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Imaging the Phase Transformation in Single Particles of Lithium Titanate Anode for Lithium-ion Batteries

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ABSTRACT

Lithium uptake and release in lithium titanate (LTO) anode materials during a discharge and charge cycle is one of the fundamental processes of a Lithium-ion battery (LIB), still not fully understood at the microscopic level. During discharge cycle, LTO undergoes a phase transformation between $\text{Li}_4\text{Ti}_5\text{O}_{12}$ and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ states within a cubic crystal lattice. To reveal the details of the microscopic mechanism, it is necessary to track the sequence of phase transformations at different discharge/charge states under operating conditions. Here we use *in-situ* Bragg Coherent Diffraction Imaging (BCDI) and *in-situ* X-ray

diffraction (XRD) experiments to examine the lithium insertion-induced materials phase transformation within a single LTO particle and a bulk battery analogue, respectively. BCDI analysis from (111) Bragg peak, shows the two-phase transformation manifesting as a distinct image phase modulation within a single LTO nanoparticle occurring in the middle of the discharge region then subsiding towards the end of the discharge cycle. We observe the biggest phase variation at the two-phase stage, indicating the formation of phase domains of 200 nm in size during the discharge process. We also observe a lattice contraction of >0.2% in a single LTO nanoparticle at the (400) Bragg peak measurement, larger than that in the corresponding bulk material. Our observation of this phase transformation at a single particle level has implications for the understanding of the microscopic/mesoscale picture of the phase transformation in anode and cathode LIBs materials.

INTRODUCTION

As the world population grows, its demand for energy grows, leading to the depletion of non-renewable energy sources. The use of these energy sources leads to a detrimental impact on the environment, and this leads to the search for clean, renewable energy sources. Solar energy is the most abundant source of energy, but much of it is not used due to lack of efficient energy storage devices. Tremendous progress has been made in search and developing electrochemical storage devices; one example is rechargeable lithium-ion batteries (LIBs), which show excellent cycling stability and high-rate capability¹⁻³. The performance of LIBs depends on the cathode, anode, and electrolyte/binder choices⁴ which can be optimized for different applications. In LIBs, the

cathode electrode is the source of Li^+ ions, while the electrolyte serves as a conductive medium that helps Li^+ ion- and electron-transport, the separator prevents direct contact between anode and cathode electrodes, and the anode electrode stores and releases Li^+ ions from the cathode⁴. Since the introduction of the first commercialized rechargeable Li-ion battery in 1991 by Sony, different anode and cathode materials have been explored to improve the power density, cyclability, and safety of the battery⁵. Cathodes such as LiFePO_4 and LiCoO_2 and anodes such as Li, graphite (by forming intercalation compound LiC_6), Si, and $\text{Li}_4\text{Ti}_5\text{O}_{12}$ have been used in different combinations⁶. Li metal has been used as anode material that could provide high specific capacity (3860 mAh/g) and lowest overall anode potential. However, cycling of Li anode leads to the formation of dendrites, which could eventually penetrate the separator and cause a shortage and fire in the battery⁷⁻⁹. Also, graphite has high energy density and long cycle life, which makes it the most popular anode in commercial LIBs, but the low lithiation voltage (~ 0.1 V vs. Li/Li^+) can result in a short-circuit, and possibly fire. Although a Si anode has the highest known theoretical capacity (4200 mAh/g), its huge volume expansion, over 300%, occurring during the lithiation/delithiation cycles leads to electrical shorting/disconnection between the active materials and the current collector^{5,10-12}. Li, graphite, and Si anodes have failed so far to satisfy the fast charging capabilities in LIBs. In contrast, $\text{Li}_4\text{Ti}_5\text{O}_{12}$ (lithium titanate or LTO) has long-cycling stability and high-rate capability because it has insignificant volume change ($<0.2\%$) between fully lithiated and delithiated end members¹³⁻¹⁵; this makes LTO a promising anode material for use in commercial fast-charging batteries, such as Toshiba's SCiB™ rechargeable batteries¹⁶.

$\text{Li}_4\text{Ti}_5\text{O}_{12}$ and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ are two end-members with a defective spinel (space group $Fd\bar{3}m$) and rock-salt (space group $Fm\bar{3}m$) structure, respectively^{17–19}. The crystal structure of LTO has cubic symmetry with lattice constants of $a = b = c = 8.3595\text{\AA}$ and $\alpha = \beta = \gamma = 90^\circ$. When LTO is fully discharged, three Li-ions are inserted per unit cell, forming $\text{Li}_7\text{Ti}_5\text{O}_{12}$ with lattice constants of $a = b = c = 8.3538\text{\AA}$ and $\alpha = \beta = \gamma = 90^\circ$ ¹⁷ showing a 0.07% lattice contraction. Compare to $\text{Li}_4\text{Ti}_5\text{O}_{12}$, the octahedral (16c) positions in $\text{Li}_7\text{Ti}_5\text{O}_{12}$ are occupied by Li, and the octahedral (16d) sites maintain the same configuration (occupation by Li and Ti). The electrochemical reaction in LTO proceeds via two-phase transformation between $\text{Li}_4\text{Ti}_5\text{O}_{12}$ (with the octahedral (16d) sites randomly occupied by Li or Ti atoms and the tetrahedral (8a) positions occupied by Li atoms) and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ (the octahedral (16c) positions occupied by Li atoms and the occupancy of the octahedral (16d) identical to that in $\text{Li}_4\text{Ti}_5\text{O}_{12}$) end-member LTO phases¹⁷. During Li^+ uptake, the Li-ions at the 8a sites move to the empty 16c sites, and the new Li ions from electrolyte also occupy the 16c sites. In the fully lithiated state, three additional Li^+ ions are added per unit cell. However, visualization of structural phase transformation and tracking the associated lithium migration directly with X-ray based techniques is experimentally challenging since the lattice constant change between the two endmembers is very small (< 0.1%), in addition to the low atomic scattering factor of Li compared to that of O and Ti atoms. Different experimental techniques such as X-ray absorption spectroscopy²⁰, electron energy-loss spectroscopy (EELS)²¹, neutron powder diffraction²², and nuclear magnetic resonance (NMR)^{23–25} have been implemented to identify the structural phase transformation in LTO anode materials. A Ti K-edge X-ray absorption spectroscopy study supported by ab initio calculations by Zhang *et al.* proposed a new quasi solid-solution

phase, which plays an instrumental role in facilitating Li^+ transport. Using STEM-EELS spectrum imaging of the lithiated $\text{Li}_7\text{Ti}_5\text{O}_{12}$, Kitta *et al.* showed that the two separate phases coexist inside the specimen with defined phase boundaries, and no apparent misfit strains or misorientations are detected²⁶. Monte-Carlo simulation combined with density functional theory (DFT) on a large LTO system showed that the three energetic parameters, namely the potential energy of the 8a site, the difference in the site energy between the 8a and 16c sites and the repulsion between two Li atoms situated at the adjacent 8a and 16c sites and the topology of the Li sites determine the system²⁷. For the electrochemical potential profile, the difference in the site energy between the 8a and 16c sites play a crucial role and in the plateau region in the potential profile two structural phases coexist²⁷. Recently, operando EELS on Li K-edge combined with first-principle calculations identified a metastable state, which consists of distorted Li polyhedra, and is proposed to play an essential role to mediate Li^+ -ion migration²¹.

The success of the LTO material for repeated cycles of charge and discharge arises from the small lattice constant change between the two LTO phases. This is its biggest advantage over alternative intercalating anode systems because it allows rapid charging and many cycles. As we will show, the small change in the lattice parameter is significant enough to cause apparent patterns of strain in micron-sized grains of LTO with a sufficiently sensitive measurement, such as our *in-situ* Bragg Coherent Diffraction Imaging (BCDI) probe a single LTO particle presented here. We observe the development of local displacement field in 3D by BCDI under *in-situ* conditions while discharging the LTO anode material. BCDI is a lensless imaging technique and, with a suitable choice of iterative algorithms, allows one to get three-dimensional shapes of the nanocrystals as

complex 3D density maps²⁸. In the images, the amplitude represents the ordered electron density of the particle, and the 3D phase distribution, $\phi(r)$, is due to local displacements in strained nanocrystals. The mapping is fully quantitative and given by the relationship $\phi(r) = \mathbf{Q} \cdot \mathbf{u}(r)$, where \mathbf{Q} is the momentum transfer vector of the reciprocal lattice point measured, and $\mathbf{u}(r)$ is the vector displacement field representing the distortions from the ideal crystal lattice²⁹. The spatial derivative of the displacement field is a component of the formal strain tensor. Since only one component of the displacement field is measured at each Bragg reflection, phase images from three or more independent \mathbf{Q} -vectors must be combined to obtain 3D spatial reconstruction of the vector field $\mathbf{u}(r)$. As we report here, the small difference in lattice parameter between the two structures is sufficient to produce an observable local displacement field: if side-by-side regions of $\text{Li}_4\text{Ti}_5\text{O}_{12}$ to $\text{Li}_7\text{Ti}_5\text{O}_{12}$ were to be present within 800 nm LTO nanoparticle, there would be a phase difference of 0.89 rad, using the lattice parameters reported in ref.¹⁷ (details of derivation is given in SI). This provides the BCDI method with plenty of sensitivity to observe nanoscale displacement fields in these materials.

In this manuscript, we present 3D renderings of phase transformation of lithium titanate (LTO) anode material during discharging, obtained using the BCDI technique. Inverted images show the formation of nano-domains with a size of about 200 nm as positively and negatively shifted image phases in the reconstructed image in the two-phase region of discharging cycle. This result is an experimental confirmation of the domain model proposed on the same material³⁰. A slice through the rendered inverted images show displacement field inhomogeneities in the sample but does not show

dislocations. In addition, the total d-spacing change estimated in a fully discharged single LTO particle is much larger than the bulk.

MATERIALS AND METHODS

A. Sample Preparation:

The samples for the *in-situ* BCDI experiment were prepared using the conventional 2032-type coin cells. $\text{Li}_4\text{Ti}_5\text{O}_{12}$ (LTO) active material from Primet Precision Materials, Inc. was mixed with a conductive agent (Super P conductive carbon black) and polyvinylidene fluoride (PVDF) binder in N-Methyl-2-pyrrolidone (NMP) solvent with 2:1:1 ratio, which was further coated on carbon paper current collector and dried in a vacuum oven. The thin LTO electrode was then punched into a 10-mm disk and assembled in an Ar-filled glove box with Li metal as the counter electrode and a Clegrad polypropylene (PP) separator. The electrolyte was prepared by mixing a 1.0M lithium hexafluorophosphate (LiPF_6) solution in ethylene carbonate ($(\text{CH}_2\text{O})_2\text{CO}$): dimethyl carbonate ($\text{C}_3\text{H}_6\text{O}_3$), with 1:1 volume ratio. For the *in-situ* characterization, an X-ray-transparent Kapton window was fabricated in the stainless-steel case by sealing it with epoxy. Before the coin-cell was assembled, we characterized the density and morphology of the electrodes with a scanning electron microscopy (SEM) (Fig. 1(C)). The SEM result shows faceted 400–800 nm sized LTO particles with a characteristic elongated ellipsoidal shape, which are ideal for BCDI experiments. We assembled several coin-cells in the Argon glove box.

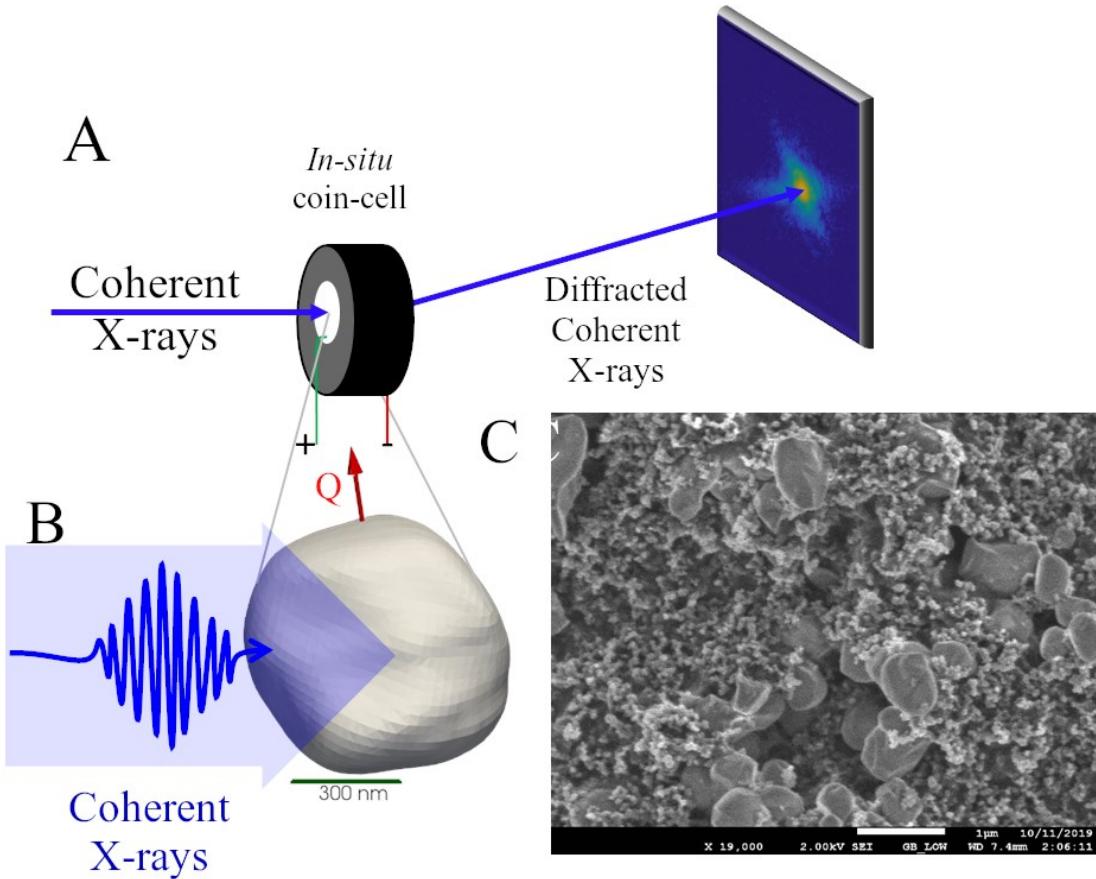


Figure 1: Experimental setup of the *in-situ* Bragg Coherent Diffraction Imaging (BCDI). (A) *In-situ* BCDI setup at 34-ID-C beamline, APS. (B) 3D reconstructed image of a single LTO crystal, shown using isosurface rendering representation. For the reconstruction, two-dimensional coherent diffraction patterns were collected by rocking the sample in and out off the Bragg condition. The arrow labeled as Q is the scattering vector. (C) SEM images of newly prepared LTO crystals reveal a clear faceted morphology of the specimen. White scale bar is 1 mm.

B. *In-situ* Powder X-ray Diffraction

In-situ X-ray diffraction data were collected at the 28-ID-2 XPD beamline of the National Synchrotron Light Source (NSLS-II). The *in-situ* XRD testing electrodes were prepared by coating the electrode slurry, a mixture of LTO, conductive carbon black, and polyvinylidene fluoride with 8:1:1 volume ratio, on a current aluminum collector. The experiment under *in-situ* conditions was conducted using 67 keV ($\lambda = 0.185 \text{ \AA}$) X-ray

beam $0.5 \times 0.5 \text{ mm}^2$ in size. X-ray diffraction patterns were acquired using a 2D PerkinElmer detector (2048 x 2048 pixels with 200 mm^2 pixel size) located 1.49 m from the sample, calibrated using CeO_2 standard powder. The 2D diffraction images were integrated azimuthally using pyFAI and GSAS-II was used for Rietveld refinement^{31,32}.

C. *In-situ* Bragg coherent diffraction imaging

Bragg coherent diffraction data were collected at the 34-ID-C beamline of the Advanced Photon Source (APS) (see Fig.1). For the coherent diffraction analysis shown in this manuscript, we used 11.2 keV and 9 keV monochromatic beams in two independent experiments. The coherent X-ray beam was focused using a pair of Kirkpatrick-Baez (KB) mirrors to $\sim 1 \times 1 \text{ }\mu\text{m}^2$ illuminating the LTO nanocrystals. As shown in Fig.1(A), the measurement was done on a $10 \text{ }\mu\text{m}$ thick LTO electrode in transmission geometry. We used the same *in-situ* coin-cell setup, which was implemented in the previous experiments³³ with a window opening of 1 mm. The coin cell was mounted on a coin cell holder vertically with the LTO electrode located downstream to minimize the absorption of the diffracted X-rays. Prior to experiment, the electrodes were tested via galvanostatic cycling and show a flat plateau region at about 1.5 V with a sharp voltage reduction in the early discharge. In addition, the coin cells were tested by a multimeter to ensure the open circuit voltage is 1.5 V (i.e., a fully functional coin cell). The particle density and morphology of the LTO particles were further examined with SEM, shown in Fig.1(C), finding 400-800 nm sized LTO particles. From the fringe-spacing in the diffraction patterns we estimated that the measured LTO nanoparticle size is approximately 800 nm. Coherent X-ray Diffraction (CXD) patterns were acquired using a

Timepix photon-counting detector mounted $D = 1.95$ m away from the sample. We obtained full rocking curves around the (400) and (111) Bragg reflections, each in a separate measurement, and collected 2D CXD patterns using a 2D detector at two-theta angles of 38.5° and 16.4° ($\Delta\theta = \pm 0.15^\circ$) respectively. Although the full sensor of the detector has 512×512 pixels with 55 mm^2 pixel size, the coherent diffraction patterns were collected utilizing just the first quadrant sensor, which has fewer bad pixels. Automatic background subtraction is implemented within the detector. For the (400) set of data, we collected 120 CXD patterns by rocking the sample in 0.0025° steps around the Bragg peak while we were cycling the coin-cell at C/2 discharge rate (C/2 rate is the current value discharge a battery in 2 hour). For the second set of data, we collected 60 CXD patterns while rocking the sample in increments of 0.005° around the (111) Bragg peak while we were cycling the coin-cell at C/2 discharge rate. For both sets of measurements, the full discharge took roughly four hours. A full 3D diffraction pattern took about 2 minutes to measure. Between consecutive scans, we optimized the sample position on a piezo scanning stage, in order to maintain the Bragg condition and avoid sample misalignment. The coin-cell was discharged using an 8-channel MACCOR battery cycler while the series of measurements progressed.

D. Phase retrieval

Before feeding the 3D diffraction data to an iterative phasing algorithm, implemented in Matlab^{29,34–37}, both a white-field correction and a hot pixel removal were applied for each frame of the 3D data stack. For the phase retrieval, we used a total of 620 iterations with 20 iterations of error-reduction (ER), and 180 iterations of a Hybrid-input-output (HIO) algorithms^{38,39}, used alternately, with the iteration starting and ending

with ER. We used the partial-coherence correction since this was found in the past to improve the reconstructed image quality³⁴. The reconstructed 3D phase images shown throughout this manuscript were shifted with respect to the center phase value for display purposes.

E. Pearson cross-correlation calculation

The Pearson cross-correlation coefficient (r_{AB}) between diffraction patterns A and image B was calculated the following formula:

$$r_{AB} = \frac{\sum_{i=1}^n (A_i - \bar{A})(B_i - \bar{B})}{\sqrt{\sum_{i=1}^n (A_i - \bar{A})^2} \sqrt{\sum_{i=1}^n (B_i - \bar{B})^2}}$$

Where the sum in i is over the 2D region of n pixels containing signal, A_i and B_i are the pixel intensity values and $\bar{A} = 1/n \sum_{i=1}^n A_i$, $\bar{B} = 1/n \sum_{i=1}^n B_i$ are the mean intensity values of the 2D diffraction patterns A and B, taken from the slice at the center of the rocking curve, respectively. The correlation coefficient r_{AB} by definition ranges from 0 to 1, where $r_{AB} = 1$ implies a perfect correlation between A and B diffraction patterns and $r_{AB} = 0$ means no correlation between them.

F. Extracting *in-situ* single-particle lattice constant changes

As a first step, we looked at the position of the (400) Bragg peak as a function of discharge voltage. We correlated the voltage cycler timestamp with each rocking scan timestamp. Since the cycler timestamp is relative to the start time, it must be corrected with the known precise starting time. Using these, we obtained the cell voltage readout of each rocking scan. The diffractometer motor positions δ , γ , χ , ϕ , and θ are all recorded in a scan file. The detector angles γ and δ are related to the total diffraction 2θ angle to a very good approximation as:

$$\cos(2\theta) = \cos(\delta) \cos(\gamma) \quad (1)$$

We calculated the γ_i and δ_i angles from the pristine state diffraction peak positions p_{x0} , p_{y0} and detector angles γ_0 , δ_0 and the diffraction peak positions on the detector, p_{xi} , p_{yi}

$$\gamma_i = \frac{(p_{xi} - p_{x0})P}{D} \left(\frac{180}{\pi} \right) + \gamma_0 \quad (2)$$

$$\delta_i = \frac{(p_{yi} - p_{y0})P}{D} \left(\frac{180}{\pi} \right) + \delta_0, \quad (3)$$

where $P = 55 \text{ } \mu\text{m}$ is the detector pixel size, and $D = 1.95 \text{ m}$ is the sample-to-detector distance. Next, we can calculate $2\theta_i$ for each scan using Eq.1 and d-spacing, d_i for each peak using Bragg's law. Finally, we calculate the relative d-spacing change, which is $\Delta d/d = (d_i - d_0)/d_0$, where d_0 is the d-spacing for the pristine state. The quantities calculated using this procedure are shown in Fig.2.

RESULTS AND DISCUSSION

A. (400) Bragg Peak Results

During the *in-situ* BCDI experiments, we cycled the electrode at C/2 discharging and charging rate while tracking a single LTO particle. In the past, similar experiments were carried out on battery cathode materials^{40,41}. Figure 2(A) shows an intensity map on the logarithmic scale of the area surrounding one of the (400) coherent diffraction peaks measured from a single LTO particle in its pristine state, and at 1.44V and 1.3V discharge states. During the discharging process (from left to right shown in Fig.2(A)) a two-step evolution of the diffraction peak signal was observed: (i) initially the peak moves inwards to a lower 2θ angle indicating a slight lattice expansion and it moves around the powder ring which is due to the rotation of the LTO particle, and (ii) in later discharging states it

moves radially outward to a higher 2θ angle indicating a lattice contraction. In addition, our *in-situ* XRD analysis showed an initial small lattice expansion during the early discharge cycle, followed by a $\sim 0.066\%$ of lattice contraction shown in Figure S5 of the SI.

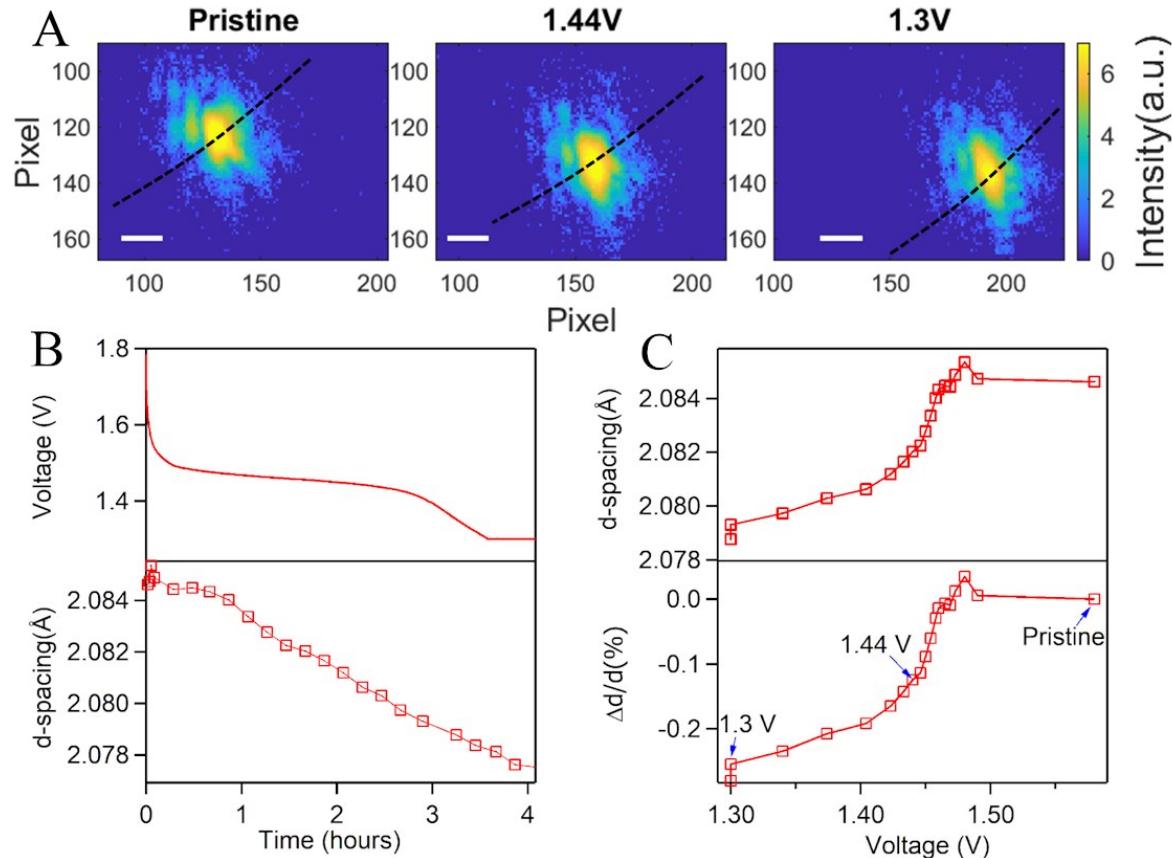


Figure 2. *In-situ* Bragg Coherent Diffraction results during a discharge cycle on the (400) Bragg peak. (A) The logarithm of the diffraction intensity at three different discharge states measured from a single LTO particle. (B) C/2 discharging cycle of coin-cell for the experiment (upper) and d-spacing changes as a function of time extracted from the shift of (400) diffraction peak on the detector (lower). (C) The extracted lattice d-spacing (upper) and relative changes (lower) of the same LTO particle during the discharge cycle as a function of the voltage of the coin-cell. The white scale bar is 20 pixels, which corresponds to $\sim 3 \times 10^{-3} \text{ \AA}^{-1}$.

In the later stage Li^+ migration from the 8a sites to the 16c sites leads to lattice contraction. The lattice contraction observed in our bulk XRD measurement is consistent with earlier reports¹⁷. The voltage profile of the discharging cycle of the electrode during *in-situ* BCDI measurements and the calculated d-spacing changes as a function of time

are shown in upper and lower panels of Fig. 2(B), respectively, the d-spacing decreases continuously, with discharge cycle consistent with lattice contraction. However, the lattice contraction measured in our BCDI experiment is larger than in earlier reports for bulk LTO samples¹⁷, where a difference of $\Delta a/a = 0.07\%$ ^{18,42} was reported. In top and bottom panels of Fig. 2(C) we plot the d-spacing and d-spacing changes with the discharging voltage, respectively. During the early stage of discharge, both the d-spacing and d-spacing stay constant, but below 1.48V, we see more lattice contraction. However, the d-spacing observed for our 400nm particles, $\Delta d/d \cong 0.2\%$, while larger than the bulk, is smaller than the $\Delta a/a = 0.81\%$ lattice contraction reported in even smaller size nanosized materials⁴³. The lattice parameter changed from 8.3526(1)Å to 8.4203(1)Å, and from 8.3585(1)Å to 8.3852(4)Å for the 44nm and 120nm nanosized samples for $\text{Li}_4\text{Ti}_5\text{O}_{12}$ and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ structural phases, respectively. This implies during a discharge the 44nm nanosized samples showed $\Delta a/a = 0.81\%$ change, whereas the 120nm showed $\Delta a/a = 0.32\%$. Systematic decrease of the lattice contraction effects with increasing particle size is to be expected due to decrease in surface area to volume ratio, which is the influence of the interface energy, which would be enhanced in smaller sized LTO samples. We note that density functional theory based first principles calculations predicted lattice parameter contraction between $\text{Li}_4\text{Ti}_5\text{O}_{12}$ and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ structural phases⁴⁴ which is much larger than the experimental results¹⁷.

The measured diffraction pattern intensity is related to the squared amplitude of the Fourier transform of the electron density of the nanocrystal. It is expected that, during the discharging process Li-ions insert into the LTO particle, which can result in local displacement field in the particle. To explore this phenomenon, we calculate the Pearson

cross-correlation coefficient between all the different coherent diffraction patterns done at different discharge voltage with the pristine coherent diffraction pattern. As shown in Fig. 3(A), the result highlights the differences between two diffraction patterns as changes in the correlation coefficient. Interestingly, the correlation plot shows two distinct regions, the first two hours of the discharging process we observe a strong correlation >0.6 and the correlation drops appreciably below 0.5 in the last two hours. Since, the change of diffraction pattern is the reflection of phase transformation within the scattering volume of the LTO particle. Therefore, the regime between 2.5 to 4 hours of the correlation plot shown in Fig. 3(A) is due to the structural transformation of the particle, called two-phase regime³⁰. This implies that partial uptake of 1 Li-ion per unit cell does change the structure locally; however, the diffraction pattern does not change significantly, which is why we get a strong correlation for the first two hours. An uptake of more than one Li-ion per unit cell leads to decorrelation in the diffraction pattern due to a change in the structure as we can see in the correlation plot. Since radiation damage can also change the diffraction pattern, to explore this possibility we measured a single pristine LTO nanocrystal for 2 hours without any voltage cycling⁴⁵. As shown in Figure S1, its shows strong correlation at all time, which indicated no evident aging or apparent beam damage in the single LTO

particle. This implies that the effect observed in Fig. 3(A) can only be due to discharge related changes in the LTO nanocrystal.

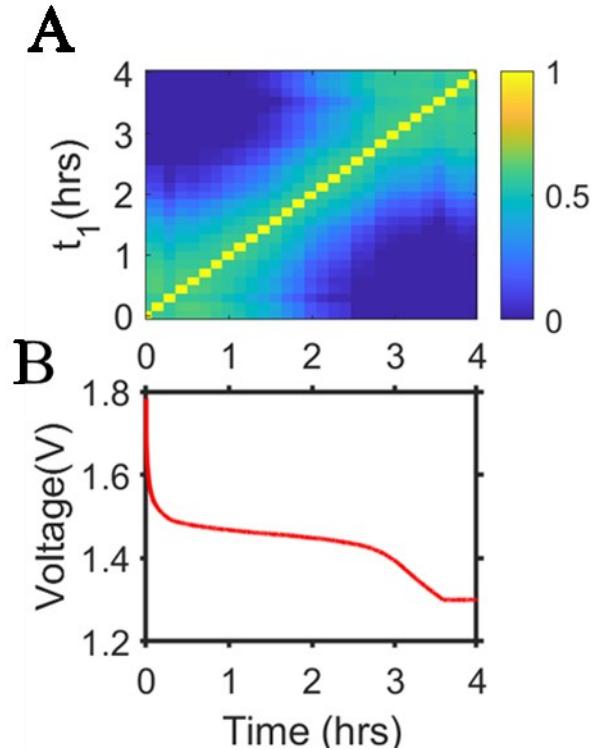


Figure 3. Monitoring structural phase transformation in LTO using (400) Bragg peak during C/2 discharge cycle. (A) Pearson Cross-Correlation of the (400) diffraction peak between different times during discharge. The correlation is calculated on the central diffraction pattern of the rocking data. (B) The voltage across the coin-cell as a function of time.

B. (111) Bragg Peak Results

On a different sample, we measured the (111) Bragg peak and calculated the Pearson correlation coefficient. The results in Fig. 4(A) show a clear phase transformation of the LTO sample during the discharge, similar to what was seen for the crystal in Fig 3. Figure 4 (C) shows reconstructed images of the LTO particle, where a clear transformation from a more uniform state to one with a stronger phase variation on its surface. The shape of the crystal was not constrained in the analysis and appears to show some small

variations; we interpret this as coupling of the surface displacement field with the isosurface contour level, and not a physical change. We attribute this phase variation to the coexistence of Li-poor and Li-rich regions, consistent with previously reported X-ray spectroscopic results^{30,46}. To visualize how the image phase transformation develops inside the LTO particle, we examine slices parallel to the Q vector of the reconstructed images shown in Fig. 5. In the early stage of the discharge state (down to 1.44V), there are no significant changes in the image phase variation within the crystal. However, below 1.35V (after 1 hr), there is a build-up of different sets of “nano-domains” seen as positively (red) and negatively (blue) shifted regions. Since this is the discharge stage where the sample is expected to be in transition between the two LTO phases⁴⁷, these nano-domains are attributed to Li-rich and Li-poor regions within a single nanocrystal. It appears that an intense image phase difference develops on one side of the LTO particle and propagates into the internal part of the particle. Although the observed image phase change is a consequence of Li-insertion, our observation does not provide direct evidence of Li-ion migration, for example, whether Li enters the core of the particle or remains confined to the outer edges. However, since the diffusion coefficient, D, of Li⁺ ions in LTO is $10^{-11} \text{ cm}^2/\text{s}$ ⁴⁸, the average distance that the Li⁺ ions can travel in the LTO material, calculated using Fick’s law $\langle x^2 \rangle = 4D\tau$, where τ is the diffusion time, is 2 μm in 1 hour. Because this distance is larger than the nanoparticles we are measuring, this means the Li⁺ ions should invade the whole LTO nanocrystal within an hour, and the changes we observe are then due to the local Li distribution.

To quantify the displacement field inside the reconstructed images, we calculated the standard deviation of the image phase over a range of different volumes selected in the

center of the particle as a function of discharge voltage, shown in Fig. 5(C). Similar trends during discharge are seen for all the choices of volume. The significant increase of standard deviation starting in the middle of the discharge plateau quantifies the appearance of inhomogeneous displacement fields within the particle, which is presumably due to the coexistence of $\text{Li}_4\text{Ti}_5\text{O}_{12}$ and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ regions within the sample.

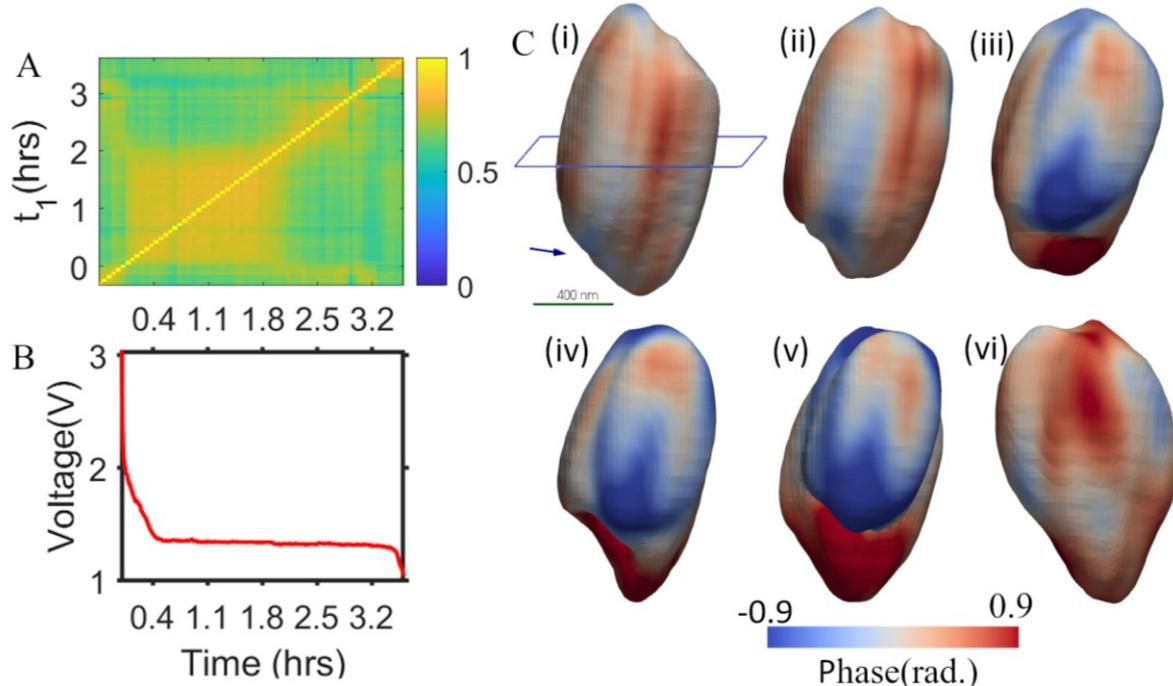


Figure 4. Evidence of the structural phase transformation in LTO during the discharging cycle. (A) Pearson cross-correlation plot of the central (111) diffraction peak of the different discharge states. (B) C/2 discharging cycle of coin-cell for the experiment as a function of time. The (i) –(vi) indicated in the curve are points for which the images of the LTO single particles were reconstructed. (C) (i) –(vi) *In-situ* BCDI images of the 3D LTO particle using the isosurface representation, colored by the phase value, measured at 3 V(pristine), 1.69 V(20 minutes), 1.36 V(40 minutes), 1.35 V(50 minutes), 1.35 V(1 hr), and 1.3 V (3 hrs). The blue arrow indicated the scattering vector (Q), and the green scale bar is 400nm. The blue plane shown in the pristine state shows the spatial location of the slices shown in Fig.5.

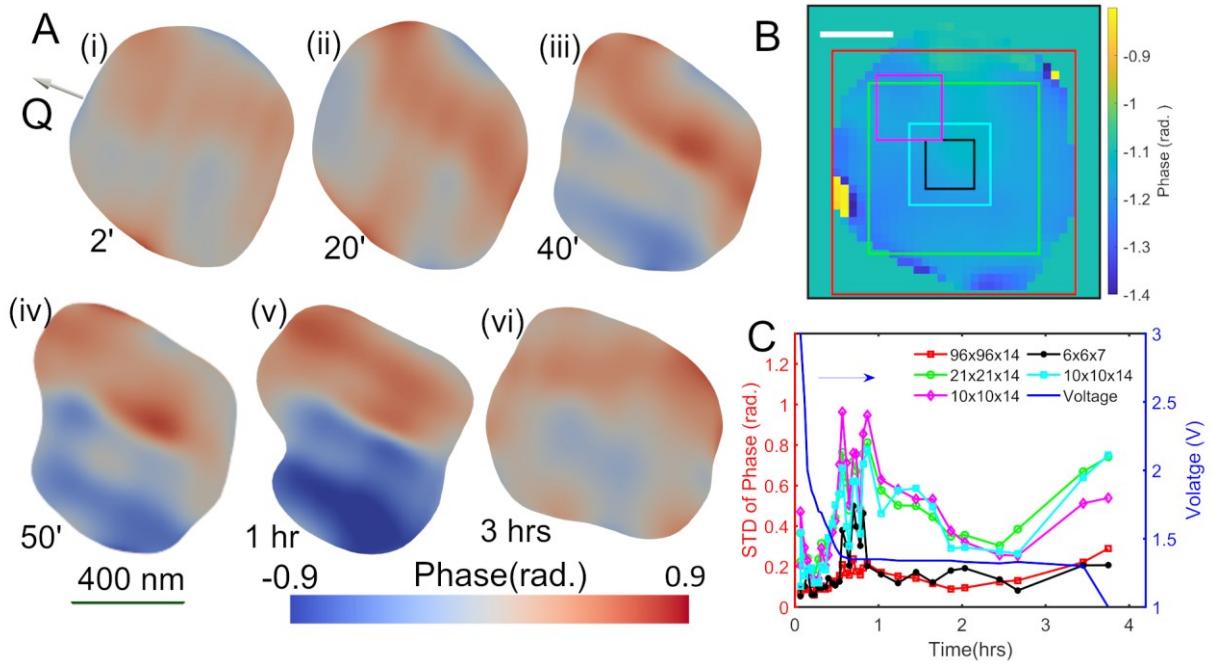


Figure 5. (A) (i)- (vi) Slices of the *in-situ* BCDI reconstructed image from its (111) Bragg peak at 3 V, 1.69 V, 1.36 V, 1.35 V, 1.35 V, and 1.3 V, respectively to show the internal displacement field development in the particle, and approximate time of the measurement is also indicated. The biggest image phase variation and relative image phase change is clear in the plateau region of the discharging voltage. (B) Central xy-slice of the reconstructed image (in the detector coordinates) with the selected voxel volumes shown as rectangles 96x96x14 (red), 21x21x14(green), 10x10x14(cyan), 6x6x7(magenta), and 10x10x14(black) measured at different discharge time. The white scalebar shown is 10 voxels and each voxel is about $(25 \text{ nm})^3$. (C) Calculated standard deviations (STD) and the discharge voltages for the corresponding different voxel volumes of the reconstructed image.

At its maximum, there is about 0.4 radians of inhomogeneity seen in the images, extending over regions of $\sim 100 \text{ nm}$ lateral dimension. This is consistent with the displacement field expected from side-by-side domains of the two material phases, estimated in the Introduction of this paper. A 3D image is shown in Fig. 6(A) in the middle of the inhomogeneous discharge state. The grey-colored background uses a low contour level to highlight the particle shape. Two clear domains with a gap in between can be seen at a higher amplitude contouring level with their surface image phases coded using

the color scheme shown. The apparent gap between the domains is due to the finite resolution and the chosen contour level. The phase difference of 1.12 rad. between the domains corresponds to a lateral shift (along the Q-vector) of about 1/6 of a (111) d-spacing, as illustrated by the atomic models shown in Fig. 6(B).

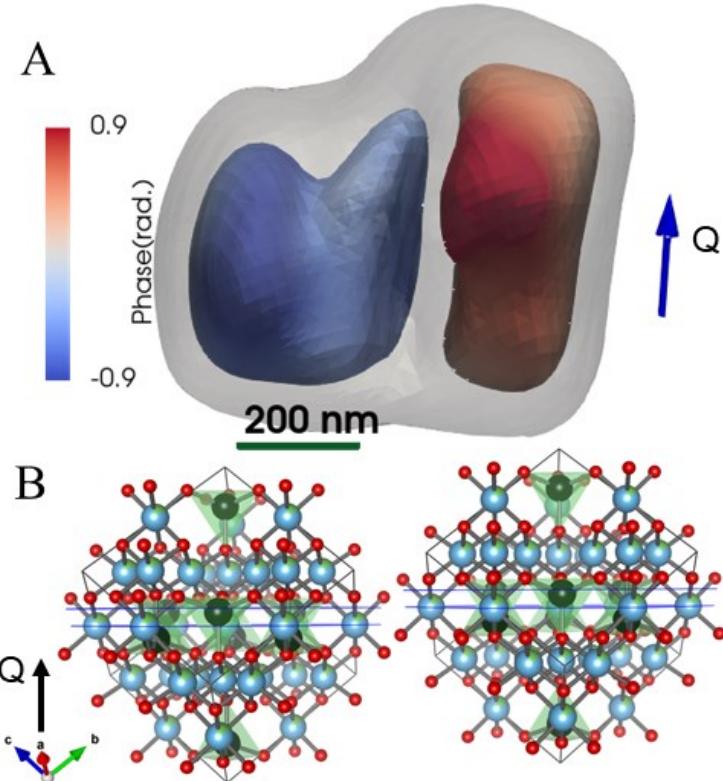


Figure 6. Material phase separation and relative image phase change in the two-phase stage. (A) Reconstructed (111) LTO single-particle image at 1.35 V, resulting in two-phase domains in the single LTO particle due to material phase separation. (B) The relative image phase change is 1.12 rad, which corresponds to the 0.8 Å translation along the Q-vector shown, which is equal to the Li-Ti plane separation shown.

CONCLUSION AND OUTLOOK

We studied the structural phase transformation of the anode material $\text{Li}_4\text{Ti}_5\text{O}_{12}$ during its lithiation process using *in-situ* BCDI measurements. When observing the (400) Bragg peak of a single LTO particle, we observed more than a 0.2% change in the d-spacing when comparing the pristine and fully lithiated states, which is significantly larger

than previously reported results from powder XRD experiments^{14,15,30,49}. This apparent discrepancy must be considered in the context of each measurement: the powder XRD yields the ensemble average lattice constant change of an ensemble average of a large volume of anode material, whereas the BCDI data result from a single LTO nanocrystal. Nanoparticles are well-known to respond differently from bulk material due to their larger relative surface area. The two-time correlation plots of the recorded coherent X-ray diffraction patterns demonstrate a clear structural evolution during the lithiation process. BCDI reconstruction, around the middle of the discharge plateau, shows the onset of Li-insertion is found to leads to a heterogeneous image phase distribution extending throughout the volume of within the nanocrystal which disappears again as the discharge completes. This informs us the formation of a lateral domain structure, implying a non-uniform lithium concentration between nano-domain regions lying side-by-side within the nanocrystal, assumed to have the $\text{Li}_4\text{Ti}_5\text{O}_{12}$ and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ compositions. The reconstructed image in Fig. 6 in the middle of the voltage plateau region shows a $2\pi/6$ image phase change between the two spatially distinct regions of the single LTO nanocrystal. This image phase shift corresponds to $d/6$ ($=0.8$ Å) rigid-body shift between the two regions, corresponding to a Li-Ti layer-spacing, illustrated in Fig. 6(B).

By monitoring the standard deviation of the image phase, we observed the biggest image variation in the middle of the plateau of the discharge where the transformation from $\text{Li}_4\text{Ti}_5\text{O}_{12}$ and $\text{Li}_7\text{Ti}_5\text{O}_{12}$ would be half-completed. The heterogenous local displacement field distribution permeating the LTO nanocrystal investigated was seen to build up during discharge and then taper off at the end of the cycle. At no point in any of our *in-situ* BCDI experiments did we detect the presence of any crystal defects, such as

dislocations, which are commonly observed in other battery material systems^{40,41}. Dislocation-mediated transformations are not expected to reverse perfectly upon charge cycling and would be expected to lead eventually to crystal degradation processes. Both our observations of lateral strain domain formation and the absence of mobile dislocations, with Burgers vector accessible in our experimental geometry, imply that the LTO crystal lattice remains completely intact during the charge-discharge cycle. This may explain the excellent cycling stability of LTO over other battery materials, which makes it suitable for durable, long-life batteries.

ASSOCIATED CONTENT

Supporting Information. Checking X-ray Beam Damage, Two-phase transformation of LTO particle, Displacement Projected along Q-vector, Lattice Constant Gradient, and In-situ X-ray Diffraction Results. This material is available free of charge via the Internet at <http://pubs.acs.org>

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AUTHOR CONTRIBUTIONS

T.A.A, F.W., and I.K.R designed the research project; T.A.A, L.W, A. S, W.C, L. X. L, R. H, F.W, and I. K. R conducted the BCDI experiments at the 34-ID-C, APS, T.A.A and I.K. R. analyzed the BCDI data, R.K analyzed XRD data with inputs from E.S.B. The

manuscript was written by T.A.A. and I.K.R., with contributions from all authors. All authors have given approval to the final version of the manuscript.

COMPETING INTERESTS

The authors declare no competing interest.

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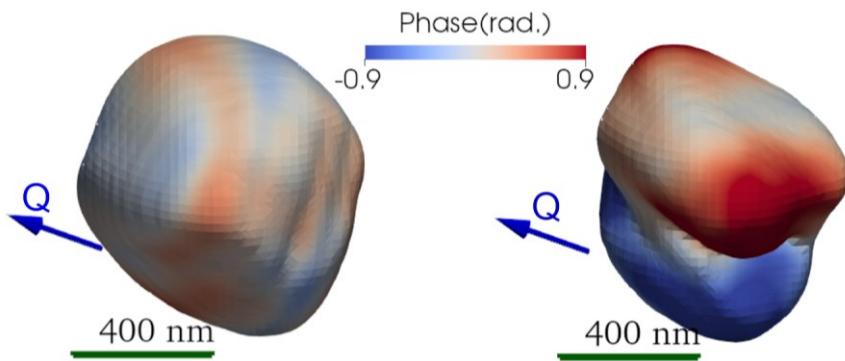
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SYNOPSIS

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SYNOPSIS

In-situ Bragg Coherent Diffraction Imaging (BCDI) and *in-situ* X-ray diffraction (XRD) experiments have been used to image the structural phase transformation within a single LTO particle during the battery discharge cycle of a lithium titanate (LTO) anode material. Analysis of the BCDI data revealed that the biggest image phase variation occurs at the two-phase stage, indicating the formation of structural phase domains during the discharge process. We also observed a lattice contraction in a single LTO nanoparticle larger than in the corresponding bulk material. This determination had not been possible using other experimental techniques (e.g., X-ray diffraction), demonstrating the potential for *in-situ* Bragg Coherent Diffraction Imaging (BCDI) methods to address scientific questions that are difficult to resolve by other means.