The Effects of Carbon Fibre and Carbon Fibre Composite Dusts on Bronchoalveolar Lavage Component of Rats

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—Carbon fibre (CF) and carbon fibre composite (CFC) as new materials have increasing industrial application. The People’s Republic of China now manufactures CF and CFC. This paper predicted their potential toxicity to human using bronchoalveolar lavage technique on the basis of comparisons with positive reference materials (quartz and chrysotile) and negative reference materials (titanium dioxide and alumina (SAFFIL) fibre). All rats dosed with dust showed some increase in lung weight relative to the saline control, though the only significant differences were seen between the rats dosed with quartz or chrysotile and those dosed with saline. From the morphological observation of lavage cells, a benign reaction of macrophages to CF and CFC was observed, whereas a series of changes in macrophages was involved in rats dosed with quartz and chrysotile. CF and CFC did not induce a significant increase in the total cell count or percentages of neutrophils and lymphocytes in bronchoalveolar lavage. The two materials tested had much lower toxicity than that of quartz and chrysotile, and were comparable with the effect induced by titanium dioxide and SAFFIL fibre which had minimal toxicity. The present work provides a scientific basis for the setting of occupational health standards for carbon fibre and carbon fibre composite dust in the workplace.

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Key words: Carbon fibre dust, Carbon fibre composite, Bronchoalveolar lavage, Cell type

Carbon fibre composite (CFC) is a new material which, by virtue of being chemically inert and having additional advantages of high strength, light weight and thermal resistance, is finding increasing industrial application. The People’s Republic of China now manufactures carbon fibre (CF) and CFC. During the manufacturing and processing of both CF and CFC, because dust may be released into the workplace air, there is an understandable concern that these processes may present an occupational risk. Because of the short history of production of both CF and CFC, epidemiological investigation could provide only limited confidence that there were no potential adverse effects. Bronchoalveolar lavage is a well established technique for predicting the toxicity of various dusts in animals. We report a study in which this technique is used to examine the in vivo cytotoxicity of CF and CFC dust samples as an aid to predicting their potential human hazard.

Materials and Methods

Rats

Twenty-one male and twenty-one female Wistar strain rats (weight range 200–300 g) were randomly assigned to seven test groups (6 in each group; 3 of each sex per group).

Test samples

The CF sample used was obtained from a factory in Ji Lin City of the People’s Republic of China producing PAN-based carbon fibre. A bundle of carbon fibres (3 cm in length) was packed into a container filled with distilled water. The whole was then frozen at -80°C. The ice embedded CF blocks were cut along the long axis of the fibres with a microtome. The sections were allowed to thaw and the water was evaporated to release residual CF. These were then sieved through a 65 µm (250 mesh) sieve. Microscopic analysis showed that the product had fibre diameters in the range of 6–8 µm with...
a median length of 37.5 $\mu$m. 97% of the fibres were ≤200 $\mu$m long. CFC was obtained from an aircraft factory in Harbin City of the People’s Republic of China. A coarse sample was produced by drilling the composite. This sample was then further reduced by grinding in an electrically operated agate pestle and mortar to produce a respirable fraction. 93% of the particles in the final product were <5 $\mu$m in diameter.

Control samples
Quartz and chrysotile were used as positive controls and TiO$_2$ and SAFFIL fibre as negative controls. A saline control was also included.

A sample of quartz was obtained from the Institute of Occupational Medicine, Chinese Academy of Preventive Medicine. The sample was analysed as 98% silica and 99% of particles were <5 $\mu$m in diameter. A sample of the UICC chrysotile A standard international reference sample was supplied by Dr. J.M.G. Davis, Institute of Occupational Medicine, Edinburgh, UK. Titanium dioxide is a commercially supplied commodity reagent (analytical grade) from Beijing Chemical Plant. The sample was ground to a fine powder in an electrically operated agate pestle and mortar. 96% of the particles in the ground sample were <5 $\mu$m in diameter. SAFFIL fibres in an inorganic refractory fibre consisted of 95% alumina with 4.3% silica and a few trace impurities. A ground sample was supplied by Dr. G.H. Pigott, Central Toxicology Laboratory, Imperial Chemical Industries plc, UK. The fibres supplied have a median diameter of 3.2 $\mu$m with 98% <5 $\mu$m in diameter. The fibres have a median length of 62 $\mu$m with 95% <211 $\mu$m.

Preparation and administration of dust suspensions
Saline was the vehicle for all dusts. Suspensions were prepared by manual shaking. All test samples, except chrysotile, were prepared at 50 mg/ml. Chrysotile was prepared at 12.5 mg/ml because of the mortality associated with the higher dose. The suspensions were sterilised by autoclaving at 120°C. Just prior to administration penicillin was added at a concentration of 400 iu/ml. The samples were administered by intratracheal injection under light anaesthesia. The volume of suspensions, including saline, administered was 1 ml for each animal, which contained 50 mg different samples (12.5 mg for chrysotile sample), respectively, for the groups.

Brochoalveolar lavage
One month after administration all the animals were anaesthetised with 2% pentobarbital and sacrificed by exsanguination via the abdominal aorta. The thorax was opened, the left primary bronchus ligatured and the right lung lavaged four times with approximately 2.5 ml physiological saline. External massage was used to ensure recovery of cells in the airways; 10 ml of lavage fluid was collected from each animal.

Post mortem examinations
For observation, lavage fluid cells were recovered by centrifugation at 2,000 rpm for 10 min. Cell pellets were resuspended in physiological saline. Total cell counts were performed in a haemocytometer. Aliquots were also sedimented in a cytocentrifuge and stained with Giemsa for differential cell counts.

The left lung of each rat was removed and weighed.

Statistical treatment of results
Intergroup comparisons were made by analysis of variance with SYSTAT software, comparisons were by Duncan’s test with a P value <0.05 considered significant.

Results
Figure 1 shows the intergroup comparison of the wet weight of the left lungs. All rats dosed with dust showed some increase in lung weight relative to the control, though the only significant differences were between the rats treated with quartz or chrysotile and those treated with saline. As similar results were observed when the wet lung weight was expressed as a percentage of the rat body weight, this effect does not reflect differences in the size of the animals concerned.

An increase in the size of particle containing cells was evident under the optical microscope for rats treated with carbon fibre or SAFFIL fibres, but the cell membranes remained intact. Some fibres were apparently phagocytosed by 2 or more macrophages, with the longer fibres sometimes giving the appearance of a string of beads. This was more remarkable in the rats treated with CF. In rats treated with CFC and TiO$_2$ black dust, particles within macrophages were seen in the bronchoalveolar lavage fluid. In some instances the macrophage nucleus was not easy to identify due to the large number of particles in the cell. Even in such particle laden cells the outer membrane remained visually intact.

In rats treated with quartz macrophages some had one or more of a series of morphological changes such as enlarged size, poor staining of the cytoplasm and nucleus, cytoplasmic vacuolation and even apparent enucleation. In addition cell debris and fragments could be seen in the lavage fluid indicating cellular disruption. In rats treated with chrysotile, similar changes were observed but to a lesser extent.

Figure 2 shows the results of total cell counts. The maximum cell count was obtained from rats treated with quartz, followed by that in rats treated with chrysotile. Cell counts in the other groups were much lower. Rats treated with TiO$_2$ or SAFFIL had counts slightly lower, but none of these changes attained statistical significance.
Differential cell counts are shown in Fig. 3. In most of test samples macrophages were the predominant cell type, accounting for 92.6%–98.8% of cells counted in all groups except those dosed with quartz or chrysotile. Statistical analysis showed the proportion of macrophages was significantly reduced in rats treated with quartz (53%) and chrysotile (76.3%) compared with those treated with other test samples.

The percentage of neutrophils reflected the differences seen in macrophages in the quartz group (41.75%) and the chrysotile group (22.0%), both significantly higher than controls. There were no other significant intergroup differences. The percentage of lymphocytes (5.25%) in the quartz group was significantly above controls. The only other group to exceed 1% was chrysotile (1.5%), though the difference was not statistically significant. Analysis of the absolute values for each of the cell types examined confirmed the increase in those treated with quartz. In rats treated with chrysotile the absolute values for each cell type were also increased compared to the other groups (excluding quartz) but the changes did not attain statistical significance. There were no other significant intergroup differences.
Discussion

The two samples tested in this study have different physical properties. CF was in fibrous form whereas the CFC sample was a non fibrous particulate. For this reason different control materials were selected for more relevant comparisons. Quartz and chrysotile are the two most widely accepted standard references as positive controls in studies of cytotoxicity and fibrogenicity. TiO₂ is also generally accepted as the standard inert particulate showing little or no cytotoxicity and fibrogenicity in a number of studies²). SAFFIL fibres were designed to minimise biological activity and their inert nature has been confirmed in a number of experiments both in vivo and in vitro³).

On the basis of the lung wet weight only the animals treated with quartz and chrysotile could be considered to show a significant reaction to dust. The slight increase in wet lung weight in the animals treated with CF, CFC, TiO₂ and SAFFIL over those treated with saline alone is too small to be considered significant, but it should be noted that the wet lung weight in rats treated with either CF or CFC was slightly below that recorded for the negative control samples.

From the morphological observation of lavage cells it was evident that the dusts persisted in the lung for at least one month after injection, but CF and CFC dusts were apparently phagocytosed by macrophages and the CF or CFC laden macrophages were morphologically intact and showed no evidence of an adverse reaction to the dust content. Similar changes were seen with TiO₂ dust and SAFFIL fibres and are considered typical of biologically inert dust. By contrast both quartz and chrysotile resulted in obvious morphological changes in macrophages and by virtue of the cellular debris seen could be considered to be cytotoxic to the phagocytic cells.

In previous studies total and differential counts of lavage cells have been used as indices to evaluate the toxicity of dusts to the lung. An increase in the total number of cells in lavage fluid has been observed in animals treated with various insoluble dusts⁴). From our observations the total numbers of cells in rats treated with CF or CFC dust were significantly smaller than those seen with quartz or chrysotile and comparable with the negative controls used. This is considered to confirm the very weak nature of the biological response to CF and CFC dust.

The only test samples to show significant changes in differential cell counts were quartz and chrysotile. In rats treated with either of these dusts, the percentage of macrophages was significantly lower than that for the other test samples and control. This mainly reflected an increase in the number of neutrophils in the lavage fluid. The presence of neutrophils in lavage fluid is believed to be a marker of pulmonary fibrosis and is considered a
Sensitive indicator of the acute effects of dust on the lung. It is suggested that this influx into the alveolar spaces reflects cell injury. In this context the lack of any significant increase in neutrophil levels in rats treated with CF or CFC is further evidence of the inert nature of these materials.

Significant increases in the number of lymphocytes occurred only in rats treated with quartz. There was no increase in the number of lymphocytes in rats treated with CF or CFC. This observation is in accord with that of Martin et al. in which five carbon fibre composite samples and one glass fibre composite sample were compared to quartz by using the bronchoalveolar lavage technique. These authors found that three of these five composite samples showed consistently low toxicity but two provoked an increase in alveolar macrophages and caused significant accumulation of neutrophils in rat lungs, but they were all much less toxic than quartz. This is in accord with the inhalation test performed by Owen et al. who exposed rats to polyacrylnitrile based carbon fibres (20 mg/m$^3$) for 16 wk (6 h/day, 5 d/wk). The only histological finding in the lungs was the accumulation of alveolar macrophages. These findings agree well with observations in the short term tests, though the size of the fibres tested suggests that respirability in the rat (and therefore the lung burden) would be low.

The present investigation shows that in vivo toxicity of the CF and CFC dust samples tested was comparable to materials of known low biological activity and distinct from that caused by quartz or chrysotile. It should be noted that no clear difference in toxicity was demonstrated between CF and CFC. Whereas the fibre is almost pure carbon, the composite uses a binding agent to trap the carbon fibre filaments, so that the chemical composition of CFC dust is more complicated. The fact that the biological effects of the two dusts showed no significant difference indicates that in this instance the binding agent did not influence the toxicity of the composite.

This investigation provides some experimental evidence for the setting of occupational exposure limits for CF and CFC dusts. Based on the in vivo cytotoxicity data available in this study, the maximum allowable concentrations of carbon fibre and carbon fibre composite dusts in the air of the workplace would be similar to that for TiO$_2$ dust.

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References