Determinant factor for clearance of ceramic fibres from rat lungs

Hiroshi Yamato, Isamu Tanaka, Toshiaki Higashi, Masamitsu Kido

Abstract
Deposition, clearance, retention, and durability of inhaled particles from lungs are known to be important factors for induction of pulmonary fibrosis or lung cancer. To study the deposition and clearance of alumina silicate ceramic fibres from the lung, male Wistar rats were exposed to ceramic fibre particles with a mass median aerodynamic diameter (MMAD) of 4.2 μm, for six hours a day, five days a week for two weeks. The average exposure concentration was 20.7 (standard deviation (SD) 4.5) mg/m³. The rats were killed at one day, three months, and six months after the end of exposure, and the fibre numbers and dimensions were measured with a scanning electron microscope. No significant differences in number and geometric mean length of residual ceramic fibres in the lungs were found between the groups. The geometric mean diameter, however, decreased according to the duration of the clearance period. These findings suggest that solubility of fibre contributes most in the clearance of the inhaled ceramic fibres from the lungs.

Asbestos has been used in various industrial products because of its physical and chemical properties. It is known, however, that occupational exposure to various types of asbestos causes lung fibrosis (asbestosis), bronchial cancer, and pleural and peritoneal mesotheliomas, and that the use of asbestos fibres has decreased greatly in recent years. On the other hand, because of substitutes for asbestos, and new industrial materials, the global production of manmade mineral fibres (MMMFs) has increased from 4.5 million tonnes in 1973 to six million tonnes in 1985.1

The safety of MMMFs has been questioned because of their similarity in shape and chemical composition to asbestos. Some in vitro and in vivo studies have considered the biological effects of MMMFs and asbestos.2

Physical dimension, clearance, and durability of fibres in the lungs were reported as important factors in predicting the biological effect of inhaled fibres.3,4 To clarify the durability and size changes of manmade ceramic fibres we conducted an inhalation exposure study of ceramic fibres in rats, and determined the changes of numbers, lengths, and diameters of residual ceramic fibres in rat lungs after various clearance times from the end of exposure.

Materials and methods
TEST SUBSTANCES
Ceramic fibres (alumina silicate refractory fibres with a nominal diameter of 2–3 μm) used in this study were made by one of the largest Japanese manufacturers. The chemical composition of the fibres is 54% SiO₂, 46% Al₂O₃. Bulk ceramic fibres were disintegrated three times with an ultracentrifugal mill (Retch Co, Germany) at a speed of 15 000 rpm to obtain the materials for exposure.

EXPOSURE SYSTEM AND EXPOSURE CHAMBER
Figure 1 presents a schematic diagram of the experimental apparatus. The dust generator is the same as that previously reported.7,8 The special feature of the dust generator is the mixing of ceramic fibres with the fluidising particles (small glass beads, diameter 210–297 μm and density 2.5 g/cm³) before use. The uniform mixing of the fibres with fluidising particles is important to keep the generation rate of fibrous dust constant. In our experiment, the mixing ratio of the ceramic fibres to the fluidising particles was 0.003. The ceramic fibres were mixed with the fluidising particles by hand. Two kg of the fluidising particles were put into a glass bottle and 6 g of ceramic fibres were added. The bottle was shaken until the colour of the mixture was judged visually to be uniform. The mixing time was about 10 minutes. After mixing, the particles were fed into the hopper and smoothly transported via a continuous screw feeder into the fluidised bed. Dry air passed upwards
through the fluidised bed in which the fibrous particles were dispersed. The air velocity was high enough to transport the fibrous particles, but not the fluidising particles, into an exposure chamber (volume 0.48 m$^3$). The fluidising particles fell back into the bed and maintained fluidisation until discharged via the overflow pipe.

CONCENTRATION AND SIZE DISTRIBUTION IN THE EXPOSURE CHAMBER

The mass concentration of ceramic fibres was measured gravimetrically at daily intervals by the isokinetic suction of air through a glass filter. The size distribution of the aerosol in the exposure chamber was determined with an Andersen cascade impactor (air sampler, Andersen type, model AN-200, Shibata Scientific Technology Ltd, Japan). The mass median aerodynamic diameter (MMAD) and the geometric standard deviation (GSD) of the ceramic fibres in the chamber were 4.2 μm and 2.0 respectively. Because the sampler is unsuitable for obtaining the full size distribution of fibrous particles, it was only used in this study to check whether the test sample was respirable or not.

RATS

Male Wistar rats nine weeks old at the start of exposure were used. The rats were exposed to ceramic fibres daily for six hours a day, five days a week, for two weeks and killed at one day, three months, and six months after the end of the exposure. Table 1 summarises the experimental conditions.

Table 1  Experimental conditions for the study

<table>
<thead>
<tr>
<th>Animals</th>
<th>Male Wistar rats</th>
</tr>
</thead>
<tbody>
<tr>
<td>Test material</td>
<td>Alumina silicate ceramic fibre</td>
</tr>
<tr>
<td>MMAD</td>
<td>4.2 μm</td>
</tr>
<tr>
<td>GSD</td>
<td>2.0</td>
</tr>
<tr>
<td>Exposure:</td>
<td></td>
</tr>
<tr>
<td>Exposure period</td>
<td>Two weeks (6 h a day, five days a week)</td>
</tr>
<tr>
<td>Exposure concentration</td>
<td>20.7 (SD 4.5) mg/m$^3$</td>
</tr>
<tr>
<td>Clearance time</td>
<td>One day, three months, six months</td>
</tr>
</tbody>
</table>

*Mass median aerodynamic diameter.
†Geometric standard deviation.

DETERMINATION OF CERAMIC FIBRES IN RAT LUNGS

The rats were anaesthetised by intraperitoneal injection of pentobarbital and exsanguinated via the heart at the end of the appropriate experimental period. The lungs were subjected to low temperature ashing (150°C) in a plasma asher (LAT-25N Yanaco, Japan) in weighing bottles. The residues were washed in a small amount of 6N HCl. After gentle mixing, each sample was diluted to volume and decanted into a nucleopore filter apparatus containing a 0.45 μm pore size filter (Toyo Filter, Japan). The weighing bottles were rinsed with distilled water several times with all fluid being passed through the filter.

Numbers, length, and diameter of fibres were determined with a scanning electron microscope (SEM:Hitachi S-700) according to the reference methods for measuring airborne MMMFs proposed by the World Health Organisation. The filters were cut into squares (roughly 7 mm x 7 mm) after

Figure 1  Schematic diagram of exposure chamber.
Table 2  Total number, length, and diameter of inhaled ceramic fibres in rat lungs

<table>
<thead>
<tr>
<th>Clearance time</th>
<th>No of rats</th>
<th>Mean no of fibres (×10³) (SD)</th>
<th>GML* (GSD) (μm)</th>
<th>GMD† (GSD) (μm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>One day</td>
<td>4</td>
<td>4.9 (0.4)</td>
<td>5.4 (2.0)</td>
<td>0.88 (1.4)</td>
</tr>
<tr>
<td>Three months</td>
<td>3</td>
<td>5.6 (0.4)</td>
<td>6.0 (2.2)</td>
<td>0.78 (1.7)</td>
</tr>
<tr>
<td>Six months</td>
<td>3</td>
<td>3.3 (1.2)</td>
<td>5.5 (1.8)</td>
<td>0.56 (1.8)</td>
</tr>
</tbody>
</table>

*Geometric median length.
†Geometric standard deviation.

Results
Fibres longer than 50 μm were found in the lungs of each group. Recovered fibres had a log normal distribution. Table 2 shows the changes in numbers and dimensions of fibres in the lungs. No significant difference was found between each clearance period in terms of the residual numbers of inhaled ceramic fibres. There was also no significant difference in the GML between each clearance period. The GMD decreased, however, according to the duration of the clearance time (figs 2 and 3).

Discussion
Previous reports suggested the following clearance mechanisms of inhaled particles from lungs: mucociliary transport, translocation to lymph nodes by alveolar macrophages, and dissolution by extracellular fluid.

Bernstein et al examined the clearance of radioactive glass fibres that had a known size distribution (GMD 1.5 μm, length of either 5 or 60 μm) from the lungs of Fisher 344 rats. There was little difference in clearance between long (half life 35.0 days) and short (half life 38.5 days) fibres at 19 weeks after instillation, but the short fibres were apparently successfully phagocytised by alveolar macrophages and cleared to the lymph nodes whereas the long fibres were not. The long fibres also produced a granulomatous response in giant cells.

Morgan et al reported that 5 × 1.5 μm fibres were 90% cleared and 10 × 1.5 μm fibres were 80% cleared from lungs during the 12 months after administration. Neither 30 × 1.5 nor the 60 × 1.5 μm fibres were cleared to any significant extent over the 12 month period. They also showed that the count median diameter (CMD) of 10 × 1.5 μm fibres was reduced by 28%. The
30 × 1·5 and 60 × 1·5 μm fibres dissolved much more rapidly and the CMD of the residual populations after 18 months was less than half that of the original. Their study concluded that longer fibres were more soluble than shorter ones.

Abraham et al.1 conducted inhalation exposure experiments with chrysotile or crocidolite asbestos that had similar size distributions. After 90 days, 95% of the chrysotile fibre was cleared but no measurable clearance of crocidolite was seen. A decrease in mass was shown by x ray diffraction and could be calculated from changes in fibre dimensions. The chrysotile that was retained appeared to be the longer, thinner, fibres. This type of selective clearance was not found with crocidolite.

Morgan and Holmes4 found no significant change in the GMD of instilled rockwool fibres in rat lungs. Even after six months, it was apparent that the fibres became thinner at their ends than in the middle, and that slow dissolution occurred. Leineweber6 has shown that the solubility of MMMFs in vitro varied over a wide range and depended on their composition.

Timbrell10 compared the length distribution of anthophyllite fibres in the lungs of Finnish workers with that of fibres that had been airborne, and concluded that the critical length above which fibres could not be cleared from the lung was about 17 μm. The studies mentioned indicate that many mechanisms can be used to eliminate inhaled fibres.

The present study showed that the critical factor for clearance of ceramic fibres is the decrease in fibre diameter, and that it depends on the solubility of the fibre. According to the Stanton-Pott hypothesis, the fibres longer than 8 μm and thinner than 0·25 μm have high carcinogenicity. Our study suggested that even the original diameters of inhaled ceramic fibres are thicker than Stanton fibres, and could be the size of Stanton fibres in the process of clearance time. The relation between the findings in the present study and the biological effects of MMMFs will be considered in future studies.

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