Needles in haystacks: using fast-response LA chambers and ICP-TOF-MS to identify asbestos fibres in malignant mesothelioma models†

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Malignant mesothelioma is an aggressive cancer associated with exposure to asbestos. Diagnosis of mesothelioma and other related lung diseases remains elusive due to difficulties surrounding identification and quantification of asbestos fibres in lung tissue. This article presents a laser ablation-inductively coupled plasma-mass spectrometry (LA-ICP-MS) method to identify asbestos fibres in cellular models of mesothelioma. Use of a high-speed laser ablation system enabled rapid imaging of the samples with a lateral resolution of 3 μm, whilst use of a prototype time-of-flight ICP-MS provided pseudo-simultaneous detection of the elements between mass 23 (Na) and mass 238 (U). Three forms of asbestos fibre (actinolite, amosite and crocidolite) were distinguished from a non-asbestos control ( wollastonite) based on their elemental profile, which demonstrated that LA-ICP-MS could be a viable technique for identification of asbestos fibres in clinical research samples.

Introduction

Malignant pleural mesothelioma (MPM) is an aggressive cancer whose pathogenesis is linked with occupational and environmental exposure to asbestos fibres.1 The prognosis of mesothelioma is poor due to the difficulty of making diagnosis in the early stage, the rapid progression, high invasiveness and lack of effective treatment.2 As the global incidence of mesothelioma is expected to increase over the next two decades,3 difficulties remain in relation to diagnosis, notably due to the lack of biomarkers and impeded identification and quantification of asbestos fibres in lung tissue.

In the UK, asbestos can be found in many industrial or residential buildings built or refurbished before the year 2000. The term “asbestos” is a general term and refers to silicate minerals that are composed of polyfilamentous bundles of long, flexible fibres with a small diameter.4 These can be easily inhaled when broken into fibrils. The mineral fibres are classified into two subgroups: (i) the amphiboles, a group of rod-like fibres including amosite, crocidolite, anthophyllite, actinolite and tremolite; and (ii) the serpentine group, consisting of chrysotile.4 Their potency of inducing asbestos-related diseases such as pleural plaques, asbestosis and MPM reportedly differs from one fibre to another due to differences in chemical structure, and therefore in solubility of the fibres in the lung.5

Detection of asbestos fibres using microscopic approaches is often inadequate. Sometimes fibres are undetectable, even in patients with heavy asbestos exposure.5 The routine quantification method uses phase contrast microscopy to count the mineral fibres that are longer than 5 μm, thinner than 3 μm, and have aspect ratios larger than 3 : 1.6 However, this method cannot detect shorter or thicker fibres and cannot distinguish between asbestos and other natural-occurring or man-made fibres, making some fibres considerably hard to detect and quantify.6 Another established method involves the use of transmission electron microscopy, following the International Standards Organisation (ISO) method for characterisation of airborne asbestos fibres.7 However, differentiation between fibre types in more complex matrices, such as biological tissue, remains challenging because fibre visibility and orientation may be hindered.

Asbestos fibres are composed of silicon and other elements, such as Mg, Ca and Fe, in varying proportions depending on fibre type. Consequently, there have been a small number of attempts to use elemental analysis, rather than optical properties, to distinguish between asbestos fibres. Pascolo et al., demonstrated the use of X-ray fluorescence (XRF) to analyse asbestos exposed lung tissue for identification of sub micrometre diameter fibres based upon Ca, Fe and Si elemental profiles.8 X-ray absorption near edge spectroscopy (XANES) has
also been used to complement XRF analysis. Transmission electron microscopy has been used in conjunction with energy dispersive X-ray (EDX) spectroscopy to identify fibres with a length and diameter as low as 0.1 μm, which optical microscopy could not detect. Based upon multielement analysis of Si, Fe, Mg and Al the specific form amosite was identified in...
ultrathin histological sections of lung tissue. In another analytical approach, secondary ion mass spectrometry was able to detect fibres as small as 1 × 5 μm that had been collected on a membrane, providing spatial information on Mg, Si and Ca.

Laser ablation inductively coupled plasma mass spectrometry (LA-ICP-MS) provides information on the spatially resolved elemental distribution within a sample. The technique has been utilised for a broad range of biological applications; from the imaging of endogenous elements, such as Ca, Zn, Cu, and Fe in human and animal tissues, to revealing the distribution of platinum anticancer drugs in tumours and tissue. In the last decade, a focus on ablation cell designs has led to dramatic improvements in transport of the ablated aerosol. Improved cell geometries have given rise to increases in absolute sensitivity and faster analysis times. The latest generation systems also offer improved spatial resolution, with lateral resolutions of a few μm now common, compared to typical resolutions of 5–100 μm achieved with older designs. The drive towards use of higher speed LA systems has contributed to a rise in the popularity of time-of-flight (TOF) ICP-MS systems for imaging applications.

The analysis was performed using FastScan mode where the magnet mass position is fixed and electrical deflectors change the measured mass position over −15% to +25% of the parked mass position in 70 μs per jump, with a dwell time of 250 μs per isotope. Using this data acquisition method a range between 23Na and 33Si was scanned every 20 ms for an average of 10 scans per jump, with a dwell time of 250 μs per isotope. The cultures were grown and harvested according to established cell culture protocols. All cell culture materials were supplied by Gibco, Life Technologies, CA, USA. All cells were confirmed to be negative for mycoplasma every three months using the MycoAlert™ Mycoplasma Detection Kit (LONZA, Switzerland).

Experimental

Cell culture
Human mesothelioma cells, MSTO-211H (ATCC, UK), were cultured and maintained at 37 °C in a humidified atmosphere with 5% CO2 in RPMI 1640 medium supplemented with 10% (v/v) heat-inactivated foetal bovine serum and 1% (v/v) penicillin/streptomycin. The cultures were grown and harvested according to established cell culture protocols. All cell culture materials were supplied by Gibco, Life Technologies, CA, USA. All cells were confirmed to be negative for mycoplasma every three months using the MycoAlert™ Mycoplasma Detection Kit (LONZA, Switzerland).

Asbestos fibres
Actinolite, amosite, crocidolite and wollastonite fibres were supplied by Health and Safety Laboratories (UK) and were kindly donated by Santia Asbestos Management Ltd. They were prepared in PBS solution to a stock concentration of 1 mg mL−1. For health and safety reasons, fibres were handled only in specially designed laminar flow hoods (Santia Asbestos Management Ltd. laboratories) until they were mixed into solutions. The fibre solutions were sterilised at 121 °C in an autoclave and stored at room temperature in PBS solutions. Prior to treatment, the solutions were passed through a 22-gauge needle 5 times for separation of the fibrils.

Cell preparation for LA-ICP-MS
For LA-ICP-MS analysis of 2D cultures, aliquots of 10 × 104 MSTO-211H cells per well were seeded in 6 well plates, with each well exposed to 3 μg mL−1 solutions of a different fibre type. After 24 h, cells were harvested and 200 μL of the cell suspensions (5 × 104 cells per mL) were cytopsunned at 100 g for 3 minutes onto plastic slides. For the 3D cell plugs, 1 × 107 cells were harvested, centrifuged and treated with 100 μL of the fibre solutions, as required, before embedding in hydroxypropyl methylcellulose (HPMC)/polyvinylpyrrolidone (PVP) media (3 : 1 ratio) and flash frozen in liquid nitrogen. Finally, the material was cryosectioned onto plastic slides to produce 12 μm sections, before introduction into the LA-ICP-MS instrument.

LA-ICP-TOF-MS
Preliminary experiments were performed using a NWR193 laser ablation system (Elemental Scientific Lasers, Bozeman, Montana), coupled to an AttoM ES sector-field ICP-MS (Nu Instruments, Wrexham, UK). For the initial experiments, the laser ablation system was equipped with the standard TwoVol3 chamber which provided a single laser shot signal width of approximately 400–600 ms at full width 10% maximum. A helium gas flow of 1–1.2 L min−1 was used through the ablation chamber, with an argon make-up flow of 0.6–0.8 L min−1 introduced through a t-piece downstream of the cell and approximately 5 cm from the connection to the ICP torch. A laser spot diameter of 3 μm, 10 Hz repetition frequency, 30 μm s−1 scan speed and 1 J cm−2 fluence were used during ablation. The analysis was performed using FastScan mode where the magnet mass position is fixed and electrical deflectors change the measured mass position over −15% to +25% of the parked mass position in 70 μs per jump, with a dwell time of 250 μs per isotope.

A NWR193 laser ablation system (Elemental Scientific Lasers, Bozeman, Montana) was coupled to a Vitesse time-of-flight ICP-MS (Nu Instruments, Wrexham, UK). The LA system incorporated a prototype TwoVol3 chamber configured with the imaging cup and was fitted with a Dual Concentric Injector (DCI) for fast washout. Helium gas flows of 1.2 L min−1 and 0.55 L min−1 were used through the cup and the outer chamber respectively, with an argon gas flow of 0.7 L min−1 introduced through the DCI. The laser ablation parameters were as follows: 3 μm spot diameter, 50 Hz repetition frequency, 150 μm s−1 scan speed, ~4 J cm−2 fluence. These parameters were chosen to ensure baseline separated signals and provide sampling of
Fig. 2  LA-ICP-MS images of the cell samples shown in panels (A–D) of Fig. 1, showing the sodium (A–D), calcium (E–H) and iron (I–L) distribution across the samples. A 400 × 200 μm area was sampled for each fibre type. The sodium signal shows the location of cells in the images, whilst the needle shaped fibres are clearly visible in the iron and/or calcium images. The circled region in the lower right panel is unlikely to be a wollastonite fibre because it lacks the characteristic silicon signal. Based on the elements found in this region, this is likely to be a piece of debris resulting from the sample preparation process.
Images of the sector-counts per second data was exported in CSV format for image shot, with no blurring between pixels. The resulting element data. The data collected before the detector, this delay was applied to the demarking values in the display as markers in the data. A delay was attributable ms displayed on a chart (ESI summed within the so as follows: groups of 40 consecutive 25.5 µs scans were initially summed within the software to give saved raw data points at ~1 ms displayed on a chart (ESI) with the correlating laser triggers displayed as markers in the data. A delay was attributable between the firing of the laser and the arrival of ions at the detector, this delay was applied to the demarking values in the data. The data collected before the first trigger signal was used to provide the gas background. The data between each delay corrected trigger was then averaged and background subtracted to provide a data point for each individual pixel. The ICP-MS software was able to demark the start and end of each line of the image by the difference in timing between laser triggers. Therefore, the pixels could be directly associated with each laser shot, with no blurring between pixels. The resulting element counts per second data was exported in CSV format for image visualisation and further processing.

Results and discussion

**Visualisation of fibres in biological matrices**

Identification of fibres within biological samples is a complex challenge, since these matrices often contain background levels of the metals of interest. Initial studies sought to distinguish whether a sector-field instrument could provide multi-element analysis of a mixture of actinolite and amosite, sufficient to distinguish them from a typical cellular background. A standard laser ablation system was used in order to provide peak widths that were considerably longer than the time taken to scan across the selected mass range. The cells contained detectable levels of endogenous 23Na, 24Mg and 31P, which facilitated registration of the LA-ICP-MS images to the microscopy data (see ESI†). Signals for 24Mg, 27Al and 28Si in the fibres were above the cellular background, confirming that LA-ICP-MS is a viable technique for visualisation of the fibres within complex matrices. However, the instrumentation used for the feasibility experiment may be unable to meet the demands of large-scale research studies into asbestos in the long-term. The sequential nature of magnetic scanning in sector-field instruments restricts their ability to perform multi-elemental analysis on transient signals. This can be somewhat alleviated through the use of FastScan mode, which uses the ion optic deflectors to provide faster electric scanning. However, this mode comes at the expense of a restricted mass range (up to 40% of the parked mass of the magnet). In the present study, this meant that several major components of asbestos, notably 40Ca and 56Fe, were not detected. The mode is also less suited to the very fast transient signals (~10 ms) from the latest generation LA systems. Clinical research requires high-throughput screening of tissues, which can be more readily achieved using these fast washout systems. The capability to differentiate between fibre subsets could therefore be enhanced by utilisation of a TOF spectrometer to provide pseudo-simultaneous detection of the full mass range.

**Differentiation of fibres**

Differentiation between fibre subsets was performed using a high-speed LA chamber, which produced peak widths of ~10 ms, along with a prototype time-of-flight mass spectrometer. Fig. 1 and 2 show the resulting data for several characteristic elements in actinolite, amosite, crocidolite and wollastonite fibres. From their chemical formulae (Table 1), it can be seen that all four of the fibre types contain silicon as a matrix element. Thus, as was expected, strong 28Si signals were observed for all four fibre types (Fig. 1 panels (I–L)), enabling rapid differentiation of the fibres from the cellular background. Magnesium was found to varying extents in all three of the asbestos fibres studied (Fig. 1 panels (E–G)), but only traces

| Table 1 | Nominal composition of the four fibre types studied |
|---------|---------------------------------------------------|
| Fibre name | Fibre type | Nominal chemical composition* |
| Actinolite | Asbestos | [Ca$_2$(Mg,Fe$^{2+}$)$_3$]Si$_2$O$_5$(OH)$_$_3$ |
| Amosite | Asbestos | [(Mg,Fe$^{2+}$)$_3$]Si$_2$O$_5$(OH)$_$_3 |
| Crocidolite | Asbestos | [Na$_x$Fe$^{3+}$Fe$^{2+}$Si$_2$O$_5$(OH)$_$_3$ |
| Wollastonite | Control | CaSiO$_3$ |

* Natural asbestos may deviate from this composition. For example, in some varieties of amphibole, the elements may be partially substituted by K, Al, Ti, Mn etc.
were observed in the wollastonite control (Fig. 1 panel H). The $^{24}\text{Mg}$ signals were strongest for amosite, where Mg accounts for >20% of the nominal composition of its cummingtonite form. During analysis, a collision–reaction cell was used to reduce nitrogen and argon related interferences, enabling $^{40}\text{Ca}$ to be measured without interference from $^{40}\text{Ar}$ and $^{56}\text{Fe}$ to be measured without polyatomic interference from $^{40}\text{Ar}^{16}\text{O}$ (see ESI†). This proved to be advantageous because comparison of iron and calcium signals was highly effective in enabling differentiation between certain fibre types. Iron is present at varying percentage levels in the three asbestos fibre types, but it is not a matrix component of the wollastonite control. The distribution pattern of iron (Fig. 2 panels (I–L)) therefore enabled rapid discrimination between the three asbestos fibres and the control sample. Furthermore, calcium is a major matrix component of wollastonite, representing ~35% of its nominal composition. Calcium is present to a much lesser extent in actinolite (~9% of its nominal composition), but it is not a matrix component of the other two fibre types studied. Thus, crocidolite and amosite (Fig. 2 panels F and G) can be easily distinguished from actinolite and wollastonite (Fig. 2 panels E and H) through visual comparison of the calcium images. The differences between amosite and crocidolite across Fig. 1 and 2 were more subtle. Sodium is a matrix component of crocidolite but not amosite, so analysis of sodium was also evaluated as a way to visually distinguish these two fibre types (Fig. 2 panels B and C). However, the very high level of sodium in biological matrices e.g. the dense cluster of cells in image B, along with the common use of PBS to prepare cell samples, may interfere with the interpretation of fibres. This serves to confirm that analysis of a single element would be insufficient to distinguish between these two fibre types. Principal component analysis was therefore employed as an additional approach to provide more in depth analysis of the differences across a range of elements. This revealed that the fibre types separate into 5 distinct groups.
(Fig. 3). The wollastonite control fibres were well separated from all three asbestos fibre types in the PCA plot. Actinolite and crocidolite were also well separated from each other. However, the amosite fibres formed two distinct groups. Amosite is found in two forms (grunerite and cummingtonite), thus it is possible that a mixture of the two forms were present in the fibres analysed in this study. Acquisition of a larger data set would be required to verify this hypothesis.

**Analysis of 3D cultures**

After demonstrating that fibres could be distinguished in monolayer cultures, a cell pellet containing amosite fibres was prepared, sectioned and provided to the analyst as a ‘blind’ sample. The densely packed nature of the 3D cultures better mimics the conditions under which patient tissue are analysed. Fibres embedded within the cells are invisible to low penetration depth techniques, such as microscopy. However, the 193 nm laser successfully penetrated through the tissue, revealing one large fibre of approximately 120 μm in length and some smaller fragments of below 50 μm in length (Fig. 4). Magnesium, iron and silicon were observed in high amounts in these images (panels (A), (B) and (D)) with an absence of calcium signal (panel (C)). Analysis of the full range of characteristic elements enabled the analyst to correctly identify the fibres as amosite, demonstrating the potential of the technique for identification of fibres in complex tissue matrices.

**Conclusions**

This is the first study to use LA-ICP-TOF-MS to provide pseudo-simultaneous multi-elemental analysis of fibres amongst a cellular background. The use of a low-dispersion ablation cell provided high-resolution, high-speed analysis, demonstrating that LA-ICP-MS has the potential to screen for the presence of fibre anomalies in clinical samples. It was apparent from the initial experiments that time-of-flight technology offers an advantage over sector-field technology for this application, particularly when used with high-speed laser ablation platforms. Use of the prototype ICP-TOF-MS system provided detection of the full range of significant elements within asbestos, which enabled an ‘unknown’ fibre, not visible under the microscope, to be correctly identified as amosite. This proof-of-concept data presents an alternative to existing approaches for fibre characterisation and offers the possibility of conducting complementary LA-ICP-MS measurements within the same specimen. For example, identification of asbestos fibres in lung tissue may complement existing LA-ICP-MS measurements of candidate metallo drugs in lung tumours.\(^\text{16,28}\) It is estimated that only 6 of approximately 400 fibre types present in nature are regulated under the generic name of asbestos.\(^\text{29}\) Many more potentially carcinogenic fibres are unregulated and continue to cause human exposure and mesothelioma. Future multi-elemental LA-ICP-MS analysis of previously unstudied, candidate fibres would help to elucidate their role in the development of mesothelioma and other asbestos-related lung diseases.

**Conflicts of interest**

P. S. and A. D. are employees of Nu Instruments.

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