TUBERCULOSIS

Investigation of mycobacterial colonisation and invasion of the respiratory mucosa

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Background: The pathogenesis of Mycobacterium avium complex and Mycobacterium tuberculosis in the respiratory tract is poorly understood, as are the reasons for their differing virulence. We have previously shown that their initial adherence to the mucosa is identical.

Methods: The interaction of M avium complex, M tuberculosis, and M smegmatis with human respiratory tissue was investigated in an organ culture model with an air interface. Tissue was infected for intervals up to 14 days and assessed by scanning electron microscopy for adherent bacteria or cultured for recoverable bacteria.

Results: The mean number of adherent bacteria/mm² (and the viable count of macerated tissue, cfu/ml) at 15 minutes, 3 and 24 hours, 7 and 14 days were: M avium complex 168 (153), 209 (136), 289 (344), 193 (313), 14140 (16544); M tuberculosis 30 (37), 39 (23), 48 (53), 1 (760), 76 (2186); M smegmatis 108 (176), 49 (133), 97 (81), 114 (427), 34 (58), (n=6). There was no significant change in morphology between infected and uninfected tissue or tissue infected with the different species over 14 days. The number of M avium complex on the mucosa and recovered from tissue increased over time (p=0.03). M tuberculosis decreased on the surface, but recoverable bacteria increased (p=0.01). M smegmatis numbers on the mucosa and recovered from tissue decreased. Sectioned tissue showed M avium complex and M tuberculosis in submucosal mucus glands and M tuberculosis penetrating epithelial cells in one experiment.

Conclusions: The initial adherence to the mucosa of the three species was similar, but after 14 days they varied in their interaction with the tissue in a manner compatible with their pathogenicity.

Mycobacterium avium complex and Mycobacterium tuberculosis both cause pulmonary disease in humans. M avium complex is generally referred to as an opportunistic pathogen since it usually infects patients with established lung damage, including scarring from previous tuberculosis, cystic fibrosis and other forms of bronchiectasis, pneumococcal lung damage, including scarring from previous tuberculosis, and chronic obstructive pulmonary disease. M avium complex can also cause primary infections in patients without any history of respiratory illness, and these have been reported with increasing frequency in recent years. In developed countries endobronchial or mediastinal infections and solitary pulmonary nodules are increasingly common. Chronic infection of the airway causes bronchiectasis. In addition, parenchymal infections manifesting as pneumonitis or cavitary disease have been reported.

M tuberculosis is an obligate human pathogen transmitted from patient to patient in respiratory secretions and aerosols. Interaction with the alveolar macrophage is the most important mechanism in the disease process but there may be others because, in one study, resident macrophages did not bind M tuberculosis efficiently. Endobronchial tuberculosis may develop via mucosal surface infection in smear positive cases by lymphatic extension from a peribronchial focus or by myeliar spread. Thompson and Kent concluded that most cases of endobronchial disease were the result of the surface implantation of M tuberculosis and that lack of apparent damage seen on bronchoscopy is possibly due to rapid healing and repair of the mucosal surface. An alternative explanation might be uptake of M tuberculosis by specialised epithelial cells which might not necessarily result in damage to the cell.

The pathogenetic mechanisms of M avium complex and M tuberculosis in the respiratory tract are poorly understood, as are the reasons for their differing virulence. We have previously reported the initial interaction of M avium complex, M tuberculosis, and M smegmatis with the respiratory mucosa of human respiratory tissue organ cultures having an air interface. These experiments showed similar adherence of the three species, predominantly to mucus with a fibrous appearance and to fibronectin in extracellular matrix (ECM) exposed when the epithelium was damaged. The results of these studies showed that the virulence of the mycobacterial species did not influence their initial adherence to the respiratory mucosa. We have therefore investigated the adherence to and recovery from organ cultures of M avium complex, M tuberculosis, and M smegmatis at intervals over 14 days incubation.

METHODS

Bacteria
A clinical strain of M avium complex previously described, M tuberculosis H37 RV (ATCC 27294), and M smegmatis strain Mc155 were used in the study. The mean viable counts of the inocula (2 µl) were calculated as previously described. The sizes of the inocula for M avium complex, M tuberculosis, and M smegmatis were 9.3 × 10⁹ cfu/µl (range 6.6 × 10⁸–1.3 × 10¹), 7.8 × 10⁹ cfu/µl (range 7.1 × 10⁸–9.1 × 10⁸), and 3.8 × 10⁹ cfu/µl (range 3.1 × 10⁸–5.0 × 10⁸), respectively. There was no significant difference between the number of bacteria in the inocula of the different series of experiments.

Organ culture
The organ culture with an air interface used in these experiments has been described previously. Human nasal turbinate was obtained from patients undergoing surgery for nasal obstruction and human bronchial tissue from patients undergoing tumour resection. Tissue was used in experiments with the written permission of the patients and the protocol of tissue removal and its use in this study was approved by the Royal Brompton Hospital ethics committee.
Experimental design

Each experiment required 80 pieces of dissected tissue. Organ cultures were studied at five time intervals with the three species matched with an uninfected control; separate organ cultures were processed for scanning electron microscopy (SEM), culture, light microscopy, and transmission electron microscopy (TEM). Two µl of a washed bacterial suspension (*M. avium* complex, *M. tuberculosis* or *M. smegmatis*) in phosphate buffered saline (PBS) were gently pipetted onto the surface of the organ culture. An uninfected control organ culture, inoculated with PBS alone, was also harvested at each time interval. Uninfected and infected tissues were placed at 37°C in 5% CO₂ for 15 minutes, 3 hours, 24 hours, 7 days, and 14 days. After harvesting from the organ cultures the tissue was washed by pipetting 100 µl (×3) of sterile PBS onto the mucosal surface in order to remove unattached bacteria.

Preparation and assessment of tissue for SEM

The method used for tissue preparation and assessment by SEM has been described previously.²¹,²² The tissue was assessed for the percentage of the surface occupied by four main mucosal features: mucus, ciliated cells, unciliated cells, and damaged mucosa. The mucosal morphology was compared over the five time intervals and between infected and uninfected tissue. The number of bacteria seen on the tissue was counted for the entire mucosa. The actual area of the mucosa was calculated taking into account the µ25 magnification of the microscope and the µ2.5 visual display unit magnification. Using this figure and the number of bacteria adherent to the mucosa, the number of bacterial cells per mm² was calculated for each species at each incubation time.

Calculation of viable counts from macerated tissue

Tissue was removed from the organ culture taking care to remove all surrounding agar. One hundred µl of sterile PBS was placed in a sterile graduated centrifuge tube and the meniscus was marked on the side of the tube. This procedure was carried out for all centrifuge tubes and for each separate experiment to compensate for variation in tubes and ambient temperature. Tissue was placed in the PBS and the displacement volume was estimated by the amount of PBS required to fill the tube. Tissue was placed in the PBS and the displacement volume was estimated by the amount of PBS required to fill the tube. There was no significant difference between organ cultures inoculated with the three species of mycobacteria, nor between infected tissue and uninfected controls (data not shown).

Preparation and assessment of tissue for TEM

Tissue was initially fixed in 2.5% glutaraldehyde and then post fixed in 1% osmium tetroxide, dehydrated, and embedded in araldite. Ultrathin (80 nm) sections were then cut and used to look for mycobacteria on the surface and within the mucosa by TEM.

Statistical analysis

Comparison of the percentages of the surface area occupied by mucosal features for infected and control tissue, the number of adherent bacteria, and the number of bacteria recovered from macerated tissue at different time intervals were analysed using Wilcoxon’s matched signed rank test. p values of ≤0.05 were judged to be significant.

RESULTS

Comparison of the number of mycobacteria observed on the mucosal surface by SEM with the number recovered from macerated tissue

Table 1 shows the morphometric analysis of the four main mucosal features: mucus, damaged epithelium, ciliated, and unciliated cells. Incubation of the organ cultures for up to 14 days had no significant effect on the morphological features of the tissue. There was no significant difference between organ cultures inoculated with the three species of mycobacteria, nor between infected tissue and uninfected controls (data not shown).

Figure 1 shows the mean number of bacteria adhering to the mucosal surface of organ cultures (bacteria/mm²) and the

### Table 1

<table>
<thead>
<tr>
<th></th>
<th>Mucus</th>
<th>Damaged mucus</th>
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<th>Unciliated mucus</th>
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<tr>
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<tr>
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<td>9.2</td>
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<td><strong>TB</strong></td>
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<tr>
<td>Mean</td>
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<td>18.5</td>
<td>17.4</td>
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<tr>
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<td>16.4</td>
<td>13.6</td>
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<tr>
<td><strong>SM</strong></td>
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Figures are the percentage of the mucosal surface surveyed occupied by a particular feature. n=number of experiments in which the mucosal feature was observed out of six. 15 m=15 minutes; 3 h and 24 h=3 and 24 hours; 7d and 14 d=7 and 14 days incubation time.
mean number of bacteria recovered from macerated tissue (cfu/μl) incubated for 15 minutes to 14 days. At 15 minutes all three mycobacterial species adhered in similar numbers to the same mucosal features, as previously described.21 22 Bacteria adhered predominantly to ECM, mucus with a fibrous appearance, and were occasionally seen adhering to areas where extruded epithelial cells had degenerated exposing their contents, open tight junctions between cells, and to healthy unciliated epithelium. Bacteria did not adhere to mucus with a globular appearance, intact extruded cells, collagen fibres, or ciliated cells.

Table 2 shows the number of experiments in which *M avium* complex, *M tuberculosis*, and *M smegmatis* were observed on the mucosal surface by SEM and were recovered from macerated tissue. *M tuberculosis* was seen by SEM in two out of six experiments at 7 days incubation. However, the mean number of *M tuberculosis* (1.2) does not register on fig 1 because of the scale of the y axis. One experiment out of six analysed by SEM showed a large amount of mucosal damage (*M tuberculosis*, 58.3% of the mucosal surface at 14 days incubation; *M smegmatis*, 62.4% at 14 days incubation) which exposed a large amount of ECM (*M tuberculosis*, 19.4% of the mucosal surface at 14 days incubation; *M smegmatis*, 17.3%), and this was the one experiment in which *M tuberculosis* and *M smegmatis* were seen on the mucosal surface at this time point and the one experiment in which *M smegmatis* was recovered from macerated tissue.

*M avium* complex increased in numbers on the mucosal surface with time of incubation. There was a major increase in the numbers of *M avium* complex seen by SEM after 14 days incubation compared with other time points. There was no significant difference in numbers observed by SEM when comparing counts between 15 minutes and 7 days incubation. *M avium* complex recovered from macerated tissue increased with time of incubation. There was an increase in the number of bacteria after 7 and 14 days incubation compared with earlier time points.

Table 2 Number of experiments in which *Mycobacterium avium* complex (MAC), *Mycobacterium tuberculosis* and *Mycobacterium smegmatis* were observed by scanning electron microscopy (SEM) and recovered from macerated tissue

<table>
<thead>
<tr>
<th>Time</th>
<th>MAC</th>
<th><em>M tuberculosis</em></th>
<th><em>M smegmatis</em></th>
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<tr>
<td>SEM</td>
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<td>14 days</td>
<td>5/6</td>
<td>1/6</td>
<td>1/6</td>
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<tr>
<td>Macerated tissue</td>
<td></td>
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<td>15 minutes</td>
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<td>6/6</td>
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<td>3 hours</td>
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<td>14 days</td>
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Data provided by a series of six separate experiments using tissue from a different donor for each experiment.

There was a decrease in the number of *M smegmatis* seen on the mucosal surface and recovered from macerated tissue after 14 days compared with other time points.

The numbers of bacteria observed by SEM and recovered from macerated tissue at different times of incubation were compared for *M avium* complex, *M tuberculosis* and *M smegmatis*. The significant differences were: *M avium* complex was seen on the mucosal surface in greater numbers than *M tuberculosis* and *M smegmatis* after 14 days incubation; *M avium* complex was recovered from macerated tissue in greater numbers than *M tuberculosis* and *M smegmatis* after 14 days incubation; and *M tuberculosis* was recovered in greater numbers than *M smegmatis* from macerated tissue after 14 days incubation.

The numbers of bacteria observed by SEM were compared with numbers recovered from macerated tissue for each mycobacterial species. There were more *M tuberculosis* recovered from macerated tissue than adhering to the mucosal surface after 7 and 14 days, but not at earlier time points. There was no significant difference between numbers adhering to the mucosal surface and recovered from macerated tissue for *M avium* complex and *M smegmatis* for all incubation times.
Mycobacterium avium complex, Light microscopy sections
large numbers of organisms (fig 2). as the bacteria were seen in sporadic "colonies" containing numbers of bacteria. This latter suggestion seems more likely features from observation by SEM because of the increased mucosal surface independently of them or obscured the features from observation by SEM because of the increased numbers of bacteria. This latter suggestion seems more likely as the bacteria were seen in sporadic "colonies" containing large numbers of organisms (fig 2).

Light microscopy sections
M avium complex, M tuberculosis, and M smegmatis were seen adhering to the mucosa at 15 minutes, 3 and 24 hours, and 7 days. M tuberculosis and M smegmatis were not seen on the mucosal surface after 14 days incubation, whereas M avium complex was seen in areas of damaged mucosa in increased numbers compared with other incubation times. M avium complex was never observed adhering to intact mucosa in these experiments. M avium complex and M tuberculosis were seen within submucosal glands underlying damaged mucosa after 7 and 14 days incubation (fig 3). Figure 3 also shows invasion of the epithelium by M tuberculosis within the gland.

They were, however, not seen in submucosal glands underlying intact mucosa. M smegmatis was never seen within submucosal mucus glands.

TEM sections
The solitary experiment in which mycobacteria were seen by TEM showed M tuberculosis entering a healthy epithelial cell (fig 4). The observation of M tuberculosis penetration of epithelial cells was not repeated in paraffin embedded light microscopy sections or by SEM analysis.

DISCUSSION
Interaction with the respiratory mucosa is likely to be a key event in the pathogenesis of M avium complex which causes chronic airways infection and granulomatous inflammation leading to bronchiectasis, whereas in tuberculosis the primary pathology is in the alveoli and endobronchial disease is less common as the incidence of prolonged smear positive cases declines. M smegmatis is an environmental species which is generally considered non-pathogenic. Infection of organ cultures with M avium complex and M tuberculosis did not affect mucosal morphology over 14 days incubation despite, in the case of M avium complex, a large increase in the numbers of bacteria on the mucosal surface and, in the case of M tuberculosis, invasion of the tissue. This observation might be because the mycobacteria do not secrete exotoxins, endotoxins, or tissue necrotising enzymes to aid their infectivity. However, it does raise the question as to how M tuberculosis in particular invades the tissue. The adherence of M avium complex and M tuberculosis to open tight cell junctions may represent a potential route of infiltration into submucosal tissues. Since this site of adherence was observed at 15 minutes, it seems likely that mycobacteria took advantage of open tight junctions already present on the mucosa. Phagocytic cells are another potential route of invasion but were not observed on the surface of organ cultures. This could reflect the lack of host circulation from which these cells could migrate, or they may have been washed from the surface during preparation of tissue for microscopic analysis.

Mucosal penetration was not observed by SEM analysis of the mucosal surface of organ cultures, but mycobacteria did adhere to normal unciliated epithelial cells and, in a single experiment (fig 4), endocytosis was observed by TEM. Clark et al used confocal microscopy to investigate the invasion of Yersinia pseudotuberculosis through murine Peyer's patch M cells with the aim of quantitating the relative contribution of M cells to intestinal invasion by Y pseudotuberculosis. We propose
that future studies of mycobacteria could employ a similar technique to investigate invasion of human airway tissue. Labeling of the mycobacteria with luciferase reporter constructs, as described by Snwin and colleagues,20 may also help to visualize bacteria penetrating the tissue.

The SEM surface counts and recoverable bacteria from macerated tissue showed that there was no difference in the numbers of mycobacteria counted on the mucosal surface and recovered from macerated tissue for the three species at early time points up to 24 hours incubation. This shows that the technique of counting bacteria over the entire mucosal surface correlated well with the total number of viable bacteria on the organ culture, and that the different pathogenicity of the three species was not related to their initial interaction with airway mucosa. The significant increase in numbers of M avium complex on the surface over time indicated that it survived and multiplied on the mucosal surface. Bacterial numbers calculated from macerated tissue exceeded SEM counts at 7 and 14 days, which suggests that some invasion of the tissue had also occurred. M avium complex formed isolated colony-like patches and did not spread over the surface. These “colonies” were probably located at the sites of initial adherence. This suggests that there were few exposed binding sites on the mucosa for mycobacterial adherence. M tuberculosis was recovered from macerated tissue in significantly increased numbers over time, but was seen on the surface of only one of six experiments after 14 days, which shows that it did not survive on the mucosal surface but had invaded the mucosa and multiplied within the tissue. M smegmatis did not survive in most organ cultures; it was only seen and recovered from one organ culture after 14 days in small numbers. This organ culture, and the organ culture on which M tuberculosis was observed by SEM at 14 days, contained large amounts of damaged mucosa so that large amounts of ECM were exposed, which may have provided a site for more mycobacteria to adhere than was usual.

A large increase in M avium complex on the mucosal surface occurred between 7 and 14 days, suggesting that a critical number of these bacteria are required before rapid multiplication occurs, similar to the principle of “quorum sensing” reported for other bacteria.20,21 Further work is required to investigate the relationship between the initial numbers of M avium complex adhering to the mucosa and infection of the mucosa since the infecting dose used in this study was high. This relationship may be important in understanding which patients are vulnerable to infection, since the greater the airway damage, the greater the number of exposed adherence targets and hence the greater the risk of infection with M avium complex. Such a study may also offer answers as to why M smegmatis was unable to survive in the organ culture after initially adhering in similar numbers to M avium complex and M tuberculosis. This might relate to nutritional factors, host defences of the tissue or, possibly, the strength of adherence to the tissue.

In our previous studies we have reported that mycobacteria adhere to mucus with a fibrous appearance under SEM,19 which was usually eradicated. An understanding of these differences may give important insights into the pathogenesis of mycobacterial disease. In condition with the clinical features of infection by the three species, this model will be useful in further investigating the virulence of mycobacteria and determining why species differ despite identical initial interactions with the respiratory mucosa.

ACKNOWLEDGEMENT
This research was funded by a grant from the Hawley Legacy.

REFERENCES
Mycobacterial infection of mucosa

LUNG ALERT

Higher influenza-related paediatric admission rates in the tropics than in temperate regions

This retrospective population-based study calculated the rate of excess hospital admissions due to influenza in children aged 13 or under with acute respiratory disease. The data were collected from the Hospital Authority of the Hong Kong Special Administrative Region, China. The excess admissions were calculated using an excess hospital admission model from periods of influenza peaks and baseline respiratory syncytial virus rates (7 weeks each in 1998 and 1999). The adjusted rates of influenza-related excess hospital admissions for 1998 and 1999 per 10 000 children were 278.5 and 288.2 for children aged <1 year, 218.4 and 209.3 for those aged 1—<2 years, 125.6 and 77.3 for those aged 2—<5 years, 57.3 and 20.9 for children aged 5—<10 years, and 16.4 and 8.1 for children aged 10—15 years. Influenza-related excess admissions accounted for 38 606 extra bed-days (5.5–8.2% of all paediatric bed-days). These influenza-related hospital admission rates for children are much higher than previously published figures. Possible explanations suggested are the inclusion of high risk patients, differences in admission practice, and higher influenza attack rates in crowded communities.

The reasons for higher influenza-related hospital admission rates in this subtropical climate compared with temperate regions need further investigation. The study highlights the rationale for continued worldwide influenza surveillance, the major impact of this common infection, and the need for effective prevention.

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