

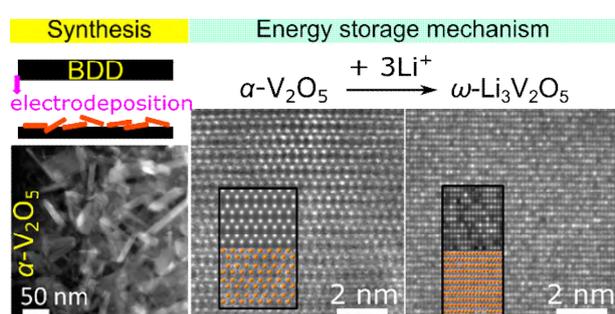
Atomic-Scale Investigation of the Reversible α to ω -Phase Lithium Ion Charge – Discharge Characteristics of Electrodeposited Vanadium Pentoxide Nanobelts

Haytham E. M. Hussein,^{†*} Richard Beanland,^{†*} Ana Sanchez,[‡] David Walker,[‡] Marc Walker,[‡] Yisong Han,[‡] and Julie V. Macpherson^{†*}

[†]Department of Chemistry, University of Warwick, Coventry, CV4 7AL, UK

[‡]Department of Physics, University of Warwick, Coventry, CV4 7AL, UK

*Corresponding authors: j.macpherson@warwick.ac.uk, R.Beanland@warwick.ac.uk, and Haytham.hussein@warwick.ac.uk



ABSTRACT

Using an electrochemical potential pulse methodology in a mixed solvent system, electrochemical deposition of amorphous vanadium pentoxide (V_2O_5) nanobelts is possible. Crystallisation of the material is achieved using in air annealing with the temperature of crystallisation identified using *in-situ* heating transmission electron microscopy (TEM). The resulting α -phase V_2O_5 nanobelts have typical thicknesses of 10-20 nm, widths and lengths in the range 5-37 nm (mean 9 nm) and 15 - 221 nm (mean 134 nm), respectively. One-cycle reversibility studies for lithium intercalation (discharge) and de-intercalation (discharge) reveal a maximum specific capacity associated with three lithium ions incorporated per unit cell, indicative of ω - $Li_3V_2O_5$ formation. Aberration corrected scanning TEM confirm the formation of ω - $Li_3V_2O_5$ across the entirety of a nanobelt during discharge and also the reversible formation of the α - V_2O_5 phase upon full charge. Preliminary second cycle studies reveal reformation of the ω - $Li_3V_2O_5$, accompanied with a morphological change in the nanobelt dimensions. Achieving α - V_2O_5 to ω - $Li_3V_2O_5$ phase reversibility is extremely challenging given the large structural rearrangements required. This phenomenon has only been seen before in a very limited number of studies also employing nanosized V_2O_5 materials and never before with electrodeposited nanocrystals.

INTRODUCTION

Improvements in lithium-ion based rechargeable batteries are essential for new technological developments and low-carbon energy technologies. Faster charge/discharge times, better capacity retention, and recycling are all under investigation, both in incumbent and new materials technologies.¹ Energy density is currently limited by the capacity of the cathode material² and in this regard, vanadium pentoxide (V_2O_5) is an interesting candidate as it can achieve a theoretical capacity of ~ 440 mAh/g provided three lithium ions (Li^+) can be intercalated reversibly per unit cell.³ A recent study by Liu et al. suggested that the capacity could be even higher due to incorporation of more than three Li^+ .⁴ The use of such materials, is thus an attractive replacement for existing, lower capacity cathodes such as $LiCoO_2$ (~ 240 mAh/g) and $LiNi_xCo_yMn_{1-x-y}O_2$ (~ 280 mAh/g).⁵ Progress in this field requires an atomic-scale understanding of structural changes such as conversion, insertion, *etc* of the cathode that occur during Li^+ ion intercalation and de-intercalation. Furthermore, defining the sites that act as either host(s) for ion insertion and extraction, or facilitate the process, can be crucial to optimising performance of the material.

The ability of vanadium (V) to exist in oxidation states from V^{2+} to V^{5+} results in many stable V-O phases and polymorphs.^{6,7} The fully oxidised V^{5+} crystalline form, V_2O_5 , has a layered crystal structure that allows rapid intercalation of Li^+ and other suitably sized ions such as Zn^{2+} , Mg^{2+} , Al^{3+} etc.,⁸⁻¹⁰ which has made it a material of interest in Li^+ and metal-ion battery cathode technology for some time.¹¹⁻¹⁵ The performance of V_2O_5 as a cathode material is strongly dependent on crystal size, with Li^+ diffusion coefficients, and hence charge/discharge times, increasing by over five orders of magnitude as the crystal size drops below 100 nm.^{13, 15-18} V_2O_5 nanostructures have been produced via a variety of synthetic methods including hydrothermal/solvothermal, electrospinning, wet-chemical, sol-gel, polyol, atomic layer deposition, and chemical vapor deposition (CVD).^{3,4,19,20} Whilst little used for V_2O_5 synthesis,^{16, 18, 21} electrodeposition has the inherent advantage that the material is, by virtue of the deposition process, electrically contacted to the electrode surface. This is an advantage over other methods which often require further processing, *e.g.* the use of binders and conductive polymer pastes²² to form an electrical contact and improve the adhesion of the particles to the collector surface.

During Li^+ intercalation, bulk crystalline V_2O_5 undergoes a sequence of transformations with the phases $\alpha \rightarrow \varepsilon \rightarrow \delta \rightarrow \gamma \rightarrow \omega$, each accommodating an increasing number of intercalated ions.^{23,24} In the absence of Li^+ , the starting, α -phase V_2O_5 structure is comprised of chains of

edge-sharing square-based pyramids in which each V is bonded to five O atoms.²⁴ Adjacent chains form sheets held together by van der Waals forces, between which Li⁺ ions are readily intercalated.²⁵ Progressive Li⁺ intercalation induces increasing buckling in the sheets to form the ϵ , δ and γ phases.²⁵ In contrast, the ω -phase is very different; the square-based pyramids convert to VO₆ and LiO₆ octahedra in a disordered rock-salt structure.^{23, 26} In bulk material, this transformation is relatively slow and subsequent delithiation does not result in complete removal of Li⁺ and restoration of the α -phase.²⁷ Instead, a disordered rock-salt or amorphous material is formed.^{4, 26}

An intriguing observation made by several research groups is that the α to ω phase transformation during Li intercalation/deintercalation can be made *reversible*, by employing V₂O₅ nanostructures.^{14, 24, 27, 28} These studies indicate that the dimensions of the nanostructure play a crucial role in achieving reversibility.^{14, 29} For example, a recent *in-situ* transmission electron microscopy (TEM) observation of current-driven lithiation of V₂O₅ nanowires (5-15 nm in thickness and 20-60 nm in width), grown by hydrothermal synthesis, showed a reversible $\alpha \rightarrow \omega \rightarrow \alpha$ transformation,²⁸ over one cycle. This study also reported formation of an amorphous surface layer ~2 nm thick on the crystalline V₂O₅ after one cycle.²⁸ Earlier work using TEM with V₂O₅ nanowires, grown by CVD, also observed single-cycle reversibility (with insertion and removal of Li⁺ chemically driven), although no information was given on the specific nanowire dimensions.¹⁴ In contrast, a complementary *in-situ* TEM study also employing V₂O₅ nanowires, grown by hydrothermal synthesis (50 – 130 nm in width, no reports on thickness) showed progressive transformation to the intermediate γ phase only,³⁰ and no observation of the ω phase. This study also reported the observation of a Li oxide shell which formed during Li⁺ intercalation.³¹

In this work, we first aim to demonstrate that electrically contacted V₂O₅ nanostructures can be produced using electrodeposition procedures, without the use of electrode templates³² or surfactants.²¹ An atom-level understanding of the resulting crystalline material and its properties in relation to the first electrochemically driven Li⁺ discharge/charge process is obtained using *ex-situ* using aberration-corrected scanning TEM (ac-STEM). Complementary electron energy-loss spectroscopy (EELS), X-ray diffraction, and X-ray photoelectron spectroscopy (XPS) measurements are also recorded. Preliminary data on the second cycle of lithiation is also presented.

EXPERIMENTAL DATA

Solutions and Chemicals

Vanadium oxide electrodeposition solutions consisted of 0.1 M vanadium (IV) oxide sulfate hydrate ($\text{VO}_2\text{SO}_4 \cdot \text{H}_2\text{O}$, 99%, STREM chemicals, UK) in 0.5 M sulfuric acid (H_2SO_4 , 99.99%, Sigma-Aldrich, UK) ($\text{pH} = 1.87$), 15 mM H_2O_2 , and 0.3 M ZnSO_4 dissolved in Milli-Q water (18.2 M Ω cm, Millipore Corp., U.S.) and *N,N*-dimethylformamide (DMF, Sigma-Aldrich, UK) at 20 °C. The ratio of water to DMF was 1: 0.33. All chemicals were used as received without further purification. For all experiments, the solutions were left aerated.

Materials and electrode fabrication

Polycrystalline 5×10 mm BDD, 80 μm thick, with a boron dopant level of $\sim 3 \times 10^{20}$ B atoms cm^{-3} was used as the starting material for production of the BDD TEM substrates (cut to a disk diameter = 3 mm, using laser micromachining).^{33, 34} The BDD was grown using microwave CVD by Element Six (Harwell, Oxford, UK) and both surfaces were mechanically polished to ~ 1 nm surface roughness. The fabrication procedure for the production of the BDD-TEM substrates is described in reference ³⁵. For electrochemical characterisation, BDD material of the same dopant density but 200 μm thick and polished to the same surface finish (on the growth surface only) was employed. A Ti/Au ohmic contact was made to the BDD electrode as previously described.^{33, 35} The geometry of the electrode area exposed to solution was defined using adhesive Kapton tape, which contained a laser-micromachined circular hole, 5 mm in diameter. The backface was also insulated using adhesive Kapton tape to prevent solution access. The electrode surface was cleaned by first polishing with alumina powder (0.05 μm particles, micropolish, Buehler, Germany) and then further polished on a water-saturated polishing pad (micropolish, Buehler, Germany). The electrode was then cycled in 0.5 M HCl for 5 minutes to verify a clean surface via the cyclic voltammetric (CV) response *i.e.* CV cycles which are reproducible with a low background current. The electrode surface was further imaged using optical microscopy (Leica DM4000M upright microscope) and the scanning electron microscopy (SEM: Zeiss Gemini) to confirm the surface was clean.

Electrochemical synthesis of vanadium oxide nanobelts using pulse-reverse anodic electrodeposition

V_2O_5 nanobelt electrosynthesis was achieved using electrochemical potential pulse (chronoamperometry) methodology from a solution containing 0.1 M $\text{VO}_2\text{SO}_4 \cdot \text{H}_2\text{O}$.³⁶⁻³⁸ First a

(i) cathodic potential was applied -0.35 V vs. SCE for 50 s, then (ii) $+1$ V vs. SCE for 5 s then (iii) $+2$ V vs. SCE for 50 s, then (iv) $+1.5$ V vs. SCE for 100 s followed by (v) $+2$ V vs. SCE for 50 s. All five steps are referred to as a “potential pulse sequence”. The rationale behind the pulse potentials chosen is described in the results and discussion section. For the majority of experiments one potential pulse sequence was carried out unless otherwise stated in the text. After electrodeposition, the electrodes were gently rinsed in deionized water for 60 s. The electrodes were stored in a vacuum desiccator for 1 hour before being annealed in a tube furnace under an air atmosphere for 2 hrs at 350 °C to crystallize the V_2O_5 (*vide infra*) The loading mass of the V_2O_5 on the BDD electrode was measured to be $\sim 1.84 \times 10^{-3}$ g cm^{-2} , and assumed consistent between different electrode geometries by keeping the charge passed constant ($= 0.097$ C) for each electrodeposition. The mass of deposit was measured using a Metrohm precision microbalance with an accuracy of 0.1 μ g.

Electrochemical measurements

All electrochemical experiments were carried out using a three-electrode set-up controlled by a potentiostat (Metrohm Autolab PGSTAT128N and CompactStat Ivium Technologies, Netherlands) using a dipped electrode arrangement (5 mm diameter electrode or the BDD-TEM electrode set-up).^{35, 39-41} A commercial SCE attached to a luggin capillary filled with the electrolyte solution was used as the reference electrode, a helical Pt wire served as the counter electrode, and BDD, in either the BDD disk electrode or BDD-TEM electrode configuration, was employed as the working electrode. The electrochemically deposited and thermally annealed V_2O_5 on BDD was used for the Li^+ ion intercalation/deintercalation studies. The electrochemical performance of the V_2O_5 -BDD electrodes was assessed in a three-electrode electrochemical cell using CV and galvanostatic discharge-charge curves. Both were recorded in a mixed water-acetonitrile (MeCN) (1:3) electrolyte containing 1 M LiCl and 1M $LiClO_4$, the latter using a potential window of 1.5 – 3.6 V vs. $Li|Li^+$ with an applied current of 0.00572 mA (active mass = 1.3×10^{-4} g) over a period of 10 hours discharge (-0.00572 mA) and 10 hours charge ($+0.00572$ mA). This corresponds to a C/10 current rate with respect to $3Li^+$ per V_2O_5 and rated capacity of 440 mAh/g. The mixed solvent electrolyte was used to widen the electrochemical stability window.⁴² Additionally, since MeCN is characterised by a low viscosity and its ability to solvate the Li^+ , Li^+ diffusion through the V_2O_5 material is promoted,⁴³ resulting in improved capacity and cyclability.⁴⁴

Scanning transmission electron microscopy tracking of Li intercalation behaviour and *in-situ* TEM heating study

To study Li⁺ ion intercalation/deintercalation at the atomic scale, V₂O₅ was electrodeposited directly onto the BDD TEM disk electrodes, which decrease in thickness from the edge of the disk to a hole at the centre ~100 μm in diameter.³⁵ The region up to ~30 μm from the edge of the hole is electron transparent. The BDD TEM electrode was electrically connected using a fine metal clamp attached to the BDD Ti/Au contact and dipped into the appropriate electrolyte solution using a manual *x,y,z* micropositioner (Newport, Oxford, UK) such that the central hole was in the solution but the Ti/Au contact remained dry. The setup is shown in **SI1**, Fig. S1. Prior to STEM imaging the BDD TEM electrode was stored under vacuum at 50 °C for 6 hrs in order to ensure the surface remained clean and free of contamination.

For TEM evaluation of the high temperature annealing process, the BDD TEM electrode was mounted in a Gatan 652 double tilt heating holder and selected area electron diffraction (SAED) patterns were collected from a region of the V₂O₅ as the temperature was slowly increased at a rate of 3.18 °C min⁻¹ to 380 °C under vacuum (**SI2**, Fig. S2).

Surface characterisation and TEM

Field emission (FE) SEM was used to obtain low magnification images of the V₂O₅ NBs after electrochemical deposition. FE-SEM images were recorded using an in-lens secondary electron detector on a Zeiss Gemini operating at 2 kV. Higher resolution images were recorded using a JEOL-JEM 2100 (LaB₆) microscope operated at 200 kV. The impact of Li intercalation/deintercalation on the shape, structure, size, chemical composition, and evolution of the V₂O₅ NBs was investigated using ac-STEM in a double aberration-corrected JEOL JEM-ARM200F TEM/STEM, operating at 200 kV equipped with a Gatan Quantum EELS. Annular dark field (ADF)-STEM and bright field (BF)-STEM images were recorded simultaneously with a probe current of 23 pA and a convergence semi-angle of ~16 mrad. The inner collection semi-angle for ADF-STEM imaging was 38 mrad. Since V₂O₅ is known to be electron beam sensitive⁴⁵ great care was taken to minimise electron dose. Crystals already aligned to the incident beam were imaged. To confirm the chemical nature of the V₂O₅, EELS spectra were recorded, at a probe convergence semi-angle of 32 mrad, a spectrometer collection semi-angle of 25 mrad and a dispersion of 0.1 eV per channel. The energy resolution of the EELS measurements was 1.2 eV, estimated from the full-width-half-maximum of the zero-loss peaks. For statistical

analysis, $n = 150$ V_2O_5 NBs were analysed in terms of length and width, from three different regions of a BDD TEM electrode.

Phase identification of the BDD- V_2O_5 electrodes prior to and after lithiation were investigated using grazing incidence X-ray diffraction (XRD; Malvern Panalytical Empyrean 3 with multicore optics (iCore/dCore), giving Cu K_α radiation). The incident beam was narrowed using a $1/16^\circ$ slit to minimise beam spread at low angle. A parallel plate collimator with a solid state Pixcel 3D detector operating as a point detector (0D) mode was used with a step size of $0.03^\circ 2\theta$. Prior to measurement, the sample surface was adjusted so that it was flat and cut the direct X-ray beam in half, assuring that the sample was in the centre of rotation of the goniometer. An incident angle of 2° was used. The grazing incidence XRD data were analysed using the Panalytical HighScore Plus software (v4.8) and the peaks matched against reference patterns from the latest ICDD database. To aid the pattern matching, the elements were restricted to just V, H, O. The diamond peaks were also accounted for.

X-ray photoelectron spectroscopy (XPS) was conducted using a Kratos Analytical Axis Ultra DLD spectrometer with a monochromated Al $K\alpha$ X-ray source (1486.69 eV) operated at 156 W in a chamber with a base pressure below 1×10^{-10} mbar. Samples were mounted on the sample bar using electrically-conductive carbon tape to ensure conductivity between the sample surface and ground, thus eliminating surface charging. Survey spectra were collected using an analyzer pass energy of 160 eV at 1 eV increments, each of which were integrated for 200 ms. High resolution core level spectra were collected using a pass energy of 20 eV (resolution of approximately 0.4 eV) using multiple sweeps in order to improve the signal-to-noise ratio. Data were analysed using the CasaXPS package, employing mixed Gaussian-Lorentzian (Voigt) lineshapes and Shirley backgrounds. The transmission function and work function of the spectrometer were calibrated using clean polycrystalline Ag foil.

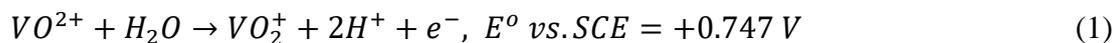
For XRD and XPS, the electrochemical potential pulse sequence was repeated five times to ensure sufficient deposit on the electrode surface such that measurable signals could be obtained.

RESULTS AND DISCUSSION

Electrochemical fabrication and characterization of high aspect ratio V_2O_5 NBs

V_2O_5 nanostructures were synthesised using a five step potential pulse sequence in a mixed water (acidified) - DMF solvent system containing vanadium sulfate ($VOSO_4$).

Thermodynamically, electrodeposition of V_2O_5 from acidic aqueous media is possible via an electrochemical (E), chemical (C)⁴⁶ mechanism, as shown in eqs 1 and 2 for $pH < 2$:



However, experimentally the process is complicated by electrochemical oxidation of water overlapping with $[VO]^{2+}$ oxidation on most electrodes.^{32, 38, 46-48} Furthermore, accumulation of protons locally (eq. 1) during electrochemical oxidation results in a further decrease in local pH at the electrode surface and re-dissolution of the deposited material becomes a possibility.^{32, 38, 46-48} Conversely, at pH values above 3.5, VO_2 will precipitate according to:^{32, 38, 46-48}



Hence there is only a very limited pH window over which V_2O_5 electrodeposition can be successfully achieved, as also shown by the Pourbaix diagram.³²

To circumvent this problem, we use a dual approach, (i) the use of BDD electrodes which, due to the catalytic inertness of the surface, kinetically retard water oxidation³³ and (ii) a mixed solvent media, here H_2O ($pH = 1.87$) and DMF, where DMF is added to the water in a ratio of (water: DMF) 1:0.33. Mixed solvents systems are useful in electrochemical systems as they alter the properties of the individual solvent. Here we use this to effect by adding the polar solvent DMF to help further increase the electrolyte/solvent window.^{49, 50} DMF has a similar dielectric constant and is miscible with water and thus also favours dissolution of inorganic compounds,⁵¹ such as VO^{2+} , but is more viscous. It also plays a role in controlling the morphology of the resulting deposit, as has been shown in the literature for other mixed solvent electrodeposition systems.^{52, 53}

The electrochemical signature of $[VO]^{2+}$ in a water ($pH = 1.87$): DMF (1.00:0.33) solution was first assessed using CV at the 5 mm diameter BDD disk electrode, as shown in Fig. 1, starting at 0.00 V (vs. SCE) and scanning positive to +2.00 V, and then negative to -2.00 V vs. SCE. In Fig. 1 a-d, one oxidation peak, at $E_p = +1.40$ V, due to the oxidation of V^{4+} (*i.e.* VO^{2+}) to V^{5+} (*i.e.* V_2O_5) is observed (Ox1), free from any interference from water oxidation, which occurs at > 2 V during the first scan (Fig. 1a). Upon reversing the scan direction, two reduction peaks are observed ($E_p = +0.3$ V and -1.5 V). At $E_p = +0.30$ V, R1 corresponds to stripping of the electrodeposited V_2O_5 (V_2O_5 to VO_2^+) followed by reduction of VO^{2+} to V^{3+} ,³⁸ at $E_p = -1.50$ V (R2). The generation of a new species, V^{3+} , is evident during the second cycle, where a second

oxidation peak is now observed (Fig. 1b), at $E_p = +0.55$ V (Ox2). This is most likely due to oxidation of V^{3+} to V^{4+} .³⁸ Starting from scan 3, all CV scans (scans 3, 4, 5, and 6) are similar to scan 2.

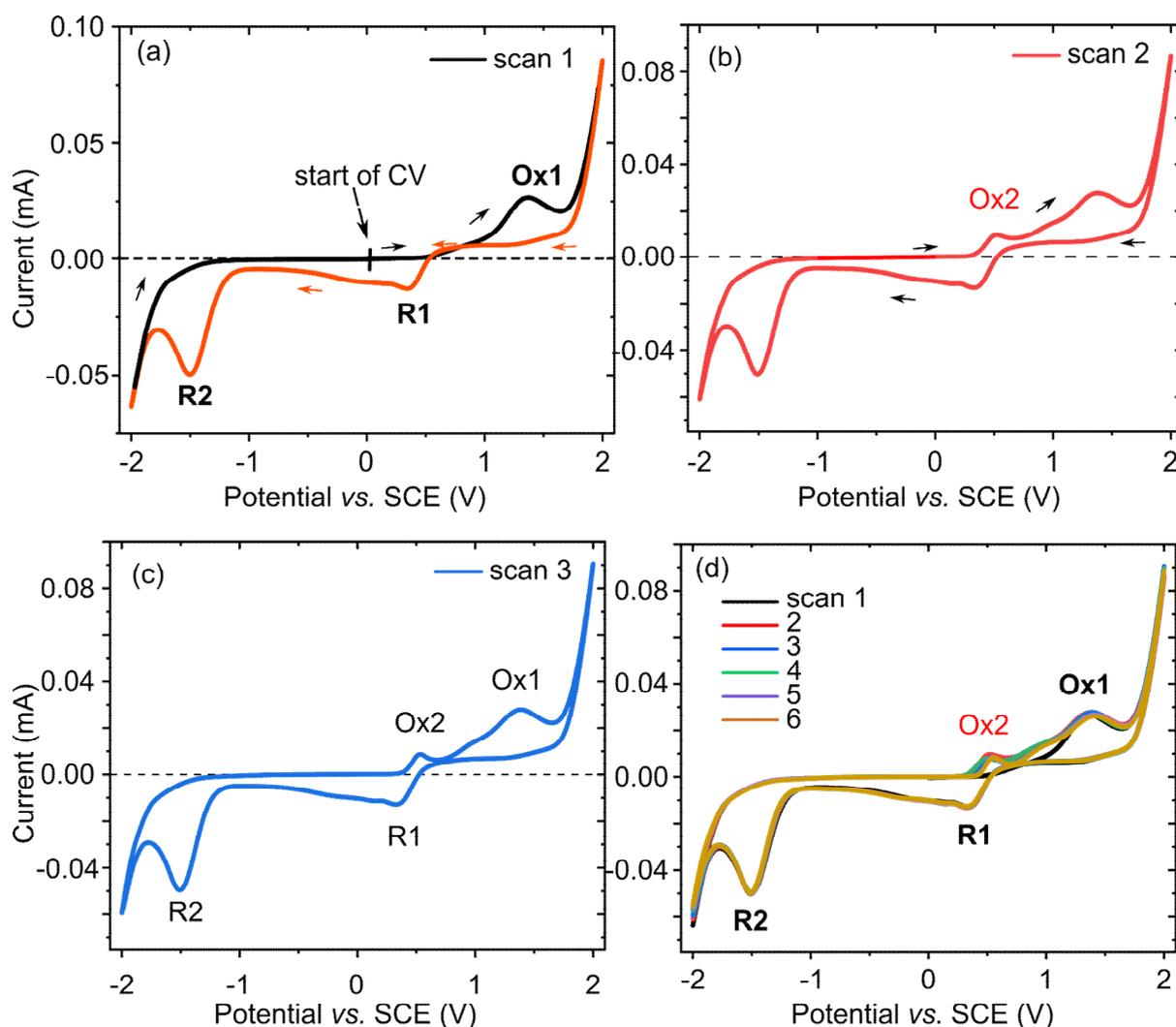


Figure 1: CV of the electrochemical behaviour of $VOSO_4$ in water: DMF (1:0.33) mixed solvent. (a) First scan indicating the starting potential and the direction of the scan sweep, scanning anodically first, (b) is the second scan, and (c) is the third scan. (d) Repetitive cycling where 6 scans were recorded at 0.1 V/s.

Electrodeposition (SI3, Fig. S3) was carried out using a series of five potential pulses. The potential was first held at a value where no current flows (-0.35 V vs. SCE) and then biased sufficiently positive to initiate oxidation of VO^{2+} (+1.00 V vs. SCE for 5 s) and form V_2O_5 nuclei (eqs 1 and 2) on the surface which can act as seeds for further nucleation and growth. By then driving the potential significantly more positive in the second oxidation step, the aim was to stimulate fast growth at these isolated centres (+2.00 V vs. SCE for 50 s). By modulating

between two high positive potentials of +1.50 V vs. SCE for 100 s and +2.00 V for 50 s, continuous growth of the nuclei centres whilst enhancing lateral growth, was ensured.

The as-deposited material exhibited no crystallinity, as evidenced by SAED, see Fig. 2a. Previous literature suggests crystalline material has a better electrical connection to an underlying electrode support.³ To crystallise the V₂O₅, thermal annealing in air was employed. Identification of the crystallisation temperature was carried out using an *in-situ* annealing SAED study employing a TEM heating holder, and heating from room temperature (RT) up to 380 °C, at a rate of 3.18 °C min⁻¹. The full sequence of the twenty-three recorded images is shown in **SI4**, Fig. S4. The first evidence of V₂O₅ crystallisation was the formation of weak intensity peaks in the SAED pattern at 350 °C (Fig. 2b); blue circles mark 111 spots from the BDD substrate. At ~365 °C the clearly defined spot pattern corresponds to the *d*-spacings of *a*-V₂O₅, Fig. 2c. The change in appearance of the deposit during annealing is apparent in Fig. 2d (as-deposited) and Fig. 2e (crystallised). These figures also indicate the approximate position of the selected area aperture used to collect the SAED patterns. The very high thermal conductivity³³ of the BDD-TEM substrate means that the temperature of the BDD electrode will be extremely close to the temperature of the micro furnace of the TEM holder.⁵⁴

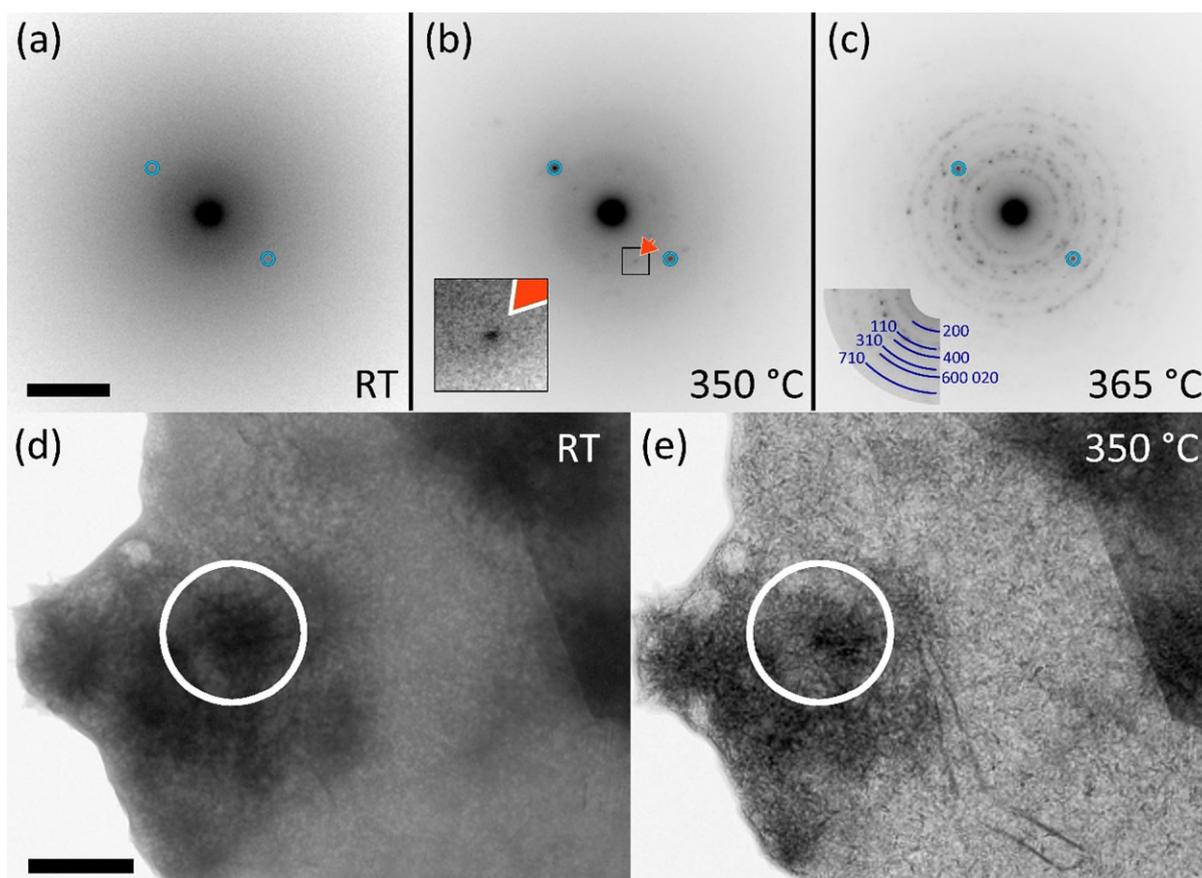


Figure 2: *In-situ* SAED heating study of the transformation of V_2O_5 from amorphous to crystalline during temperature controlled annealing. (a) As-electrodeposited amorphous V_2O_5 , at room temperature, showing only diffuse contrast (scale bar is 10 nm^{-1}). (b) The first appearance of crystalline V_2O_5 spots at $350 \text{ }^\circ\text{C}$, one of which is marked by an arrow and enlarged with contrast enhanced in the inset. (c) Rings of spots, showing full conversion to crystalline material at $365 \text{ }^\circ\text{C}$. Blue circles mark 111 spots from the BDD TEM support. (d) and (e) images corresponding to (a) and (b) with the position of the SA aperture marked. Scale bar is 500 nm .

Using the *in-situ* TEM study to guide controlled crystallisation of electrodeposited V_2O_5 on the BDD electrode, all subsequent studies used a 2 hr anneal in air at $350 \text{ }^\circ\text{C}$ followed by an extended cool-down from $350 \text{ }^\circ\text{C}$ to RT (12 hours) prior to further analysis. The resulting material was analysed using STEM, XPS, and powder XRD. ADF- and BF-STEM images of annealed V_2O_5 on a BDD TEM substrate are displayed in Figs. 3a and b and show a dense mat of NBs (acicular crystals); see also **SI5**, Fig. S5 and Table S1 for a statistical analysis of crystal size dimensions.

Surface sensitive XPS of the annealed material on a BDD-TEM substrate was consistent with formation of α - V_2O_5 (see also **SI6**, Fig. S6 for the survey spectrum). The curve-fitted XPS spectrum in Fig. 3c shows an oxygen 1s peak attributed to the V–O stretch bond at 529.27 eV and vanadium peaks $2p_{1/2}$ at 524.57 eV and $2p_{3/2}$ at 517.15 eV . The main V $2p_{3/2}$ core peak

corresponds to a valence state of V^{5+} (solid red fitted curve), associated with V_2O_5 , as does $Vp_{1/2}$, whilst the tiny shoulder on the right indicates the presence of a very small amount of V^{4+} (solid orange fitted curve).¹⁵ A small V^{4+} peak has been seen before in the XPS spectra for V_2O_5 material subject to thermal treatment in air at similar temperatures (400 °C).^{21, 55} XRD of an annealed thick film also confirms crystallisation (Fig. 3d). The ω -2 θ plot exhibits peaks at 21°, 27°, 32°, and 34° corresponding to the crystal planes (101), (110), (400), (211) of α - V_2O_5 (ICDD 00-041-1426, $Pm\bar{m}n$, $a = 11.519 \text{ \AA}$, $b = 3.563 \text{ \AA}$, $c = 4.375 \text{ \AA}$). The strong peak at 43° is from the BDD substrate.

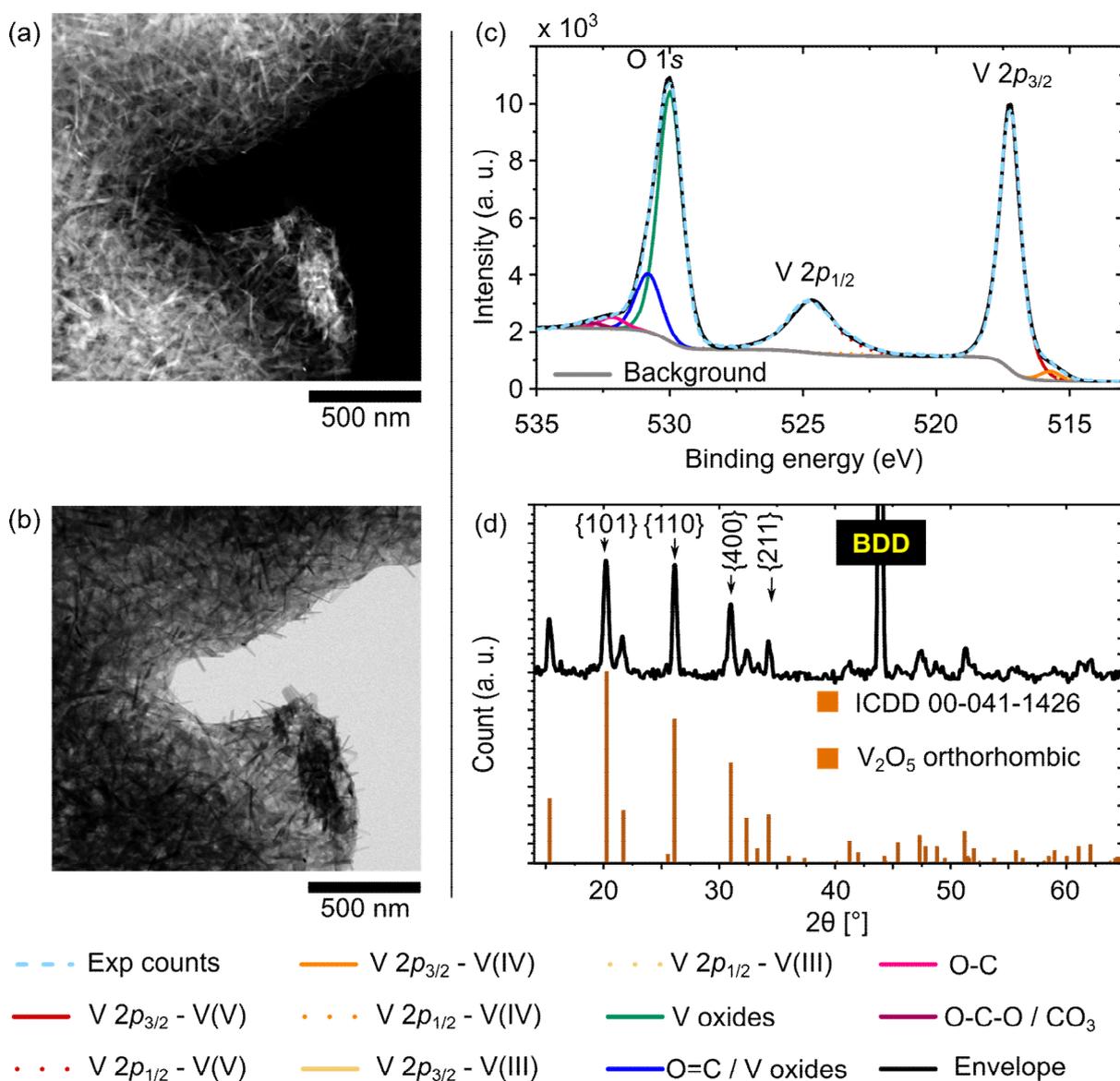


Figure 3: Morphology and analysis of electrodeposited and thermally annealed V_2O_5 . (a and b) ADF-STEM and BF-STEM images of high coverage V_2O_5 on a BDD TEM support, forming a dense mat of acicular crystals. (c) XPS spectrum and curve fitting of the O 1s and V 2p_{3/2} peaks. (d) ω -2 θ XRD pattern identifying the deposit as the orthorhombic V_2O_5 structure.

To explore the electrodeposited and annealed α -V₂O₅ in more detail, Fig. 4 shows high resolution aberration-corrected ADF-STEM images. In Fig. 4a the characteristic parallelepiped acicular shapes (also referred to as ‘calissons’¹⁷) of the V₂O₅ NBs are again evident. By taking side-on views of the NBs (e.g. Fig. 4b showing a [100] view and the (001) facets), it was possible to determine thickness values. These were found to be typically in the range 10 - 20 nm. The long axis of the NBs lies parallel to [010] and the NBs had a mean length of 134 nm (range 15 – 221 nm) and mean width 9 nm (range 5 – 37 nm), **SI5**. Occasionally other unidentified polymorphs were observed (see **SI7**, Fig. S7).

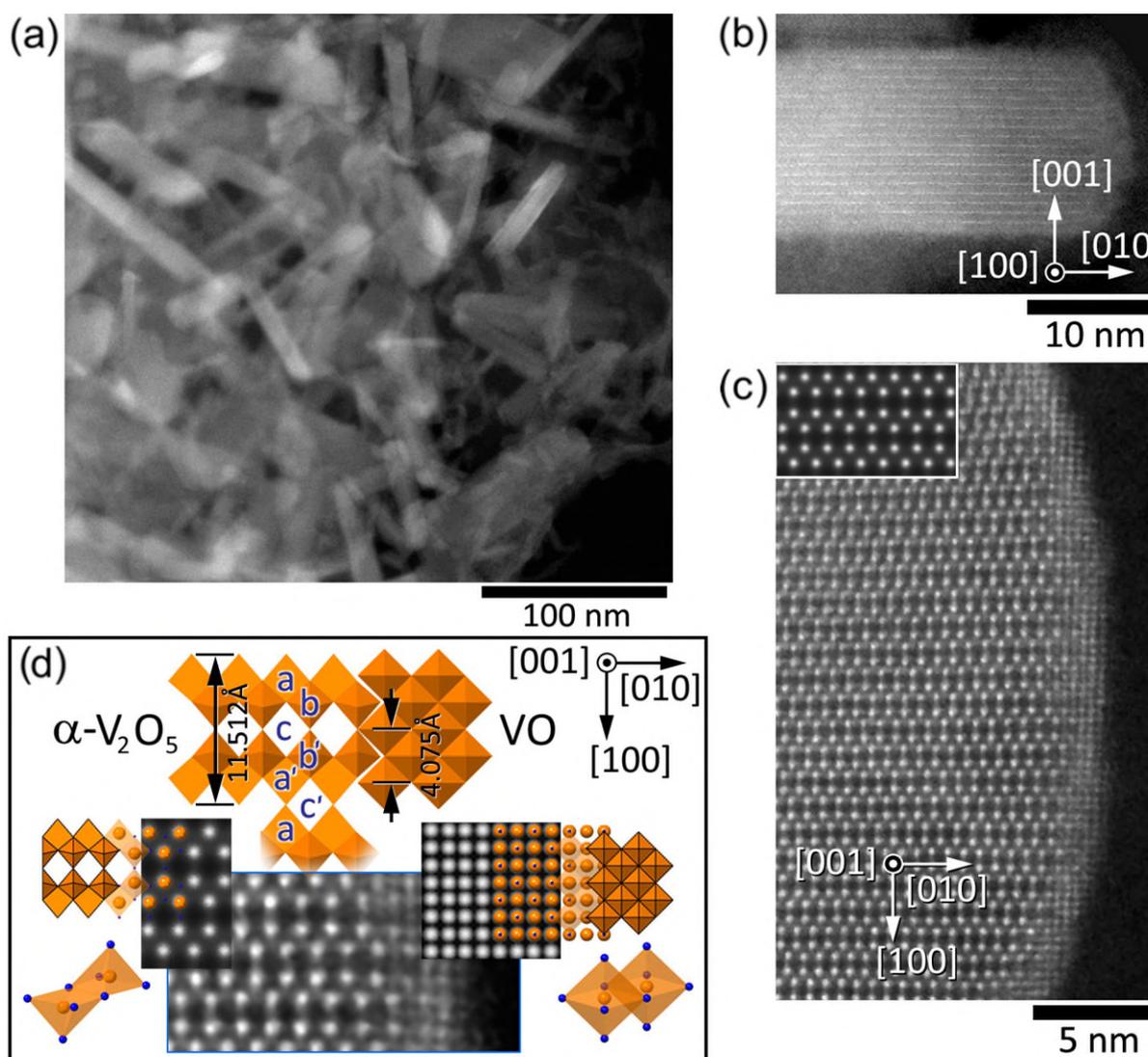


Figure 4: ADF-STEM of electrodeposited and thermally annealed V₂O₅. (a) Electrodeposited and thermally annealed α -V₂O₅ on BDD with high coverage, forming a mat of acicular crystals with lengths 15 – 221 nm. (b) A single α -V₂O₅ NB seen edge-on, with (001) facets and long axis parallel to [010]. (c) Another NB viewed face-on, close to the [001] axis, showing distinctive double-layers of vanadium (inset: multislice simulation of [001] α -V₂O₅). An altered layer 1-2 nm in thickness is visible at the (010) facet. (d) The

altered layer at the (010) facet has an appearance consistent with rock-salt VO, with octahedral coordination of vanadium (orange) to oxygen (blue).

More detailed information on atom arrangement in a NB is shown in Fig. 4c. Focusing on the region away from the edge first, the ADF-STEM image, where only the high atomic number V atoms produce contrast, confirms the material is α -V₂O₅. A schematic of the structure is shown at the top and left of Fig. 4d, where V-O polyhedra are shown in orange, individual V atoms are represented by orange circles and O atoms are represented by blue circles. α -V₂O₅ consists of (001) sheets, comprised of [010] chains of edge-sharing square-based pyramids in which each V lies close to the base and is bonded to five O atoms. For each pyramid there is an sixth oxygen atom present but this does result in an octahedron around the V atom as it is slightly too distant to form the appropriate bond.^{24, 56}

In the (001) schematic of α -V₂O₅ of Fig. 4d, half of the pyramids point down towards [00 $\bar{1}$], (marked 'a'; the flat base of the pyramid is visible), and half point upwards towards [001], (marked 'b'; apex and facets of the pyramid are visible). The horizontal [010] chains of pyramids have apices that alternate in an up-up-down-down b-b-a-a sequence. As the a-b chains connect at pyramid corners, there is also a row of empty sites, marked 'c' in Fig. 4d. This a-b-c, filled-filled-vacant stacking sequence is evident in the ADF-STEM images in Figs. 4c and 4d. The two rows of pyramids with central V⁵⁺ atoms can be seen with bright contrast, while the third, empty, row is much darker in comparison. A multislice simulation of the ideal α -V₂O₅ structure is also shown on the left of Fig. 4d connecting the atomic model to the experimental image and as an overlay in Fig. 4c. Further information on the crystal structure of α -V₂O₅ and BF-STEM images are given in **SI8**, Figs. S8 and S9.

Fig. 4c, and at higher resolution in Fig. 4d, also shows the presence of a change in structure at the edge of a (010) facet, noting it was not possible to resolve this feature on all facets of a NB or indeed on all NBs. The atom arrangement appears consistent with a rock-salt type structure, VO, where V is now octahedrally coordinated to six O atoms, forming a structure of close-packed edge-sharing octahedra as shown in the top and right schematics in Fig. 4d. Rock-salt VO can be produced in bulk form,⁵⁷ with a range of stoichiometry VO_x (0.8 < x < 1.3) and high concentrations of both V and O vacancies.⁵⁸ As is apparent from Fig. 4d, the [010] rows of atoms are continuous throughout the NB, but in the VO region there is no row of empty sites and the [001] view shows a square array of bright atom columns. This image is in excellent agreement with the multislice simulation that bridges the gap between the structural model and experimental image on the right of Fig. 4d. The VO structure often appears noticeably brighter

than the bulk of the crystal (see also SI9, Fig. S10), which is also consistent with the higher density of oxygen-depleted VO, in comparison with α -V₂O₅. From Fig. 4d information on the lattice mismatch between VO and α -V₂O₅ along [100] can also be obtained; the plane spacing of V₂O₅ is 11.512 Å while that of VO is 4.075 Å, giving a 3:1 matching with a misfit of 6%. However, despite the difference in crystal structure the VO region appears coherent with the α -V₂O₅ crystal, with an orientation relationship [100]_{VO}//[100]_{V₂O₅}, [001]_{VO}//[001]_{V₂O₅}. Other examples of VO formation on a NB edge are given in SI9, Fig. S10.

It is interesting to compare the macroscale (XRD and XPS) with the atom scale (ac-STEM and EELS) data in Figs 3 and 4. XRD which is a bulk measurement indicates the formation of α -V₂O₅ crystals. XPS which is sensitive to only the first few surface layers³² confirms this, via observation of a dominant V⁵⁺ oxidation peak, but also indicates the presence of a small amount of V⁴⁺ but no V²⁺ (as expected for VO). For crystals subject to annealing, under both UHV (thermal reduction) and in air, loss of oxygen and the formation of V⁴⁺ oxidation states has been previously documented, using XPS, EELS and DFT,^{21, 56, 59} with DFT predicting loss of vanadyl oxygens and formation of a V-O-V bond between layers upon thermal reduction.^{56, 60} However, given that XPS does not show a V²⁺ signature, VO presence must be at levels below the detection sensitivity of XPS in the area of the XPS measurement. This fits in with our TEM observations that the VO layer is not present on all NBs imaged or indeed all facets.

It is also important at this stage in the discussion to consider the possible effect of the electron beam on the ac-STEM data. In previous high resolution TEM studies, α -V₂O₅ subject to high doses of electron irradiation in UHV for long periods of time (minutes to tens of minutes), resulted in a transition to rock-salt VO^{45, 61, 62}. By lowering the flux of electrons it was possible to observe the intermediate phases V₄O₆ and V₆O₁₃ before reaching VO.⁴⁵ However, in our study, the radiation dose in our work is approximately three orders of magnitudes smaller than previous work, and imaging was for much shorter periods of time (10 μs/pixel), see SI9. When the VO structure was present it was observed immediately and noticeable structural changes in the NBs were only seen after having made several passes over the NB with the electron beam. This suggests that the thermal anneal was responsible for the observed atomic rearrangements.

Characterisation of the α -V₂O₅ NBs during one cycle Li intercalation and deintercalation

The first lithiation (discharge) and de-lithiation (charge) cycle for electrodeposited and annealed α -V₂O₅ NBs on a BDD electrode was investigated galvanostatically by cycling between +3.6 V and +1.5 V vs. Li|Li⁺ at a C/10 current rate (see experimental section). The

discharge-charge curve of the α -V₂O₅ NBs is presented as a potential (y axis) versus composition (top axis), represented as a number of intercalated Li (*i.e.* Δx_{Li} in Li_xV₂O₅)/specific capacity in mAhg⁻¹ (bottom axis) and is shown in Fig. 5 for the first discharge (lithiation - left hand side) – charge (delithiation - right hand side) cycle.

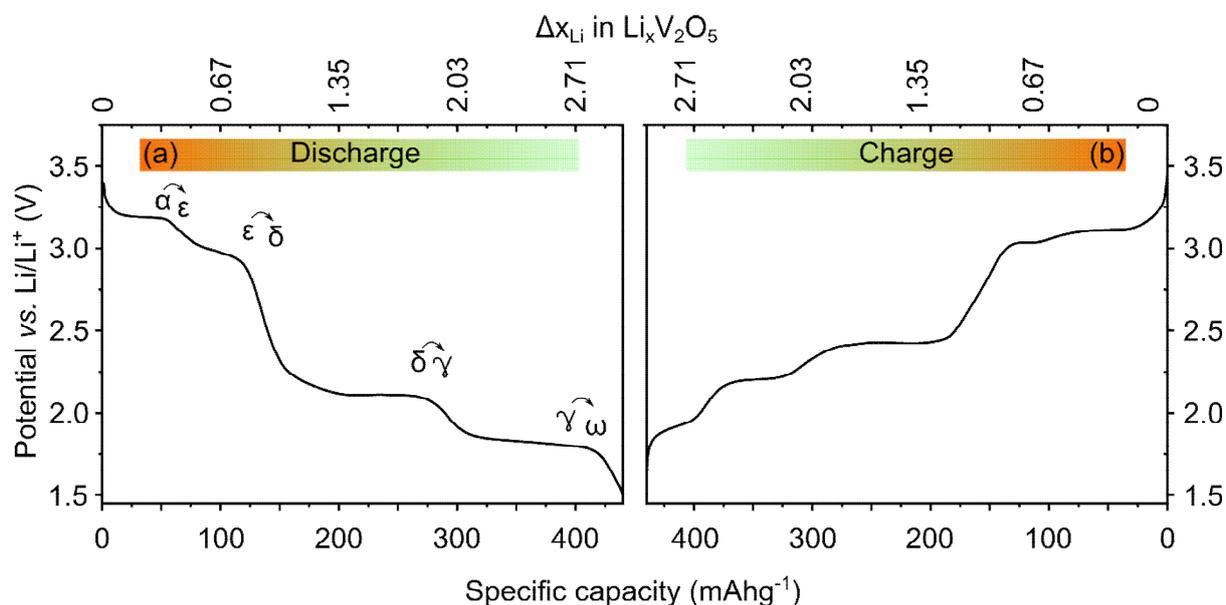


Figure 5: Galvanostatic potential profile. First discharge (left hand side) – charge (right hand side) of α -V₂O₅ NB-BDD electrode in 1 M LiCl + 1 M LiClO₄ in 1:3 mixture of water and MeCN at C/10 rate.

During discharge, the electrode is cycled to less positive potential values to promote intercalation/insertion of Li⁺ ions into the α -V₂O₅ material (lithiation). Fig. 5a shows a series of distinct potential plateaus, which demonstrate the formation of different Li_xV₂O₅ phases. The galvanostatic profile shows various sloped regions associated with the following sequence of transitions; $\alpha \rightarrow \epsilon \rightarrow \delta \rightarrow \gamma \rightarrow \omega$, Li_xV₂O₅. The first three transitions describe the Li_xV₂O₅ phases with increasingly puckered layers of VO₅ square-based pyramids, as the number of Li⁺ intercalated per unit cell increases.²⁴⁻²⁶ The fourth represents the formation of the rock salt ω -Li₃V₂O₅ phase. In a similar manner to the structural transition α -V₂O₅ \rightarrow VO, we expect this phase transition involves conversion of VO₅ square-based pyramids to VO₆ octahedra. This is accompanied by the expulsion of a fraction of V atoms into unoccupied rows, most likely driven by exchange with Li⁺, where they also become octahedrally coordinated. Complementary CV data showing the voltammetric response of the NBs during the first discharge/charge cycle, in a solution of 1 M LiCl + 1 M LiClO₄ in 1:3 mixture of water and MeCN, is shown in **SI10**, Fig. S11.

A maximum discharge specific capacity of 440 mAhg^{-1} was achieved during the first galvanostatic cycle (on discharge / lithiation) which is as expected if all the material was converted to $\omega\text{-Li}_3\text{V}_2\text{O}_5$. A number of Li^+ per V_2O_5 (ΔxLi) in $\text{Li}_x\text{V}_2\text{O}_5$ of ~ 3.0 was also achieved. The lithiated V_2O_5 material, obtained after discharging, was further investigated by STEM and macroscale XPS and XRD characterisation. The XPS data of the V region is presented in Fig. 6 and the survey spectrum as well as the XPS data of the Li and O regions are shown in **SI11**, Figs. S12 and S13. Curve fitting of the core XPS spectrum in Fig. 6 between 510 eV and 535 eV, and comparison with Fig. 3a, reveals the changes in the valence state of vanadium upon lithiation. In particular, the dominant V $2p_{3/2}$ peak at ~ 517 eV can now be deconvoluted into three peaks; the solid red, orange and yellow fitted curves indicate V^{5+} (517.5 eV), V^{4+} (516 eV) and V^{3+} (~ 515.5 eV). The data shows a substantial increase in the $\text{V}^{4+/3+}$ valence states compared to the pristine material (Fig. 3a). In addition the XPS signal for lithium (Fig. S13 in **SI11**; between 60 eV and 52 eV) confirms lithium incorporation into the material. The presence of $\text{V}^{4+/3+}$ highlights the formation of $\omega\text{-Li}_3\text{V}_2\text{O}_5$.^{4, 26} The XRD $\omega\text{-}2\theta$ plot of the material after the first discharge (lithiation) is also shown in **SI11**, Fig. S14, and is similar to what has previously been reported for $\text{Li}_3\text{V}_2\text{O}_5$ by Delmas *et al.*²³ for one cycle.

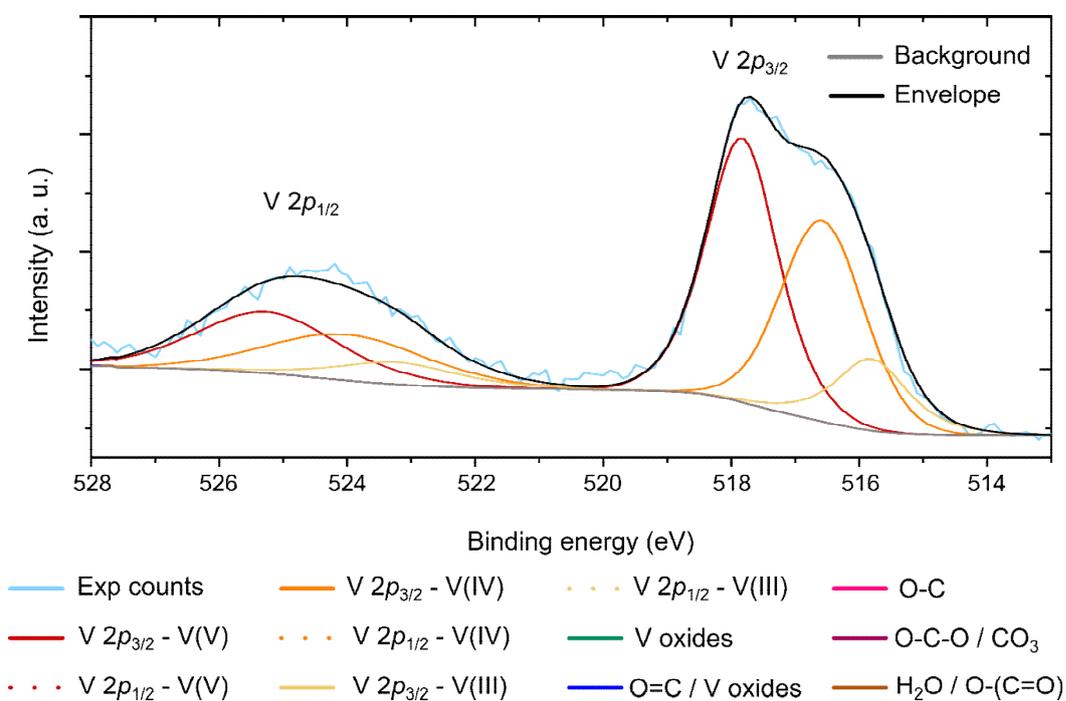


Figure 6: XPS study of the structure of the V_2O_5 NBs after lithiation (discharge). XPS spectrum and curve fitting of the V $2p$ of the V_2O_5 NBs after lithiation showing the V $2p_{3/2}$ and the V $2p_{1/2}$ regions.

To reveal the atomic structure of the resulting lithiated NBs after the first full discharge, STEM images were again recorded. Fig. 7a shows the conversion of the NB to the rock salt ω -Li₃V₂O₅ phase. In structure, this appears very similar to the VO region at the (010) edge of the α -V₂O₅ NBs (Fig. 4b-d). In this lithiated material, with a nominal Li₃:V₂ ratio, we may expect V atoms to be replaced by Li in three out of every five octahedra.⁴ There are relatively few octahedra in a crystal of only 10-20 nm thickness. Thus, statistical variations in the V occupancy of individual atom columns should also be readily visible since high atomic number atom columns give a larger contrast than low atomic number ones. This is shown in the bottom half of Fig. 7a and is consistent with a mixture of randomly arranged LiO₆ and VO₆ octahedra in the ω phase. This interpretation is also supported by multislice simulations performed using a random structure model overlaid on the images (outlined in green in Fig. 7a). The ω -Li₃V₂O₅ structure has the same orientation relationship with respect to the crystal shape as the VO region, but now extends over all of the NB. The lattice parameter of ω -Li₃V₂O₅ is 4.095 Å,⁴ only slightly larger than that of VO, giving a misfit of 7% to α -V₂O₅. EELS spectrum of the lithiated materials recorded from the centre of the lithiated NB also shows a change in vanadium oxidation state,³⁰ Fig. 7b (green line). The V 2p_{1/2} peak has moved by ~1.0 eV indicating a V³⁺/V⁴⁺ valence state, as expected for formation of ω -Li₃V₂O₅^{4,26} and consistent with the XPS data (Fig. 6 and SI11, Fig. S13).

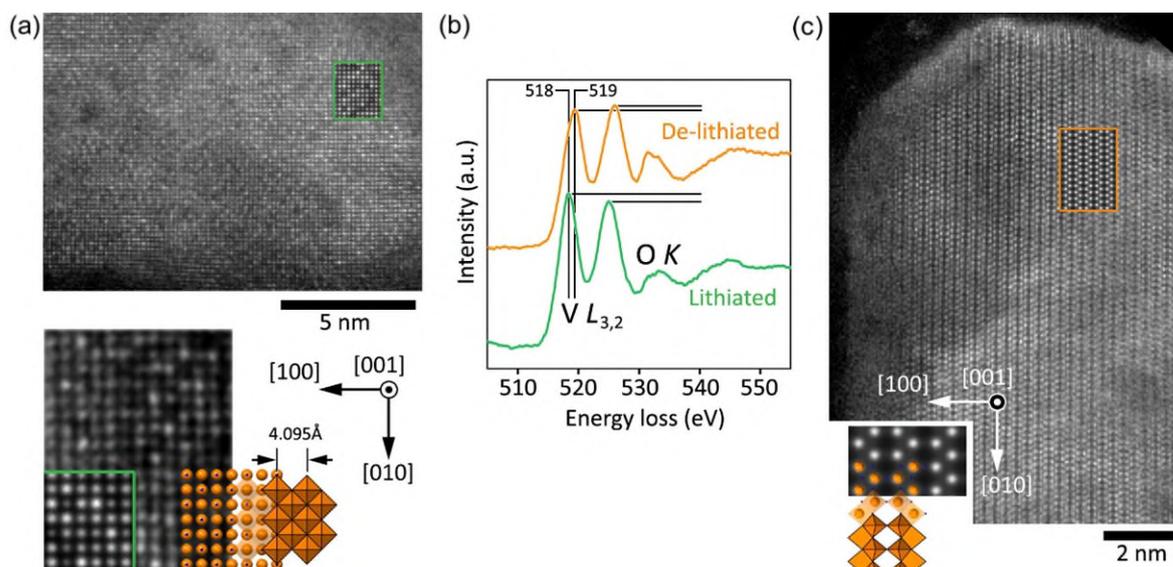


Figure 7: ADF-STEM of lithiated-delithiated vanadium oxide NBs. (a) after first lithiation, showing the rock-salt ω -Li₃V₂O₅ phase. The variable occupancy of the Li/V octahedral site in the rock-salt ω -Li₃V₂O₅ phase is apparent from the highly variable intensity of the atom columns in the enlarged ADF-STEM image. Overlays show a multislice simulation of a

random disordered rocksalt structure model. (b) Vanadium and oxygen EELS spectra for lithiated and de-lithiated material showing changes in vanadium valence state (see text). (c) After one delithiation, showing reversion to the α -V₂O₅ phase and retainment of the VO surface layer. Multislice simulation shown as an inset.

The galvanostatic charge profile in Fig. 5b shows several plateaus during the delithiation process at slightly different potential values in comparison with the first discharge process suggesting differences in the dynamics of the mechanism of Li⁺ intercalation/ deintercalation during a single charge-discharge cycle. A maximum charge specific capacity of 439 mAhg⁻¹ was achieved, which indicates recovery of the starting material, α -V₂O₅.¹ To further investigate the status of the material after one discharge/charge cycle, ADF-STEM was employed to image individual NBs of the fully charged material (after the first delithiation), Fig. 7c, whilst XPS and XRD measurements were made on the bulk material (SI12, Figs. S15 and S16). The ADF-STEM image of a typical NB after the first delithiation process reveals restoration of the original α -V₂O₅ structure but also shows retainment of the thin rock-salt surface layer (VO) (visible at the (010) facets in Fig. 7c, c.f. Fig. 4c and 4d). Additionally, EELS spectrum of the delithiated materials recorded from the centre of the V₂O₅ NB also shows a change in vanadium oxidation state, Fig. 7b (orange line). The V 2p_{1/2} peak appears at 519 eV indicating a V⁵⁺ valence state, as expected for formation of α -V₂O₅. Both the macroscale XPS data (SI12, Fig. S15) and XRD (SI12, Fig. S16) data are also extremely similar to the pristine α -V₂O₅ XPS and XRD data recorded in Figs. 3c and 3d, indicating no remaining lithiated phases and removal of Li⁺.

Finally, whilst the focus of the work was primarily on the first cycle discharge/charge response of electrodeposited α -V₂O₅ we also recorded preliminary STEM experiments and electrochemical data, exploring the second discharge process, shown in Fig. 8 and SI13, Figs. S17 - S19.

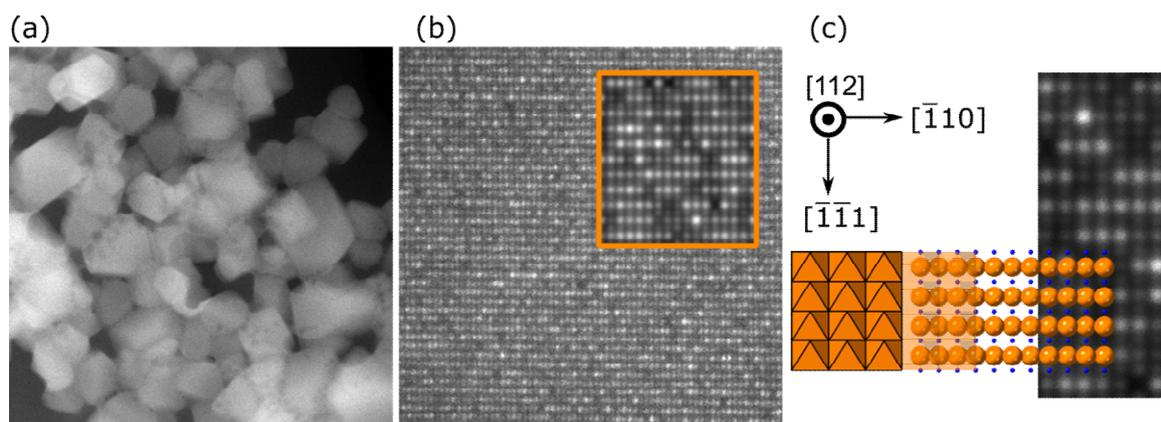


Figure 8: ADF-STEM of vanadium oxide NBs after a second lithiation. (a) Crystal morphologies become noticeably more equiaxed (see also **SI13** Figs. S18). (b) The material has the rock-salt ω - $\text{Li}_3\text{V}_2\text{O}_5$ phase and as before the variable occupancy of the Li/V octahedral site is apparent from the variable intensity of the atom columns, Multislice simulation shown as an inset. (c) Orientation of LiO_6/VO_6 octahedra in this $[211]$ view and correspondence with the simulated disordered ω - $\text{Li}_3\text{V}_2\text{O}_5$ structure.

The second discharge curve shows a similar behaviour to the first discharge profile (**SI13**, Fig. S17), with a discharge capacity of 439 mAh/g was achieved. However, the morphology of the NBs has changed (Fig. 8a-b and **SI13** Figs. S18-S19), with the crystals becoming more equiaxed and having lower aspect ratios. A high magnification of the crystals after lithiation, Fig. 8b, indicates that the cubic ω - $\text{Li}_3\text{V}_2\text{O}_5$ phase is once again obtained and represents the majority of NBs investigated with STEM. The change in morphology is quantified in **SI13**, Figs. S20 and S21. The variable occupancy of the Li/V octahedral sites is again apparent from the variable intensity of the atom columns in Fig. 8b. The multislice simulation is shown as an inset. Fig. 8c shows the orientation of the LiO_6/VO_6 octahedra in the $[211]$ view of the lithiated material and correspondence with the simulated disordered ω - $\text{Li}_3\text{V}_2\text{O}_5$. Whilst this observation is interesting, it is clear further investigation is required to understand the impact this morphology change has on the ability of the material to undergo repetitive reversible cycling between α - V_2O_5 and ω - $\text{Li}_3\text{V}_2\text{O}_5$. Furthermore, some unidentified ordered structures were occasionally seen (see **SI13**, Fig. S19). The presence of such structures could also be the reason why the capacity started to decrease.

CONCLUSIONS

A mixed solvent method, in combination with an electrochemical potential pulse technique, was developed to deposit amorphous V_2O_5 NBs, with thermal treatment used to transform the amorphous V_2O_5 into crystalline α - V_2O_5 . The optimal temperature for this process was determined by *in-situ* heating TEM. The resulting α - V_2O_5 NBs had thicknesses of 10-20 nm,

lengths 15 – 221 nm (mean 134 nm) and widths 5-37 nm (mean 9 nm). For successful electrodeposition, it was necessary to suppress the oxidation of water, achieved through the use of BDD electrodes and a mixed aqueous – aprotic (DMF) solvent, and employ a potential pulse sequence. STEM of the pristine material revealed that the NBs were (in the majority) a single crystal of α -V₂O₅ with a [010] long axis and (100) and (001) faces. A thin surface layer, 1-2 nm in thickness, was found on some NBs in edge regions, with an appearance consistent with a few monolayers of rock-salt VO coherently strained to match the underlying α -V₂O₅ crystal. It was however undetectable in XPS suggesting it was not a significant presence. Further work to understand its significance and origin is underway.

Single cycle charge–discharge data indicated a specific capacity consistent with the reversible incorporation of three Li⁺ per unit cell *i.e.* formation of rock-salt ω -Li₃V₂O₅. This was confirmed by ac-STEM images of fully lithiated material, which showed not only the rock-salt structure but also variable intensities of atom columns in annular dark field images, indicating a disordered and locally variable composition. Upon full delithiation, α -V₂O₅ was recovered, also evidenced using XPS and XRD, highlighting the one cycle $\alpha \leftrightarrow \omega$ phase reversibility of this nanostructured electrodeposited material. VO surface layers were also observed in the delithiated material. Preliminary second cycle ac-STEM lithiation studies again revealed formation of the ω phase but interestingly this was accompanied by a morphology change, with the NBs becoming more equiaxed. Further studies are required to establish whether this electrodeposited nanomaterials is a viable battery cathode, when run over many cycles.

This investigation both demonstrated the importance of STEM as an analysis technique to better understand material transformation in hybrid aqueous/nonaqueous batteries and the potential for using electrodeposition as a synthesis methodology for battery related materials.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge. Additional information and figures as (PDF) contains; Experimental set-up for vanadium oxide electrodeposition and Li intercalation using the BDD-TEM electrode, Illustration of the *in-situ* TEM heating set-up, Chronoamperometric response of vanadium sulphate solution using the BDD electrode in a water-DMF mixed solvent, In-situ TEM thermal annealing of V₂O₅ from amorphous to crystalline, Statistical analysis of the V₂O₅ NBs dimensions and size, XPS survey spectrum of

the crystalline V_2O_5 NBs electrodeposited on BDD, Unknown vanadium oxide polymorphs, The crystalline structure of α - V_2O_5 , Electron dose and additional observations of VO surface layers, CV data of the first discharge/charge cycle in a solution of 1 M LiCl, XPS and XRD of the V_2O_5 structure post Li intercalation (Li insertion), XPS and XRD of the V_2O_5 structure post Li deintercalation (Li removal), Electrochemical response during the second discharge and charge cycle and the formation of ordered $Li_3V_2O_5$ structure. Low magnification STEM images of V_2O_5 structures after lithiation and delithiation cycles, and References

AUTHOR INFORMATION

Corresponding Authors

* Prof. Julie V. Macpherson: Department of Chemistry, University of Warwick, Coventry, CV4 7AL, UK. Tel: +44 (0) 2476 573886. Fax: +44 (0) 2476 524112. E-mail: j.macpherson@warwick.ac.uk, and

* Prof. Richard Beanland: Department of Physics, University of Warwick, Coventry, CV4 7AL, UK. Tel: +44 (0) 24 765 73884. E-mail: R.Beanland@warwick.ac.uk, and

* Dr Haytham Hussein, : Department of Chemistry, University of Warwick, Coventry, CV4 7AL, UK. E-mail: Haytham.hussein@warwick.ac.uk.

ORCID

Julie V. Macpherson: 0000-0002-4249-8383

Richard Beanland: 0000-0003-1749-4134

Ana M. Sanchez: 0000-0002-8230-6059

Haytham E. M. Hussein: 0000-0002-5238-4590

David Walker: 0000-0002-8583-4279

Marc Walker: 0000-0002-5522-0516

Yisong Han: 0000-0002-7638-6785

Author Contributions

H.E.M.H devised the procedure and concept for electrochemical fabrication of the V_2O_5 NBs and the use of BDD TEM platforms for ac-STEM monitoring of the Li intercalation and deintercalation process in hybrid battery platform. H.E.M.H. carried out the electrochemical

experiments and prepared the BDD substrates. H.E.M.H. and R.B. carried out the *in-situ* TEM heating. H.E.M.H. and A. S. carried out the STEM imaging. EELS measurements were carried out by Y.H., and A. S. D.W. carried out the XRD experiments and M.W. carried out the XPS experiments. H.E.M.H, R.B., D.W., M.W. and J.V.M. were involved in analysis and interpretation of the data. H.E.M.H, R.B. and J.V.M. wrote the manuscript with input from the other authors.

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

H.E.M.H. thanks the University of Warwick and the Research Development Fund for funding. We thank Element Six Ltd. for the provision of high grade BDD material, Profs. Louis Piper (WMG) and Richard Walton (Chemistry), University of Warwick, for useful discussions and Josh Tully (Department of Chemistry, University of Warwick) for laser cutting and ohmically contacting the BDD TEM disks.

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