Multivariate hyperspectral data analytics across length scales to probe compositional, phase, and strain heterogeneities in electrode materials

Highlights
- Phase inhomogeneities and stress gradients govern electrode performance in batteries
- Hyperspectral imaging allows for the mapping of compositional variations
- Spectral databases are critical to investigating compositional heterogeneities
- Chemistry-geometry-mechanics coupling is crucial to lithiation phenomena

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In brief
In battery electrode materials, compositional heterogeneities give rise to stress gradients that ultimately result in degenerative failure. Progress in hyperspectral imaging has enabled an unparalleled view of multiphysics processes. Data science methods hold promise for deciphering mechanistic understanding from high-dimensional data. In this work, the utility of X-ray spectromicroscopy coupled with multivariate data analytics is demonstrated through multi-scale investigation of intercalation phenomena in a canonical cathode material aided by curation of a spectral standards database.
Multivariate hyperspectral data analytics across length scales to probe compositional, phase, and strain heterogeneities in electrode materials

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SUMMARY
The origins of performance degradation in batteries can be traced to atomistic phenomena, accumulated at mesoscale dimensions, and compounded up to the level of electrode architectures. Hyperspectral X-ray spectromicroscopy techniques allow for the mapping of compositional variations, and phase separation across length scales with high spatial and energy resolution. We demonstrate the design of workflows combining singular value decomposition, principal-component analysis, k-means clustering, and linear combination fitting, in conjunction with a curated spectral database, to develop high-accuracy quantitative compositional maps of the effective depth of discharge across individual positive electrode particles and ensembles of particles. Using curated reference spectra, accurate and quantitative mapping of inter- and intra-particle compositional heterogeneities, phase separation, and stress gradients is achieved for a canonical phase-transforming positive electrode material, $\alpha$-V$_2$O$_5$. Phase maps from single-particle measurements are used to reconstruct directional stress profiles showcasing the distinctive insights accessible from a standards-informed application of high-dimensional chemical imaging.

INTRODUCTION
Lithium-ion (Li-ion) batteries have expanded from consumer electronics to become a mainstay in electromobility applications and are starting to claim a growing share of the grid-level storage market.1,2 The stringent performance and safety requirements of medium- and large-area energy storage formats have spurred a greater focus on the atomistic mechanisms and multiphysics
coupling underpinning degradation phenomena. In many cases, the origins of performance degradation are traceable to multi-field and multiphysics coupling originating at atomistic scales, manifested at mesoscale dimensions, and compounded up to the level of electrode architectures. A detailed understanding of electrochemistry-mechanics coupling and resulting emergent phenomena across many decades of length scales is imperative to unlock unexploited performance from existing battery chemistries, develop dynamic process controls, and design next-generation materials and architectures that are purpose-built to alleviate common modes of degradation.

Degenerative capacity loss in batteries can derive from many sources, including loss of active material, stoichiometric drift, SEI formation, and rising internal resistance. For the active electrode components of Li-ion batteries, mechanical stresses developed during electrochemical cycling play a key role in many degradation mechanisms. The accumulation of stress underpins delamination, crack formation, exfoliation, and even accelerated dissolution/migration reactions at interfaces. In insertion electrodes, the insertion (extraction) of Li-ions drives successive dilation (contraction) of the host framework. The close coupling between mechanics and electrochemistry plays an especially prominent role in phase-transforming electrode materials. In these materials, intercalation gives rise to multiphase coexistence regimes, which result in significant coherency strains at the interfaces between differently lithiated phases. Direct observations of phase separation demonstrate a hierarchy of lithiation heterogeneities at the single-particle level, which are propagated across extended length scales, resulting in increased interfacial resistance, degraded rate performance, and loss of effective capacity. Despite a growing appreciation of the link between lithiation inhomogeneities and battery degradation, the mechanisms governing electrochemistry-mechanics coupling remains underexplored, especially with regard to localized phenomena. A primary impediment lies in probing these processes with adequate resolution and chemical sensitivity across length scales.

Increasingly, state-of-the-art electron microscopy and X-ray spectromicroscopy methods are being used to probe these processes using various contrast methods. At the micron and nanometer length scale, for example, cross-section analyses of diffractions electrons from electron backscatter diffraction demonstrate the role of grain orientation on electrode degradation. At the atomic scale, the transmission of electrons through electrode materials has been used to measure structural changes during cycling using transmission-electron microscopy. In X-ray spectromicroscopy, brilliant and highly tunable X-rays from a synchrotron source are used to map pixel/voxel-wise X-ray absorption signatures across a heterogeneous sample affording insights into the composition and critical atomistic and geometric correlates of intercalation phenomena. Compared with electron microscopy, X-ray methods offer high penetration power and a significant degree of tunability (enabling the collection of chemical information over a range of length scales) while minimizing sample damage. In a typical X-ray spectromicroscopy experiment, transmission intensities are recorded for a predefined field of view (typically between 4 and 100 µm²) over one or more elemental X-ray absorption edges; this technique thus provides spatially resolved chemical information with element and orbital specificity. A representative spectromicroscopy dataset can comprise as many as 1 million individual spectra and 250 images containing complex information about the chemical properties of a material mapped with nanometer-scale spatial resolution. A key challenge is thus to extract from these complex datasets a chemically intuitive understanding of compositional heterogeneities and their relation to particle dimensions and geometry.

A critical imperative for understanding degradation mechanisms is to map compositional variations to resulting stress gradients. Thus, for X-ray spectromicroscopy measurements that rely on absorption contrast, stress information needs to be derived from compositional gradients extrapolating from known structure parameters and stress tensors. Statistical regression and machine learning approaches hold promise for more effective utilization of hyperspectral datasets to correlate observed lithiation heterogeneities to geometric features and for the reconstruction of stress gradients. However, it is essential that such approaches be grounded in a physically meaningful interpretation of X-ray absorption spectra.

In this article, we demonstrate a workflow wherein multivariate statistical methods are used in conjunction with a library of reference spectra curated for samples with well-defined extents of lithiation to develop accurate chemical maps of lithiation gradients within individual particles and across ensembles of particles. At the individual particle level, composition maps from scanning transmission X-ray microscopy (STXM) and X-ray ptychography are further correlated with stress gradients with the help of finite element analysis (FEA). Using orthorhombic V₂O₅ as a model system, we examine phase evolution upon Li-ion intercalation and construct a library of X-ray absorption standards for samples that have been rigorously characterized by multimodal methods. By combining statistical approaches with curated reference spectra, we demonstrate a generalizable approach to composition, phase, and stress mapping, which provides critical insights into modulating battery chemistry and electrode architecture to alleviate common degradation mechanisms.

RESULTS AND DISCUSSION

A curated X-ray absorption spectroscopy reference database for Li₃V₂O₅

Orthorhombic α-V₂O₅ is a layered compound with multiple accessible redox couples (V⁵⁺/V⁴⁺ and V⁴⁺/V³⁺) and a framework that can support intercalation of up to three Li-ions per formula unit (i.e., up to a final composition of Li₅V₂O₅). The promise of this material as an intercalation electrode stems from a high theoretical capacity (up to 441 mAh/g) and the potential for multielectron redox. A growing appreciation for the geographical diversity of vanadium deposits and the maturity of extraction methods with low-carbon footprints has driven further interest in V₂O₅ in light of growing concerns about greenhouse emissions from battery manufacturing and the resilience of supply chains. Despite current use as a positive electrode (cathode) for small-format commercial cells, widespread adoption of V₂O₅ in larger formats has been stymied by a series of intercalation-induced phase transformations, which gives rise to stress and diffusion impediments and limits the accessible energy density and cycle life. Thus, while
specific emphasis is placed on the methodology of interpreting hyperspectral imaging data in this work, in light of increasing material criticality challenges faced by incumbent positive electrode chemistries, the insights obtained here the role of defects and particle geometry on electrochemistry-mechanics coupling in V$_2$O$_5$ have practical implications for next-generation batteries.\textsuperscript{32,40}

To develop a rigorous set of standard X-ray absorption spectra for use in hyperspectral image analysis, phase-pure stoichiometric Li$_x$V$_2$O$_5$ materials (where $x$ denotes the extent of lithiation) were prepared by reacting V$_2$O$_5$ powders with stoichiometric amounts of lithium iodide (LiI), as described in the experimental procedures.\textsuperscript{24} As V$_2$O$_5$ incorporates Li, its crystal lattice and electronic structure are strongly modified, as shown in Figure 1.\textsuperscript{11} Powder X-ray diffraction (PXRD) patterns for each phase are shown in Figure 1A, and the corresponding Rietveld refinements and structure solutions are shown in Figures S1 and S2, respectively. Tables S1–S6 show the Rietveld refinement statistics and atom positions; the evolution of lattice parameters is summarized in Table S7. Upon intercalation, V$_2$O$_5$ undergoes several discontinuous changes to its crystal structure, stabilizing $\alpha$-, $\epsilon$-, and $\delta$-phases with increasing lithiation, corresponding to lithium stoichiometries of ca. $0.0 < x < 0.1$, $0.3 < x < 0.7$, and $0.9 < x < 1.0$, respectively.\textsuperscript{34} With the exception of the highly lithiated $\delta$-phase, which necessitates a half-layer shift in the ab plane (Figure 1E), contraction of the lattice along the a axis, and expansion of the lattice along the b axis can be observed with increasing lithiation, as shown in Figures 1B and 1C, respectively. With increasing lithium content, electron density becomes localized on specific vanadium sites, the nearest apical oxygens become canting toward intercalated lithium-ions (Figure 1D), and the V$_2$O$_5$ layers are increasingly puckered (Figure S3; Table S8).\textsuperscript{41} These profound changes in bonding motifs coupled with the structural distortions give rise to inhomogeneous stress gradients as a result of anisotropic lattice contraction and expansion along different crystallographic directions.

Figure 1. Li-ion intercalation-induced evolution of crystal lattice and electronic structure
(A–C) (A) Powder X-ray diffraction patterns collected for phase-pure stoichiometric Li$_x$V$_2$O$_5$ materials demonstrate intercalation-induced (B) contraction of the lattice along the a axis and (C) expansion of the lattice along the b axis. Rietveld refinement of the powder diffraction data shown in (A) is shown in Figure S1. Structure solutions obtained by Rietveld refinement are shown in Figure S2.
(D) Li-ion intercalation further induces monotonically increasing canting of the apical oxygens toward the intercalated lithium-ion and puckering of the V$_2$O$_5$ layers (Figure S3).
(E) Finally, the VO$_{5+1}$ quasi-octahedron evolves as the sixth long bond gradually lengthens. In the highly lithiated $\delta$-phase, a half-layer shift in the ab plane gives rise to a purely VO$_5$ square pyramid coordination environment.
(F) Closely overlapping V L- and O K-edge X-ray absorption spectra obtained by analysis of STXM data for individual particles (Figures S4 and S5) demonstrate lithiation-induced modification of local coordination environments.
(G and H) Expanded regions of the (G) V L$_3$- and (H) O K-edges are provided. Shifts of absorption features to higher energies and a monotonic diminution in the intensity of pre-edge features are demonstrated with increasing lithiation. Linear curve fitting of these spectra is shown in Figures S6 and S7.
The phase transformations and changes in lattice symmetry observed by XRD are accompanied by changes in local vanadium and lithium coordination environments. These alterations are demonstrated in X-ray absorption near-edge structure (XANES) spectra, which depend sensitively on the local coordination geometry and ligand field. Interpretation of XANES spectra is complicated by multiplet effects, known limitations of density functional theory in predicting the electronic structure of complex oxides, and challenges in treating core holes. First-principles approximations, including multiple scattering, time-dependent density functional theory, and local pair natural orbital coupled cluster methods, and self-consistent-field methods coupled with many-body perturbation theory, have been used to simulate XANES spectra with varying degrees of success. First-principles approximations provide insight into the physical origins of spectral features but accurate treatments of core holes are often challenging to implement for many-body correlated systems, such as typical positive electrode materials, especially upon incorporation of defects and alloying. We present here a database of experimental spectra derived from STXM measurements (Figure 1F) acquired across V L- and O K-edges of phase-pure individual particles of \( \alpha \), \( \varepsilon \), and \( \delta \)-Li\( \text{V}_2\text{O}_5 \) (Figures S4). Principal-component and k-means clustering analyses have been performed for each measurement to validate the phase purity of the samples, as shown in Figure S5. The spectra in Figure 1F can thus be unambiguously assigned as spectral signatures of the differently lithiated phases with variations manifested in Figure S5 solely from detector non-linearity at higher thicknesses.

**Evolution of spectral features at V L- and O K-edges as a function of lithiation**

As shown in Figure 1F, the V L-edge (~510–525 eV) comprises two dominant features centered at ~518 (V L\( _3 \)) and ~525 eV (V L\( _2 \)). The V L\( _2 \)-edge results from transitions from V 2p\( _{1/2} \) \( \rightarrow \) V 3d states and remains relatively consistent across all phases. A Coster-Kronig Auger process that broadens spectral states and remains relatively consistent across all phases. A Coster-Kronig Auger emission process that broadens spectral states and remains relatively consistent across all phases. A principal-component and k-means clustering analysis has been performed for each measurement to validate the phase purity of the samples, as shown in Figure S5.

The relative intensities of the t\( _{2g} \) and e\( _{g}^* \) features in O K-edge XANES spectra in Figure 1H are also greatly modulated by the lithiation extent. With increasing lithiation, the t\( _{2g} \) feature is diminished in intensity with respect to the e\( _{g}^* \) absorption. This modulation of relative intensities is a consequence of electron correlation (since lithiation introduces V 3d\( _1 \) states), the loss of degeneracy between spin-up and spin-down states upon lithiation, and the concomitant reduction of vanadium centers (which modifies V 3d-O 2p hybridization), and the elongation and canting of V-O bonds (Figure S3; Table S9). The changes in V-O hybridization cause a shift in the spectral weight of the e\( _{g}^* \) and O 2p-V 4sp hybridized states to higher energies (Figure S7; Tables S9–S11 and S13). Notably, the pronounced structural rearrangement across the \( \varepsilon \rightarrow \delta \) phase transition does not follow this trend. Instead, a shift from a distorted octahedral (i.e., V\( \text{O}_5^{1+} \) environment) to an unambiguously square pyramidal coordination results in a change in the crystal field splitting marked by a re-emergence of a sharp 3d\( _{xy}^{2}\) feature centered near 517 eV at the V L\( _3 \)-edge and renormalization of the t\( _{2g} \)-to-e\( _{g}^* \) intensity at the O K-edge.

**Using reference spectra to guide composition and phase mapping by X-ray spectromicroscopy**

Access to validated reference Li\( \text{V}_2\text{O}_5 \) XANES spectra (Figure 1) enables accurate and efficient interpretation of lithiation heterogeneities imaged by high-dimensional X-ray imaging techniques, such as STXM and X-ray ptychography. In STXM, transmission images are obtained over a series of image positions, \( p = 1 \ldots P \), along a series of energies \( n = 1 \ldots N \). During pre-processing, the incident flux \( I_0 \) is utilized to calculate an optical density matrix to establish a linear relationship between the absorption coefficient (\( \mu \)), which is dependent on the extent of lithiation and thickness (\( t_0 \)). For a heterogeneous sample comprising a set \( s = 1 \ldots S \) number of spectroscopically distinct constituents, the optical density, \( D_{N \times P} \), can be expressed as a spectrum for each pixel as:

\[
D_{N \times P} = \mu_{N \times S} \cdot t_{S \times P}.
\]  
(Equation 1)

where \( \mu_{N \times S} \) represents the set of exact spectra for all the \( s = 1 \ldots S \) and their corresponding weighting at every pixel is denoted by \( t_{S \times P} \). Thus, provided \( D_{N \times P} \) from X-ray spectromicroscopy measurements, the goal is to find solutions for the set of spectra (\( \mu_{N \times S} \)) and weighting maps (\( t_{S \times P} \)) which accurately represent the contributions of each component at individual pixels. A workflow for these processes and the construction of stress maps is illustrated in Figure 2. When the exact set of absorption spectra (\( \mu_{N \times S} \)) is well defined, spectral maps (\( t_{S \times P} \)) can be obtained by matrix inversion using singular value decomposition (SVD) as follows:

\[
t_{S \times P} = (\mu_{N \times S})^{-1} \cdot D_{N \times P}.
\]  
(Equation 2)

Furthermore, when spectral signatures are known for each composition, hyperspectral maps can be used to visualize composition gradients. This scheme, where \( \mu_{N \times S} \) is directly defined using the curated spectral library acquired for standards, is described in this work as a direct deconvolution, which is represented in the overall workflow sketched in Figure 2 by the direction of the orange arrows. Alternatively, when a direct convolution with reference spectra is not possible (as a result of the presence of impurities, energy calibration errors, or excess noise), additional steps are required to find a solution for \( \mu_{N \times S} \). An example of this scheme is represented in Figure 2 by the direction of the teal arrows (i.e., principal-component analysis (PCA) → reduced-dimensionality search space → cluster analysis → \( \mu_{N \times S} \)). A more detailed presentation of the matrix notation.
A region of interest is imaged across a range of incident photon energies spanning one or more elemental absorption edges. A frame stack can be generated by appropriately aligning the images to account for beam drift and stacking them along the energy dimension. A background spectrum is utilized to transform transmission images to optical density data represented as a matrix, $D_{NKP}$. If a direct analysis with reference spectra is possible, spectral maps can be generated by SVD (indicated by orange arrows). Alternatively, PCA can be utilized to aid cluster analyses of pixels based on spectral similarity; comparison of the cluster spectra against reference spectra informs accurate assignments of spectra to $Li_xV_2O_5$ phases (indicated by teal arrows). Composition maps can be further used to map intercalation-induced stresses by FEA to extract insight into electrochemistry-mechanics coupling. The latter is a purely physics-based calculation and solves for linear elastic mechanics in 2D.
for this process is available from Lerotic et al.50,51 and Kopri- 
narov et al.52

Initially, we describe these approaches using the synthetic data-
tset shown in Figure 3A to facilitate discussion and validation of 
each methodology. In particular, we seek to effectively utilize the 
measured spectral signatures of distinctly lithiated phases (Fig-
ure 3B) to develop a transparent and physically meaningful 
framework for interpreting large hyperspectral datasets. The 
synthetic dataset shown in Figure 3A consists of three spectro-
scopically distinct (Figure S8) regions comprising phase-pure 
\( x \)-Li\(_{0.1}\)V\(_2\)O\(_5\) and \( \epsilon \)-Li\(_{0.45}\)V\(_2\)O\(_5\) in addition to a Li\(_{0.35}\)V\(_2\)O\(_5\) 
a/\( \epsilon \) phase 
mixture oriented in a core-shell motif, 53 a common multiphasic 
lithiation pattern observed during ion intercalation. In a direct 
deconvolution of the synthetic dataset, a set of images (\( tS \)) 
representing the weighting of each spectrum in the sample 
(Figure S9), is obtained by SVD as described in Equation 2 , 
wherein \( m \) has been defined using the library of XANES stan-
dards. When plotted using a uniform grayscale, the spectral 
maps corresponding to the phases present in the dataset are 
clearly represented (S9B and S9D) in sharp contrast to the re-
mainig spectral maps corresponding to V\(_2\)O\(_5\) phases that are 
not present in the sample (S9A, S9C, S9E, and S9F). In this 
example, spectral maps are calculated for the entire library of 

XANES spectra, and the relative weight-
ings of each component are used to 
deduce which phases are present in the 
sample. As shown in the composition 
map in Figure 3D, this approach captures 
the artificial dataset’s ground-truth, a 
core-shell phase separation motif, with excellent fidelity (Fig-
ure S8). It is worth noting, however, that prior knowledge 
about the sample, such as depth of discharge, can be utilized 
to narrow the selection of spectral signatures considered for 
SVD (i.e., to visualize the spatial orientation of known phases in 
the sample).

It is important to note here that the success of a direct decon-
volution with standards stems in some measure from the fact 
that the validation dataset was initially generated using reference 
spectra; this represents the best-case scenario wherein the input 
spectra used for deconvolution are an excellent match to the 
measured data. While the great utility of a direct analysis cannot 
be understated, a direct spectral deconvolution with standards 
comes with some caveats. For example, while the matrix inver-
sion process described in Equation 2 can be applied to any 
combination of input spectra, the fidelity of the spectral 
maps depends strongly on the extent to which the reference 
spectra are a one-to-one representation of the measured data 
(\( D_{\text{inv}} \)).52 Here, sample thicknesses above a certain threshold 
(inducing non-linearities), parasitic reactions generating non-
vanadium oxide phases, environmental noise, errors in the energy 
calibration, potentially polarization dependences of spectral 
intensities arising from particle orientation,54 and contributions
from the halo of the incident beam can lead to erroneous results. An illustration of this is provided in Figure S10, which shows invalid results produced by a direct SVD, using the library of standards applied to a modified dataset incorporating a $-0.4$ ev energy calibration error and the addition of Gaussian noise.\(^56\) One approach to minimize this limitation is a collection of internal standards from known phase-pure regions; however, the complexity of this approach greatly increases with the number of chemical constituents. In this work, a considerable amount of effort has been placed on minimizing many of these sources of error during data collection, for example, by limiting sample thickness and mitigating the formation of non-vanadium oxide phases, as discussed in the experimental procedures.

We have examined an alternative approach to analyzing hyperspectral datasets acquired for complex samples, which is the most likely scenario for realistic energy storage materials. This approach, illustrated by the direction of the teal arrows in Figures 2 and 3, fuses knowledge of standards (Figure 3B) with PCA and cluster analysis (Figure 3C) to find spectrosopically similar groupings of data that represent distinct $Li_3V_2O_8$ phases. The first step in this approach aims to provide a reduced dimensionality search space that retains all of the meaningful variations in the data—this can be accomplished by PCA, as shown in Figure S11. Next, pixels with similar spectral signatures can be grouped according to their Euclidean distance in the eigenvector search-space provided by PCA; here, the average spectral response of every pixel belonging to a cluster can be used to define the set of spectra, $\mu_{\text{HXS}}$, as shown in Figure S12 for the test set. In this approach, the number of selected principal components and clusters is user adjustable and critical to an accurate determination of $\mu_{\text{HXS}}$. Without access to standards, selecting the number of principal components can be aided by referencing their cumulative variance.\(^56\) Indeed, as shown in Figure S11 for the test dataset, the majority of the variance in the data is captured by the first few components (>99.9% from the first two components in the test set). Subsequently, during clustering analyses, color-coded scatterplots of clusters, as shown in Figure S12A, can be utilized to investigate how clusters are separated in eigenvector space to inform the number of user-selected clusters that form $\mu_{\text{HXS}}$. In an ideal execution of k-means clustering via PCA, the number of partitions and their spectral identity should match the chemically distinctive constituents present in the sample (contrast Figure S8 with Figure S12)—in practice, this can be difficult to achieve by solely monitoring the statistical properties of the data. In this work, signatures of the eigenspectra (which are often reminiscent of the difference spectra between standards) and cluster spectra are validated using the library of X-ray absorption standards to enable an electronic-structure-informed selection of the number of principal components and clusters selected for $\mu_{\text{HXS}}$; Figures S11 and S12 show that this supervised approach, guided by standards to PCA and clustering, respectively, can successfully partition data based on statistical deconvolution to obtain a set of spectra $\mu_{\text{HXS}}$, even in the presence of Gaussian noise and energy misalignment in the modified dataset.\(^56\) A combined PCA and k-means clustering approach thus represents a powerful means of determining $\mu_{\text{HXS}}$ of the spectral components based on statistical deconvolution of datasets.\(^50,52,67-68\) Notably, the cluster analysis in Figure S12 delineates three distinct clusters despite the presence of only two $Li_3V_2O_8$ phases in the generated dataset. These results thus highlight a crucial distinction between direct deconvolution aided by spectral standards and deconvolution based on clustered spectra. The former can disentangle the individual contributions of multiple phases. In contrast, the latter must assign regions of mixed composition to either a separate cluster or as part of a closely related cluster. One approach to resolving this uncertainty is to over-cluster the data to achieve greater segmentation and enable interactive curation by analyzing dendrograms (based on Euclidian distance) or comparison with standards.\(^58,59\) An additional method for refining the search space has been highlighted by Lerotic and co-workers, wherein cluster-similarity regularization is incorporated with non-negative matrix analysis to introduce a physically meaningful bias toward properties of known spectra.\(^60\)

In all instances, it is crucial to be cognizant of each approach’s limitations and monitor any residual errors to avoid erroneous results. Using PCA, for example, has been shown to produce negative weighting values, which are unphysical and represent a limitation to its utility for quantitative interpretation.\(^50\) Although the fine spectral features of the X-ray absorption spectra provide the physical basis for distinguishing components in a heterogeneous material, spectra should be placed on an absolute mass-absorption scale to avoid fitting peak intensity differences stemming from thickness effects—this can introduce systematic errors resulting from uncertainties in the mass-absorption coefficients.\(^52\) These and other sources of error can be monitored by tracking residual maps (Figure S9G) and residual plots, as shown in a subsequent section. Whether employing a direct deconvolution of datasets using standards or a combined approach using PCA and cluster analysis, access to a library of reference spectra enables user-supervised validation to ensure physically meaningful interpretation of the data.

The spatially resolved weighting of each phase, either determined by direct deconvolution or abetted by the use of reference spectra and statistical methods, can be utilized to generate composition/concentration maps, providing a quantitative representation of the extent of lithiation (or depth of discharge in the case of electrochemically lithiated samples).\(^75\) To reduce noise and enable a continuous representation of the data, spectral maps are subjected to a Gaussian filter and bilinear interpolation step (see experimental procedures). The reconstructed composition map for the test particle, shown in Figure 3D, thus reproduces the spatial localization of the $\alpha$-$Li_01V_2O_8$ and $\epsilon$-$Li_{0.45}V_2O_8$ phases, their phase mixtures, and the relative thickness gradient within the particle (Figure S8) with fine detail and excellent fidelity to the original dataset. Figure 3E shows von Mises stress values obtained for the test particle by translating composition gradients to stress via FEA as described in the supplemental experimental procedures.\(^24,34\) The process of translating composition maps to stress gradients is physics-based and involves FE calculations, solving 2D linear elastic mechanics with corresponding boundary conditions. The commonly used von Mises stress (2D) criterion is applied to compute an equivalent stress of the Cauchy stress tensor at every point within the particle, thereby providing a snapshot of the overall stress state resulting from lithiation heterogeneities.\(^61\)
In general, regions of high von Mises stress correlate with highly lithiated domains as a result of compressive stresses. The most significant stress experienced by the test particle coincides with phase boundaries, reflecting the sharp changes in lithiation and elastic misfit strain at the $\alpha$-Li$_{0.1}$V$_2$O$_5$ and $\epsilon$-Li$_{0.45}$V$_2$O$_5$ interface. A complete picture of lithiation-induced stresses is provided in Figure S13, wherein the complex stress state is deconvoluted into the von Mises, normal, and shear stress components under fixed and free boundary conditions. Here, the choice of boundary conditions is intended to capture the likely varying interactions, in a real electrode, between the active primary particles and the surrounding electrode, which can significantly influence the stress state of a given particle. It is important to note here that, while linear elasticity does not directly predict fracture, it provides an overall snapshot of the stress state, which captures the intercalation-induced stresses that typically precede failure. Delineation of displacement, von Mises stress, nominal stress along $x$, and shear stress along $y$ for fixed boundary conditions in Figure S13 provides an unprecedented view of chemistry-mechanics coupling in insertion cathodes.

In summary, we have demonstrated that spectral standards can enable accurate chemical mapping using a synthetic hyperspectral dataset generated from reference spectra. In the absence of spectral contamination, reference spectra can be used directly to deconvolute contributions from different phases within a heterogeneous sample. Alternatively, reference spectra enable a supervised approach to spectral deconvolution in conjunction with statistical clustering approaches in a more general scenario. The composition maps generated from the accurate assignment of the spatial disposition of lithiated phases can furthermore be used to reconstruct a detailed picture of stress gradients within individual particles. We now focus on applying standards to experimental STXM data collected for heterogeneous Li$_x$V$_2$O$_5$ nanoparticle samples to demonstrate the utility of this approach. Phase distributions are investigated across length scales with the help of our curated reference spectra (Figure 1F). In subsequent sections, lithiation heterogeneities are mapped at the level of individual particles, small ensembles, and particle networks.

### Evaluating lithiation heterogeneities and stress gradients within individual particles

We start at the smallest length scale with hyperspectral data of the highest spatial resolution. The lithiated V$_2$O$_5$ nanowire fragments shown in Figure 4 were initially screened by STXM (Figure 4A) at 50 nm resolution and subsequently by X-ray ptychography at 10 nm resolution (Figure 4B). X-ray ptychography is a coherent diffraction imaging technique capable of making hyperspectral measurements, bearing resemblance to STXM as an element-specific probe, with marked improvements in spatial resolution, as is readily apparent by contrasting Figure 4A (STXM) with 4B (ptychography). As a lens-less technique, the spatial resolution of X-ray ptychography is limited only by the wavelength of the incident photon energy. Given the massive datasets generated by spectroptychography (on the order of terabytes), STXM can be utilized as a preliminary screen to identify a limited set of photon energies that best capture chemical contrast. Figures 4C and 4D show ptychography maps constructed from coherent diffraction patterns at 518.2 and 519.4 eV, respectively, which correspond to the $d_{xy}$ and $d_{xzy}$
absorption features at the V L_{3}-edge as shown in the integrated absorption spectrum derived from STXM in Figure 4E. Optical density maps at 518.2 and 519.4 eV have been normalized by an optical density profile collected at the post-edge (555.0 eV) to eliminate mass thickness contrast. This enables the comparison of relative d_{xy} and d_{xy/yz} absorption intensities, which, as observed in Figure 1G, are powerful indicators of the extent of intercalation. The inferred lithiation gradients from individual d_{xy} and d_{xy/yz} images are consistent and broadly depict core-shell behavior wherein nanowire edges are sparsely lithiated relative to a Li-rich core. A linear combination fit (LCF) with energy-aligned standards has been performed on the STXM data to enable a more quantitative look at phase separation, as shown in Figure S14. The results from LCF indicate a nearly evenly mixed phase of ε-Li_{0.1}V_{2}O_{5} and ε-Li_{0.30}V_{2}O_{5}. The concentration map shown in Figure 4F has been computed based on pixel intensities from Figure 4G (indicative of lithiation) within lower and upper bounds for lithiation set by the ε-Li_{0.1}V_{2}O_{5} and ε-Li_{0.30}V_{2}O_{5} phase. Notably, the tips of both nanowire fragments appear heavily lithiated, likely due to preferential nucleation at fragmented edges with a large density of defects, which lower the barrier to nucleation. The spatial resolution of X-ray ptychography enables visualization of additional Li-rich channels originating from the nanowire edges and running along the length of the nanowire fragments. Here, the insertion of lithium ions between specific layers likely drives further insertion into the expanded layer, a phenomenon akin to staging, thus driving preferential diffusion along locally expanded layers.

A high-resolution composition and phase map enables visualization of stress through finite element simulations, as shown in Figure 4G. Here, we primarily discuss the von Mises stress since it provides, at a glance, an expression of the overall stress state of the material. It should be noted here that stress values are calculated (not directly imaged) utilizing the composition map combined with knowledge of the partial molar volumes of the individual variously lithiated phases. The most significant regions of von Mises stress coincide with areas of high lithiation, reflecting the compressive strain induced by a locally expanded Li-rich core against a sparsely lithiated shell. Striations oriented perpendicular to the nanowire length are observed due to boundary constraints that account for additional compression or tension due to volumetric expansion or contraction against the substrate. Secondary areas of elevated von Mises stress coincide with highly lithiated nanowire tips; lithiation-induced local compression of the crystal lattice likely alters the local diffusion rate, exacerbating lithiation differences and driving additional coherency strain.

A full profile of stress calculations from FEA is shown in Figure S15—which provides a rich picture of lithiation-induced stress gradients. For the particle fragments shown in Figure 4, interparticle diffusion is limited, and the thermodynamic driving forces for phase separation are accommodated by intraparticle lithiation gradients. Furthermore, defects at frayed nanoparticle tips strongly modify nucleation and subsequent phase propagation. As such, Figure 4 provides a compelling demonstration of the use of single-particle X-ray spectroscopy and X-ray ptychography aided by reference spectra to image phase separation with <10 nm spatial resolution. Future work will focus on acquiring and utilizing phase images provided by X-ray ptychography with greater spectral resolution to maximize contrast in individual particles and cross-sectioned single crystals.

**Imaging particle-by-particle lithiation in small ensembles of V_{2}O_{5} nanoparticles**

We next turn our attention to a small ensemble of V_{2}O_{5} nanoparticles to examine the role of interfaces between active electrode particles. Figure 5 shows interparticle heterogeneity in two pairs of interfacing nanowires following “on-substrate” lithiation described in the experimental procedures. This approach to chemical lithiation enables the investigation of interactions between adjacent or solid-solid interfaced particles during Li-ion insertion. The spatial resolution in Figures 5A and 5B is 30 and 50 nm, respectively, which is sufficient for investigating interparticle and intraparticle lithiation inhomogeneities. Spectral deconvolution by SVD reveals comparably low and high lithiated phases, demonstrating the strongly preferential lithiation of one particle at the expense of the other during early (Figure 5A) and later stages of lithiation (Figure 5B).

The spectrum for the low lithiated nanowire in Figure 5A (orange) is shown in Figure 5C and is characterized by sharp d_{xy} and d_{xy/yz} features at the V L_{3}-edge, as well as a relatively strong t_{2g} feature compared with the e_{g} feature at the O K-edge. These features collectively suggest little-to-no lithium insertion. These features collectively suggest little-to-no lithium insertion. The blue spectrum is diminished and has broader lineshapes, indicating some reduction of the formal vanadium oxidation state. Furthermore, the O K-edge exhibits a lower t_{2g}-to-e_{g} ratio, demonstrating a relatively higher degree of lithiation. Particle-by-particle lithiation is driven by a strong energetic preference for diffusion across an interface, which is preferred over nucleation of a lithiated phase. Analogous mosaic instabilities have been modeled by Delmas et al. and experimentally shown by Chueh et al. for another phase-separating positive electrode material, LiFePO_{4}. This mechanism is characterized by phase coexistence within particle ensembles but not within single particles since diffusion across interconnected particles can mitigate the formation of phase boundaries. Indeed, PCA and k-means clustering analysis performed for isolated segments from each nanowire suggest little-to-no intraparticle lithiation inhomogeneities (Figure S16).

Unlike LiFePO_{4}, which has a singular miscibility gap and shows two-phase separation between Li-rich and Li-poor phases, lithiation of orthorhombic V_{2}O_{5} progresses through a sequence of Li_{x}V_{2}O_{5} phases (Figures 1 and S17), thereby introducing multiple energetic barriers for phase nucleation. The spectrum shown in Figure 5D for the lowly lithiated nanowire (yellow, Figure 5B) is most similar to the ε-Li_{0.3}V_{2}O_{5} phase. In sharp contrast, the adjacent particle (blue) shows a much greater extent of lithiation corresponding to ε-Li_{0.9}V_{2}O_{5}. Notably, lithiation beyond x > 1 in V_{2}O_{5} leads to the irreversible formation of a γ-phase requiring considerable distortion and inversion of the square pyramids in addition to the sliding of layers. Here, in a particle-by-particle sequence of lithiation, nucleation of the ε-Li_{0.3}V_{2}O_{5} phase in a previously unlithiated particle is energetically preferred over nucleation of the highly incommensurate γ-phase in the Li-rich particle. Such a sequence of phase transformations demonstrates an interplay between the thermodynamic transformation barrier and a barrier to interparticle transport. At the electrode scale, interparticle heterogeneities can...
lead to significant variations in current density, driving high local current hotspots that can serve as a stress concentrator, nucleate crack formation, and accelerate degradation. Approaches for mitigating interparticle differences include the formation of mesoscale architectures that maximize concurrent nucleation events and modifying intercalation phase diagrams to stabilize extended solid-solution lithiation regimes. As such, analysis of STXM composition and phase maps aided by reference spectra provide insight into the nature of interfaces between positive electrode particles, illustrating a preference for interparticle heterogeneity over intraparticle phase boundaries when particles are connected through interfaces that enable facile Li diffusion. The heterogeneous transformation of individual particles drives a large fraction of the current to a small fraction of the active material, thereby contributing to local thermal and stress gradients.

**Quantitative maps of mesoscale lithiation heterogeneities**

At mesoscale dimensions, the spatiotemporal evolution of phase propagation fronts strongly depends on particle dimensions and geometry, their interfaces, (dis)charge rates, and their relative position in the electrode architecture. An example of this is highlighted in Figure 6A, which shows the galvanostatic discharge/charge profiles of nanosphere (0D), nanowire (1D), nanoplatelet (2D), and micron-sized platelet bulk (3D) geometries (corresponding SEM images shown in Figures 6B–6E, respectively). Here, several voltage plateaus, corresponding to the individual phase transitions in the V₂O₅ phase diagram crucial to investigate how these irregularities are manifested across length scales from ensembles to entire porous electrodes.

**Figure 6F**, coincide with multiphase coexistence. Most notably, in this series, a size dependence on specific discharge capacity is observed, which reflects the higher internal resistance and diffusion limitations of larger particles. Not surprisingly, several efforts to mitigate transport limitations in positive electrode materials include a reduction in particle size. Nevertheless, a large fraction of commercial cathodes, prepared via slurry comprising the active material, binder, solvent, and other conductive additives, contain several geometric irregularities that can engender large heterogeneities. To this end, it is crucial to investigate how these irregularities are manifested across length scales from ensembles to entire porous electrodes.

**Figure 5. Particle by particle lithiation at V₂O₅ particle interfaces**

(A–D) Nanowires lithiated by an “on-substrate” method exhibit particle-by-particle lithiation. Two sets of nanowires are shown in (A and B), wherein the colored regions in the composite maps correspond to the spectral components shown in (C and D), respectively, as identified by SVD with spectra from a combined principal-component and k-means clustering analysis informed by comparison with reference spectra. Orange (C)/yellow (D), and blue (C/D) regions demarcate locally low and highly lithiated wires and their corresponding spectral identity. Scale bars, 1 μm.
Figure 6. Quantitative analysis of lithiation heterogeneities on an ensemble of Li$_x$V$_2$O$_5$ nanoparticles

(A–E) (A) Discharge/charge profiles of various α-V$_2$O$_5$ particle geometries at a C-rate of C/5 for the first cycles between 2.0 and 4.0 V. The morphology of the (B) nanospheres, (C) nanowires, (D) nanoplates, and (E) micron-sized “bulk” platelets is depicted by SEM. Scale bars, 100 nm (B) and 1 μm (C–E), respectively.

(F) A simplified phase diagram showing the evolution of lithiation-induced Li$_x$V$_2$O$_5$ phases is included in (F).

(legend continued on next page)
Notably, the mode values for the surface area of the closely lithiated \( \varepsilon \)-Li\(_{0.3}\)V\(_2\)O\(_5\) (12.4 \( \mu \)m\(^2\)) and \( \varepsilon \)-Li\(_{0.45}\)V\(_2\)O\(_5\) (11.9 \( \mu \)m\(^2\)) phases are in proximity, and in contrast to the formation of \( \delta \)-Li\(_{0.3}\)V\(_2\)O\(_5\), which is observed primarily in particles with a 6.8 \( \mu \)m\(^2\) mode surface area. It should be noted here that the calculated surface area for smaller particles, in particular, is an overestimation due to feathering at particle edges, resulting from limited spatial resolution and the presence of agglomeration.

Interestingly, a significant fraction of the larger particles in this distribution show regions that can best be represented by either \( \varepsilon \)-Li\(_{0.3}\)V\(_2\)O\(_5\) or \( \varepsilon \)-Li\(_{0.45}\)V\(_2\)O\(_5\), demonstrating compositional differences within individual particles (albeit within a single solid-solution regime) above a critical size threshold. This heterogeneity is likely driven at least in part by geometric asphericity\(^{15}\) as well as the presence of extended defects within particles.\(^{10,34}\) This quantitative view of phase coexistence and its size dependence provides a compelling illustration of the use of reference spectra acquired for well-defined standards. Through the lens of electrochemistry-mechanics coupling, scaling down particle dimensions increases the relative penalty associated with phase-boundary-induced interfacial mismatch, thus suppressing phase separation, an origin of stress and battery degradation.\(^{24}\) Indeed, this quantitative mapping underscores the interplay between crystallite geometry and lithiation heterogeneities; whereas micrometer-sized platelets of V\(_2\)O\(_5\) exhibit considerable phase heterogeneities across the thickness of the electrode; electrodes comprising homogenous-nanometer-sized spherical V\(_2\)O\(_5\) show remarkably homogeneous lithiation with extended solid-solution lithiation.\(^{16}\) Together, the results in Figure 6 attest to the importance of controlling electrochemistry-mechanics coupling to enhance electrode performance.

A closer investigation of the size distribution of the three phases in Figure 6 reveals that, following lithiation (discharge), a limited number of large and small particles remain lithiated (discharged) and delithiated (charged), respectively, reflecting a minority population that shows deviations from the norm. Indeed, this suggests that particle size is not the sole determinant of the sequence of intercalation events. As shown in Figures 4 and 5, for example, the presence of defects and interparticle connectivity can significantly alter phase separation patterns. In view of this, local considerations of electrode materials, proximity and accessibility of Li-ion exchange with adjacent particles, and geometric asphericity play a significant part in driving the observed intercalation phenomena.\(^{15,30,34}\) This coupling between geometry, electrochemistry, and mechanics remains inadequately understood. However, it is clearly critical to realizing the full potential of intercalation electrodes,\(^{84}\) such as V\(_2\)O\(_5\) as demonstrated by recent studies leveraging geometry\(^{16,34}\) strain\(^{24,85–87}\) and atomistic modifications\(^{35,88}\) as levers to modulate intercalation phase diagrams to circumvent significant phase inhomogeneities and stress gradients. Pint and co-workers demonstrated that straining V\(_2\)O\(_5\) thin films through coupling with shape-memory alloys modulated intercalation potentials by as much as 40 mV and provided a 2.5× increase of the Li-ion diffusion coefficient.\(^{85,86}\) Electrochemistry-mechanics coupling in cathode materials is a multiphysics problem that occurs over decades of length scales from atomic to mesoscale dimensions; visualizing these interactions through hyperspectral imaging, as shown in this work, is a step toward deciphering design principles for mitigating the consequences of battery degradation.

**Conclusions**

The fundamental mechanisms of chemistry-geometry-mechanics coupling in Li-ion batteries remain inadequately understood but continue to be of pivotal importance to the lifetime of commercial batteries. Atomistic phenomena propagate across mesoscale dimensions, manifesting as phase inhomogeneities and stress gradients, which strongly impact the performance and longevity of electrode architectures. This work demonstrates the remarkable utility of synchrotron-based hyperspectral X-ray spectromicroscopy experiments, which, when coupled with spectral standards, enable accurate and quantitative mapping of inter- and intraparticle compositional heterogeneities, phase separation, and stress gradients in the canonical positive electrode material, \( \varepsilon \)-V\(_2\)O\(_5\). A detailed account of Li-intercalation-induced changes in the crystal lattice and electronic structure is developed to curate a robust set of physically interpretable spectral standards. Distinctive workflows, including SVD, PCA, k-means clustering, and linear combination fitting, are deployed in conjunction with reference spectra to develop high-accuracy quantitative maps.

Soft X-ray nanotomography, informed by standards, enables high-resolution visualization of lithiation gradients, illuminating the role of defects and substrate coupling during nucleation-limited lithiation. Experimentally determined phase and composition maps at the level of individual particles are translated to stress gradients, enabling a direct link between chemistry and mechanics and providing key insights into the role of particle geometry in stress concentration. STXM imaging of interconnected particles reveals the delicate balance between barriers for phase nucleation, the energetic penalties associated with stabilizing phase boundaries, and Li-ion transport across interlinked particles. Finally, quantitative analysis of phase separation across an ensemble of nanoparticles demonstrates the role of particle size and geometry in governing phase separation and phase evolution. Deterministically controlling the trajectories of intercalation reactions remains a grand challenge for battery diagnostics and prognostics. Future efforts will focus on combining the chemical and spatial resolution of hyperspectral imaging with...
temporal resolution within operando cells to track intercalation phenomena under an electrochemical stimulus with an eye toward developing design principles for real-time process control of key processes. In addition to the challenges of extracting correlators from dynamic imaging data, it is key to map laboratory measurements and the phenomenology observed in imaging experiments to data steams available in real-world applications. The latter are predominantly electrochemical impedance and current-voltage inputs monitored in battery management systems (BMSs). It is thus imperative to establish rigorously validated correlations and transfer functions between information-rich hyperspectral imaging data and impedance signals.24,34,41 Such a capability could have tremendous potential for integrating meaningful performance descriptors into existing BMSs to monitor battery state-of-health and prolong battery service life.

EXPERIMENTAL PROCEDURES

Resource availability

Lead contact
Further information and requests for resources should be directed to and will be fulfilled by the lead contact, Sarbajit Banerjee (banerjee@chem.tamu.edu).

Materials availability

This study did not generate new materials.

Data and code availability

• The hyperspectral datasets, PXRD diffraction data, and X-ray absorption spectroscopy reference database can be found in a Mendeley Data repository: Santos, David (2022), “Multivariate Hyperspectral Data Analytics Across Length Scales to Probe Compositional, Phase, and Strain Heterogeneities in Electrode Materials,” Mendeley Data, V1, https://doi.org/10.17632/79j72k9gbp.1.
• The original code used for generating composition maps and stress maps (via FEA) from hyperspectral imaging is included in the Mendeley Data repository.
• Access to the Analysis of X-ray Images and Spectra (aXis 2000) software is provided, free of charge, by Adam Hitchcock: http://unicorn.chemistry.mcmaster.ca/axis/aXis2000-download.html.
• Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

Synthesis of various V2O5 particle morphologies

α-V2O5 nanowires were synthesized by a hydrothermal growth process adapted from a previously reported method and range from 50 to 400 nm in width and thickness, respectively, and span several microns in length—the <400 nm thickness of the nanowires, parallel to the direction of the beam, ensures sample thickness is within the detector linearity region during STXM measurements.24,44,41 In a typical reaction, 300 mg of commercially sourced V2O5 (Sigma Aldrich, 99.5%) was treated hydrothermally under reducing conditions (5.2 [v/v] mixture of water and 2-propanol; typically, the total reaction volume is 16 mL) to obtain green/blue V2O5-H2O nanowires. V2O5 nanowires were obtained by calcination of the recovered V2O5-H2O nanowires in static air at 350 °C for 48 h.

The micron-sized platelet bulk (3D) geometries, shown in Figure 6E, were obtained from Alfa Aesar, α-V2O5 nanospheres (Figure 6B) were synthesized based on a procedure outlined in previous work.46 For the preparation of the nanoplatelet morphology (Figure 6D), 10 mL H2O2 (35 wt %) was slowly added into 0.36 g bulk V2O5 (Alfa Aesar) with magnetic stirring and reacted for 10 min, followed by addition of 30 mL nano-pure water. The suspension was stirred continuously for 10 min. Next, 30 mL isopropanol was added, stirring the solution for another 10 min. The brown suspension was collected, transferred into a Teflon autoclave, and secured into a stainless-steel reactor. The reactor was placed into an oven and maintained at 180 °C for 24 h. After cooling to room temperature (24 h), a precipitate was obtained through vacuum filtration. The precipitate was washed with nano-pure water and iso-propanol and subquabity dried under ambient conditions. Finally, the powder was annealed at 300 °C for 24 h in the muffle furnace. For all morphologies, phase purity of the resulting orthorhombic V2O5 (Pmmn) was established by PXRD.

Synthesis of stoichiometric LiV2O5 phases

V2O5 nanowires (300 mg) were initially dispersed in 20 mL of dry acetonitrile under an inert Ar atmosphere. A 1.25 M excess of the desired stoichiometric amount of anhydrous LiI (i.e., 0.1, 0.3, 0.45, 0.7, and 0.9 mol of LiI per mol of V2O5) was added to the acetonitrile/V2O5 mixture, which was allowed to stir gently for 7 days. Over the course of the reaction, the supernatant developed a brown/red color indicative of I2 byproduct formation as per:

\[
V_2O_5(s) + xLiI(MeCN) \rightarrow Li_xV_2O_5(s) + \frac{1}{2}I_2(MeCN). \quad \text{Equation 3}
\]

Notably, I2 acts as a mild oxidant and is sufficiently oxidizing so as to act as a chemical deintercalation agent. Consequently, reaction 2 is best characterized as reversible, and phase purity of the powder can be achieved at equilibrium. The stabilization of phase-pure Li1V2O5 nanowires is key to construction and curation of a spectral database for well-characterized materials (Figures S6 and S7). This is in direct contrast to the use of the more highly reducing n-butyllithium (vide infra) as a lithiating agent, which is not reversible. The resulting Li1V2O5 powders were separated from the supernatant by centrifugation and washed three times with dry acetonitrile to remove I2 and excess unreacted LiI. After drying under an inert Ar atmosphere, phase purity was confirmed by PXRD (Figure 1A). X-ray absorption spectra obtained for each phase were collected in transmission mode (STXM) across V L- and O K-edges, as shown in Figures 1 and S4.

PXRD

PXRD patterns were collected on a Bruker D8-focus diffractometer in Bragg-Brentano geometry (Cu Kα, λ = 1.5418 Å; 40 kV; 25 mA current). Rietveld refinement of the X-ray diffraction patterns was performed using GSAS-II.41

Chemical lithiation of V2O5 nanowires via n-butyllithium

Chemical lithiation of all samples not used as X-ray standards was achieved via immersion in a solution (molar excess of ~4:1 Li/V2O5 of 0.025 M n-butyllithium (Sigma, Aldrich) in n-heptane under an inert atmosphere (argon, <0.1 ppm O2 and <0.1 ppm H2O). Lithiation of the V2O5 nanowires proceeded according to:

\[
V_2O_5(s) + xC4H9Li(C4H9)2 \rightarrow Li_xV_2O_5(s) + \frac{x}{2}C_4H_9H_2. \quad \text{Equation 4}
\]

The samples shown in Figures 4, 5A, 5B, and 6 were lithiated for 5, 5, 30, and 5 min, respectively. Relatively short lithiation times are employed to prevent saturation (i.e., complete lithiation) and to further reflect the early stages of lithiation (low depth of discharge). In contrast, longer lithiation times reflect a deeper depth of discharge.

The particles shown in Figure 6 were lithiated as a powder, utilizing a previously detailed procedure.41 Here, the lithiated V2O5 powders were dispersed via ultrasonication in a 2-propanol (5 mg/mL) solution and subsequently drop cast onto a silicon nitride substrate.

The nanoparticles shown in Figures 4 and 5 were drop cast prior to lithiation.44,41 An on-substrate lithiation protocol allows for the effects of interconnectivity on lithiation to be probed. For all samples, the as-prepared grids were immediately sealed under an inert argon atmosphere for transport to the Canadian Light Source and the Advanced Light Source for STXM and ptychography measurements, respectively.

Galvanostatic measurements

Galvanostatic measurements were carried out using coin cells (CR2032) assembled in a glove box filled with argon gas. The working electrode was prepared by casting a mixture of active material (V2O5 powder), conductive carbon black (Super C45, MTI Corp.), and binder (poly(vinylidene fluoride), with a weight ratio of 70:20:10 (w/w/w) in N-methyl-2-pyrrolidinone solution onto an aluminum foil as a current collector. A lithium foil (Sigma-Aldrich)
was utilized as the reference electrode. A Celgard 2500 membrane and a 1 M solution of LiPF$_6$ in ethylene carbonate and dimethyl carbonate (Sigma Aldrich) were used as the separator and electrolyte, respectively. Galvanostatic discharge-charge measurements of the coin cells were performed on a Land (CT2001A) system under a voltage window between 2.0 and 4.0 V at 25°C.

**STXM**

The STXM measurements in Figures 5 and 6 were performed at the 10ID-1 beamline of the Canadian Light Source in Saskatoon, SK, CA. The 10ID-1 beamline is equipped with an elliptically polarized Apple II type undulator, which provides an intense beam in the 130–2,700 eV energy range. State-of-the-art Fresnel zone plate optics are utilized in conjunction with an order-sorting aperture to achieve a focused X-ray beam. The regions of interest shown in Figures 4A, 5A, 5B, and 6 were imaged with a spatial resolution of 50, 30, 50, and 80 nm, respectively. Details of phase identification were based on experimental goals, i.e., the study of single particles, interconnected particles, or particle ensembles. Typically, ROIs are located using single-energy X-ray microscopy measurements. Before spectromicroscopy experiments, an average composition is estimated from a line-scan (ensemble X-ray spectroscopy). For the STXM measurements shown in Figures 4, 5, and 6, the V L- and O K-edges were collected in a single stack ranging from 508 to 560 eV with energy steps of 0.2 eV in the regions of spectral interest (Figure 1) and 1 eV in the continuum regions before and after the specific elemental edges. In a typical STXM measurement, a region of interest is raster-scanned while transmission intensities are recorded in a stepwise fashion by a charged-couple device detector. A 500-line mm$^{-1}$ monochromator was used to tune the incident photon energy to the V L- and O K-edges for imaging the Li$_x$V$_2$O$_5$ samples; here, a 1 ms pixel$^{-1}$ energy$^{-1}$ dwell time was used for signal acquisition.

Image registration was achieved by a cross-correlation analysis using the “Jacobsen stack analyze” routine in the aXis2000 software suite (version updated May 28, 2022). An incident spectrum was collected from a fully transmitting region in the sample (i.e., the silicon nitride substrate) to convert the transmission measurements to an optical density matrix. Preliminary screenings of the data were recorded by integrating the spectra across all pixels to obtain an averaged spectrum that could be compared indirectly or directly to energy-calibrated reference spectra. Spectral shape, peak positions, and relative peak intensities are qualitatively assessed in an indirect comparison, whereas a direct comparison fits a linear combination of the standard spectra to experimental spectra in the Athena suite offered by the Demeter software package (http://bruceravel.github.io/demeter/).

PCA and subsequent k-means clustering were carried out using the “PCA GUI” routine (version 1.1.1) within aXis2000. The number of principal components considered for clustering was chosen based on their eigenvalues and their corresponding eigenspectrum/eigenimage representation of the data. Access to the analytical standards aids in a preliminary interpretation of the eigenspectra, which considerably improves the selection of significant components. The number of clusters sought by a subsequent k-means search was refined by comparison with standards and by linear combination fitting of experimental spectra with standards. The averaged spectrum for each cluster was then assigned a lithium stoichiometry of $x = 0$, $x = 0.1$, $x = 0.3$, $x = 0.45$, or $x = 0.7$ in Li$_x$V$_2$O$_5$ to obtain a set of phase-specific signatures from which composition maps could be generated by SVD (vide infra).

**X-ray ptychography**

The X-ray ptychography measurements shown in Figure 4 were performed at the coherent scattering and microscopy beamline (COSMIC) 7.0.1.2 of the Advanced Light Source in Berkeley, CA. The recursive translation function of the X-ray ptychography plugin within ImageJ (https://imagej.nih.gov/ij/) was utilized to register ptychography images generated in a stepwise fashion along the V L-edge. The particles in Figure 4 were initially imaged by STXM with a spatial resolution of 50 nm, and an energy resolution of 0.2 eV in the regions of spectral interest (Figure 1) and 1 eV in the continuum regions before and after the specific elemental edges—their averaged spectra were compared against standards to fingerprint Li$_x$V$_2$O$_5$ phases. Single-energy X-ray ptychography images were collected with 10 nm spatial resolution at energies corresponding to the d$_{20}$, d$_{100}$, and post-edge features (energy positions were directly informed by the averaged spectrum from STXM). Relative intensities at the d$_{20}$ and d$_{100}$ were assessed to infer lithium gradients after normalizing for thickness effects based on pixel intensities at the post-edge.

**Construction of a test STXM dataset**

A synthetic stack can be generated (as opposed to experimentally collected) from a single image (Figure S1A) and spectrum (Figure S1B) in the aXis2000 software suite. In brief, for a given spectrum with N number of absorption measurements (determined by the incident photon energy range and step size [eV]), the resulting stack comprises N images containing pixel values that result from the product of the pixel intensity in the original image and the energy-specific absorption intensity as shown in Figure S1C.

Intraparticle heterogeneity was simulated by creating a synthetic stack dataset comprising three spectroscopically distinct regions. The images in Figures S8A and S8C were combined with the reference spectra (Figure 1F) in Figures S8D and S8F, which represent phase-pure $\alpha$-Li$_x$V$_2$O$_5$ and $\varepsilon$-Li$_x$V$_2$O$_5$, respectively. Figures S8B and S8E correspond to a phase mixture comprised of 70% $\varepsilon$-Li$_x$V$_2$O$_5$ and 30% $\alpha$-Li$_x$V$_2$O$_5$ (generated from a weighted average of the spectra in Figures S8D and S8F). The importance of accounting for regions with intermediate concentration (within the spinodal region) is 2-fold; first, thin metastable interfaces between Li-rich and Li-poor domains have been experimentally observed, and, second, phase boundaries oriented along the direction of the incident X-rays are possible based on the orientation of the nanowire to the substrate as shown in Figure S20.

**Generation of composition maps from STXM data**

Prior to performing SVD, spectra obtained directly from standards or a standards-informed PCA selection and k-means analyses were converted to a mass absorption scale by matching the pre- and post-edge absorption intensities to the sum of the tabulated atomic mass attenuation coefficients for each material composition and density to enable quantitative analysis. The results of applying SVD to STXM data include a set of phase maps that describe the spectral weighting of each component and thickness in every pixel, as shown in Figure S9. In this work, the generation of high-resolution composition maps has been adapted from a previously reported procedure that directly uses the spectral maps to visualize composition (vide infra). To reduce noise and enable a continuous representation of the data, the spectral maps generated from the SVD process were subjected to a Gaussian filter $G$, according to:

$$G(x, y) = \frac{1}{2\pi \sigma} \exp \left(-\frac{x^2 + y^2}{2\sigma^2} \right),$$

where $\sigma$ denotes the filter width—increasing the value of $\sigma$ leads to a broader “smearing” of the input data over the image plane. This work uses the multidimensional Gaussian filter from the SciPy package for smoothing. The $\sigma$ hyperparameter of the Gaussian filter is chosen based on the spectral resolution of the original image ($\sigma = 1$, mode = “reflect”). The remaining default settings were retained from the SciPy package. A bilinear interpolation enables a continuous representation of the data to better visualize the spatial orientation of distinctly lithiated domains. This work uses Omnic’s calculator (https://www.omnicommlator.com/math/bilinear-interpolation) to perform bilinear interpolation on each spectral map. Subsequently, the superposition of the pixel intensities contained in the spectral maps are weighted by the corresponding stoichiometric fraction $x_i$ for each spectral map (informed by standards), yields a composition map according as:

$$C(x, y) = \sum \phi_i(x, y) \times x_i.$$

where $\phi_i$ denotes the intensity value at a given pixel after Gaussian transformation, and $i$ denotes the number of phases in Li$_x$V$_2$O$_5$. The composition C at each pixel is thus derived from a superposition of the various lithiated phases present in each pixel, as inferred from a standards-informed deconvolution of the phase mixtures. To avoid the introduction of errors during bilinear interpolation, composition maps have been contrasted before and after the interpolation step—no notable inconsistencies were identified.
To calculate stress maps, a finite element mesh is first generated from the composition map using the open-source mesh generator GMSH. The find - contour function within the scikit-image library applies the marching square algorithm to extract the particle’s geometry, which aids in mesh generation. Subsequently, local lithium stoichiometries are mapped onto the nodes of the planar triangle mesh utilizing a linear interpolation according to:

\[ c = C_{\text{min}} + \left( x - C_{\text{min}} \right) \frac{C_{\text{max}} - C_{\text{min}}}{x_{\text{max}} - x_{\text{min}}} \].

(Equation 7)

where \( C_{\text{min}}, C_{\text{max}} \) denote the lower and upper bounds of the lithium concentration at corresponding composition bounds \( x_{\text{min}}, x_{\text{max}} \) according to literature data (Table S14).

For a general case, nonlinear kinematics and inelasticity are expected. Linear elasticity is assumed wherein a constitutive relationship links stresses and strains according to:

\[ \sigma_{ij} = C_{ijkl} \varepsilon_{kl} \].

(Equation 8)

with a fourth-order elasticity tensor, \( C_{ijkl} \), that describes the linear dependence between the symmetric second-order stress tensor \( \sigma_{ij} \) on the elastic strain \( \varepsilon_{kl} \), which can be further defined according to:

\[ \varepsilon_{ij} = \frac{1}{2} \left( u_{ij} + u_{ji} \right) \].

(Equation 9)

where \( u_{ij} \) denotes the components of the displacement vector. To visualize the resulting stress states and their proximity to failure in the present material system, the von Mises stress is calculated as

\[ \sigma_{\text{M}} = \sqrt{\sigma_{ij}^2 + \sigma_{ij}^2 - \sigma_{ij}\sigma_{kl} + 3\tau_{ij}^2} \].

(Equation 11)

where the equivalent von Mises stress \( \sigma_{\text{M}} \), and \( \sigma_{ij}, \sigma_{kl}, \tau_{ij} \), denote the normal and shear stress components in Cartesian coordinates (see supplemental information and Figures S21–S23). For simplicity, isotropic material behavior and a concentration-independent Young’s modulus of \( E = 43 \text{ GPa} \), Poisson’s ratio of \( \nu = 0.3 \) were assumed for the finite element simulations.\(^{14,16}\)

Both fixed and free boundary conditions were considered to account for the constrained and unconstrained volumetric expansion of the particle boundaries, resulting from varying degrees of particle–particle or particle–substrate surface interactions. The fully constrained and unconstrained conditions examined here enable a quantitative estimation of the upper and lower bound of von Mises stress developed within single particles. The overall stress level for the constrained conditions is nearly 2- to 3-fold greater when compared with the unconstrained alternative. Constraining the boundaries affects a stress state shift toward the compressive regime, an overall increase in the magnitude of stresses, and \( \tau_{ij} \) "hotspots" at phase interfaces, as shown in Figures S13 and S16.

SUPPLEMENTAL INFORMATION

Supplemental information can be found online at https://doi.org/10.1016/j.patter.2022.100634.

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AUTHOR CONTRIBUTIONS

D.A.S., conceptualization, formal analysis, investigation, data curation, methodology, writing – original draft, visualization. J.L.A., conceptualization, formal analysis, investigation, data curation, visualization, writing – original draft. B.L., formal analysis, methodology, visualization, writing – original draft. L.R.D.J., conceptualization, investigation. Y.L., investigation, formal analysis. S.P., formal analysis. M.A.G., investigation. L.C., investigation. P.S., methodology. Y.D., supervision. B.-X.X., supervision, funding acquisition. S.B., supervision, funding acquisition, writing – review & editing.

DECLARATION OF INTERESTS

The authors declare no competing interests.

INCLUSION AND DIVERSITY

One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in their field of research or within their geographical location. One or more of the authors of this paper self-identifies as a gender minority in their field of research. One or more of the authors of this paper self-identifies as an underrepresented minority in their field of research or within their geographical location. One or more of the authors of this paper self-identifies as a gender minority in their field of research. One or more of the authors of this paper self-identifies as a member of the LGBTQIA+ community. One or more of the authors of this paper received support from a program designed to increase minority representation in their field of research. While citing references scientifically relevant for this work, we also actively worked to promote gender balance in our reference list.

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