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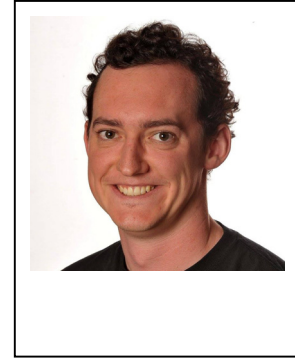
# **MODERN DEVELOPMENTS AND APPLICATIONS IN MICROBEAM ANALYSIS**

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**EXPLORING MARS AT THE NANOSCALE: APPLICATIONS OF TRANSMISSION ELECTRON MICROSCOPY AND ATOM PROBE TOMOGRAPHY IN PLANETARY EXPLORATION**

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## 1. INTRODUCTION

Rocks from Mars are among the rarest meteorites and are the only physical samples we currently have of the red planet. They represent a total of 277 individual stones, with a combined weight of 162.5031 kg [2] split between five main lithologies (four igneous and one breccia) the shergottites, nakhlites, chassignites, black beauty paired stone, and the anomalous Alan Hills (ALH) 84001 [2, 4-6]. Cosmogenic exposure ages indicate that they sample 4 - 8 ejection sites on Mars [5]. A Mars sample return mission is being planned to recover samples that had been previously collected and stored by the Mars 2020 rover; these samples will add a further ~ 0.5 kg of fresh oriented rocks to our collection - potentially including sedimentary lithologies [9, 10].

Martian meteorites have proven invaluable in providing insights into volcanic activity, the action of water and impact processes on Mars [4, 6, 11-15]. However, the rarity of Martian samples means that as analytical scientists we are duty bound to maximise the amount of data collected from the smallest sample volume to provide the greatest insight into Martian planetary processes. Here we describe our recent application of transmission electron microscopy (TEM) and atom probe tomography (APT) techniques to Martian meteorites. TEM and APT are two nanoscale techniques that provide the high-resolution culmination of a correlative analytical pipeline (Fig. 1) that maximises data output from the minimum sample volumes to address research questions pertinent to the red planet.

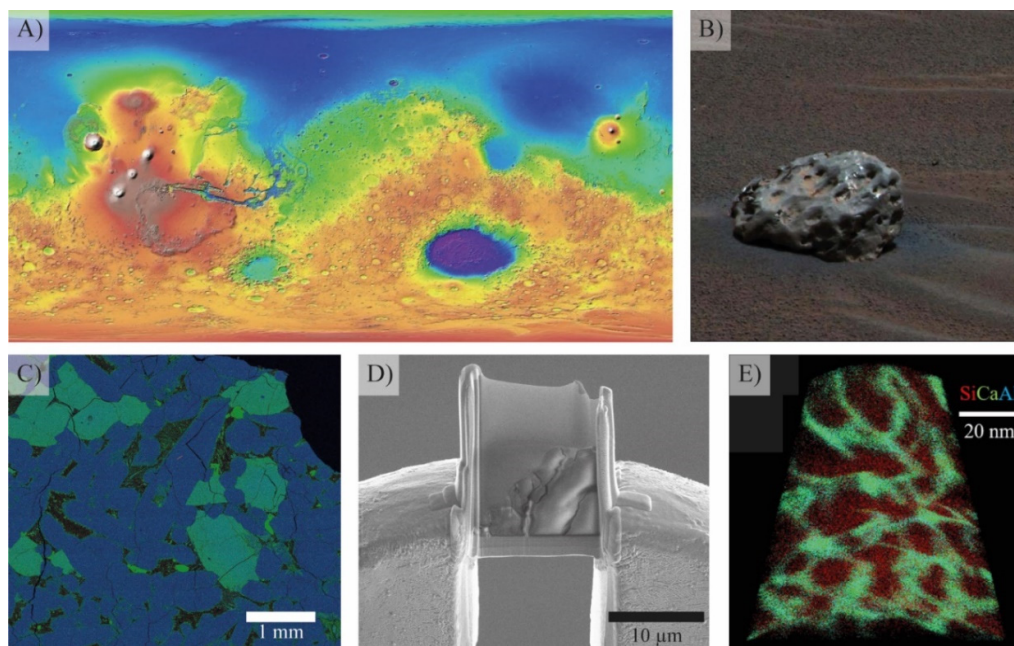


Figure 1. A correlative pipeline across a broad array of length scales to interpret the geological history of the planet Mars. a) An elevation map of the Martian surface from the NASA MOLA science team. b) An image of an iron meteorite on the surface of Mars taken by NASA's Opportunity rover. c) A red (Al), green (Fe), blue (Mg), EDS image of Martian meteorite Yamato 000802. d) A TEM lamella of a section through a magnetite grain in the Tissint meteorite. e) A APT analysis of an atom probe needle produced extracted from the mesostasis of Martian breccia North west Africa 11522 (after [3]).

## 2. FOCUSED ION BEAM TEM/APT SAMPLE PREPARATION

The preparation of geological and extraterrestrial samples for TEM and APT usually involves extraction from a polished section using a focussed ion beam (FIB) instrument. Typical TEM sample preparation requires extracting a 1 - 2  $\mu\text{m}$  thick bar of material that is then thinned to electron transparency, usually  $< 100 \text{ nm}$  [16]. TEM samples produced by Ga-FIBs are generally  $15 \mu\text{m} \times 20 \mu\text{m}$  as larger samples would be too time intensive to extract. However, with the development of Xe/plasma-FIB instrumentation that can remove ‘vast’ volumes of material rapidly, the width and depth of a TEM sample are limited only by the cohesive strength of the material. It is now feasible to produce  $100 \mu\text{m}$  wide TEM sample. To avoid bending/warping during thinning of wider samples the structural integrity can be maintained by only thinning ‘windows’ at regular intervals, with structurally sound ‘posts’ in between. Wider TEM samples are no longer limited to small regions of interest and can now evaluate heterogeneities in mineralogy, density, porosity, etc., across  $100 \mu\text{m}$  lengths. This is incredibly powerful in meteoritics because most meteorites are heterogeneous over these length scales.

Typical APT sample preparation approaches involve milling out a triangular prism and attaching segments to pre-prepared posts [17]. Sample mounts for APT are Cu or Mo five post grids otherwise used for TEM analysis that have been FIB milled into needles or pre-grown arrays of Si posts. Once attached, the APT samples are thinned using an annular milling pattern to produce a needle with a  $100 \text{ nm}$  diameter tip [17]. APT samples can also be prepared in situ using the ‘satellite dish’ method ([1] Fig. 2a). This approach involves cutting a  $150 \mu\text{m}$  diameter wide  $50 \mu\text{m}$  deep circular depression in the sample with APT needle in the centre [1]. A wedge shape ‘window’ is also cut to view the needle in the atom probe ([1] Fig. 2a). This approach maintains the orientation of the specimen relative to the sample. The satellite dish method combined with electron backscatter diffraction (EBSD) would be a powerful correlative analytical approach. However, this approach is only viable using a Xe-FIB due to the ‘vast’ amounts of material that need to be excavated and where most of the sample surrounding the region of interest is forfeit and as such is only suitable for specific applications. The ‘*button method*’ of FIB APT/TEM sample preparation enables sub-micrometre grains, interfaces, defects and twin boundaries to be targeted ([8] Fig. 2b). This approach becomes particularly powerful when combined with SEM, EBSD, FIB-TOF-SIMS characterisation prior to FIB preparation to pinpoint the region of interest [8]. A ‘*button*’ of electron beam deposited platinum is deposited to highlight the feature and locate it after covering with the ion beam protective Pt strip ([8] Fig. 2b). These buttons permit accurate extraction and thinning of TEM/APT samples to ensure the region of interest is contained within the sample. This approach has been successfully applied to extract  $100 \text{ nm}$  diameter refractory metal alloys in meteorites [18, 19].

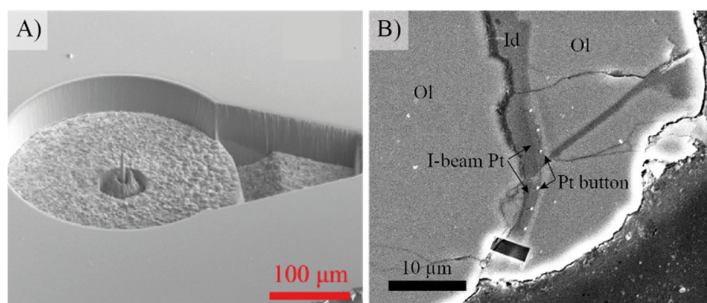


Figure 2. Images of a suite of FIB sample preparation approaches to both TEM and APT. a) APT sample preparation using the in situ ‘satellite dish’ approach, adapted from [1]. b) The ‘button method’ of APT sample preparation of dish from Rickard *et al.* [8] used here on to decorate and target the olivine-iddingsite interface in the Lafayette meteorite.

## TRANSMISSION ELECTRON MICROSCOPY

TEM has been used extensively to characterise water-bearing minerals in Martian meteorites (e.g., [20, 21]), and is complementary to APT. TEM works by irradiating a thin sample – ideally < 100 nm – with a beam of high energy electrons (typically 200 kV) under vacuum. These thin samples are often referred to as ‘electron transparent’. Those electrons that pass through the sample can be used to form images, down to atomic resolution. If the beam is relatively broad and static, bright- and dark-field images are formed using the unscattered and scattered electrons, respectively. By scanning a focused beam over the sample, scanning TEM (STEM) images are formed, again in bright-field (using unscattered electrons) and dark-field (using scattered electrons) modes. Electron diffraction patterns represent the angular range of scattering, and the d-spacings obtained from them can be used to identify minerals and determine their crystallographic orientation. The energy that the electrons lose as they pass through the sample can also be measured to obtain information on chemical composition and bonding (i.e., electron energy loss spectroscopy, EELS). An energy-dispersive X-ray (EDX) detector located above the thin sample enables quantitative and qualitative chemical analysis in the same manner as EDX in an SEM. For a more detailed description of TEM and STEM in planetary science see Lee [22].

### 3. ATOM PROBE TOMOGRAPHY

The APT technique is described in detail in Kelly and Larson [23] and Gault *et al.* [24]. Therefore, we only provide a brief overview of the technique here. APT is a high-resolution technique capable of analysing small cylindrical volumes (up to approximately  $100 \times 100 \times 1,000$  nm) with a spatial resolution of 0.3 nm in three dimensions [23]. The required specimen geometry for APT analysis is needle-like, with a 100 nm diameter tip and a half shank angle <  $10^\circ$  [23]. The needle-like specimen is loaded in the atom probe under high vacuum and situated near an annular electrode [25]. A voltage of  $\sim 1,000$  V is applied to the specimen by the electrode. If the specimen is conductive, the voltage can be pulsed to facilitate field evaporation of individual atoms from the surface of the needle [23, 24]. However, if the sample is non-conductive (i.e., most geological and extraterrestrial samples) field evaporation can be initiated through the addition of heat by applying a high frequency (100 - 250 kHz) ultraviolet

laser to the tip of the sample to achieve single atom ionisation [26]. It was the addition of the laser to atom probe instruments that enabled the analysis of non-conductive geological samples [26].

Field evaporated ions are accelerated across a potential difference where they impinge upon a position sensitive detector [23, 24]. The time of flight from laser/voltage pulse to impact provides the mass-to-charge-state-ratio of the ion and thus its elemental and isotopic identity, while the x- and y-coordinates on the detector can be used to reconstruct the initial position of the atom within the sample and thus reconstruct the 3D atomic structure of the sample [23, 24]. The detection efficiency of the detector is dependent on the available surface area on the detector surface, and is generally between 50 - 65 % [23, 24]. However, new generation detectors in atom probes have an increased surface area available for detection and can now collect over 80 % of the ions produced. In principle, the energy imparted to the sample should ionise all elements of the periodic table with equal efficiency resulting in 100 % ionisation of the sample [27, 28]. Therefore, APT does not, in theory, suffer mass dependant fractionation of isotopes or produce biases in the elements detected. The only sources of uncertainty are then derived from counting statistics based on the actual number of ions measured [26]. As such, APT has a high sensitivity and spatial resolution, and so can in principle be used to determine crystallography and atomic lattice spacings, complete geochemical characterisation including trace elements [19, 29-31], depict nanoscale geochemical heterogeneities and clustering of atoms [29, 32-34], and isotope analysis across all elements in the periodic table [35-40].

To extract elemental and isotope abundances from the mass-to-charge-state-ratio-spectrum resulting from the analysis, the total number of counts beneath each peak must be calculated. This is done by ‘ranging’ the counts beneath each peak [24] using the IVAS software package over a specific unified atomic mass unit ( $u$ ) width followed by a local background subtraction from a nearby region that contains no peaks over the same  $u$  width. Details about ranging methods and advantages and disadvantages of different approaches can be found in Daly *et al.* [38] and Parman *et al.* [25]. In general, for geochemical analysis the full width of the peak should be used whereas for isotope analysis slightly less than the width of the smallest peak should be used consistently for all peaks isotope peaks of interest [38].

Analysis of geological and extraterrestrial samples is complicated by the formation of multi-ion complexes, i.e., OH or Si<sub>2</sub>O<sub>3</sub>. These result in isobaric interferences that need to be deconvolved to extract robust elemental or isotopic abundances. APT can detect trace abundances of elements down to 10 atomic parts per million (appm) for sample volumes over 10<sup>6</sup> nm<sup>3</sup> [23] while newer atom probe instruments are capable of detection limits as low as 1 appm. This means APT can be utilised to analyse the trace element chemistry of mineral phases. Furthermore, the 3D reconstruction of the atomic positions and the size of the sample mean that it is possible to ensure only atoms emitted from the phase of interest are included in the spectrum. This means an accurate bulk chemistry of the phase of interest can be derived (e.g. [19, 29, 30, 41]). Therefore, APT is ideal for analysing sub-micrometre size phases whose trace element

chemistry cannot be obtained through conventional techniques, or where the signal from the phase of interest is contaminated by contributions from surrounding phases. In addition, it is possible to extract nanostructural geochemical information such as planar or point cluster concentrations of elements even in trace amounts within samples. This approach has been demonstrated many times for the geochronological studies of Pb clusters in zircon, and baddeleyite [32, 33, 35-37] and gold in arsenopyrite [29].

The mass resolving power ( $m/\Delta m$ ), where  $m$  is mass and  $\Delta m$  is the variation in mass between atoms, of the atom probe is better than 1,000 [26, 42]. Therefore, APT is capable of discerning peaks separated by 0.15  $u$  [26, 42]. Thus, isotopic peaks from ions of different charge states (i.e.,  $^{94}\text{Mo}^{3+}$  and  $^{63}\text{Cr}^{2+}$ ) can be distinguished, but APT cannot currently determine the mass difference between ions that have the same mass (i.e.,  $^{54}\text{Cr}^{2+}$  and  $^{54}\text{Fe}^{2+}$ ) [26, 42].  $^{54}\text{Cr}^{2+}$  and  $^{54}\text{Fe}^{2+}$  have a difference in mass of 0.0005  $u$ , which requires a mass resolving power  $> 1.5 \times 10^5$  [26, 42] to distinguish these ionic species. These isobaric interferences are beyond the mass resolving power of any technique currently available [26, 42] and require peak deconvolution to estimate the contribution of each isotope to the total peak. Field evaporation during APT analysis is ideal for isotopic analysis of nanoscale materials as the whole sample is ionised, and it is assumed that there is no bias in the detection of elements or between isotopes, i.e., all species are ionised and detected equally [27, 28]. However, unique factors in each analysis, such as voltage, tip shape and temperature are not constant between analyses and their effect on the resulting mass spectrum is not quantified and potentially not quantifiable [25, 33]. Despite these challenges a robust standardised approach for isotope analysis has been proposed for Re/Os isotope analysis and is readily adaptable to other isotope systems and can distinguish isotopic ratio variations between 0.5 - 1 % with substantial room for improvement [38].

#### 4. A BRIEF HISTORY OF ATOM PROBE TOMOGRAPHY IN PLANETARY SCIENCE

The first paper on non-conductive geological materials using a laser assisted atom probe microscopy was by Valley *et al.* [35] who demonstrating that the  $^{207}\text{Pb}/^{206}\text{Pb}$  isotopic abundances can be measured in zircon using APT. The APT  $^{207}\text{Pb}/^{206}\text{Pb}$  isotope ratio recorded an age consistent with measurements of the same zircon using secondary ion mass spectrometry (SIMS) [35]. This study put atom probe on the map as a powerful geochronological tool for geoscience applications that is capable of reliably reproducing results of other more established high precision isotopic techniques but using a much smaller sample volume.

One of the first uses of APT on extraterrestrial samples was Heck *et al.* [39] who reported APT data for C isotopic ratios in nanodiamonds within primitive chondritic meteorites. These results indicated that the  $^{12}\text{C}/^{13}\text{C}$  ratio is anomalous relative to Solar System values [39]. However, these anomalous isotopic ratios may instead reflect contributions of  $^{12}\text{C}$ -hydrides to the  $^{13}\text{C}$ -peak (Heck *et al.* 2014) and possible underrepresentation of the  $^{12}\text{C}$  peak due to molecular ion formation and detector dead times [41]. This work has been followed by continued APT work

on pre-solar grains in Lewis *et al.* [40], unpicking the nanostructures and geochronological information trapped in shocked zircon, baddeleyite and Lunar meteorites [34, 36, 37], unpicking the magnetic properties of iron meteorites [31] and geochemical analysis of refractory metal alloys [18, 19]. We have only scratched the surface of the insights of APT in planetary science.

The real interpretive power of APT and TEM is through correlative analysis combining several techniques to provide context to these nanoscale analyses (e.g., [8]). Below is an example workflow showing how light microscopy, SEM, EBSD, FIB, TEM, STEM and APT can be successfully combined to locate and characterise water-bearing minerals in Martian meteorites.

## 5. WATER ON MARS

Ashworth and Hutchison [11] first described evidence for water-bearing material in the nakhlite group of Martian meteorites (see [12] for a recent review). These authors described the water-bearing material as “iddingsite”, which they interpreted to have formed by hydrothermal alteration of olivine. This fluid that produced the iddingsite assemblage was oxidising in nature, low temperature (< 140 °C) [15] and affected these rocks approximately 633 Ma [43]. The heat source that melted these subsurface ices is not currently known, but the primary theories are impact generated hydrothermal activity [44] or a magmatic event [43].

The iddingsite veins can be readily identified using transmitted light microscopy (Fig. 3) and are a few micrometres in thickness. They resemble narrow veins of ‘rust’ in transmitted light (Fig. 4a) and although more detail can be observed by SEM (Figs. 4b and 4c) their properties remain difficult to determine. EDS shows that these veins have a Mg-Fe silicate composition (~ 40 wt% SiO<sub>2</sub>, 40 wt% Fe<sub>2</sub>O<sub>3</sub>, 5 wt% MgO) and their low analytical totals (~ 85 - 90 wt%) indicate the presence of unanalysed OH/H<sub>2</sub>O (i.e., Martian water). EBSD demonstrates that the veins have a preferred crystallographic orientation parallel to [010], and this crystallographic information can be used to help guide the orientation in which the electron-transparent samples can be cut and extracted using FIB (Figs. 2c and 2d). Bright-field TEM images show that the host olivine is microstructurally complex, probably reflecting mild shock metamorphism [21] whereas the iddingsite veins appear to be homogeneous and featureless (Fig. 4d).

To characterise OH/H<sub>2</sub>O-bearing Fe-Mg silicates, compositional maps were obtained from the area in Fig. 4d using EDX and EELS (Fig. 5). Results highlight the location of siderite and parts of the carbonate have altered to ferrihydrite (Fe<sup>3+</sup><sub>8.2</sub>O<sub>8.5</sub>(OH)<sub>7.4+</sub> 3H<sub>2</sub>O) (Fig. 5). As well as elemental analysis, EELS is used to determine the valence state of Fe (and other elements) [45]. A Fe valence map of the vein shows that the OH/H<sub>2</sub>O-bearing Fe-Mg silicate and ferrihydrite both contain Fe<sup>3+</sup> whereas the siderite and olivine have Fe<sup>2+</sup> as expected (Fig. 5).

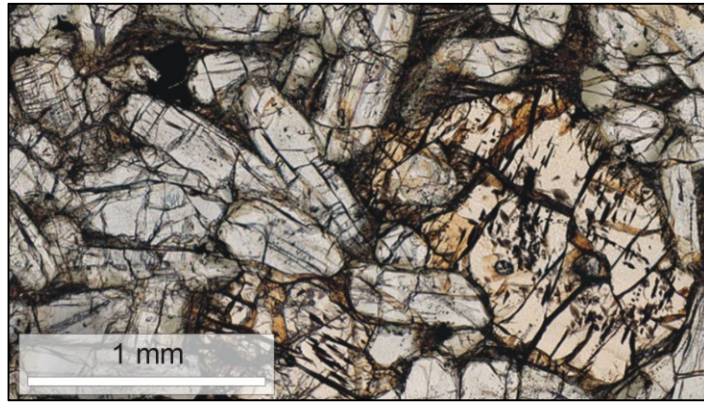


Figure 3. Transmitted light image of a thin section of the Nakhla meteorite. An olivine grain in the centre of the field of view has a light brown colour and is cut by several very thin veins of rust-coloured iddingsite. The clear tabular mineral grains are augite. Image from the Open University virtual microscope ([www.virtualmicroscope.org/content/nakhla-2](http://www.virtualmicroscope.org/content/nakhla-2)).

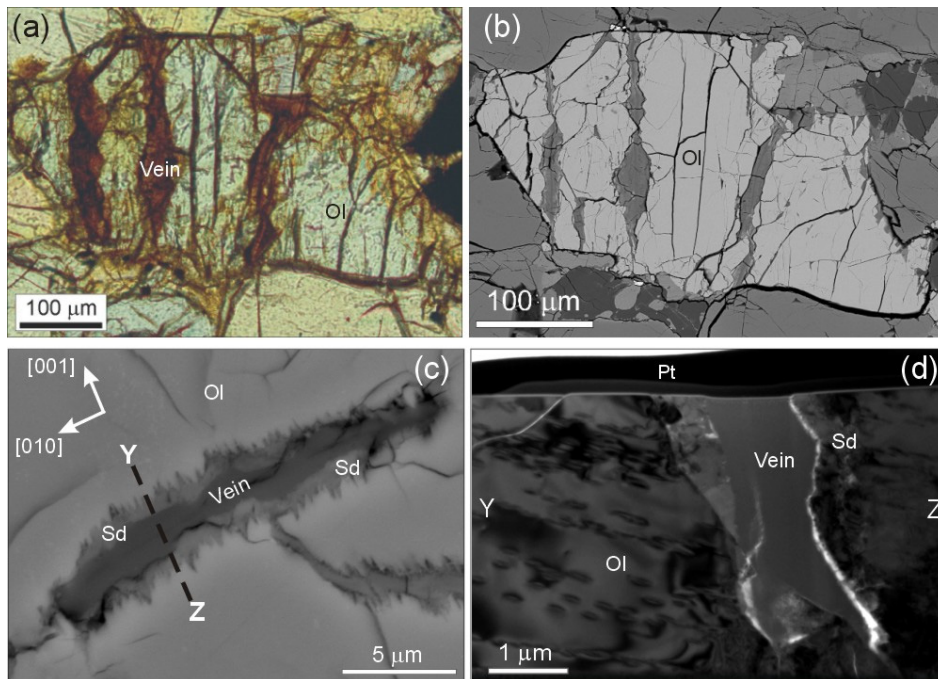


Figure 4. Iddingsite in the Lafayette meteorite, (a & b), and the Nakhla meteorite (c & d), hosted by olivine (Ol). a) Transmitted light image, and b) backscattered electron SEM image of an olivine grain that is cross-cut by three rust-brown coloured iddingsite veins. c) Backscattered electron SEM image of an iddingsite vein. Between the iddingsite and olivine is a selvage of siderite (Sd) with a finely serrated contact with olivine. The crystallographic orientation of the vein relative to the host olivine grain is indicated, as determined by EBSD. The dashed black line Y-Z denotes where a FIB section was extracted. d) Bright-field TEM image of the sample extracted from (c). The olivine is various shades of grey as its defect-rich microstructure is diffracting electrons to different intensities. At this magnification the vein is homogeneous and apparently free of diffracting crystals. Siderite (Sd) is difficult to distinguish from olivine. The black band along the top of the image is the platinum (Pt) strap that was deposited by the FIB to protect the sample from ion beam damage.

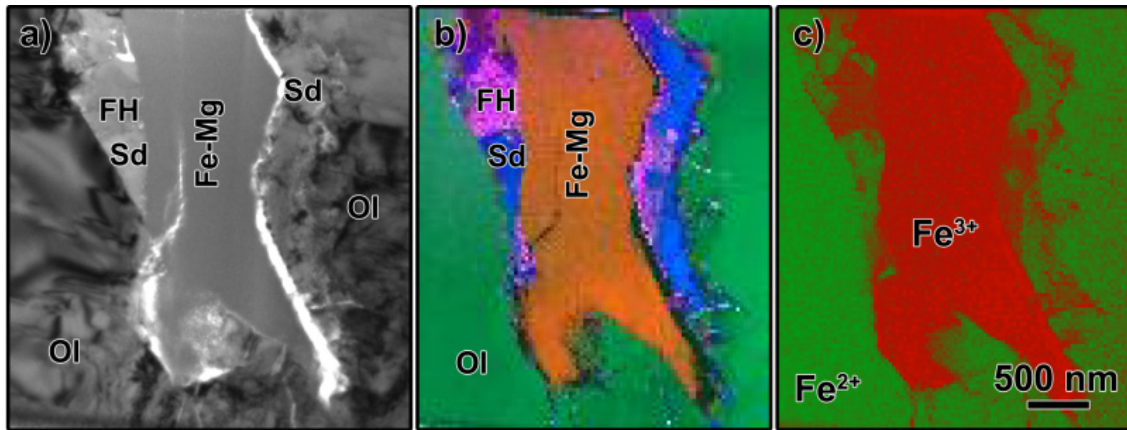


Figure 5. Iddingsite veins from the Nakhla meteorite. a) Bright-field TEM image of a vein of OH/H<sub>2</sub>O-bearing Fe-Mg silicate (Fe-Mg) that is flanked by siderite (Sd) and hosted by olivine (Ol). The siderite also contains ferrihydrite (FH). b) EDS map of (a) with element combinations coloured so as to render olivine green, siderite blue, ferrihydrite pink, and the Fe-Mg silicate orange. d) EELS valence map of (a) and (b) showing that olivine and siderite both contain Fe<sup>2+</sup> (coloured green) and the Fe-Mg silicate and ferrihydrite have Fe<sup>3+</sup> (red).

The identity of the OH/H<sub>2</sub>O-bearing Fe<sup>3+</sup>-Mg silicate could not be determined by TEM imaging or EELS alone. Thus, higher resolution STEM images and EDS maps were obtained using an aberration-corrected TEM with a probe size of < 2 nm (Fig. 6). STEM images show that the Fe-Mg silicate is composed of a closely packed mass of particles that are circular/oval and a few nanometres in diameter (Fig. 6a). EDS maps show that the particles are composed of Si with a Fe-rich rim (Fig. 6b). Given that this material contains OH/H<sub>2</sub>O and its Fe<sup>3+</sup>, the Fe-Mg silicate is interpreted to be composed of nanoparticles of opaline silica (amorphous SiO<sub>2</sub>·nH<sub>2</sub>O) that are rimmed by Mg-bearing ferrihydrite.

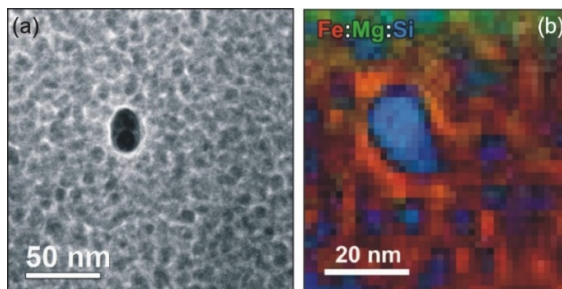


Figure 6. Iddingsite veins from the Nakhla meteorite. a) High-angle annular dark-field (HAADF) image of the OH/H<sub>2</sub>O-bearing Fe-Mg silicate showing that it is composed of a tightly packed mass of nanoparticles. b) EDS element map of one area of (a) demonstrating that the nanoparticles are composed of silica and have a Fe-rich rim.

APT samples have been extracted across the naxhlite olivine-iddingsite interface (Fig. 7) to observe nanoscale geochemical gradients and corroborate TEM observations. APT analysis of iddingsite reveal it had a heterogeneous density consistent with the HAADF images obtained by TEM (Figs. 6a and 7) but homogeneous chemistry. As such on the length scales of an APT needle in this region, we only identified clay minerals. No oxides, carbonates, sulphates or

halides [4] or nanoscale chemical domains were found. The density variations in iddingsite are interpreted to represent the preferential ionisation of individual phyllosilicate platelets (high density) followed by ‘bursting’ removing the clay mineral to leave a void (low density). APT analysis of across the iddingsite-olivine interface reveal that this boundary is gradational at the nanoscale. Towards the interface over 30 - 20 nm steady depletions in the major cations such as Mg are observed. These depletions are complemented by a steady enrichment in H towards the interface (Fig. 7). EBSD analysis of NWA 817 indicate the iddingsite vein is parallel to the (021) plane of the olivine consistent with previous petrographic descriptions of crystal orientation relationships in these meteorites [14]. However, the observation of a 20 nm wide diffuse boundary not a sharp boundary (Fig. 7) is inconsistent with recent models of fluid alteration of glasses, such as interfacial dissolution-reprecipitation [46]. Our APT data agree better with a diffusion coupled hydration model where inward diffusion of hydrogen species is coupled with outward diffusion of cations [47]. In addition, this 20 nm layer may store H more securely than iddingsite whose hydrogen isotopic composition is strongly perturbed by terrestrial weathering [48, 49] making inferences about the fluids origin challenging. However, directly measuring the D/H ratio of hydrogen contained within the olivine across these interfaces may reveal the D/H ratio of the original fluid, and thereby constrain its source (e.g., Martian crust versus atmosphere).

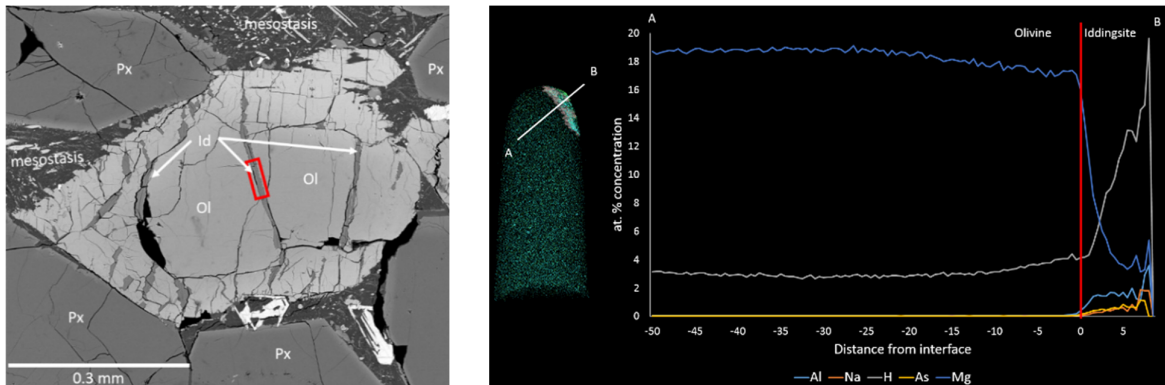


Figure 7. APT analysis of an olivine-iddingsite interface in nakhlite meteorite NWA 817. Image to the left is a backscattered electron (BSE) image of an olivine phenocryst with cross-cutting crystallographically controlled iddingsite veins. A set of APT samples were extracted from the red box so as the iddingsite-olivine interface was present at the apex of the needle. Ol = Olivine, Id = Iddingsite, Px = Pyroxene. Image below is an APT analysis across an olivine-iddingsite interface. The atom map represented by Al (blue dots) and Na (green dots) atoms defines the interface (grey iso-concentration surface). The graph shows a concentration profile of the major elements across the boundary from iddingsite (b) into olivine (A) where a clear depletion of Mg and enrichment of hydrogen is observed up to 20 nm into the olivine and may provide a secure reservoir for Martian Hydrogen after [7].

## 7. CONCLUSION

We stand upon the edge of a nanorevolution in planetary science. Through measurements at the atomic-scale using TEM and APT we can provide new insights into Martian and Solar System processes that act on length scales over 20 orders of magnitude larger than the sample volume analysed such as action of water on Mars that has implications for past and future habitability of the red planet. APT combined with improvements to the resolution of more traditional analytical approaches like TEM is about to open a whole new suite of nanoscale observations. This is particularly important for samples from the red planet where meteorite samples are rare and planned Mars sample return mission will provide at most a kilogram of material. Therefore, it is critical to maximise the research output from these small sample volumes. The combination of APT and TEM form a final set of techniques at the end of a correlated pipeline of analysis that will maximise the information gathered from such small sample volumes.

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