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# Investigating structural properties and reaction mechanism of non-stoichiometric spinel LNMO via solid state NMR†

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In this study, <sup>6,7</sup>Li solid state nuclear magnetic resonance (ssNMR) has been used to characterize local disorder in LiNi<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub> (LNMO) spinel cathodes and to understand their implications for the phase transformation mechanism during electrochemical cycling vs. lithium. By comparing NMR spectra of LNMO samples with slightly varying transition metal stoichiometries, we demonstrate the high sensitivity of ssNMR in the indirect detection of transition metal order and its ability to identify disorder induced by stoichiometric variations. Secondly, we investigate the lithiation reaction mechanism of the non-stoichiometric transition metal-ordered LNMO spinel cathode by *ex situ* <sup>7</sup>Li NMR and highlight that its reaction process involves primarily two successive biphasic reactions. However, our results suggest that stoichiometrically induced transition metal disorder promotes a mixed solid solution and biphasic reaction mechanism. Besides an extended solid solution region, we evince that the biphasic reaction involves the presence of intermediate transient species that make it possible to bridge the miscibility gap between the Li<sub>1</sub> and Li<sub>0.5</sub> phases. These results go beyond our *operando* XRD results, underlining that local structural analysis by <sup>7</sup>Li ssNMR can provide valuable additional insights into the complex reaction mechanism that are difficult to access by other techniques.

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## 1 Introduction

There has been a recent resurgence of interest in the spinel LNMO positive electrode, specifically aiming to establish a systematic comprehension of the relationship between its structural properties and electrochemical performance.<sup>1</sup> At the same time, the study of phase transitions during charging and discharging, and the role of structural order and disorder therein, are of great relevance in the field of electrochemical energy storage materials.<sup>2</sup>

These are crucial not only for understanding the prevailing redox mechanism but also for determining the battery performance, including rate capability and cyclability. Despite more than a decade of research on high-voltage spinel, LiNi<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub>

(LNMO) has yet to unleash its inherent advantages of high energy density (464 W h kg<sup>-1</sup>), high operating potential of 4.7 V (vs. Li/Li<sup>+</sup>) and high rate capability in commercial battery applications.<sup>3,4</sup>

LNMO adopts a spinel arrangement, where Li<sup>+</sup> ions are situated at the tetrahedral sites, and the transition metal ions occupy the octahedral sites within an oxygen cubic close-packed lattice. This configuration promotes rapid lithium diffusion in all three dimensions within the framework.<sup>5,6</sup> The stoichiometric ideal composition of the material entails Ni and Mn to be in the +2 and +4 oxidation states, respectively. In this ideal stoichiometry the extraction of Li ions is achieved through the oxidation of Ni<sup>2+</sup>/Ni<sup>3+</sup>/Ni<sup>4+</sup>, while Mn<sup>4+</sup> ions remain electrochemically inactive. The arrangement of transition metal (TM) cations in the octahedral sites of LiNi<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub> can vary depending on the synthesis conditions.<sup>7,8</sup> In fact, Ni and Mn can either exhibit ordering in the 4a and 12d sites, respectively within a P<sub>4</sub>32 space group or be randomly distributed in the 16d sites of a Fd $\bar{3}m$ -type unit cell.<sup>9,10</sup> The transition metal ordering has been reported to take place when calcining at  $\approx 700$  °C.<sup>10,11</sup> Previous studies have evinced that elevating the calcination temperatures above the transition temperature results in the partial reduction of Mn(IV) to Mn(III), that can be observed from a progressively growing Mn<sup>3+</sup>/Mn<sup>4+</sup> redox plateau

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between 4–4.3 V vs. Li<sup>+</sup>/Li in the electrochemical signature curve.<sup>12–16</sup>

Recent consensus is that the reduction of Mn<sup>4+</sup> to Mn<sup>3+</sup> in LiNi<sub>(II)</sub><sub>0.5–y</sub>Mn<sub>(IV)</sub><sub>1.5–y</sub>Mn<sub>(III)</sub><sub>2y</sub>O<sub>4</sub> serves as a compensatory mechanism for the formation of a secondary rock salt phase resulting in Ni<sup>2+</sup> loss in the spinel, which also explains the release of oxygen, all without necessitating the presence of oxygen defects in the spinel phase.<sup>1</sup>

Previous studies from our group showed that TM ordering in LNMO leads to the formation of nanosized antiphase domains that are detrimental to Li<sup>+</sup> transport, while site mixing in ordered samples leads to performance comparable to that of fully disordered samples.<sup>15,16</sup> Recently, Stüble introduced a more accurate categorisation of “ordered” and “disordered” phases in LNMO spinel including both TM order and TM stoichiometry, rectifying the misconception present in some studies that cation order and Mn(III) content are strictly correlated.<sup>1</sup>

It is general consensus that electrochemical delithiation reaction of LNMO occurs *via* at least three thermodynamically stable phases, which are the fully lithiated LiNi<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub> (Li<sub>1</sub>, Ni<sup>2+</sup>), the half-lithiated Li<sub>0.5</sub>Ni<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub> (Li<sub>0.5</sub>, Ni<sup>3+</sup>), and the fully delithiated Ni<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub> (Li<sub>0</sub>, Ni<sup>4+</sup>) compositions. The ordered Li<sub>x</sub>Ni<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub> undergoes two distinct phase transitions within the range of 1 ≥ x ≥ 0, with their redox potential at around of 4.70 V (Li<sub>1</sub>/Li<sub>0.5</sub>) and 4.74 V (Li<sub>0.5</sub>/Li<sub>0</sub>) versus Li/Li<sup>+</sup>, whereas the disordered spinel only undergoes one phase transition from Li<sub>0.5</sub> to Li<sub>0</sub> and shows solid solution behaviour between Li<sub>1</sub> and Li<sub>0.5</sub>.<sup>6,9,17,18</sup> This is believed to be one of the reasons for the superior rate performance and/or cycle life of disordered LNMO in comparison to the ordered LNMO, as reported in previous studies.<sup>1,14,16,18,19</sup> By varying the synthetic atmosphere and annealing temperature, it was previously shown that small amount of atomic mixing in a partially ordered spinel, was sufficient to significantly improve electrochemical performance aligning to those of completely disordered spinel.<sup>20</sup> Similar observations were also reported by Stüble *et al.* and DeGennaro *et al.* that the partially ordered and disordered shows almost similar electrochemical performance.<sup>1,21</sup> The reaction mechanism and kinetics of LNMO (de)lithiation are significantly affected by various factors, particularly Mn/Ni ordering and the content of Mn<sup>3+</sup>, *i.e.* TM stoichiometry, which are not necessarily coupled as recently shown.<sup>1</sup>

Neutron powder diffraction (NPD) and Raman spectroscopy are well established techniques to detect TM order in LNMO spinel, nevertheless these techniques are unable to precisely quantify their proliferation within the material. While NPD allows quantitative estimation of TM stoichiometry as well as their position within the lattice, it does not provide information on the disorder of local environments experienced by Li nor on the TM oxidation state. Over the past decade, solid-state NMR spectroscopy has emerged as a valuable technique for studying the chemical environments of lithium (Li<sup>+</sup>) and sodium (Na<sup>+</sup>) in cathode materials for batteries.<sup>22–27</sup> The sensitivity of <sup>6,7</sup>Li magic-angle spinning (MAS) NMR is widely recognized for its ability to detect and characterize the local surroundings of lithium atoms. This technique enables direct assessment of

various transition metal (TM) arrangements around lithium, allowing for the quantification and analysis of these distinct environments. In a previous study, Cabana *et al.* utilized <sup>6</sup>Li MAS NMR spectroscopy to examine LiNi<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub> samples synthesized at different temperatures, aiming to discern varying levels of ordering. Similarly, Duncan *et al.* investigated the impact of Ni deficiency in a class of spinel LNMOs LiNi<sub>x</sub>Mn<sub>2–x</sub>O<sub>4</sub> (0.3 ≤ x ≤ 0.5) employing <sup>6</sup>Li NMR to analyze the effect in local ordering of transition metal ions.<sup>10,14</sup> The <sup>6,7</sup>Li MAS NMR technique has also been extensively employed to characterise the chemical structure of positive electrode materials in their pristine, partially lithiated, and delithiated states. This technique provides direct insights into the behaviour of mobile Li<sup>+</sup> ions, which are directly involved in electrochemical processes. Additionally, this technique is quantitative, enabling the tracking and quantification of the species that are either eliminated or retained in the structure during battery cycling.

In our recent *operando* X-ray absorption study we have demonstrated that Ni-deficient partially TM ordered and disordered LNMO undergo similar sequences of redox reactions but differences in their local structure evolution were evinced. The aim of this work is to investigate the local and long-range structure of spinel LNMO in order to understand the role of stoichiometric induced TM disorder on the electrochemical reaction (phase transformation) mechanism. With this aim in mind, we have studied a non-stoichiometric (Ni-deficient) “ordered” spinel LNMO alongside with a near stoichiometric “ordered” LNMO sample by *ex situ* <sup>7</sup>Li NMR and *operando* XRD to understand the phase transformation mechanism at different lithiation stages during charging and discharging and to gain insight into the local disorder induced by Ni deficiency. Neutron powder diffraction (NPD) was used to confirm the cation arrangements, while <sup>6,7</sup>Li magic angle spinning (MAS) NMR was used to study the local environments.

## 2 Experimental

For the labelling of the LNMO samples investigated in this study we use a nomenclature in which the description of LNMO is followed by a letter D or O indicating whether the sample is predominantly transition metal disordered or ordered, this is followed by a two digit suffix stating the Ni stoichiometry in the formula unit per 1 Li, according to the classification recently proposed by Stüble *et al.*<sup>1</sup> The description of LNMO-O-45 would hence translate to a predominantly transition metal ordered spinel with LiNi<sub>0.45</sub>Mn<sub>1.55</sub>O<sub>4</sub>. Two high-purity samples of TM ordered LNMO with two different Mn/Ni stoichiometries (LNMO-O-45 and LNMO-O-48) were provided by Topsoe.

### 2.1 Material characterisation

Neutron powder diffraction (NPD) of pristine and *ex situ* cycled TM ordered (LNMO-O) were carried out at D1B and D2B diffractometer at the Institute Laue-Langevin (ILL), Grenoble France<sup>28–30</sup> using a wavelength of 1.288 and 1.595 Å, respectively. The use of different beamlines resulted in different angular resolution. The samples were mounted on a 6 mm diameter



vanadium sample holder and placed on a carousel that allows automatic sample change. The neutron diffraction patterns were collected at RT with a high statistic to enhance the data accuracy. Rietveld analysis was carried out using the FullProf suite.<sup>31</sup>

Raman spectra of the material was recorded with a Renishaw spectrometer (Nanonics Multiview 2000) operating with an excitation wave-length of 532 nm. Spectra were acquired with 15 s of exposition time of the laser beam to the sample.

The magnetic properties have been characterised using the ACMS option of a PPMS Evercool II instrument from Quantum Design. Magnetic susceptibility has been performed in AC mode, from 5 to 300 K, with an AC field of 10 Oe (see Fig. S1a (left)†). Magnetisation *versus* magnetic field has been performed in DC mode at 5 K, scanning DC magnetic field from -4 to 4 T (see Fig. S1b (right)†); the magnetisation at 4 T is considered to be the value at saturation. Please see SI for more detail on the method used for extracting Mn<sup>3+</sup> excess from magnetisation at saturation.

Micrographs were taken on a Thermo Fisher Quanta 200 FEG high-resolution Scanning Electron Microscope (SEM). The working voltages of the Quanta 200FEG 20 kV and Everhart-Thornley detector (ETD) was used for imaging. Particle sizes were evaluated with a MasterSizer 3000 (Malvern Panalytical, Netherlands).

The samples were analyzed with inductively coupled plasma - atomic emission spectroscopy (ICP-AES) using a Horiba Ultima 2 (Jobin Yvon, Longjumeau, France) in conjunction with a AS500 autosampler and Activanlyst software (version 5.4). The ICP-AES operating conditions were as follows: 1.0 kW of RF power, 13 l min<sup>-1</sup> of a plasma-gas flow rate, 0.2 l min<sup>-1</sup> of a sheath-gas flow rate and 0.25 l min<sup>-1</sup> of a nebulizer-gas flow rate. Solutions were introduced into the plasma torch using nebulizer and a cyclonic type of spray chamber at a flow rate of 0.87 ml min<sup>-1</sup>. Calibration solutions were prepared using a commercial calibration standards of Li, Ni and Mn (Scharlab, Barcelona, Spain) at concentration of 1000 mg l<sup>-1</sup>. Nitric acid 69% analytical grade, Hydrochloric acid 37% Ultratrace from Scharlab (Barcelona, Spain) and Ultrapure Water from Fischer Scientific (Waltham, Massachusetts, USA) were used for dilutions. The most prominent analytical lines of Li 670.784 nm, Ni 216.556 nm and Mn 257.610 nm were selected for measurements. Concentrations of these elements were quantified using four-point external calibration curve within the concentration range of 0.01–100 mg l<sup>-1</sup>.

<sup>6,7</sup>Li Magic Angle Spinning solid state Nuclear Magnetic Resonance (MAS NMR) experiments were performed on a Bruker 300WB spectrometer charged to a field of 4.69 T equipped with a standard 1.3 mm MAS probe. Spinning frequencies were set to 50 kHz. A rotor synchronised spin-echo pulse sequence was used with typical 90 and 180° pulses of 1.3 and 2.6 μs, respectively. A recycle delay of 0.7 s and 2 s was used and around 89k and 1k scans were typically acquired in a <sup>6</sup>Li and <sup>7</sup>Li NMR experiment, respectively. The recycle delays were set to ensure full relaxation of the spins. The spectra were referenced to a 1 M solution of LiCl (set at 0 ppm). The spectra

were analysed and deconvoluted using the Dmfit software package.<sup>32</sup>

<sup>6</sup>Li exhibits smaller quadrupolar and dipolar couplings to the nuclear and electronic spins compared to <sup>7</sup>Li due to its smaller quadrupole moment and gyromagnetic ratio. As a result, <sup>6</sup>Li typically produces higher-resolution NMR spectra with fewer spinning sidebands, enabling a clearer understanding of the local environment. Therefore, we measured <sup>6</sup>Li NMR spectra in our studies of the pristine materials. However, one drawback of <sup>6</sup>Li is its significantly lower natural abundance. Typically for the small sample volumes taken from the cycled cells (*i.e. ex situ* samples), <sup>6</sup>Li enrichment is necessary. Furthermore, no distinction was observed in the NMR resonance for the pristine material between <sup>7</sup>Li and <sup>6</sup>Li, which had no impact on the analysis. Consequently, <sup>7</sup>Li NMR was utilized for the *ex situ* investigation of LNMO-O-45.

## 2.2 Electrode formulation and cell assembly

For the *ex situ* <sup>7</sup>Li NMR and *operando* XRD experiments, LNMO powder samples were mixed in NMP based slurry together with PVdF binder (Kynar HSV900; Arkema, France) and carbon additive (C65; Imerys, Switzerland) in ratio of 93:3:4 and deposited on single side carbon coated Al foil (Armor, France) with active material loading around 10.6 mg cm<sup>-2</sup>. After cutting, pressing and drying, electrodes were mounted in CR2032 coin cells for *ex situ* NMR samples. All reported C-rates are relative to theoretical capacity of 148 mA h g<sup>-1</sup>. The cells were recycled for two complete cycles and then stopped at the desired SoC, the corresponding electrochemical cycling profiles of these *ex situ* samples are depicted in Fig. S6 and S7 (ESI)†. The voltage of the cells were held at the desired SoC until they were rapidly disassembled and the recovered electrode material was transferred to the NMR rotor, to minimise relaxation previously reported.<sup>33</sup> For the *operando* XRD measurements, the slurry was prepared as for NMR in ratio of 84:8:8 and coated on 6 μm Al foil current collector with active material loading around 6.1 mg. After cutting and drying, the positive electrode was assembled in specially designed *in situ* cell for *operando* XRD measurements, recently described elsewhere.<sup>34</sup> The cells were assembled in an argon-filled glove-box (≤0.1 ppm H<sub>2</sub>O and O<sub>2</sub>) with a 16 mm diameter LNMO positive electrode, a quartz fiber separator (QM-A; Whatman) and a 16 mm diameter lithium disc counter-electrode, using LP30 electrolyte.

## 2.3 Operando XRD

*Operando* XRD measurements were performed on a lab-scale Bragg-Brentano diffractometer (Bruker D8 advance, with Cu K<sub>α1,2</sub> tube source and LYNXEYE 1D detector), using a custom-made *in situ* electrochemical cell equipped with Beryllium Windows, previously described elsewhere.<sup>34</sup> Galvanostatic cycling with potential limitation was performed using a Bio-Logic ST-150 potentiostat at a C/n rate (expressed as 1 mol of Li reacted in n hours per mole of LNMO). The electrochemical cycling during *operando* measurements were carried out at C/20 rate within voltage window of 3.5 to 5 V vs. Li<sup>+</sup>/Li in a 2θ range of 8–22° and 35–52°. 20 min patterns were collected sequentially



with an angular step size of  $0.1^\circ$ . The electrochemical tests were carried out at room temperature.

### 3 Results

In the first part, see Section 3.1, two pristine LNMO samples are compared by different material characterisation techniques (NPD, ICP, SEM, Raman, magnetic measurements and  $^6\text{Li}$  NMR) to highlight the influence of slight deviations of the TM stoichiometry on the TM order and its effect on the electrochemical properties. In the second part we characterize the lithiation reaction mechanism of one of the previously examined samples from local and crystal structure perspective by employing *ex situ*  $^7\text{Li}$  NMR and *operando* diffraction analysis, see Section 3.2. These findings are further discussed in Section 4.

#### 3.1 Comparison of two transition metal ordered pristine LNMO

In a first step, two predominantly TM ordered LNMO samples were analyzed by NPD. It is a highly informative technique since it allows not only to determine phase purity and crystal phases but, more importantly, to elucidate the presence of TM ordering and estimate TM stoichiometries, thanks to the sufficiently different neutron cross section of Ni and Mn. In Fig. 1(a) the NPD patterns and Rietveld refinements of the two pristine materials are compared. Selected fitting parameters are presented in Table 1. Single phase TM ordered space group  $P4_332$

Table 1 Selected results of the Rietveld refinements of the NPD data<sup>a</sup>

Sample	Lattice const. [ $\text{\AA}$ ]	$\theta$ crystal size [nm]	Mn/Ni ratio
LNMO-O-45	8.1833(1)	124/6 <sup>b</sup>	3.405(3)
LNMO-O-48	8.16789(7)	420/16 <sup>b</sup>	3.1442(7)

<sup>a</sup> Single phase, space group  $P4_332$  and antiphase model have been used for both samples. <sup>b</sup> Corresponds to antiphase domain size.

with antiphase model as previously proposed<sup>16</sup> was used for the refinement of both samples. The good agreement of the calculated with the experimental pattern underlines the phase purity and the predominantly TM-ordered nature of both LNMO samples. Nevertheless, faint features at  $35.0$  ( $d = 2.14$ ) and  $45.2^\circ$  ( $d = 2.07$ ) which are not comprised in the spinel phase refinement can be made out in the bottom pattern (indicated by arrow markers) which could hint presence of a rocksalt impurity. For both samples, a Ni-deficient stoichiometry was determined from these NPD refinements, resulting in Mn/Ni ratios higher than stoichiometrically expected value of  $3/1$ . In accordance with the nomenclature proposed by Stüble *et al.*,<sup>1</sup> the Ni-deficient LNMO samples with Mn/Ni ratios of 3.41 and 3.14 will be referred to as LNMO-O-45 and LNMO-O-48, respectively. The refinement also reveals larger crystal domains and larger antiphase domains for the LNMO-O-48 sample, which is consistent with the fact that a higher degree of order is achieved

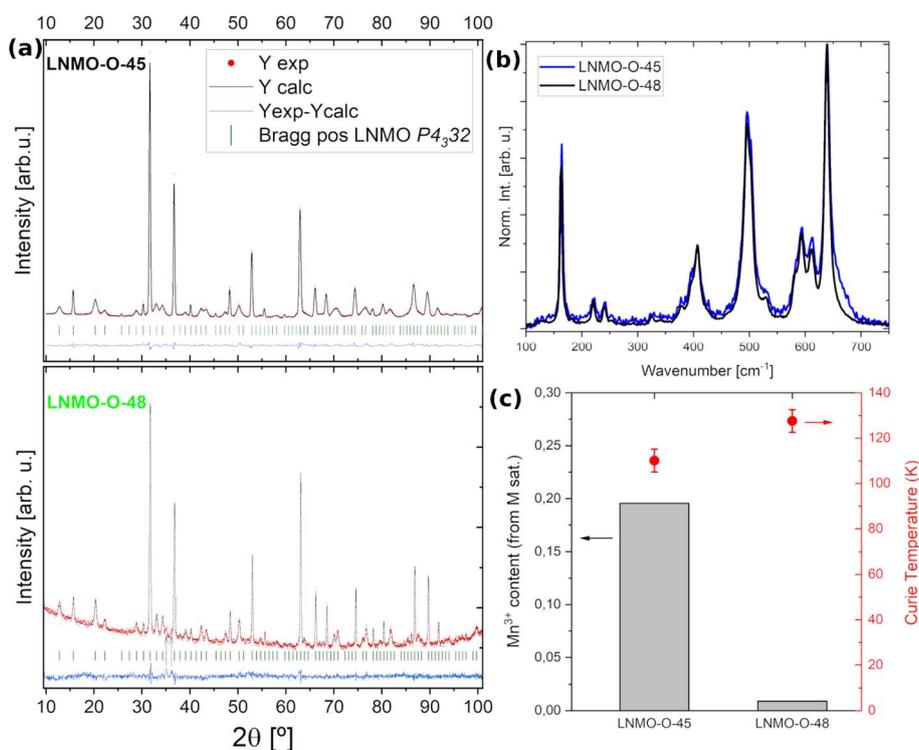


Fig. 1 (a) Rietveld refinement of Neutron diffraction pattern for LNMO-O-45 (top) and LNMO-O-48 (bottom), respectively. Arrow markers in the bottom plot indicate features not comprised in the spinel phase refinement. (b) Raman spectra of pristine LNMO-O-45 and LNMO-O-48. (c) Comparison of  $\text{Mn}^{3+}$  content based on magnetic saturation (grey bars) and Curie temperature (red markers) of LNMO-O-45 and LNMO-O-48.



in a more stoichiometric compositions. The full set of refinement parameters are reported in Table S2.†

ICP analysis was used to support stoichiometry determination. The analysis revealed that the ratio of sum of both transition metals to Li is close to the expected value of 2/1 for both samples, which underlines the assumption that the Ni deficiency is compensated by a Mn excess to maintain charge neutrality. Furthermore, ICP analysis yields to Mn/Ni ratios of 3.4(2) and 3.0(2), for LNMO-O-45 and LNMO-O-48, respectively. This former value corresponds to an excess of Mn of  $y = 0.05$  in  $\text{LiNi}_{(ii)0.5-y}\text{Mn}_{(iv)1.5-y}\text{Mn}_{(iii)2y}\text{O}_4$  for sample LNMO-O-45, in agreement with NPD data. For sample LNMO-O-48, no Ni deficiency was evinced in the overall sample, which could be due to the presence of minor Ni rich rocksalt segregation besides the LNMO-O-48 spinel phase, as hinted by the faint additional NPD feature, *vide supra*.

Scanning Electron Microscopy (SEM) has been carried out to compare particle morphology and size of the two pristine LNMO materials. The micrographs depicted in Fig. S2† illustrate that both materials have similar primary and secondary particle size and shape, consisting of spherical particles with an average diameter of 8  $\mu\text{m}$  which consist of smaller polygonal shaped particles with average size of 1  $\mu\text{m}$ .

Raman spectra of the two pristine TM ordered LNMO samples are depicted in Fig. 1(b) showing that both spectra largely superimpose in peak position and shape. This structural fingerprint indicates that both LNMO samples have same local structure as well as similar degree of structural order. Moreover, both spectra reveal the unique features of TM ordered LNMO at 223, 242 as well as the 593  $\text{cm}^{-1}$ .<sup>35</sup> In the case of the LNMO-O-48 these features stand out more clearly than in the LNMO-O-45. Another noteworthy difference between the two spectra is the more pronounced shoulder peak at  $\approx 665 \text{ cm}^{-1}$ , which has been attributed to bending vibrations of the TM in the 16d sites of the disordered  $Fd\bar{3}m$  phase.<sup>33</sup> The discrepancy between NPD and Raman are a result of the different length scales probed, moreover it should be noted that Raman spectroscopy does not provide quantitative information on TM order.

To maintain the charge neutrality, the Ni-deficient stoichiometry entails an excess of Mn in combination with the

reduction of  $\text{Mn}^{4+}$  to  $\text{Mn}^{3+}$ , resulting in the following formula;  $\text{LiNi}_{(ii)0.5-y}\text{Mn}_{(iv)1.5-y}\text{Mn}_{(iii)2y}\text{O}_4$ . For each deficient  $\text{Ni}^{2+}$ , two  $\text{Mn}^{3+}$  are formed. The evaluation of the  $\text{Mn}^{3+}$  content is hence an indirect way of probing the TM composition. Due to its difference in magnetic moment, the  $\text{Mn}^{3+}$  can be probed *via* its magnetisation saturation. We found  $\approx 0.01 \text{ Mn}^{3+}$  content in the LNMO-O-48 sample and 0.19 of  $\text{Mn}^{3+}$  for the LNMO-O-45, see Fig. 1(c). This difference is in qualitative agreement with the NPD and ICP results, which point towards stronger Ni deficiency and hence more elevated  $\text{Mn}^{3+}$  content in the LNMO-O-45 compared to LNMO-O-48. While for the LNMO-O-48 the determined Mn excess (0.01) lies within stoichiometry determined by ICP and NPD, the Mn excess determined based on Mn saturation of LNMO-O-45 exceeds those indicated by ICP and NPD ( $y = 0.05$  yielding to expected  $\text{Mn}^{3+}$  content of 0.1). This deviation could be due to the fact that the orbital moment is not entirely quenched or to the presence of amorphous  $\text{Mn}_{(iii)}$ -oxide segregation.<sup>36</sup> Interestingly, these magnetic measurements also reveal an elevated Curie temperature  $T_C$  for the LNMO-O-48 which suggests a higher degree of TM ordering, according to previous study.<sup>37</sup>

The impact of TM stoichiometry and ordering can also be observed in the electrochemical signature of the LNMO sample. The electrochemical cycling curve of the LNMO-O-45 along with the corresponding derivative ( $dQ/dV$ ) are shown in Fig. 2. Two processes can be easily observed: a shorter one centered at 4.1 V, referred to as the 4 V capacity, contributing approximately 15  $\text{mA h g}^{-1}$ , and a longer process at  $\approx 4.75 \text{ V}$ , which accounts for the majority of the capacity.

According to previous works, the first minor region at around 4.0 to 4.3 V can be attributed to the  $\text{Mn}^{3+}/\text{Mn}^{4+}$  redox couple.<sup>18,21</sup> This can be used as an indication of  $\text{Mn}^{3+}$  presence in the spinel structure. This is in agreement with the absence and presence of this plateau in this voltage range for the LNMO-O-45 and LNMO-O-48 samples, respectively (see Fig. 2). Continuing the charge reaction above 4.3 V for LNMO-O-45, a slope leads up to an extended plateau, which can be subdivided into two subsequent plateaus which are highlighted in the derivative curve, see Fig. 2 (right). These two plateaus are expressed by two consecutive peaks at 4.72 and 4.76 V vs.  $\text{Li}^+/\text{Li}$

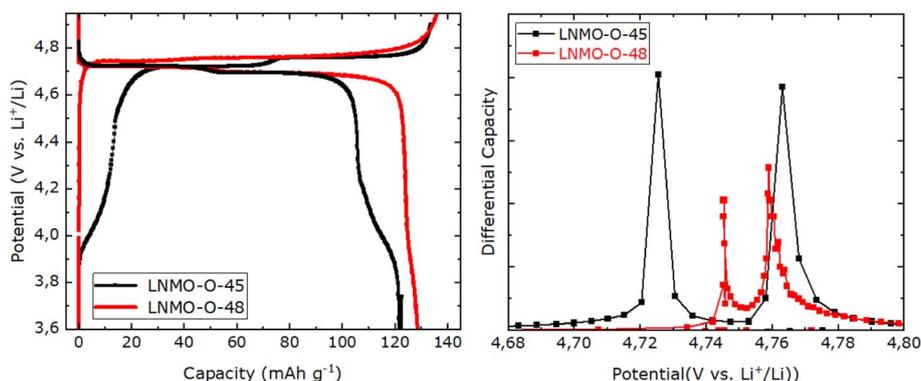


Fig. 2 (Left) Electrochemical cycling curve of LNMO-O-45 (black) and LNMO-O-48 (red) at C/10 vs. Li metal in the third cycle. (Right) Corresponding derivative of the Ni charge redox reaction.



and which are attributed to the redox couples  $\text{Ni}^{2+}/\text{Ni}^{3+}$  and  $\text{Ni}^{3+}/\text{Ni}^{4+}$ , respectively. In contrast, in the charge/discharge curves of near stoichiometric LNMO-O-48, the Ni plateaus merge into a single long flat plateau at 4.7 V reflected by a much narrower voltage step between the  $\text{Ni}^{2+}/\text{Ni}^{3+}$  and  $\text{Ni}^{3+}/\text{Ni}^{4+}$  redox couples.

In the case of the non-stoichiometric TM ordered LNMO (LNMO-O-45), based on capacity values obtained in Fig. 2 (left), the 4 V capacity in the third cycle at 0.1C is around 12% of the total capacity following eqn (1). This is in general agreement with 5% Ni deficiency evinced by NPD and ICP which entail a 10%  $\text{Mn}^{3+}$  content for the LNMO-O-45.

$$\frac{Q_{\text{cha}}^4 \text{V} + (Q_{\text{dis}}^{\text{tot}} - Q_{\text{dis}}^4 \text{V})}{2Q_{\text{dis}}^{\text{tot}}} \quad (1)$$

As reported by Song *et al.*, TM cation ordering influences the voltage of the redox plateaus of nickel, or, more precisely, on the voltage of the  $\text{Ni}^{2+/3+}$  plateau.<sup>12</sup> This effect is difficult to spot in the voltage vs. capacity profile but is clear in the  $dQ/dV$  profile shown in Fig. 2 (right). According to Song and co-workers, the presence of TM order is reflected by the difference in potential between the two Ni redox plateaus. An increase in TM order leads to elevation of the  $\text{Ni}^{2+}/\text{Ni}^{3+}$  redox plateau and hence results in a comparatively smaller voltage gap to the  $\text{Ni}^{3+}/\text{Ni}^{4+}$  plateau which remains unaltered regardless of the TM stoichiometry, as recently demonstrated by Stüble *et al.* by comparing LNMO samples of varied TM order and stoichiometry.<sup>1</sup> For the LNMO-O-45, we observe a gap of 38 mV between the  $\text{Ni}^{2+/3+}$  and the  $\text{Ni}^{3+/4+}$  plateaus in the charge curve. Within the compositions screened by Stüble, this value is in best agreement with the one of their partially ordered LNMO (LNMO-43). Our observed value falls in between their reported values of 20 and 60 mV for the ordered and disordered LNMO, respectively. This is an indication for the coexistence of different TM ordering in the LNMO-O-45. The degree of TM ordering in spinel LNMO can

affect the electronic band structure of the Ni ions, which in turn influences their redox behaviour as observed by different Ni redox plateaus in the voltage curve. A potential reason for this could be the differences in orbital overlap of the Ni ions as function of TM ordering. In this regard a more distinct electronic states of  $\text{Ni}^{2+}$  and  $\text{Ni}^{3+}$  within TM ordered LNMO would result in a higher driving force (voltage) needed for the delithiation compared to TM disordered structure in which a larger orbital overlap of Ni is prevalent.

Moreover, their reported 4 V capacity contribution attributed to  $\text{Mn}^{3+}$  of around 13.7% is in good agreement with our findings for LNMO-O-45 sample underlining similar TM stoichiometry. Similarly, Duncan *et al.* showed the effect of  $\text{Mn}^{3+}$  presence due to Ni deficient TM stoichiometry on the TM ordering by comparing voltage gap of samples with varying degree of Ni deficiency.<sup>14</sup> Based on their voltage gap classification, the expected Ni stoichiometry in our LNMO-O-45 sample falls within the range of 0.4 to 0.45, which aligns well with our results obtained from ICP and NPD analyses.

In Fig. 3 the  $^6\text{Li}$  NMR spectra of the pristine non-stoichiometric LNMO-O-45 and the near-stoichiometric LNMO-O-48 are shown for comparison. In both spectra, a set of signals are observed between 700 and 1100 ppm. These shifts are clearly out of the range of the shifts typically observed in the  $^6\text{Li}$  solid-state NMR spectra of diamagnetic compounds (10 to -10 ppm). The large shifts observed in the spectra of LNMO are typically observed in paramagnetic solids and are due to the Fermi contact interactions between Li ions and the unpaired electrons in the  $t_{2g}$  and  $e_g$  orbitals of the transition-metal ions in the local coordination shell.<sup>10</sup> The magnitude and sign of the induced paramagnetic shifts of the NMR signals are determined by the bond distances and angles between the metals and Li centres, as well as oxidation states and types of orbitals overlapping along the pathway. In this 3d-transition-metal system, we assume the hyperfine shifts are additive, a consequence of minimal spin-orbit coupling; as such, one can decompose the

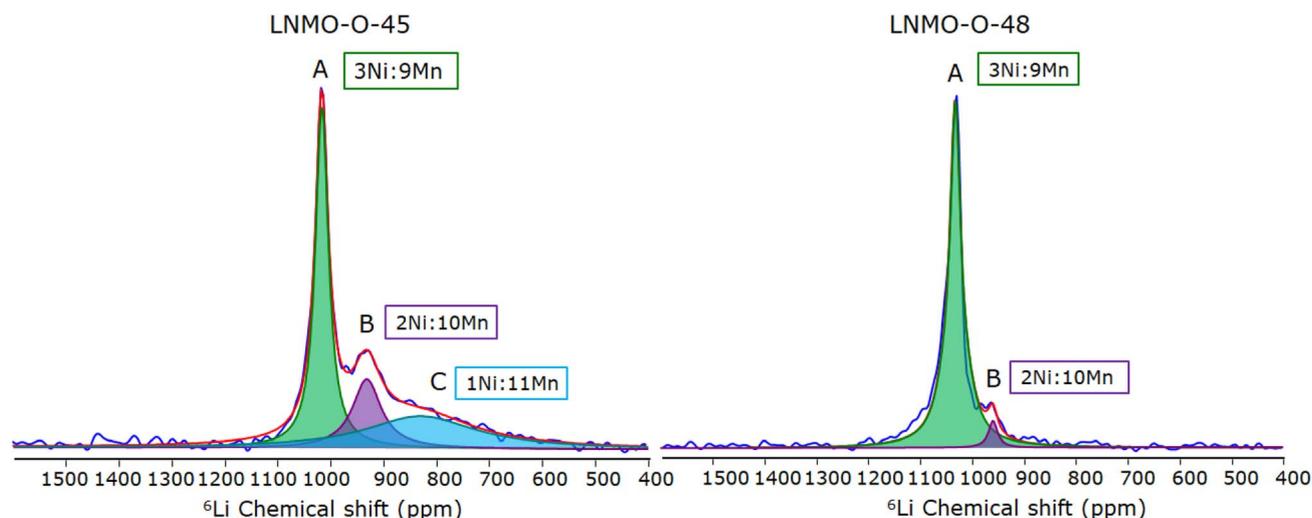


Fig. 3  $^6\text{Li}$  NMR spectra of pristine TM ordered LNMO samples: non-stoichiometric LNMO-O-45 (left) and near-stoichiometric LNMO-O-48 (right).



Table 2 Peak intensity of deconvoluted  ${}^6\text{Li}$  NMR spectra for LNMO-45 and LNMO-48 attributed to different Li environments

TM number <sup>a</sup>		LNMO-O-45			LNMO-O-48		
#Mn	#Ni	${}^6\text{Li}$ shift [ppm]	Int. <sup>b</sup> [%]	LW [ppm]	${}^6\text{Li}$ shift [ppm]	Int. <sup>b</sup> [%]	LW [ppm]
9	3	1017	48	29	1032	95	29
10	2	932	20	73	960	5	20
11	1	830	32	256	—	—	—

<sup>a</sup> Refers to the number of TM atoms in the first shell around Li. <sup>b</sup> Values are not T2-weighted, which could influence absolute integration values.

shift into a set of pathways between Li and nearby transition metal ions.<sup>38</sup> The spectra of LNMO-O-48 and LNMO-O-45 shown in Fig. 3 are both characterised by a dominant signal observed at 1017 and 1032 ppm for LNMO-O-45 and LNMO-O-48, respectively, in agreement with previous  ${}^6\text{Li}$  solid state NMR characterisations of ordered LNMO samples.<sup>10,14,39,40</sup> These signals are assigned to lithium environments with Mn/Ni ratios in the first coordination shell of 3/1. The slightly lower chemical shift of this dominant feature in the LNMO-O-45 sample has its origin in the less oxidised Mn, *i.e.* higher  $\text{Mn}^{3+}$  content.<sup>14</sup> Perfectly TM ordered spinel LNMO would only contain a single lithium environment, hence its NMR spectra should consist of a single signal. Indeed, a dominant single peak at 1032 ppm is observed here for the LNMO-O-48 sample shown in Fig. 3 (right), indicating the presence of a nearly ideal TM order and stoichiometric structure (9Mn : 3Ni). Additional signals at lower ppm values are also observed in both spectra. The chemical shifts and relative intensities of all the signals obtained by deconvolutional fit are summarised in Table 2.

The additional minor signal observed around 962 ppm in LNMO-O-48, indicates that not all the Li in the spinel are surrounded by 3  $\text{Ni}^{2+}$  and 9  $\text{Mn}^{4+}$ . This minor resonance feature can hence be attributed to a Mn-rich Li environment (10Mn : 2Ni; *i.e.* presence of  $\text{Mn}^{3+}$ ) in the first TM coordination shell of Li as indicated in the Table 2. This local Mn excess evinced by Li NMR is congruent to the global Mn excess observed by NPD and magnetic measurements.

Compared to the LNMO-O-48, the spectrum of the LNMO-O-45 shown in Fig. 3 (left) has a lower relative intensity for the signal ascribed to the ideal structure (9  $\text{Mn}^{4+}$  and 3  $\text{Ni}^{2+}$  in the first sphere of coordination) at 1017 ppm, as well as at least two strong additional signals are observed. The presence of multiple strong resonances is a clear indication of TM disorder, as previously demonstrated.<sup>15</sup> As a higher concentration of Mn is present in LNMO-O-45, these additional features observed at lower ppms are assigned to Li ions in a Mn-rich environment. In particular, the signal at 932 ppm is in agreement with Li ions coordinated to 10Mn : 2Ni and the broad signal at 830 ppm to a 11Mn : 1Ni environment.<sup>10</sup> The fact that the latter is not visible in the spectrum of LNMO-O-48 indicates that a higher Mn excess stoichiometry is required to form such Mn rich local environments. However, it should be noted that the extreme case of pure Mn environment (12Mn : 0Ni) which would correspond to the Li environment in spinel  $\text{LiMn}_2\text{O}_4$  located at  $\approx 512$  ppm is not observed in either of the two samples.<sup>39</sup>

An increase in linewidth (LW) of the peaks centered at 932 and 830 ppm of the LNMO-O-45 with increasing Mn share is observed. This can be explained by either the higher number of unpaired electrons in  $\text{Mn}^{3+}$  compared to  $\text{Ni}^{2+}$  or  $\text{Mn}^{4+}$ , and/or the local heterogeneity (*i.e.* TM disorder) of Li environments at these positions.

### 3.2 Reaction mechanism of non-stoichiometric TM ordered LNMO

**3.2.1 *Ex situ* and *operando* crystal structure analysis.** To investigate the reaction mechanism of the non-stoichiometric TM ordered LNMO-O-45, *ex situ* NPD at distinct points of lithiation as well as an *operando* XRD of a complete charge and discharge were carried out. The *ex situ* neutron diffraction patterns of the LNMO-O-45 at distinct points of charge; namely pristine, fully charged (EOC), after one full cycle (EOD) and after 11 complete cycles are depicted in Fig. S4.† These neutron diffraction patterns confirm that the main spinel structure features as well as the superstructure features (see Fig. S4,† inset), are maintained during electrochemical deinsertion and insertion of lithium from the spinel host structure. The superstructural features are the result of the TM ordering, which are reflected in a reduced symmetry cell  $P4_332$  space group.

Fig. 4, left shows the *operando* XRD profiles of the LNMO electrode during charge from the fully discharged state and the following full discharge, both at 0.05C rate. A reversible capacity of  $\approx 129$  mA h  $\text{g}^{-1}$  was obtained. In the XRD profiles, three regions of change can be identified attributed to the  $\text{Li}_0$ ,  $\text{Li}_{0.5}$ , and  $\text{Li}_1$  phases respectively, following previous reports.<sup>41,42</sup> It should be noted that the two peaks observed for the (311) reflection are stemming from the bichromatic Cu  $K_{\alpha 1,2}$  radiation. Continuous peak shifts for the  $\text{Li}_1$  phase at the beginning of the charge and the end of the discharge are observed, which point towards a predominantly solid solution driven reaction mechanism at high lithiation degree. In the region between the 50 and 80 mA h  $\text{g}^{-1}$  ( $\approx 0.6 \leq x \leq 0.4$ ) the decrease of the  $\text{Li}_1$  and the rise of  $\text{Li}_{0.5}$  phase are observed which indicates a biphasic transition. A closeup of this region is depicted in Fig. S5, see ESI.† This underlines that in the  $\text{Ni}^{2+}$  to  $\text{Ni}^{3+}$  transition ( $\text{Li}_1$  to  $\text{Li}_{0.5}$  phase) two different reaction mechanism occur successively, firstly a solid solution reaction (blue) followed by primarily biphasic region (red). Upon further delithiation only the decrease of  $\text{Li}_{0.5}$  peak intensity and rise of the completely delithiated  $\text{Li}_0$  phase intensity are observed which illustrates



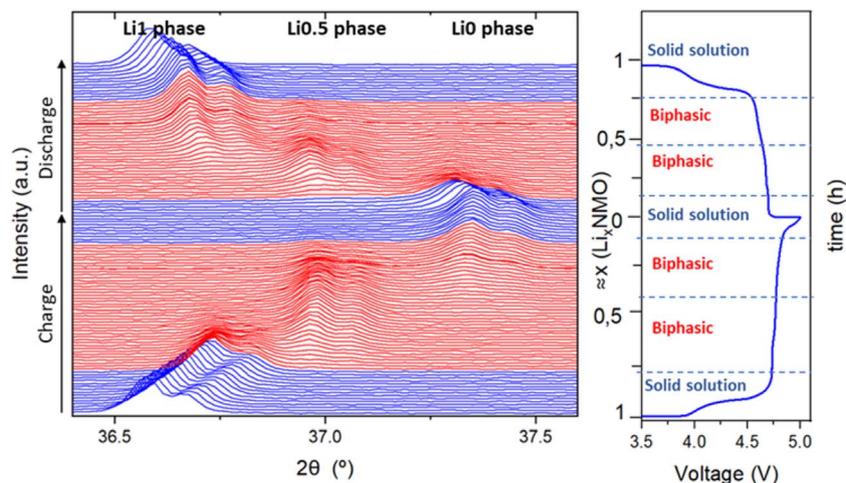


Fig. 4 (Left) *Operando* XRD profiles for diffraction reflection (311) of 125 patterns ( $\text{Cu K}_{\alpha 1,2}$ ) acquired during one complete charge and discharge at rate of 0.05C, for a LNMO-O-45 electrode. Prevalent phase reaction mechanism are indicated by blue (solid solution, single phase) and red (biphasic reaction). (Right) Corresponding electrochemical cycling curve with approximate lithiation degree.

that the  $\text{Ni}^{3+}$  to  $\text{Ni}^{4+}$  reaction is a purely biphasic transition (red). Interestingly at very high SOC a peak shift of the solely existing  $\text{Li}_0$  phase is observed. Lattice contraction of the  $\text{Li}_0$  phase has been observed previously, but its origins remain unclear.<sup>14,18,41,43</sup> Upon discharge the above described changes are reversed, resulting in pattern congruent to pristine. This accentuates the good reversibility of the structural changes of the LNMO spinel phase, in accordance with findings from *ex situ* NPD, see Fig. S4.†

**3.2.2 Ex situ solid state NMR of LNMO-O-45.** To gain complementary insights into the reaction mechanism of non-stoichiometric TM ordered LNMO-O-45 from a local Li structure perspective,  $^7\text{Li}$  NMR spectra of  $\text{Li}_x\text{NMO-O-45}$  electrodes at different lithiation state were recorded during charge and discharge. The resulting spectral evolution are depicted in Fig. 5 for charge (top) and discharge (bottom), respectively. The spectrum of the electrolyte soaked sample at 3.5 V (OCV), shows the same  $^7\text{Li}$  NMR spectrum as the pristine LNMO-O-45 powder with a main sharp signal at 1013 ppm together with two broader resonances at 932 and 830 ppm.

This first delithiation step corresponds to the plateau observed between 4.0–4.3 V in the capacity vs. voltage profile of Fig. 5(top). The  $^7\text{Li}$  NMR spectrum of a sample extracted from a cell charged to 4.05 V (*i.e.*,  $x \approx 0.95$  in  $\text{Li}_x\text{NMO}$ , corresponding to the removal of 0.05  $\text{Li}^+$  ions per formula unit), which corresponds to 7  $\text{mA h g}^{-1}$  shows a similar pattern to the one obtained at 3.5 V but with a small shift of peak position of 6 ppm towards higher frequency as well as a decay in intensity of the signals at 937 and 837 ppm (see Fig. 5(top)). This small shift on the centre of the spectrum and the decay in the two broad signals can be ascribed to changes in the first and second coordination shell of the Li ion as a result of partial oxidation of  $\text{Mn}^{3+}$  to  $\text{Mn}^{4+}$ . This is in agreement with the observed chemical shift difference between two pristine LNMO samples (*i.e.* LNMO-O-45 and LNMO-O-48), *vide supra*. As stated in Table 2, the latter two last signals are assigned to Li environments with

increased Mn content (*i.e.* presence of  $\text{Mn}^{3+}$ ) in the first Li coordination shell of TM ions. At 4.05 V, where extraction of 0.05 Li is achieved, an oxidation of about half of  $\text{Mn}^{3+}$  to  $\text{Mn}^{4+}$  as the total amount of  $\text{Mn}^{3+}$  present in the material was estimated at  $\approx 12\%$  from electrochemistry (and magnetic measurement) *vide supra*.

Upon further charging to 4.3 V (*i.e.*  $x \approx 0.9$ , corresponding to the extraction of approximately 0.1  $\text{Li}^+$  ion) also equivalently to 13  $\text{mA h g}^{-1}$ , both signals at 937 and 837 ppm are no longer observable in agreement with the total oxidation of  $\text{Mn}^{3+}$  to  $\text{Mn}^{4+}$ . In this spectrum a single resonance at 1017 ppm is observed, resembling that of an ideally TM ordered  $9\text{Mn} : 3\text{Ni}$  Li environment., see Table 2). This confirms Cabana's model for a spinel sample with complete ordering of TM ions, where the ratio of  $\text{Mn}^{4+}$  to  $\text{Ni}^{2+}$  is in accordance with the overall spinel stoichiometry  $\text{LiNi}_{0.5}\text{Mn}_{1.5}\text{O}_4$ .<sup>10</sup> This model suggests that  $\text{LiNi}_{0.5}\text{Mn}_{1.5}\text{O}_4$  would exhibit a sole lithium environment, as also observed by Duncan in fully stoichiometric LNMO ( $y = 0.0$  in  $\text{LiNi}(\text{ii})_{0.5-y}\text{Mn}(\text{iv})_{1.5-y}\text{Mn}(\text{iii})_{2y}\text{O}_4$ ).<sup>14</sup> It is noticeable that this signal at 1017 ppm is broader and has shifted back to the lower frequency by 14 ppm compared to the one observed at 4.05 V. This is likely due to a slight relaxation of the system *i.e.* rapid exchange of  $\text{Li}^+$  ions between the TM ordered (9Mn : 3Ni) and Mn rich sites.

Beyond the  $\text{Mn}^{3+/4+}$  redox plateau (above 4.6 V,  $x > 0.9$ ), five *ex situ*  $^7\text{Li}$  NMR measurement points were taken to shed light on the Ni redox activity providing insights on its reaction mechanism. The electrodes studied at the first Ni plateau (*i.e.*  $\text{Ni}^{2+/3+}$ ) are equivalent to the extraction of  $x = 0.15$  to 0.5 Li. In this range of lithiation, the resonance around 1019 ppm attributed to the Li rich ( $\text{Li}_i$ ) phase is solely present. The absence of any other NMR signal suggests no additional phase present at this SoC hinting a single phase solid solution reaction driven mechanism. This is in agreement with *operando* XRD of LNMO-O-45, see Fig. 4, showing a single phase behaviour between  $x \approx 1$  and 0.7 which also points towards a solid solution mechanism.



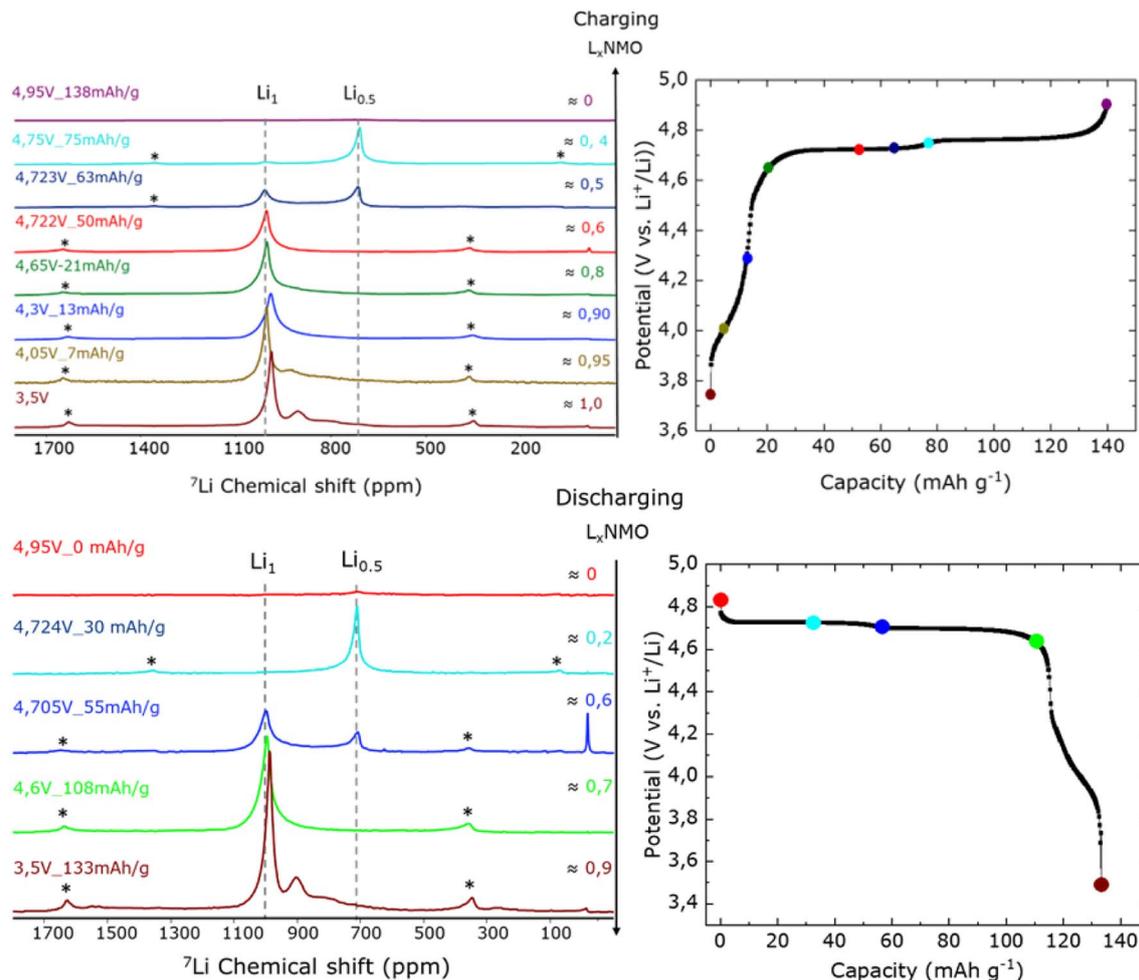


Fig. 5  $^7\text{Li}$  NMR spectra of *ex situ* LNMO samples at selected stages during (top) charge (delithiation) and (bottom) discharge (lithiation) reaction along with corresponding cycling curve in which points at which spectra were acquired are indicated by colour coded markers. \* indicates side band.

At  $x \approx 0.6$  the appearance of a new NMR signal can be observed centred at around 730 ppm. This new resonance can be assigned to the  $\text{Li}_{0.5}$  phase, in agreement with Shimoda.<sup>44</sup> The existence of two separate signals attributed to  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phases confirms a two-phase reaction during the first charge plateau corresponding to  $\text{Ni}^{2+}$  to  $\text{Ni}^{3+}$  redox reaction. Notably, the integrated intensity of the peak assigned for  $\text{Li}_{0.5}$  is comparable to that of the  $\text{Li}_1$  phase. This observation suggests that the biphasic phase transformation from  $\text{Li}_1$  to  $\text{Li}_{0.5}$  occurs within a narrow lithiation region, with  $0.7 > x > 0.4$ . Furthermore, a closer look at the NMR spectra in this lithiation region (see Fig. 6(b)) hints that additional species with chemical shift between the two thermodynamically stable main phases (*i.e.*  $\text{Li}_1$  and  $\text{Li}_{0.5}$ ) are present. The implications of such intermediate phases are further discussed in Section 4.

The appearance of a new  $^7\text{Li}$  NMR resonance at 730 ppm at lower frequency, can be explained by the oxidation of the  $\text{Ni}^{2+}$  to  $\text{Ni}^{3+}$ , which leads to the formation of the new  $\text{Li}_{0.5}$  phase clearly distinct from  $\text{Li}_1$  phase that contains Ni only in a +2 state. In this 3d-transition-metal systems, hyperfine shifts mainly

depend on the pathways between Li and nearby transition metal ions. Consequently, during delithiation when the phase transforms from  $\text{Li}_1$  to  $\text{Li}_{0.5}$ , this transformation also involves a valence change from  $\text{Ni}^{2+}$  to  $\text{Ni}^{3+}$ . Therefore, in the  $\text{Li}_1$  phase, Ni exists solely in the +2 form, whereas in the new  $\text{Li}_{0.5}$  phase, Ni is in the +3 form. This change could lead to a reduced spin density transfer between the orbital pathways of the transition metal ion and Li due to the absence of one unpaired electron in  $\text{Ni}^{3+}$  in the  $\text{Li}_{0.5}$  phase resulting in a lower frequency shift for  $\text{Li}_{0.5}$  phase compared to  $\text{Li}_1$  phase. At the end of the first Ni plateau ( $x \approx 0.4$ ), it can be observed that the new  $\text{Li}_{0.5}$  phase is prevalent with a small remnant of  $\text{Li}_1$  phase, in agreement with the almost complete transformation of  $\text{Li}_1$  to  $\text{Li}_{0.5}$  phase. Upon further delithiation ( $x \leq 0.4$ ) the latter is the only Li containing phase observed by NMR, the  $\text{Li}_0$  phase being undetectable in Li NMR experiments. However, the coexistence of  $\text{Li}_{0.5}$  and  $\text{Li}_0$  was evinced by *operando* XRD, *vide supra*.

At fully charged state  $x = 0$  (*i.e.* 4.95 V), as depicted in the Fig. 5(top), no  $^7\text{Li}$  NMR resonances of neither the  $\text{Li}_1$  nor  $\text{Li}_{0.5}$



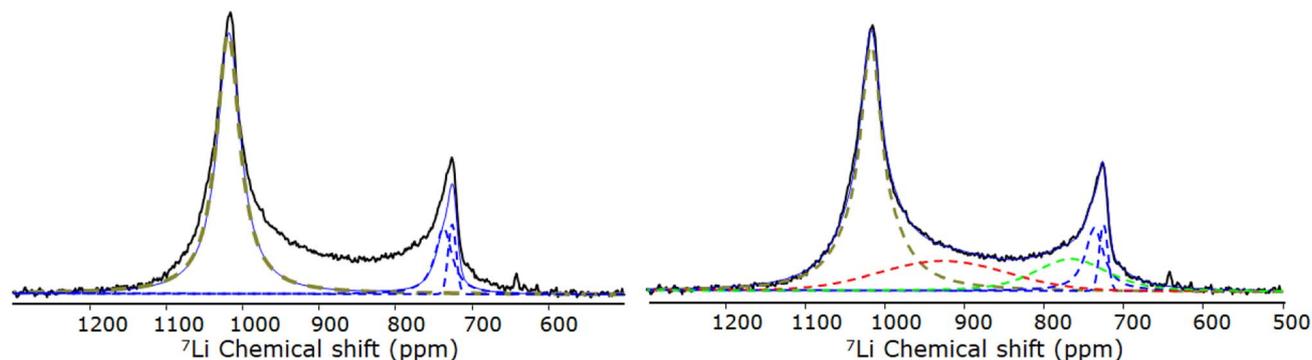


Fig. 6  ${}^7\text{Li}$  NMR spectra at  $x \approx 0.6$  (discharge) corresponding to phase transition region between  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phases. (Left) Fitted using solely signals for the  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phases. (Right) Fitted with two additional intermediate components. Black: experimental, light blue: sum of calculated, dark blue:  $\text{Li}_{0.5}$  component, olive:  $\text{Li}_1$  component and red and light green: for intermediate components.

phases are present, suggesting that the transformation to fully oxidised spinel phase ( $\text{Li}_0$ ) devoid of lithium is achieved.

During subsequent lithiation to  $x \approx 0.2$ , the reappearance of the  $\text{Li}_{0.5}$  phase at 780 ppm is found. Upon further lithiation at  $x \approx 0.6$ ,  $\text{Li}_1$  phase reemerged and coexisted momentarily with the  $\text{Li}_{0.5}$  phase suggesting a biphasic reaction, as depicted in Fig. 5(bottom) Between Li content of 0.4 and 0.7, the  $\text{Li}_{0.5}$  phase gradually vanishes, as observed during charging in a similar lithiation range. Simultaneously, the  $\text{Li}_1$  phase becomes the predominant phase, as evidenced in the  $x \approx 0.7$  sample, indicating that at higher Li content the mechanism is essentially a solid solution, in agreement with *operando* XRD findings. Upon reaching full lithiation in the range of  $0.9 < x < 1$ , the NMR features linked to the presence of the ordered (9Mn : 3Ni) at 1013 ppm as well as Mn rich local Li environments (932 ppm and 830 ppm) were observed. This observation reflects the reversible reduction of  $\text{Ni}^{4+}$  to  $\text{Ni}^{2+}$  and  $\text{Mn}^{4+}$  to  $\text{Mn}^{3+}$ , respectively. Furthermore, a shift in the center of gravity toward lower frequency was observed yielding to congruent NMR spectrum to that of the pristine state. This concordance serves as compelling evidence for the reversible relithiation of the sites within the LNMO spinel structure.

## 4 Discussion

Monitoring local Li environments at different stages of SoC upon delithiation and lithiation of the non-stoichiometric partially ordered LNMO-O-45 spinel *via*  ${}^7\text{Li}$  NMR and contrasting them with *operando* XRD resulted in plentiful insights whose implications will be further discussed in the following.

At the beginning of the charge reaction in the 4 V plateau region, we observe the preferential delithiation of the Mn-rich (Mn > 9) local Li environments. This is reflected by the disappearance of the two NMR signals (at 932 ppm and 830 ppm) assigned to these Li environments when reaching 4.3 V. This implies that the Mn oxidation and Li vacancy formation upon electrochemical charging occur in close vicinity. Albeit appearing logical, this spatial concurrence has not been shown in any previous LNMO studies to the best of the authors knowledge. Moreover, the chemical shift of the main feature observed in

this SoC region hints that a fast Li exchange occurs between the TM ordered stoichiometric (9Mn : 3Ni) and Mn-rich sites.

A second noteworthy finding is the extended solid solution region evinced for this TM ordered LNMO sample by both *ex situ*  ${}^7\text{Li}$  NMR and *operando* XRD. This finding is reflected by the absence of the  $\text{Li}_{0.5}$  phase approximately until about 1/3 of total charge is reached, well beyond  $\text{Mn}^{3+}$  related 4 V plateau. Only upon further delithiation, the biphasic reaction mechanism expected for the TM ordered LNMO in the  $\text{Ni}^{2+/3+}$  plateau is observed. These findings underline the importance of both structural parameters, TM stoichiometry and TM order to adequately describe LNMO characteristics, as the former greatly affects the latter. Although TM order features are salient for the LNMO-O-45 sample in both Raman spectroscopy and Neutron diffraction, its electrochemical mechanism strongly deviates from the expected predominantly biphasic reaction mechanism of TM ordered LNMO.<sup>17,18,44,45</sup> Indeed our findings show that already small Mn-excess stoichiometry induces TM disorder (by partially populating Ni 4b sites) which result in an extended solid solution reaction region in the phase transformation from  $\text{Li}_1$  to  $\text{Li}_{0.5}$  phase as well as the presence of intermediate species.

The existence of additional spectral features besides those of the expected  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phase towards the end of the  $\text{Ni}^{2+}$  to  $\text{Ni}^{3+}$  transition (Fig. 6) is another interesting observation revealed by the *ex situ*  ${}^7\text{Li}$  NMR. The fact that such intermediate environments were not observed in the ideal stoichiometric and fully TM ordered LNMO investigated by Shimoda *et al.*<sup>44</sup> hints that their presence could be related to the stoichiometric induced TM disorder in the LNMO-O-45. In Fig. 6 the *ex situ* NMR spectrum at SoC  $x \approx 0.6$  is depicted. It is evident that the experimental spectrum cannot be matched by solely using two signals for the expected  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phases, see Fig. 6 (left). Our Gaussian and Lorentzian fit analysis suggest that there are at least two additional broad features present at this SoC, see Fig. 6 (right). These additional broad features, exhibiting a paramagnetic shift between those of the  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phases, suggest the existence of intermediate Li environments. However, these intermediate species do not result in distinct resonances, as observed for the  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phases; instead, they represent a continuous distribution of environments between the two



equilibrium phases. We believe this intermediate region exhibits characteristics of a mix of both phases could suggest a narrow miscibility gap between the two phases with a benefit of facilitating a facile Li hopping between the  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phase, possibly correlated with a blend of  $\text{Ni}^{2+}$  and  $\text{Ni}^{3+}$  states as well as various lithiation states as suggested by *ex situ* NMR having a chemical shift between the  $\text{Li}_1$  and  $\text{Li}_{0.5}$  phases. The presence of these intermediate environments may contribute to narrowing the lattice mismatch or miscibility gap between the fully lithiated ( $\text{Li}_1$ ) and half lithiated ( $\text{Li}_{0.5}$ ) phases. This phenomenon has been reported mainly for LFP,<sup>46,47</sup> but a clear correlation between cation ordering, reaction mechanism, and rate has yet to be fully established for spinel LNMO. Battery performance is not solely constrained by Li transport in the electrolyte and electron movement within the electrodes but also by the phase transformations occurring during lithiation and delithiation. Therefore, the absence of a phase transformation or the presence of an intermediate phase that narrows the miscibility gap is generally considered a key factor in achieving high rates and better cyclability. In this regard Omenya *et al.* showed that by partially substituting Fe with V the miscibility gap in LFP can be significantly decreased. Moreover, this leads to a reduction of the solid solution formation temperature in the two-phase region and enhances Li solubility. In our spinel LNMO case, the local environment change stems from the Ni-deficient stoichiometry resulting in partial “substitution” of  $\text{Ni}^{2+}$  sites by Mn ( $\text{Mn}^{4+}$  and  $\text{Mn}^{3+}$ ). Such stoichiometrically induced partial disorder can hence be a viable approach to close the miscibility gap and enable better cycling

performance. If these intermediate species exist only on a local structural level or also exist as long range intermediate crystal phases cannot be answered with certainty at this point. The analysis of our *operando* XRD data in this region of SoC see Fig. S5 (see ESI†) doesn't provide conclusive evidence, if besides the evident features of biphasic transition additional phases are present. Nevertheless the existence of several intermediate species with different lithiation stages besides the thermodynamically stable phases  $\text{Li}_1$  and  $\text{Li}_{0.5}$  was previously suggested based on *operando* XRD by Komatsu and coworkers.<sup>41</sup> Furthermore, it should be noted that the co-existence of multiple local environments at about half lithiation was already hinted in our previous *operando* X-ray absorption study by a maxima in local disorder of the Ni–O bonds (Debye–Waller factor) for the TM ordered LNMO.<sup>15</sup>

In Fig. S8 (in ESI†), the peak intensity ratios and chemical shift of the four spectral features present during delithiation and lithiation are depicted in single graph to highlight their evolution as function of SoC. The full data set of deconvoluted spectra is provided in Table S1 in the ESI† along with a discussion of the trends observed.

On a macroscopic level the narrow co-existence range of thermodynamically stable phases  $\text{Li}_1$  and  $\text{Li}_{0.5}$  ( $0.6 \leq x \leq 0.4$ ) evinced by both *ex situ* NMR and *operando* XRD points towards the prevalence of a concurrent reaction propagation model rather than particle by particle reaction propagation.<sup>41,48</sup> In such concurrent model a quasi-simultaneous transformation occurs in all particles. A scheme of this proposed concurrent model reaction front propagation is illustrated in Fig. 7 along with

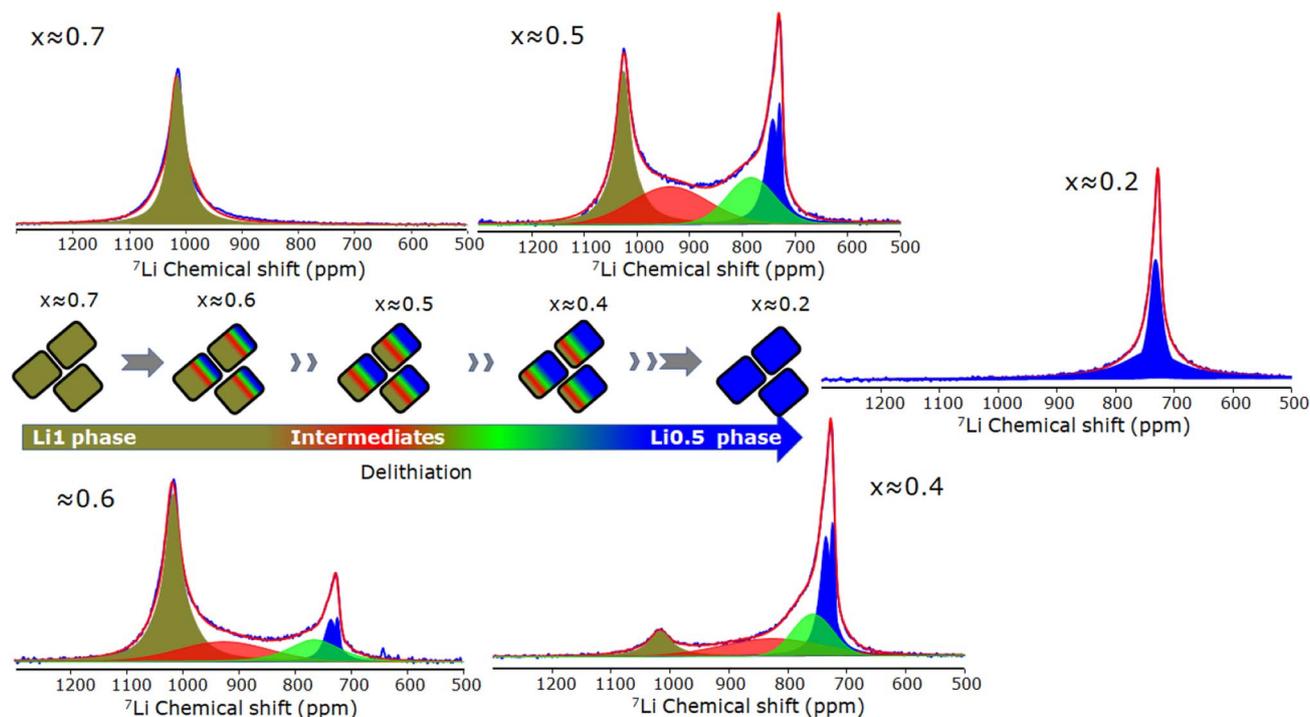


Fig. 7 Scheme of concurrent reaction propagation model showing the narrow lithiation range of coexistence ( $0.6 \leq x \leq 0.4$ ) of both thermodynamically stable phases ( $\text{Li}_1$  and  $\text{Li}_{0.5}$ ) as well as intermediates species based on corresponding *ex situ*  $^7\text{Li}$  NMR spectra.



corresponding NMR spectra. Varying charge states within the same particle, so called intra-particle state of charge heterogeneity, has been observed in several cathode active materials.<sup>49,50</sup> This heterogeneity in phases likely also exist in partially delithiated LNMO, as observed by Kuppen *et al.*<sup>48</sup>

Another remarkable finding of our *ex situ* <sup>7</sup>Li NMR study is the complete delithiation of the spinel structure at the end of the charging process, as evidenced by the absence of signals in the <sup>7</sup>Li NMR spectra. It should be considered that NMR is an element-selective technique and only detects the nuclei under investigation. Hence in this case, the Li<sub>0</sub> phase present at the lower lithiation state cannot be observed by Li NMR. Our results indicate a complete phase transformation to the Li<sub>0</sub> phase (Ni<sub>0.5</sub>Mn<sub>1.5</sub>O<sub>4</sub>), which is confirmed by our diffraction results (NPD and XRD) and is in agreement with the results of other authors.<sup>51,52</sup> However, it may contradict studies which claim that incomplete delithiation (only up to 90%) can be obtained with a full charge.<sup>43</sup> The complete delithiation of the spinel structure evinced by *ex situ* <sup>7</sup>Li here is a strong asset of the LNMO spinel which results in better use of lithium compared to other cathode materials such as NMC.

## 5 Conclusion

In this study, we have shown that NMR is highly sensitive to key structural parameter of TM ordering of the LNMO spinel cathode material. Its ability to evince and quantify TM order degree but also distinguish between thermodynamic and stoichiometry induced disorder is a unique asset. In this regard we demonstrated that even minor deviations from ideal TM stoichiometry ( $0.02 \leq y \leq 0.05$  in LiNi<sub>(II)</sub><sub>0.5-y</sub>Mn<sub>(IV)</sub><sub>1.5-y</sub>Mn<sub>(III)</sub><sub>2y</sub>O<sub>4</sub>) result in strong changes of the NMR spectra and greatly affect the electrochemical mechanism. By coupling local and long range sensitive techniques we have gained a holistic picture of the delithiation/lithiation process, revealing that Ni deficient LNMO-O-45 does not follow a simple sequence of a two biphasic reactions as previously claimed for such predominantly TM ordered LNMO spinels. Instead our *ex situ* NMR and *operando* XRD suggest that delithiation linked to Ni<sup>2+/3+</sup> redox reaction occurs firstly *via* an extended solid solution mechanism followed by the expected two phase reaction mechanism. However, we evince for the first time the presence of reaction intermediates during this later stage which suggest that the formation from Li<sub>1</sub> to Li<sub>0.5</sub> phase occurs on a local structural level *via* intermediate steps. We attribute their existence to the stoichiometric induced TM disorder and rationalise their potential for material performance enhancement by closing the miscibility gap. During the Ni<sup>3+/4+</sup> redox reaction a two phase reaction mechanism (Li<sub>0.5</sub> to Li<sub>0</sub>) is prevalent resulting in the complete delithiation of the spinel structure at the end of the charge process.

## Conflicts of interest

There are no conflicts to declare.

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