Respiratory symptoms, immunological responses, and aeroallergen concentrations at a sawmill


Abstract
After identification of a case of extrinsic allergic alveolitis due to exposure to wood dust at a sawmill, all employees at the sawmill where he worked were studied with an occupational, environmental, and symptom questionnaire, spirometry, skin prick tests, and serum specific IgG measurements. Ninety-five of current and 14 of 17 ex-sawmill workers were studied. As a basis for comparison, a group of 58 workers from a nearby light engineering factory were also studied. Few women (6) were employed and they were excluded from the analysis. Workers at the sawmill were stratified into high and low exposure groups depending on their place of work. This division was supported both by their subjective assessment of the dustiness of their environment and the results of personal dust samples. There were no significant differences between the three groups in age, height, smoking habits, exposure to other causes of extrinsic allergic alveolitis, forced expiratory volume in one second, forced vital capacity, atopic state, or cutaneous reactivity to moulds. In the high exposure group the prevalence of work-related cough and nasal and eye symptoms was higher than in the low exposure and comparison groups. The prevalence of work-related wheeze was similar in both the high exposure and comparison groups, but was lower in the low exposure group. The prevalences of chronic bronchitis and symptomatic bronchial hyperreactivity were similar in the high and low exposure groups but were lower in the comparison group. Serum concentrations of specific IgG against extracts of sawdust and *Trichoderma koningii* were significantly higher in the high exposure group than in the other two groups. The prevalence of symptoms suggestive of extrinsic allergic alveolitis was 4-4% in the high exposure group, greater than in the low exposure group (0%), and the comparison group (1-9%). In conclusion extrinsic allergic alveolitis probably occurs in British sawmills, and among the exposed population its prevalence may be as high as that reported in Sweden. The allergen responsible is likely to be from mould growing on the wood and may be from *Trichoderma koningii*.

Materials and methods
THE SAWMILL
The sawmill, which opened in 1980, cuts timber from all parts of the United Kingdom. The mill site comprises five main buildings: the office building; the green mill with attached sorting shed, saw shop, and maintenance shop; the autostacker; the kiln; and the dry mill. The types of timber cut have been described previously.1

About 3300 m³ of timber was cut in the green mill each week. Extraction of sawdust from beneath the band saws, which were enclosed, relied on conveyor belts. There was no atmospheric dust extraction system. Large pieces of wood were cleaned from beneath the machinery each evening, but dust was not regularly cleaned from inaccessible parts of the machines. The whole of the green mill was blown down with compressed air every two to three months.

About 20% of the cut wood was machined in the dry mill where some was planed or mouldings cut and the rest cut into shorter lengths. Sawdust from these blades was extracted by overhead vents. Less than 5% of the green mill's output passed through a kiln before reaching the dry mill. The kiln was manually loaded, but was otherwise automatically operated to maintain a temperature of about 60°C for three to four days. Humid air was vented from the kiln directly into the atmosphere.

About 5% of the cut planks were treated with a fungicide in an automatic process with manual loading and unloading of the treatment tank, which was run overnight. The fungicide was a mixture of sodium dichromate (44-6% w/w), copper sulphate (34-6% w/w), and arsenic pentoxide (19-6% w/w).
Saw blades were replaced every three to four months and were resharpened in the sawshop, where new blades were also made. Most band saw blades were made from steel, but some circular saw blades had tungsten carbide tips.

THE WORKFORCE
The sawmill employed a total of 100 people between the buildings and outdoors. Two shifts (0600 to 1400 and 1400 to 2200) were operated for most of the workforce. Office staff worked from 0900 to 1700 and a few employees worked a night shift from 2200 to 0600. During each of the day shifts most of the men worked in the green and dry mills, two men worked in the autotacker, and the remainder were outside, either driving vehicles or working in the log yard. Those inside the mills were subject to the highest levels of dust exposure, but although those working outside seemed less exposed to fresh sawdust, they encountered stored logs, sawdust, and wood chips, as well as mulched bark and vapore from the kiln. The maintenance engineers visited all parts of the sawmill and were exposed to old dust on and around the machines.

EX-WORKERS
The sawmill provided a list of all employees other than those employed as casual labour for periods of only a few weeks. During the nine years of operation, 17 workers had left the sawmill, which was by far the largest employer in the area. All of these workers were sent questionnaires at their last known addresses and were invited to attend during the survey for the remaining tests. Those who did not attend were visited at home where practicable.

COMPARISON GROUP
Workers at a light engineering factory producing large industrial heaters and four wheel drive vehicles were asked to volunteer as a comparison group. The factory employed 110 people and was situated in a similar rural environment to the sawmill. It comprised one large shop floor and attached offices. The workforce consisted of sheet metal workers, lathe operators, welders, paint sprayers (who used two pack polyurethane paint), tool makers, assemblers, warehousemen, and storemen, as well as office staff. Each of these departments was asked to find a number of volunteers to produce a sample with about the same distribution of occupations as at the sawmill.

EXPERIMENTAL DESIGN
A cross sectional study of the entire workforce at the sawmill and the comparison group was made during one week in October 1989. All subjects gave informed consent before taking part in the survey and the study had been approved by Brompton Hospital ethics committee. The subjects completed an occupational, environmental, and smoking questionnaire under supervision. A symptom questionnaire was presented by a doctor blinded to the answers to the first questionnaire, and subjects then had lung function and skin prick tests. Blood samples were also taken.

QUESTIONNAIRES
The questionnaires contained sections on systemic, respiratory, nasal and conjunctival symptoms, smoking, and occupational history, as well as questions about exposure to other known causes of extrinsic allergic alveolitis (mouldy hay or grain, mushroom growing, and birds). Questions on cough, phlegm, breathlessness, wheezing, and smoking were taken from the Medical Research Council's questionnaire on respiratory symptoms. Questions on nasal, conjunctival, and flu-like symptoms, weight loss, and fevers were also included. All questions on symptoms asked about the previous 12 months (or the last 12 months of employment at the sawmill). After each of the questions about symptoms were questions about the date of onset of symptoms, whether they improved at weekends or on holiday, and whether there was any seasonal variation. Further questions on wheezing during the four weeks before taking part were taken from a questionnaire that has been validated against histamine bronchial reactivity, and a positive response to one or more of these questions was used to define symptomatic bronchial hyperreactivity.

Symptoms were called work related if they improved at weekends, or on holiday and if they had started after the subject had joined the sawmill or factory. For the purposes of this survey extrinsic allergic alveolitis was defined as the combination of one or more work related respiratory symptoms (cough or breathlessness) and a constitutional symptom (work related flu like symptoms or unintentional weight loss).

LUNG FUNCTION
A Vitalograph dry wedge spirometer was used to record forced expiratory manoeuvres from each subject. Subjects were allowed to perform a maximum of six manoeuvres and readings of forced expiratory volume in one second (FEV₁) and forced vital capacity (FVC) were taken. Mean values were calculated from the best two technically satisfactory manoeuvres. From the data for men from both factories multiple regression analysis was performed with age, height, and smoking state and using the residuals for each worker from this line a normalised FEV₁ and FVC were calculated with the mean age and height as standards.

SKIN PRICK TESTS
Skin prick tests were performed with extracts of mixed grass pollen, house dust mite (Dermatophagoides pteronyssinus), cat fur, Alternaria alternata, Aspergillus fumigatus, Cladosporium herbarum, Serpula (Merulius) lacrymans (Bencard), and Mucor mucido, Aureobasidium (Pullularia) pullulans,
Penicillium notatum, and Rhizopus nigricans (Allergopharma). A test was called positive if a weal was seen of at least 2 mm diameter at 10 minutes after testing after subtraction of any weal produced by a control solution. A subject was regarded as atopic if one or more positive reactions were elicited by mixed grass pollen, house dust mite, or cat fur.

AIR SAMPLING
Airborne dust was sampled with personal dust samplers in the breathing zone of workers who were divided into different exposure groups based on the zoning method of Corn and Esmon. Concentrations of total airborne dust were measured with seven hole filter holders housing preweighed polytetrafluoroethylene (PTFE) filters (1-2 μm pore size, 25 mm diameter, Sartorius Instruments Ltd, GB-Belmont, Surrey) and Casella AFC 123 pumps (Casella London Ltd) at a flow rate of 2 l/min. Sampling times ranged from six to eight hours covering a work shift. After sampling, the filters were immediately removed and reweighed on a Sartorius R180D balance so that the dust concentration could be calculated.

DETERMINATION OF AIRBORNE FUNGI
Air was sampled for fungal spores with both personal and static area air samplers. To assess spore concentrations at different sites around the sawmill, air was sampled with an Andersen sampler. The sampler was held about 1-5 m above floor level and was operated at a flow rate of 25 l/min. At each site successive sets of six petri dishes containing 2% malt extract agar, penicillin (20 IU/ml medium), and streptomycin (40 IU/ml), or dichloran rose bengal chloramphenicol agar were loaded into the Andersen sampler and exposed for 15–60 s. One set of each medium was incubated at 25°C and another at 40°C. All samples were replicated three times and counts for each plate were adjusted for multiple deposition of viable particles beneath jets from Andersen’s tables. Isolates were identified by the appearance of colonies and microscopic morphology from standard texts.

To assess the exposure of individual workers airborne particle samples collected in the breathing zones of workers as described earlier were analysed for the presence of fungal spores. Spores were resuspended by shaking the filter on a whirlimixer (Fisons, Loughborough, Leics) with 5 ml of wash fluid (bacteriological peptone, (1 g); Tween 80, (0-5 g); inositol, (20 g); distilled water, (1000 ml). The suspension was then used to prepare logarithmic dilution series in quarter-strength Ringer’s solution and 0-1 ml quantities of suitable dilutions were spread over the agar surface of prepoured petri dishes. Six dishes of 2% malt extract agar or dichloran rose bengal chloramphenicol agar were inoculated with each dilution and three of each were incubated at 25°C and three at 40°C. Colonies were identified as for Andersen sampler isolates.

SPECIFIC IgG ASSAY
Specific IgG of subtypes IgG1, IgG2, and IgG4 against wood allergens were assayed with a solid phase radioimmunoassay based on the method described by Hay et al. with protein A as the tracer.

PREPARATION OF ALLERGENS
Samples of sawdust were collected from the floor around the green mill bandsaws and mulched bark was collected from the storage pile. Extracts were made with Coca’s solution. These were dialysed and lyophilised before reconstitution in 50% aqueous glycerol at a final concentration of 500 μg/ml. Filtrates from broth cultures of single species of moulds isolated from air samples were also dialysed and lyophilised before similar reconstitution.

SERUM SAMPLES
Venous blood (10 ml) was collected from subjects and the serum was stored at −20°C until assayed.

RADIOIMMUNOASSAY
The conditions of the radioimmunoassay were optimised with serum known to contain specific IgG before the assay of serum samples from the survey. Wells on Immulon microtitre plates (Dynatech Laboratories Inc, Chantilly, Virginia, USA) were coated by incubating 100 μl of a 1:1000 dilution of antigen in 200 mM sodium carbonate/bicarbonate buffer pH 9-2 for 20 hours in a humid chamber at room temperature. The plates were washed, 50 μl duplicate samples of 1 in 500 dilutions of serum were added to the wells, and the plates were again incubated for 20 hours in a humid chamber at room temperature. After further washing 50 μl of a 5 ng/ml solution of 125I labelled protein A (Amersham International Plc, Amersham, Bucks) was added to each well. The plates were incubated in a humid chamber at room temperature for five hours before a final wash. The amount of protein A bound to the wells was measured in a gamma counter (Packard Cobra 5 Channel, Canberra Packard, Pangbourne, Berks). Reference tubes to which 50 μl of 5 ng/ml 125I protein A had been added were also counted and the results of the assay were expressed as the percentage of total counts of 125I protein A bound to the test wells. The non-specific binding of protein A in tubes coated with the relevant antigen, which had not been incubated with serum, or which had been incubated with fetal serum obtained from placental cord blood was negligible. The non-specific adsorption of test serum antibodies on to the well surface was also negligible under these conditions and blocking of the plates did not seem to be necessary.

STATISTICAL TESTS
Data from the questionnaires, spirometric measurements, and skin prick test results were coded and entered into a computer...
Results

Ninety five out of the 100 current sawmill workers completed questionnaires and performed spirometry; 88 had skin prick tests and gave a blood sample. Fourteen of the 17 ex-workers were studied and all completed questionnaires, performed spirometry, had skin prick tests, and gave a blood sample. At the comparison factory 58 volunteers were studied and with the exception of two who declined to give blood, all completed all parts of the study.

There were six women among the current employees at the sawmill and six women among the comparison volunteers. As their number was small and they were exclusively involved in secretarial work in offices away from the sawmills and shop floors, the data relating to these women have not been included in the analysis. The data from the current and ex-workers have been combined and analysed as one group. Table 1 shows the number of men employed in different parts of the sawmill.

<table>
<thead>
<tr>
<th>Group</th>
<th>No of employees</th>
<th>Median subjective dustiness (range)</th>
<th>Mean (SEM)</th>
<th>95% CI</th>
<th>Dust concentration (mg/m³)</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sawmill</td>
<td>58</td>
<td>45</td>
<td>36</td>
<td>35</td>
<td>36</td>
<td>35</td>
</tr>
<tr>
<td>Low dust</td>
<td>78</td>
<td>0.024 (0.011-0.051)</td>
<td>0.024 (0.011-0.051)</td>
<td>0.024 (0.011-0.051)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>High dust</td>
<td>43</td>
<td>0.024 (0.011-0.051)</td>
<td>0.024 (0.011-0.051)</td>
<td>0.024 (0.011-0.051)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Comparison</td>
<td>52</td>
<td>0.024 (0.011-0.051)</td>
<td>0.024 (0.011-0.051)</td>
<td>0.024 (0.011-0.051)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 2 Median subjective dustiness and percentage reporting work related symptoms, for men in low and high dust jobs, and at the comparison factory

- No: 58
- High dust: 45
- Comparison factor: 52

Table 3 Demographic data for men in the three groups

<table>
<thead>
<tr>
<th>Group</th>
<th>Sawmill</th>
<th>Low dust</th>
<th>High dust</th>
<th>Comparison</th>
</tr>
</thead>
<tbody>
<tr>
<td>No</td>
<td>45</td>
<td>58</td>
<td>52</td>
<td></td>
</tr>
<tr>
<td>Mean age (y)</td>
<td>37.0</td>
<td>35.9</td>
<td>33.6</td>
<td></td>
</tr>
<tr>
<td>(range)</td>
<td>(20-42)</td>
<td>(16-44)</td>
<td>(20-64)</td>
<td></td>
</tr>
<tr>
<td>Mean height (cm)</td>
<td>174-8(1-1)</td>
<td>173-8(0-9)</td>
<td>174-60(9)</td>
<td></td>
</tr>
</tbody>
</table>

Previous employment:
- Farm work: 26.6%
- Mushroom culture: 0%
- Other sawmills: 24.4%
- Pet birds: 13.3%
- Domestic mould: 13.3%
- Ex-smokers: 66.7%
- Mean (SEM) cigarettes/day: 16-2(2-5) 15-3(2-1) 14-1(9)
- Mean (SEM) pack-years: 17-0(3-7) 13-7(3-1) 10-0(1-7)
There were no significant differences in the prevalence of symptomatic bronchial hyper-reactivity (table 2), or in the normalised FEV₁ or FVC (table 4). There was also no significant difference in the prevalence of atopy (those with one or more positive reactions to skin prick tests with mixed grass pollen, house dust mite, or cat fur) or cutaneous reactivity to mould extracts among high and low dust workers (table 5).

The results of the IgG binding studies of serum samples from workers against extracts of sawdust from the centre of the logs, mulched bark, Trichoderma koningii, Penicillium glabrum (frequensans), and Aspergillus fumigatus showed that the distribution of the data was skewed. Therefore, for statistical analysis the data were subjected to a log transformation to give a normal distribution. Table 6 shows the mean log, % binding for workers in the high and low exposure groups at the sawmill. All those with symptomatic extrinsic allergic alveolitis had high concentrations of antibody against fungi. Log, % binding against sawdust and Trichoderma koningii was significantly greater in men in the high exposure group than in the low exposure group. Among all sawmill workers log, % binding against sawdust, mulched bark, Trichoderma koningii, and Aspergillus fumigatus was significantly less among smokers than non-smokers (table 7).

COMPARISON OF HIGH EXPOSURE GROUP AT THE SAWMILL AND COMPARISON FACTORY

There were no significant differences between the mean ages, height, smoking habits, or exposure to other causes of extrinsic allergic alveolitis in the men in the high exposure group and men at the comparison factory (table 3). There was also no significant difference between the mean % predicted FEV₁ and FVC in the two groups (table 4). Unlike men at the sawmill, no men at the comparison factory had seen mould there.

Significantly more men in the high exposure group at the sawmill reported work related cough, breathlessness, nasal, ocular, and flu-like symptoms (table 2). There was no significant difference in the prevalence of work related wheeze. There was a higher prevalence of chronic bronchitis and symptomatic bronchial hyper-reactivity in the high exposure group than at the comparison factory, but there was no significant difference between the prevalence of unintentional weight loss. There were no significant differences between the prevalence of atopy or cutaneous reactivity to mould extracts in the high exposure group and at the comparison factory (table 5). A higher proportion of the men in the high exposure group had symptoms suggestive of extrinsic allergic alveolitis than at the comparison factory (4.4% v 1.9%).

The log, % binding values against sawdust and Trichoderma koningii were significantly higher in the high exposure group at the sawmill than in the low exposure and comparison groups (table 6). The log, % binding against Trichoderma koningii was high in all those men at the sawmill with symptoms suggestive of extrinsic allergic alveolitis.

FUNGAL SPORE COUNTS

Concentrations of fungal spores collected by personal dust sampling varied considerably between the different occupational groups (table 8). Concentrations similar to or less than those measured in the comparison factory were found in the sawmill offices, outside, autostacker, and sawshop samples. Significantly larger concentrations were found in the dry and green mills. Maintenance workers were also heavily exposed (table 8). Penicillium species were the most numerous fungi in nearly all samples taken at the sawmill but their numbers were small in the sawmill offices and in the comparison factory. Aspergillus fumigatus was also common at all sites in the sawmill but less common in the offices, the autostacker, and in the comparison factory. Trichoderma, Stilbella and Talaromyces species were numerous at some sites, although numbers of Stilbella were very variable with one sample yielding very large numbers of colonies accounting for almost all isolates in a group.

<table>
<thead>
<tr>
<th>Sawmill</th>
<th>High exposure</th>
<th>Low exposure</th>
<th>Comparison factory</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean normalised FEV₁</td>
<td>4.06(0.09)</td>
<td>4.11(0.07)</td>
<td>4.13(0.07)</td>
</tr>
<tr>
<td>Mean normalised FVC</td>
<td>4.90(0.11)</td>
<td>5.02(0.08)</td>
<td>5.01(0.08)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Antigen</th>
<th>Sawdust</th>
<th>Mulched bark</th>
<th>Trichoderma koningii</th>
<th>Penicillium glabrum</th>
<th>Aspergillus fumigatus</th>
</tr>
</thead>
<tbody>
<tr>
<td>log, % binding</td>
<td>2.49(0.18)</td>
<td>1.79(0.12)</td>
<td>1.71(0.06)</td>
<td>2.41(0.07)</td>
<td>2.51(0.06)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Antigen</th>
<th>Non-smokers</th>
<th>Smokers</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sawdust</td>
<td>2.91(0.23)</td>
<td>1.79(0.10)</td>
</tr>
<tr>
<td>Mulched bark</td>
<td>3.52(0.09)</td>
<td>3.22(0.05)</td>
</tr>
<tr>
<td>Trichoderma koningii</td>
<td>3.27(0.13)</td>
<td>2.70(0.08)</td>
</tr>
<tr>
<td>Penicillium glabrum</td>
<td>2.07(0.09)</td>
<td>1.90(0.03)</td>
</tr>
<tr>
<td>Aspergillus fumigatus</td>
<td>2.41(0.07)</td>
<td>2.33(0.04)</td>
</tr>
</tbody>
</table>
Table 8 Mean concentrations of different fungi and mean total concentration of fungi (colony forming units (cfu/m³) in the breathing zones of workers in different parts of the sawmill and in the control factory

<table>
<thead>
<tr>
<th></th>
<th>Offices</th>
<th>Outside</th>
<th>Dry mill</th>
<th>Auto-stecker</th>
<th>Green mill</th>
<th>Maintenance</th>
<th>Saws shop</th>
<th>Control</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aspergillus fumigatus</td>
<td>148</td>
<td>4501</td>
<td>9462</td>
<td>2330</td>
<td>8700</td>
<td>4920</td>
<td>4400</td>
<td>602</td>
</tr>
<tr>
<td>Mucor</td>
<td>28</td>
<td>7</td>
<td>4</td>
<td>1</td>
<td>3</td>
<td>1</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Cladosporium</td>
<td>123</td>
<td>266</td>
<td>104</td>
<td>26</td>
<td>619</td>
<td>543</td>
<td>879</td>
<td>1000</td>
</tr>
<tr>
<td>Penicillium</td>
<td>28</td>
<td>7</td>
<td></td>
<td></td>
<td>217</td>
<td>264</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Talaromyces</td>
<td>—</td>
<td>123</td>
<td></td>
<td></td>
<td>132</td>
<td>219</td>
<td>879</td>
<td>—</td>
</tr>
<tr>
<td>Rhizopus</td>
<td>—</td>
<td>212</td>
<td>6</td>
<td></td>
<td>815</td>
<td>159</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stilbella</td>
<td>40</td>
<td>1803</td>
<td>26</td>
<td>1232</td>
<td>284</td>
<td>262</td>
<td>3560</td>
<td>3120</td>
</tr>
<tr>
<td>Talaromyces</td>
<td>—</td>
<td>38</td>
<td>20</td>
<td>124</td>
<td>26</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Total</td>
<td>1009</td>
<td>8891</td>
<td>33088</td>
<td>5038</td>
<td>27120</td>
<td>18872</td>
<td>13518</td>
<td>6646</td>
</tr>
</tbody>
</table>

— = Not detectable.

Table 9 Mean concentration of airborne fungi (cfu/m³) in different parts of the sawmill as determined by Andersen sampler

<table>
<thead>
<tr>
<th></th>
<th>Log yard</th>
<th>Bark saw</th>
<th>Bark stripper</th>
<th>Dry mill</th>
<th>Green middle</th>
<th>Mill end</th>
</tr>
</thead>
<tbody>
<tr>
<td>Penicillium</td>
<td>1520</td>
<td>1440</td>
<td>16000</td>
<td>1200</td>
<td>2720</td>
<td>2880</td>
</tr>
<tr>
<td>Aspergillus fumigatus</td>
<td>60</td>
<td>380</td>
<td>160</td>
<td>120</td>
<td>80</td>
<td>800</td>
</tr>
<tr>
<td>Talaromyces</td>
<td>240</td>
<td>420</td>
<td>160</td>
<td>200</td>
<td>240</td>
<td>720</td>
</tr>
<tr>
<td>Talaromyces</td>
<td>380</td>
<td>40</td>
<td>90</td>
<td>120</td>
<td>80</td>
<td>240</td>
</tr>
<tr>
<td>Cladosporium</td>
<td>40</td>
<td>60</td>
<td>160</td>
<td>120</td>
<td>80</td>
<td>800</td>
</tr>
<tr>
<td>Acreosporium</td>
<td>—</td>
<td>40</td>
<td>20</td>
<td>120</td>
<td>240</td>
<td>80</td>
</tr>
<tr>
<td>Phellinus</td>
<td>80</td>
<td>60</td>
<td>120</td>
<td>2160</td>
<td>80</td>
<td>800</td>
</tr>
<tr>
<td>Stilbella</td>
<td>600</td>
<td>80</td>
<td>120</td>
<td>240</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Mucor</td>
<td>120</td>
<td>120</td>
<td>240</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Total</td>
<td>4210</td>
<td>3700</td>
<td>17020</td>
<td>2840</td>
<td>8960</td>
<td>7120</td>
</tr>
</tbody>
</table>

Table 9 shows the numbers of fungal spores collected by the Andersen sampler at different sites throughout the sawmill. **Penicillium** species, **Aspergillus** and **Tichoderma** were again commonly isolated at all sites. In general the prevailing concentrations were lower than those measured with personal samplers but extremely large concentrations of **Penicillium** spores were measured near to the automatic debarker.

**Discussion**

Occupational exposure to softwood dust is one of the areas still to be reviewed under the 1986 Control of Substances Hazardous to Health (COSHH) regulations and there is currently no maximum exposure limit; however, suggested limits have been published for guidance. With the exception of workers in the sawshop and in the green mill when cutting Douglas fir, dust concentrations were below the suggested exposure limit (eight hour time weighted average) for softwood dust of 5 mg/m³. It is likely that some of the dust collected by personal samplers worn by men in the sawshop was metallic, from the saws, so that their exposure to wood dust might also be less than the suggested limit.

Both the results of personal dust sampling and the workers own subjective assessment of the dustiness of their environment supported our stratification into high and low exposure groups at the sawmill.

Significantly more men reported work-related mucosal irritation (eye and nose) in the high exposure group at the sawmill than in the low exposure group or at the comparison factory and this is in keeping with other studies of woodworkers.

The prevalence of symptomatic extrinsic allergic alveolitis among the high exposure group at the sawmill (4-4%) suggests that wood associated alveolitis may occur as often in British sawmills as in Swedish sawmills. No cases of extrinsic allergic alveolitis occurred in men in low dust jobs, suggesting that exposure to wood dust was an important aetiological factor.

The finding that work-related wheeze was more prevalent in the comparison and high exposure groups than in the low exposure group suggests that there were sensitisers or irritants in the dust at both sites; at the sawmill these were likely to be in the wood dust, at the comparison factory these may have been related to paint spraying or welding. The fact that the prevalence of symptomatic bronchial hyper-reactivity was similar in both groups at the sawmill and was higher than at the comparison factory, suggests that occupational exposure to sensitisers at the sawmill may be responsible for the development of bronchial hyper-reactivity. Although certain hardwood dusts are known to be potential sensitisers, softwood dusts are not generally considered to be important causes of occupational asthma although they have been recognised as causes of occupational dermatitis. Extrinsic allergic alveolitis and wheeze caused by exposure to the same allergen has previously been described but is uncommon.

Swedish woodtrimmer’s disease is thought to be caused by exposure to moulds growing on wood, the growth of which is nurtured by kiln drying. Only a small proportion of the wood at this sawmill was kiln dried, but half of the sawmill workers had seen mould around the mill, mostly on the unsawn logs in the yard, or on boards being cut in the dry mill. Measurements of airborne spores showed that the largest concentrations occurred near to the automatic debarker and in the green mill. This is consistent with the finding that most wood mould grows in the area between the bark and the wood so that spores will be dispersed when high energy teeth rip off the bark or cut the outer layers of the stripped log. Despite the finding of spontaneous heat generation by the pile of mulched bark (presumably due to decomposition of the bark by fungi) spore concentrations nearby were no greater than in the rest of the log yard and were about half the concentrations found in the green mill. The results of personal sampling are in broad agreement with these findings.

At almost all sites the three most commonly isolated fungi were **Penicillium** species, **Aspergillus fumigatus**, and **Tichoderma** species. **Penicillium** species and **Aspergillus fumigatus** were also the two most abundant fungi at the comparison factory although fewer in number, but **Tichoderma** species were uncommon. Total concentrations of fungal colony forming units were of the same order
Respiratory immunological responses, and aeroallergen concentrations at a sawmill

as the mean daily concentration of fungal spores determined microscopically, in Cardiff, less than 50 miles from the sawmill. These are probably somewhat lower than concentrations at inland sites, but concentrations determined by culturing are also lower than those determined microscopically as not all airborne spores are viable. The composition of species in Cardiff differed greatly, however, from that found in the sawmill. In outdoor air by day, Cladosporium is usually predominant and spores of Aspergillus and Penicillium type are relatively few. At the sawmill and in the comparison factory, Penicillium predominated with Aspergillus fumigatus, Trichoderma, and, sometimes, Stilbella. The last two genera are particularly characteristic of a timber substrate. Cladosporium, Alternaria species, Aureobasidium pullulans, Penicillium species, Aspergillus species, and Trichoderma species have all been found to be numerous in the air of Canadian and Swedish sawmills. Alternaria species and Aureobasidium pullulans were not common, however, at either the sawmill or the comparison factory. Penicillium and Trichoderma species have also been found to be common aerospores in wood furniture factories.

Penicillium species are ubiquitous fungi occurring in soil as saprophytes on decaying vegetation and causing decomposition of vegetable matter such as grains and foodstuffs. They have previously been associated with both asthma and extrinsic allergic alveolitis. Trichoderma is also widespread and is commonly found on wood, wood products, and in soil.

The higher concentrations of IgG against Trichoderma koningii in men in the high exposure group at the sawmill compared with those in the low exposure group or at the comparison factory are likely to reflect exposure to the fungus rather than being indicative of disease. Nevertheless, all those workers with symptoms consistent with extrinsic allergic alveolitis had high concentrations of antibody against Trichoderma koningii. High concentrations of serum antibodies against Trichoderma have also been reported in Finnish sawmill workers.

The mean concentration of IgG against sawdust was also significantly greater in workers in high dust jobs at the sawmill than in those in low dust jobs, or at the comparison factory. The presence of high concentrations of IgG against extracts of sawdust and mulched bark probably also reflects exposure, but concentrations were again high in workers with symptoms of extrinsic allergic alveolitis. The identity of the relevant epitopes in the wood extracts remains unclear. Binding inhibition studies have shown partial antigenic identity with fungal antigens, in particular Trichoderma, present in the wood presumably as a result of infection but there also seem to be other antigens present, some of which may be constituents of wood itself. Wood is known to contain numerous chemicals capable of producing topical dermatitis and exposure to one or more of these may be responsible for inducing specific IgG production. As in other studies of extrinsic allergic alveolitis, the mean specific IgG concentrations were lower in smokers than in non-smokers.

This study has shown that exposure to wood dust at the sawmill was associated with an increased prevalence of symptoms related to nasal and conjunctival irritation and with a high prevalence of symptomatic bronchial hyper-reactivity. Sawmill workers exposed to high concentrations of dust had more work related respiratory symptoms than those exposed to lower dust concentrations and cases of extrinsic allergic alveolitis only occurred in the more heavily exposed group. As well as being exposed to high concentrations of wood dust, this group was also exposed to high concentrations of fungal spores and had high concentrations of specific IgG against extracts of sawdust and Trichoderma koningii. High exposure to fungal spores, particularly to Trichoderma koningii, seems the most likely cause of extrinsic allergic alveolitis in these sawmill workers.

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