhao, Y. W. Chen and G. R. Wang, *Solid State Ionics*, 2010, **181**, 1399.
- 38 J. N. Reimers, E. W. Fuller, E. Rossen and J. R. Dahn, *J. Electrochem. Soc.*, 1993, **142**, 3396.
- 39 J. Cho, *Chem. Mater.*, 2001, **13**, 4537.
- 40 A. Van der Ven and G. Geder, *Electrochem. Solid-State Lett.*, 2000, **3**, 301.
- 41 A. Van der Ven and G. Ceder, *J. Power Sources*, 2001, **97–98**, 529.
- 42 A. Van der Ven, J. Bhattacharya and A. A. Belak, *Acc. Chem. Res.*, 2013, **46**, 1216.
- 43 A. Urban, J. Lee and G. Ceder, *Adv. Energy Mater.*, 2014, **4**, 1400478.
- 44 R. Malik, D. Burch, M. Bazant and G. Ceder, *Nano Lett.*, 2010, **10**, 4123.
- 45 Y.-I. Jang, B. Huang, F. C. Chou, D. R. Sadoway and Y.-M. Chaing, *J. Appl. Phys.*, 2000, **87**, 7382.
- 46 W. Du, Z. Su and Y. Zhang, *Ceram. Int.*, 2016, **42**, 6500.
- 47 M. Ama, Z. Su and H. Pan, *Ceram. Int.*, 2015, **41**, 13887.

