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Association between smoking and tuberculosis infection: a population survey in a high tuberculosis incidence area

S den Boon, S W P van Lill, M W Borgdorff, S Verver, E D Bateman, C J Lombard, D A Enarson, N Beyers

Background: Associations between smoking and tuberculosis disease including death from tuberculosis have been reported, but there are few reports on the influence of smoking on the risk of developing Mycobacterium tuberculosis infection. The aim of this study was to determine the association between smoking and M tuberculosis infection.

Methods: In a cross sectional population survey, data on smoking and tuberculin skin test (TST) results of 2401 adults aged ≥ 15 years were compared.

Results: A total of 1832 (76%) subjects had a positive TST (≥ 10 mm induration). Of 1309 current smokers or ex-smokers, 1070 (82%) had a positive TST. This was significantly higher than for never smokers (unadjusted OR 1.99, 95% confidence interval (CI) 1.62 to 2.45). A positive relationship with pack-years was observed, with those smoking more than 15 pack-years having the highest risk (adjusted OR 1.90, 95% CI 1.28 to 2.81).

Conclusion: Smoking may increase the risk of M tuberculosis infection.

In a recent large epidemiological study, 50% of deaths from tuberculosis among Indian men was attributed to smoking.1 While an association between smoking and tuberculosis disease has been shown in various studies,2–7 it is less clear to what extent smoking increases the risk of M tuberculosis infection, the risk of progression from infection to disease, or the risk of death among tuberculosis patients.

We report here a population survey conducted in two urban communities in Cape Town, South Africa that provides evidence that smoking may increase the risk of tuberculosis infection. The study area in which we have performed epidemiological research for more than 10 years has a population of 36 334 and 97.9% are of mixed ethnicity (categorised as “coloured”; Statistics South Africa: Census 2001). The tuberculosis notification rate in the area is 238 new cases per 100 000 population per year (1993–8),8 and the prevalence of HIV infection is estimated to be lower than the overall rate for the Western Cape province at the time of the survey (12.4% (8.8–15.9%) in 2002).9

METHODS

A random sample of 15% of the residential addresses was selected. Of the 833 selected addresses, the head of the household of 218 (26%) declined to participate. These addresses were replaced by neighbouring addresses according to fixed rules. Written informed consent was obtained from the household of 218 (26%) declined to participate. These addresses were replaced by neighbouring addresses according to fixed rules. Written informed consent was obtained from every adult (≥ 15 years) and the study was approved by the research ethics committees of Stellenbosch University and the University of Cape Town.

Data collection methods included a tuberculin skin test (TST) and a precoded questionnaire with questions on smoking. For the TST we used an intradermal injection of 2 TU (0.1 ml) PPD RT 23 into the ventral aspect of the left forearm. The induration was measured 48–72 hours later.

Smoking was defined as having ever smoked for at least 1 year. The average number of cigarettes smoked per day was recorded for all ex-smokers and current smokers. The number of pack-years smoked was calculated as the average number of cigarettes smoked per day multiplied by the duration of smoking divided by 20. Infection with M tuberculosis was defined as a TST of 10 mm or more, referred to as a positive TST. The 10 mm cut off point is conventionally used and appeared justified in this population since intermediate reactions to TST, usually attributed to environmental mycobacteria, were uncommon (fig 1).

The following potential confounders were considered: age, sex, education level, body mass index (BMI), and individual monthly income. The possible interactions between pack-years and the variables age and sex were investigated. Odds ratios (OR) and their 95% confidence intervals (95% CI) were determined to identify risk factors, and random effects logistic regression was used to control for confounding as well as for possible clustering at the address level.

RESULTS

Of 3512 adults who completed the questionnaire, a TST result was recorded in 2443 (70%). Data on smoking habits were incomplete in 42 individuals so analyses were performed on 2401. Compared with subjects excluded from the analyses, those included were less often male (unadjusted OR 0.57, 95% CI 0.49 to 0.65), were less often current or ex-smokers (unadjusted OR 0.82, 95% CI 0.70 to 0.94), and fewer had an income above 2000 Rand (300 US$) (unadjusted OR 0.62, 95% CI 0.51 to 0.76).

Of the 2401 adults analysed, 1832 (76%) had a positive TST and 1309 (53%) were current or ex-smokers; 82% (n = 1070) of the 1309 current or ex-smokers had a positive TST compared with 70% (n = 762) of the 1092 never smokers (unadjusted OR 1.99, 95% CI 1.62 to 2.45). Men were more likely to have a positive TST than women (unadjusted OR 1.34, 95% CI 1.08 to 1.65; table 1). A positive TST was more common in the 25–44 year age group than in other age groups. The proportion of individuals with a positive TST increased with income, but not with education level or BMI (data not shown).

In the multiple logistic regression analyses, income and sex did not confound the association between smoking and infection. However, as sex was associated with smoking and with having a positive TST, it was retained in the model. The
association between infection and pack-years smoked was not significantly different for the different age and sex strata. After adjusting for age and sex, there was a significant association between smoking and a positive TST (adjusted OR 1.77, 95% CI 1.33 to 2.35). The probability of a positive TST seemed to increase slightly with the number of pack-years smoked, although the differences between the different pack-year categories were not significant (table 1).

**DISCUSSION**

This study shows that current or ex-smokers had a higher prevalence of *M tuberculosis* infection than never smokers and that there was a slightly higher risk of infection for those who smoked more than 15 pack-years than for those who smoked less, although this was not significant. This suggests that the increased risk of disease and death from tuberculosis among smokers may be due, at least in part, to an increased risk of smokers becoming infected with *M tuberculosis*.

An unexpected finding was the positive association between a positive TST and income. It should be noted, however, that the mean incomes in the study area are low, and that the categorisation threshold used in the analyses identifies only the poorest in the community. Nevertheless, the reason for their lower TST rates requires further study, including the possibility of lower risk of exposure through fewer social contacts.

Our study confirms previous studies that showed an association between smoking and tuberculosis infection in at risk groups.10–14 For example, in an immigrant population Plant et al10 reported a higher risk of infection among smokers which increased with duration of smoking. In contrast to previous studies investigating specific high risk groups,10–14 the current study is the first to investigate the association between smoking and tuberculosis infection in a cross sectional population survey in a high incidence community.

The reason for the increased risk of infection in smokers is unclear, but may be explained by the effects of smoking on pulmonary host defences. Smoking has been shown to reduce natural killer cytotoxic activity, to suppress T cell function in both lung and blood, to impair mucociliary clearance of particles, and to increase numbers of alveolar macrophages in the lower respiratory tract. Cells of the macrophage-phagocytic group influence immediate or innate immunity through their handling and elimination of mycobacteria, and products of cigarette smoke may therefore favour persistence and/or replication of ingested mycobacteria by impairing the macrophage or dendritic cell function.15 16

To take possible sources of bias into account we have considered the following. Men and persons in the highest income category are under-represented, but this is unlikely to be of significance as neither sex nor income was a confounder for the association between smoking and positive TST. Smokers and ex-smokers were also slightly under-represented but still comprised 55% of the sample, and we can see no reason to assume that the smokers who had undergone a TST might be different from those who had not. A weakness of the study is that we did not test the HIV status

![Figure 1 Distribution of induration of tuberculin skin test in 2401 adults.](image)

**Table 1 Risk factors for *M tuberculosis* infection**

<table>
<thead>
<tr>
<th>Smoking (pack-years)</th>
<th>TST $\geq$10 mm</th>
<th>Total</th>
<th>% TST $\geq$10 mm</th>
<th>Unadjusted (95% CI)</th>
<th>Adjusted* (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Never smoked</td>
<td>769</td>
<td>1102</td>
<td>70</td>
<td>1.86 (1.42 to 2.43)</td>
<td>1.77 (1.33 to 2.35)</td>
</tr>
<tr>
<td>$&lt;5$</td>
<td>443</td>
<td>550</td>
<td>81</td>
<td>2.39 (1.74 to 3.29)</td>
<td>1.77 (1.25 to 2.50)</td>
</tr>
<tr>
<td>$5$–$15$</td>
<td>343</td>
<td>407</td>
<td>84</td>
<td>2.00 (1.40 to 2.84)</td>
<td>1.90 (1.28 to 2.81)</td>
</tr>
<tr>
<td>$&gt;15$</td>
<td>236</td>
<td>288</td>
<td>82</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unknown</td>
<td>41</td>
<td>54</td>
<td>76</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>1100</td>
<td>1477</td>
<td>74</td>
<td>1.34 (1.08 to 1.65)</td>
<td>1.24 (0.99 to 1.56)</td>
</tr>
<tr>
<td>Male</td>
<td>731</td>
<td>923</td>
<td>79</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Unknown</td>
<td>1</td>
<td>1</td>
<td>100</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>15–24</td>
<td>432</td>
<td>651</td>
<td>66</td>
<td></td>
<td></td>
</tr>
<tr>
<td>25–34</td>
<td>444</td>
<td>498</td>
<td>89</td>
<td>4.47 (3.14 to 6.37)</td>
<td>4.30 (3.00 to 6.17)</td>
</tr>
<tr>
<td>35–44</td>
<td>412</td>
<td>484</td>
<td>85</td>
<td>3.16 (2.28 to 4.37)</td>
<td>2.82 (2.00 to 3.99)</td>
</tr>
<tr>
<td>45–54</td>
<td>263</td>
<td>330</td>
<td>80</td>
<td>2.11 (1.50 to 2.97)</td>
<td>1.96 (1.35 to 2.85)</td>
</tr>
<tr>
<td>$&gt;55$</td>
<td>264</td>
<td>408</td>
<td>65</td>
<td>0.88 (0.66 to 1.18)</td>
<td>0.88 (0.64 to 1.22)</td>
</tr>
<tr>
<td>Unknown</td>
<td>17</td>
<td>30</td>
<td>57</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Income (Rands)$\dagger$</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>$&lt;500$</td>
<td>804</td>
<td>1088</td>
<td>74</td>
<td></td>
<td></td>
</tr>
<tr>
<td>500–1999</td>
<td>685</td>
<td>893</td>
<td>77</td>
<td>1.18 (0.94 to 1.48)</td>
<td></td>
</tr>
<tr>
<td>$\geq2000$</td>
<td>315</td>
<td>381</td>
<td>83</td>
<td>1.93 (1.38 to 2.68)</td>
<td></td>
</tr>
<tr>
<td>Unknown</td>
<td>28</td>
<td>39</td>
<td>72</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>1832</td>
<td>2401</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Adjusted for pack-years, age, sex, and taking into account possible clustering at address level.

$\dagger$R6.50 = US$1.00.
of participants and were therefore not able to correct for HIV status. Confounding factors that were taken into consideration were individual monthly income, BMI, and education level. However, we cannot entirely discount the possibility that socioeconomic and behavioural differences other than smoking may have affected the relationship between smoking and tuberculous infection.

We conclude that smoking may increase the risk of M tuberculosis infection. We propose that further studies be conducted to investigate this association and to establish whether smoking reduction strategies contribute to tuberculous control.

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Authors’ affiliations

S den Boon, S WP van Lill, N Beyers, Centre for TB Research and Education, Department of Paediatrics and Child Health, Stellenbosch University, P O Box 19063, 7505 Tygerberg, Cape Town, South Africa

S den Boon, M W Borgdorff, Academic Medical Centre, Amsterdam, The Netherlands

M W Borgdorff, S Verwer, KNCV Tuberculosis Foundation, The Hague, The Netherlands

E D Bateman, Division of Pulmonology, Department of Medicine, University of Cape Town, Cape Town, South Africa

C J Lombard, Biostatistics Unit, Medical Research Council, Cape Town, South Africa

D A Enarson, International Union Against Tuberculosis and Lung Diseases, Paris, France

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NB, EB, Svl, DE, and MB were involved in the study design and writing of the manuscript. SdB performed the statistical analysis and wrote the paper. Advice on statistics was given by CL, SV and MB. All authors participated in interpretation of the results. NB supervised the data collection and MB the statistical analyses.

REFERENCES


