

In situ observation of compressive deformation of an interconnected network of zinc oxide tetrapods

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ABSTRACT

Zinc oxide tetrapods have remarkable functional and mechanical properties with potential applications in different fields including nanoelectronic and optoelectronic sensing, functional composites and coatings, as well as energy harvesting and storage. Based on the 3D shape of these microparticles, they can be assembled into highly porous (up to 98%) macroscopic ceramic framework structures that can be utilized as a versatile template for the fabrication of other multi-scaled foam-like materials. Here we investigated the three-dimensional structure of low density interconnected zinc oxide tetrapod networks by high resolution X-ray computed tomography. *In situ* observations during mechanical loading show inhomogeneous development of anelastic strain (damage) during compression, and homogeneous elastic recovery on unloading. Individual tetrapods are observed to deform by arm rotation to accommodate strain.

Zinc oxide (ZnO) is a semi-conducting metal oxide with remarkable functional properties including piezo-electricity and opto-electricity due to its large band gap and hexagonal-wurtzite crystal structure as well as good mechanical and thermal properties [1]. It can be grown in a variety of micro- and nanostructures of which one prominent shape is a three-dimensional (3D) tetrapod geometry of four arms connected at a central core [1,2]. It has already been demonstrated that nano and micro-scale tetrapods of ZnO (t-ZnO) have widespread potential applications in many fields, including biomedical engineering in the form of functional composites, optoelectronic and nanoelectronics sensing, energy harvesting and storage, as well as in environmental remediation [1]. Furthermore, tetrapodal particles can be further assembled into highly porous (up to 98%) macroscopic ceramic framework structures, providing a very high free volume as the spatially extended arms of the tetrapods prohibit efficient close packing. These interconnected networks offer easy access to active surfaces in a mechanically stable, compliant and low-density structure [2] and can be further utilized as a versatile template for the fabrication of other multi-scaled foam-like materials, so-called aeromaterials, from different 1D and 2D nanomaterials, such as graphene, hexagonal boron nitride, as well as carbon

nanotubes [3,4]. These freestanding aeromaterials are ultra-lightweight, with densities in the order of only a few mg/cm³ and are characterized by a unique set of mechanical, electrical and optical properties [4,5].

The low nominal elastic moduli of such interconnected networks are very favourable for flexible electronics and mechanically robust sensors [6], but there is limited knowledge of the relationships between network architecture and properties, nor how these degrade with mechanical damage such as from cyclic loading [7]. This is needed to design functional materials with optimal properties and prolonged lifetimes. Novel micromechanical studies of the hinging and buckling deformation of individual aero-graphite tetrapods [8] provided insights into how they may behave collectively as self-assembled networks [4]. However, there have been no such studies of functional ceramics such as t-ZnO, although both the elastic modulus and electrical conductivity have been observed to decrease as network connections change when the elastic limit is exceeded [9].

Here we characterise the 3D network of interconnected t-ZnO using high resolution X-ray computed tomography (μ XCT) to quantify the tetrapods' geometry and distribution. *In situ* μ XCT observations during mechanical loading allow deformation within the bulk material to be

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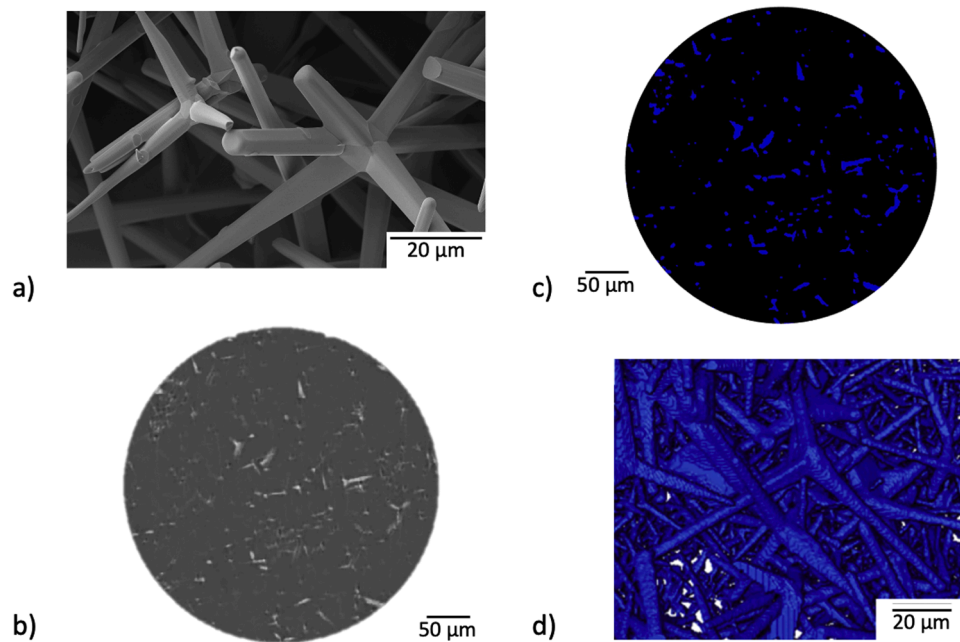


Fig. 1. Visualisation of the t-ZnO network (0.3 g cm^{-3} density): (a) scanning electron microscopy; (b) cross-sectional orthoslice of tomograph (after filters to reduce noise); (c) threshold segmented orthoslice; (d) 3D rendering of a tomographed region (after segmentation).

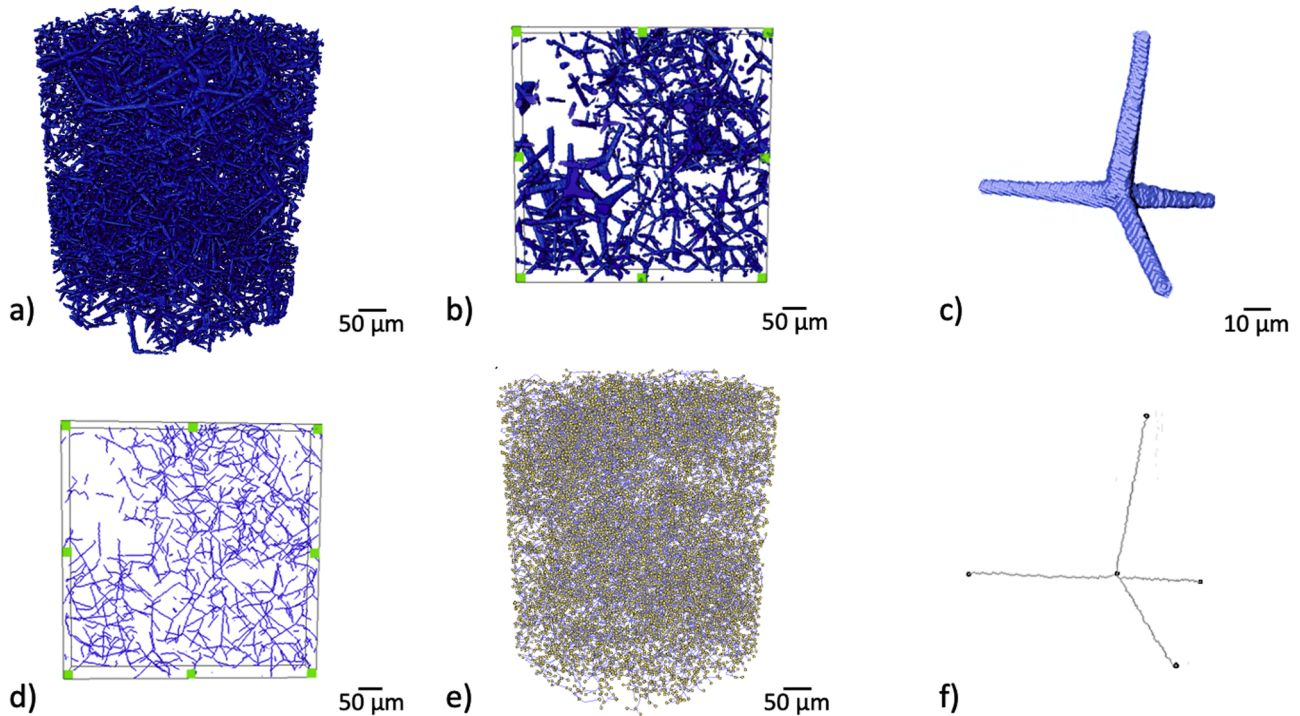


Fig. 2. Skeletonisation of the t-ZnO network (0.3 g cm^{-3} density): (a) visualisation of the tomographed region of interest after labelling; (b) tetrapods in a subregion of the same volume; (c) a single tetrapod; (d) the subregion after skeletonization (ligaments only shown); (e) tomographed region of interest after skeletonization (ligaments and nodes shown); (f) the skeletonised single tetrapod with ligaments and nodes.

examined, for the first time.

The t-ZnO material was produced via flame transport synthesis with post-synthesis compression and sintering [9]. Briefly, zinc microparticles (size 1–10 μm) were mixed with polyvinyl butyral (mass ratio 1:2) and heated for 30 min at 900 $^{\circ}\text{C}$ to produce a loose t-ZnO powder. Different masses were compressed in a ceramic mould (3 mm height, 3 mm diameter) to form pellets that were heated (unconstrained) at

1150 $^{\circ}\text{C}$ for 5 h to obtain specimens with nominal bulk densities between 0.3 and 0.9 g cm^{-3} . Scanning electron microscopy (Zeiss NVision 40 SEM, operating at 5 kV with a $\sim 10 \text{ nm}$ carbon coating) (Fig. 1a) confirmed the characteristic network architecture of interconnected tetrapods [10].

3D characterisation was done by μXCT (Zeiss Xradia Versa 610 microscope, operating between 60 and 120 kV with voxel size from 1 to

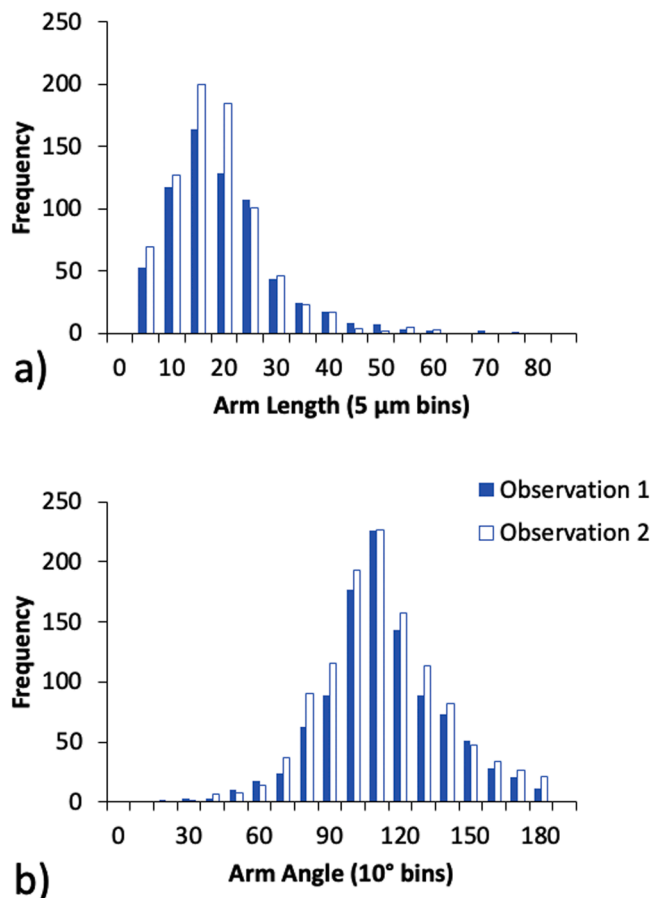


Fig. 3. Distributions of arm lengths and inter-arm angles measured by skeletonization of tomographs of t-ZnO with 0.3 g cm^{-3} density. Observation 1 and 2 were at 5 and 10% nominal strain, and show no significant difference.

1.7 μm) using the instrument software for attenuation contrast image reconstruction. Post-processing and analysis of the tomographs was done using the Avizo Fire software (v.2020.3.1). A Non-Local Means Filter was applied before intensity-based segmentation of the higher attenuation ZnO (Fig. 1b–d). Tomographs at 1.7 μm per voxel were sufficient to resolve the t-ZnO structure, though volume and surface measurements of the tetrapods by image segmentation have a subjective bias; the partial volume effect [11] is significant because the t-ZnO arms have cross-section dimensions ($\sim 1\text{--}3 \text{ μm}$) comparable to the image voxel size [3]. All samples were quite homogeneous; the observed solid fraction of ZnO within the sample volume (a moving cube of $\sim 300 \text{ μm}$ side) was constant both radially and axially (variance $< 15\%$) for all densities. Precise measurement of the total volume ($\pm 0.001 \text{ cm}^3$) via image texture classification to determine the sample's external boundaries, and of its mass using an A&D GH-202 balance ($\pm 0.01 \text{ mg}$) gave true densities of 0.29 ± 0.01 , 0.41 ± 0.01 , 0.52 ± 0.01 and $0.74 \pm 0.01 \text{ g cm}^{-3}$ for “nominal” densities of 0.3, 0.45, 0.6 and 0.9 g cm^{-3} respectively. The difference, which indicates expansion of the pellets, is greater for the higher density materials and is deduced to arise from interactions between the tetrapods after compression during processing.

To quantify the structure of the tetrapod network (Fig. 2a and b) a thinning algorithm [12] can be applied to the segmented image to determine its medial axis skeleton, which is described as a set of nodes connected by ligaments (1 voxel thickness) that are equidistant from the surfaces of the solid (Fig. 2d and e). Dead-end branches with length fewer than 5 voxels were regarded as noise and removed. This allowed measurement of arm lengths (between connected nodes) and interarm angles (between pairs of arms with common nodes). This skeletonization generally, and incorrectly, resolved the centers of tetrapods as 2 closely

adjacent nodes (1 or 2 voxel separation) instead of a single node. An example tetrapod, from a 0.3 g cm^{-3} sample tomographed at 1 μm voxel, is shown in Fig. 2c. Its skeleton (Fig. 2f) has a pair of 3-coordinated nodes at its center that have been joined for visualisation. Considering this double-counting, the number of tetrapods within a reconstructed volume of $\sim 0.05 \text{ mm}^3$ was estimated at ~ 3000 , i.e., a number density of $\sim 6 \times 10^4 \text{ mm}^{-3}$. The general network characteristics were insensitive to deformation (Fig. 3); in successive compressive observations (5 and 10% nominal strain) the mean arm lengths (Fig. 3a) were 17.4 μm (standard error in the mean 0.4 μm , standard deviation 11 μm) and 16.0 μm (standard error in the mean 0.3 μm , standard deviation 9.2 μm) respectively. The interarm angle distributions were also similar (Fig. 3b) at 107.0° (standard error in the mean 0.8° , standard deviation 25.7°) and 107.4° (standard error in the mean 0.8° , standard deviation 26.0°) respectively. The mean angle is close to the theoretical angle of $\sim 110^\circ$ [13]. The observed range illustrates the lack of perfect symmetry that arises from lattice strains during crystal growth [9] and is consistent with electron microscopy of t-ZnO structures produced by chemical vapour deposition [14].

To examine the deformation of the t-ZnO network in more detail, *in situ* XCT observations at 0.7 μm voxel were performed of a 0.45 g cm^{-3} sample, which was mounted in an X-ray transparent PMMA load-bearing tube and compressed between PTFE anvils by sequential displacements of $\sim 75 \text{ μm}$ applied via a manually controlled micrometer. The nominal strain over the sample length (3 mm) was estimated by μXCT observation of the change in anvil separation. Incremental global digital volume correlation (gDVC), which converged for mesh sizes of $100\text{--}250 \text{ μm}$, was applied using Avizo, whereby the tomograph at each step was correlated with the reference tomograph of the previous step. The incremental displacement vector fields were summed to obtain the cumulative three-dimensional displacements. Further details are summarised in the supplementary information. As an example, Fig. 4a is a vertical section of the residual field at 8.7% nominal strain. The residual field [15] quantifies the difference between the second image and the reference, where the latter has been deformed by the gDVC-calculated displacement field; large residuals (i.e., high contrast in this visualisation) indicate poor correlation, which can be caused by local changes in the image (i.e., the structure of the material). Significant residuals were observed close to the moving anvil within a region where the mean displacements were large and of high variability (Fig. 4b); the size of this region increased with applied nominal strain. This region of poor correlation and large displacements demonstrates significant and local changes to the network structure. In the lower part of the sample, where low residuals indicated accurate displacement measurement and no significant changes in structure, the displacement gradient (i.e., strain) increased with the applied strain. The measured strain in the lower region was less than the applied strain, due to accommodation of the displacement of the moving anvil in the upper region, and increased in magnitude as the high residual region was approached (Fig. 4c), showing nonlinear material behavior with increasing deformation. However, similar behavior was observed on unloading in both regions; when the 0.45 g cm^{-3} sample was unloaded by 2.4% nominal strain from a compressive nominal strain of 8.7%, the gDVC analysis measured a uniform strain relaxation of $2.5 \pm 0.2\%$ with low residuals over the entire sample. Such reversible behavior is quite like the compaction of a granular solid [16] in which *in situ* observations have also shown anelastic deformation to localize at the contacting surface [17,18].

The electrical conductivity of a t-ZnO network (density $\sim 0.115 \text{ g cm}^{-3}$) has been observed to increase and then decrease with compression [9], which was attributed to improved tetrapod contact with increasing strain and then disruption of the network; the applied deformation indicated an elastic limit of $\sim 2\%$ nominal strain. Inspection of Fig. 4c shows the transition between the high correlation and low correlation regions is at local strain of $\sim 0.5\text{--}1\%$. This lower elastic limit may be due to the higher network density of this 0.45 g cm^{-3} sample.

The mechanism by which the network structure accommodates

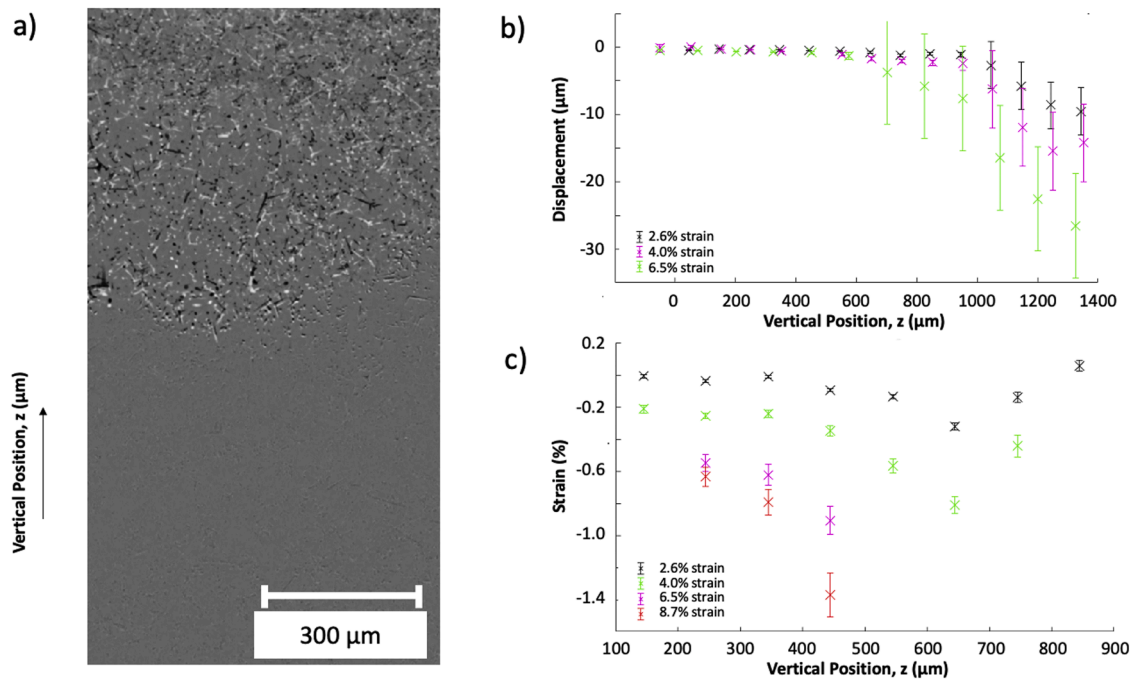


Fig. 4. Measurement of deformation in t-ZnO (0.45 g cm^{-3} density) by global digital volume correlation of computed X-ray tomographs; (a) the residual field at 8.7% nominal strain; (b) vertical displacement with distance from the bottom of the sample at different nominal strains (data at 8.7% strain are not shown due to the high residuals); (c) vertical strain (in the region of low residuals) with distance from the bottom of the sample at different nominal strains. The fixed anvil is at the lower surface ($z \sim 0$).

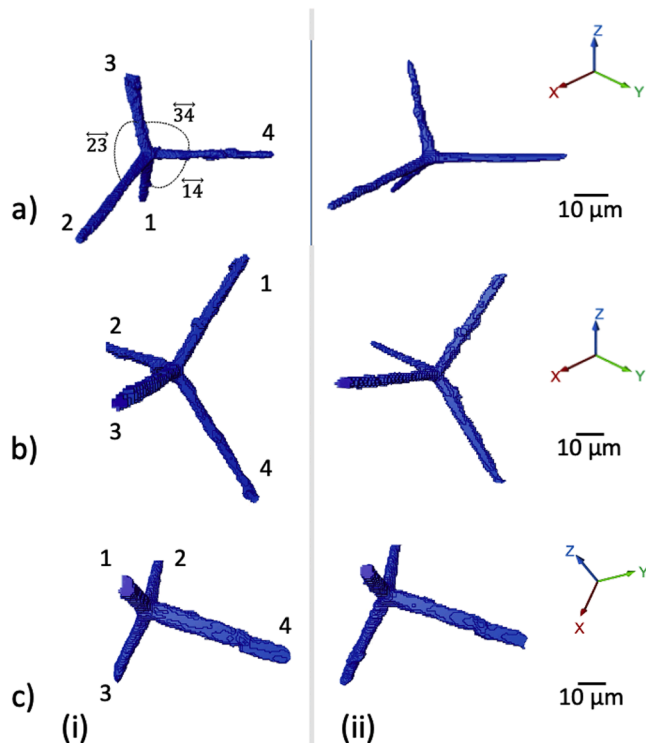


Fig. 5. Visualisation of t-ZnO tetrapods a) to c) from (i) observation 1 at 5% nominal strain and (ii) observation 2 at 10% nominal strain in a network with 0.3 g cm^{-3} density. The anvil compression direction between *in situ* observations was parallel to the Z-axis. Arms 1–4 are labelled, where 34 is the angle between arms 3 and 4, etc.

Table 1

Inter-arm angles ($^\circ$) measured for tetrapod examples (a)–(c) (see Fig. 5) at (i) observation 1 at 5% nominal compressive strain and (ii) observation 2 at 10% nominal strain in t-ZnO with 0.3 g cm^{-3} density. 12 is the angle between arms 1 and 2, etc. Significant changes are highlighted in bold.

		12	13	14	23	24	34
a)	(i)	115 \pm 2	131 \pm 5	103 \pm 1	100 \pm 1	102 \pm 1	101 \pm 1
	(ii)	101 \pm 2	102 \pm 2	117 \pm 3	102 \pm 2	127 \pm 5	103 \pm 2
b)	(i)	99 \pm 3	131 \pm 6	110 \pm 3	117 \pm 6	98 \pm 3	98 \pm 2
	(ii)	97 \pm 7	100 \pm 2	106 \pm 6	114 \pm 9	119 \pm 12	116 \pm 7
c)	(i)	93 \pm 2	128 \pm 7	124 \pm 7	109 \pm 3	92 \pm 3	102 \pm 3
	(ii)	133 \pm 13	87 \pm 5	108 \pm 7	116 \pm 4	105 \pm 3	103 \pm 2

deformation was examined for individual tetrapods (Fig. 5) in a 0.3 g cm^{-3} sample that was observed *in situ* (at $1 \mu\text{m}$ voxel). The inter-arm angles of the same tetrapods were measured in successive observations at nominal compressive strains of approximately 5 and 10% (Table 1); i.e., where the bulk of the sample was beyond the elastic limit. Angular measurement errors were estimated conservatively using $\pm 2^\circ$ and $\pm 1^\circ$ uncertainty in the location of the central node and arm tip nodes. The first example (Fig. 5a) increased the inter-arm angles 14 and 24, and decreased 12 and 13 with increasing strain. This corresponds to rotation of arms 1 and 2 away from arm 4 and towards arm 3 and shows the tetrapod was compressed parallel to the loading axis. No bending of the arms could be observed. ZnO whiskers have a very high elastic limit [19] and it is assumed the deformation of the individual tetrapod arms may be similar to the elastic hinging behavior that has been observed in micromechanical testing of single arms of aero-graphite tetrapods [8]. Other t-ZnO examples (Fig. 5b and c) exhibited changes in shape that were less clearly aligned with the applied load. This indicates a quite complex mechanical interaction with adjacent structures, as has been predicted in interconnected tetrapod networks [4]. In the second example (Fig. 5b), arm 3 moved towards arm 1 largely by rotation about arm 2. In the third example (Fig. 5c), arm 1 rotated away from arm 2 towards arms 3 and 4.

In summary, we have shown that the network architecture of sintered t-ZnO can be observed by μ XCT and quantified in a reproducible manner by image post-processing. *In situ* μ XCT with gDVC finds significant and non-uniform deformation of the network structure with changes in the geometry of individual tetrapods. This observation is relevant to t-ZnO and also to materials for which it provides a template, such as aerographene [20] and aero-hBN [3,5]. Such deformation will affect how properties change at the elastic limit and the response of functional properties to cyclic fatigue, for example. μ XCT may therefore be a very useful tool to study the interactions between deformation, microstructure and properties of functional network materials.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.scriptamat.2022.115153.

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