

# Particulate matter is associated with sputum culture conversion in patients with culture-positive tuberculosis

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**Abstract:** Emerging risk factors for tuberculosis (TB) infection, such as air pollution, play a significant role at both the individual and population levels. However, the association between air pollution and TB remains unclear. The objective of this study was to examine the association between outdoor air pollution and sputum culture conversion in TB patients. In the present study, 389 subjects were recruited from a hospital in Taiwan from 2010 to 2012: 144 controls with non-TB-related pulmonary diseases with negative sputum cultures and 245 culture-positive TB subjects. We observed that a 1  $\mu\text{g}/\text{m}^3$  increase in particulate matter of  $\leq 10 \mu\text{m}$  in aerodynamic diameter ( $\text{PM}_{10}$ ) resulted in 4% higher odds of TB (odds ratio = 1.04, 95% confidence interval = 1.01–1.08,  $P < 0.05$ ). The chest X-ray grading of TB subjects was correlated to 1 year levels of  $\text{PM}_{10}$  ( $R^2 = 0.94$ ,  $P < 0.05$ ). However, there were no associations of pulmonary cavitation or treatment success rate with  $\text{PM}_{10}$ . In subjects with TB-positive cultures, annual exposure to  $\geq 50 \mu\text{g}/\text{m}^3$   $\text{PM}_{10}$  was associated with an increase in the time required for sputum culture conversion (hazard ratio = 1.28, 95% confidence interval: 1.07–1.84,  $P < 0.05$ ). In conclusion, chronic exposure to  $\geq 50 \mu\text{g}/\text{m}^3$   $\text{PM}_{10}$  may prolong the sputum culture conversion of TB patients with sputum-positive cultures.

**Keywords:** air pollution, chest X-ray, particulate matter, sputum culture, tuberculosis

## Introduction

Tuberculosis (TB) is a communicable disease caused by inhalation of the bacillus *Mycobacterium tuberculosis*. TB is spread to others when patients with pulmonary TB expel bacteria. In 2013, the World Health Organization (WHO) reported that 9 million people developed TB, and 1.5 million people died from TB.<sup>1</sup> In Taiwan, the treatment success rate of new smear-positive cases is 70%.<sup>2</sup> More effort still needs to be put into controlling TB and increasing the awareness of other risk factors that can modulate disease treatment.

Air pollutants are thought to influence the development of pulmonary disease by interfering with nonspecific and specific lung defenses.<sup>3</sup> Many studies conducted over the past 30 years have shown an association between air pollution and TB.<sup>4,5</sup> People who are exposed to particulate matter that is  $< 10 \mu\text{m}$  in aerodynamic diameter ( $\text{PM}_{10}$ ) generated from biomass burning are more likely to be infected with *M. tuberculosis* and progress to TB disease.<sup>6,7</sup> Smith et al<sup>8</sup> observed that outdoor  $\text{PM}_{10}$  may be a causative agent for the development of TB. A Taiwan cohort study further showed that exposure to outdoor fine particulate matter was associated with an increased risk of active TB (adjusted hazard ratio = 1.39 per 10  $\mu\text{g}/\text{m}^3$ ).<sup>5</sup> Jassal et al<sup>9</sup> suggested that outdoor fine particulate matter was associated with TB lung pathology and the prevalence of

smear-positive TB.<sup>9</sup> Occupational exposure to silica particles was identified as a risk factor for TB.<sup>10</sup> Additionally, other pollutants, such as sulfur dioxide (SO<sub>2</sub>), were identified as risk factors for TB infection.<sup>11</sup> However, a possible link between air pollution and TB treatment after establishing a diagnosis has not previously been examined.

The reference standard method for diagnosing TB is sputum microscopy to directly observe bacteria. Regarding TB treatment, four standard treatment regimens of first-line TB drugs are used in clinical practice, which achieves success rates of >95%.<sup>12</sup> The period for taking different TB drugs is normally between 6 and 9 months. However, a longer period may be required if cavitation, extensive disease, immunosuppression, or a sputum culture that remains positive at 8 weeks occurs in TB patients.<sup>13</sup> The objectives of the present study were to investigate the associations of PM<sub>10</sub> with 1) the risk of acquiring pulmonary TB; 2) the severity of pulmonary TB disease as graded by a chest X-ray (CXR) appearance; and 3) the rate of sputum culture conversion.

## Materials and methods

### Study population

A retrospective case–control study of cases reported between January 1, 2010 and December 31, 2012 was conducted. Study subjects were recruited in the chest clinic of Taipei Medical University, Shuang Ho Hospital (New Taipei City, Taiwan). Included in the study were sputum culture-positive patients of both sexes, aged >16 years, who had begun treatment during the period of study. Patients who were smear- and culture-positive were included in the study. TB was diagnosed using sputum culture according to diagnostic standards.<sup>13,14</sup> The frequency of sputum culture was the 1st day, and the 2nd, 6th, 9th, and 12th months. A clinical-based control population was recruited from patients who visited the chest clinic and were diagnosed with non-TB pulmonary diseases. Additionally, a sputum culture examination was conducted in control subjects. The sputum culture-negative group served as the control group. Excluded were those for whom the mandatory report was incomplete and for whom missing data could not be retrieved. Cases of TB were assessed at the start and end of TB treatment, taking care to exclude cases ruled out by the attending physician. Personal, clinical, and epidemiological data were obtained from patients' clinical progress records. Microbiological laboratory data and medical records of patients were reviewed. The TB treatment success rate was defined as having successfully completed treatment, with or without bacteriological evidence of success. This study was approved by the Taipei Medical University Joint Institutional

Review Board (Taipei, Taiwan). Records of all patients were anonymized and de-identified before the analysis.

### Grading of CXRs

To quantify the severity of chest lesions observed in CXR images of TB patients, a scoring method was applied in the present study.<sup>15</sup> Briefly, the presence of nodules, patchy or confluent consolidation, cavitation (size), bronchial lesions or fibrosis, effusion, and lymphadenopathy were recorded. To grade the percentage of the lungs that was affected, a visual estimation of the extent of opacification, cavitation, or other pathology as a percentage of the visible lung was determined. Dense and patchy opacification of a zone was graded. The severity of TB CXRs was scored blindly by two chest physicians (KYL and PHF).

### Air pollution exposure

Measurements of air pollutants were obtained from 25 monitoring stations operated by the Taiwan Environmental Protection Administration. The monitoring stations, located throughout northern Taiwan, measured air pollutants and weather data on a daily basis. Daily concentrations of PM<sub>10</sub>, ozone (O<sub>3</sub>), nitrogen dioxide (NO<sub>2</sub>), SO<sub>2</sub>, and carbon monoxide (CO) were collected to represent exposure to air pollution among the population by indicating the nearest station within 10 km of their residence. If there was more than one station within 10 km of their residence, the participant was also assigned exposure values equal to the weighted average of all monitors in their area of residence, with weights proportional to the inverse of the square of the distance between their residence and the station. All daily air pollution measurements were matched to the date of treatment initiation for study subjects. For the control group, the previous 1 year PM<sub>10</sub> data were collected when a subject was recruited for the study. The average environmental data values for the 365 days before the matched date were used to estimate the yearly average of daily levels.

### Statistical analysis

The Shapiro–Wilk test was used to test for normality. A logistic regression model was used to assess the risk of air pollutants associated with subjects with TB-positive cultures compared to the risk associated with subjects who had negative cultures. This model produced estimates of the odds ratios and corresponding 95% confidence intervals (CIs). The multivariate Cox proportional-hazards regression model was used to assess the hazard ratio and corresponding 95% CI for sputum culture conversion in subjects with positive sputum

cultures relative to PM<sub>10</sub> exposure. A linear regression model was used to examine the correlation of quintiles of PM<sub>10</sub> with pulmonary cavity, treatment outcomes (success rate), and CXR grading. All statistical analyses for this study were performed using SPSS 15.0 software (SPSS, Chicago, IL, USA). The significance criterion was set to  $P < 0.05$ .

## Results

### Demographic characteristics of the study population

Table 1 shows the demographic characteristics of study subjects. Males made up 63% and 71% of patients with TB-negative and -positive cultures, respectively, and average ages of patients in these groups were 56 and 59 years, respectively. Values of the body mass index of all subjects ranged between 22 and 23 kg/m<sup>2</sup>. Married cases accounted for 87% and 83% of those with TB-negative and -positive cultures, respectively. Percentages of smokers were 12% for culture-negative subjects and 13% for culture-positive subjects. Alcohol consumption rates for the TB-negative and -positive groups were 46% and 27%, respectively. In the study groups, most subjects had less than high school education (46% for TB-negative subjects and 53% for TB-positive subjects), and the majority of subjects were in the middle-income socioeconomic group (95% for culture negative and 83% for culture positive). There were no human immunodeficiency virus coinfections in these subjects. In terms of CXR, 26% of subjects with a positive culture presented with pulmonary cavity lesions, which accounted for 33 in the CXR grading. The number of days for sputum culture conversion from positive to negative was 242 days (range: 32–773 days).

### Air pollution and TB

Table 2 summarizes the 1 year average data of air pollutants, and the values were 48.7 µg/m<sup>3</sup> for PM<sub>10</sub>, 3.2 ppb for SO<sub>2</sub>, 21.6 ppb for NO<sub>2</sub>, 0.6 ppm for CO, and 25.7 ppb for O<sub>3</sub>. We observed associations of TB risk with increased PM<sub>10</sub>, after adjusting for age and sex. An increase of 1 µg/m<sup>3</sup> of PM<sub>10</sub> resulted in a 4% higher odds of TB (odds ratio =1.04, 95% CI: 1.01–1.08,  $P < 0.05$ ). Incident TB was not associated with SO<sub>2</sub>, NO<sub>2</sub>, CO, or O<sub>3</sub>.

### PM<sub>10</sub>, pulmonary lesions, and treatment outcomes

One-year averages of PM<sub>10</sub> for quintiles in relation to pulmonary cavity, treatment success, and CXR grading were examined (Figure 1). We observed a correlation of PM<sub>10</sub> with CXR

**Table 1** Demographic characteristics of subjects with TB-negative (n=144) and TB-positive cultures (n=245) from 2010 to 2012

Variables	Negative culture subjects (%)	Positive culture subjects (%)
Sex, no (%)		
Female	53 (37)	70 (29)
Male	91 (63)	175 (71)
Age, years		
Mean ± SD	56±20	59±21
Range	19–96	16–94
BMI, kg/m <sup>2</sup>		
Mean ± SD	23±4	22±4
Range	14–40	12–33
Marital, no (%)		
Unmarried	19 (13)	42 (17)
Married	125 (87)	203 (83)
Smoking, no (%)		
Nonsmoker	92 (64)	146 (60)
Former smoker	34 (24)	67 (27)
Current smoker	18 (12)	32 (13)
Alcohol consumption, no (%)		
No	78 (54)	179 (73)
Yes	66 (46)	66 (27)
Education, no (%)		
Less than high school	67 (46)	129 (53)
High school or equivalent	50 (35)	77 (31)
More than high school	27 (19)	39 (16)
Socioeconomic status, no (%)		
High	3 (2)	2 (1)
Middle	137 (95)	204 (83)
Low	4 (3)	38 (16)
HIV infection		
Yes	0	0
No	144 (100)	245 (100)
Pulmonary cavity		
Yes	0 (0)	63 (26)
No	144 (100)	182 (74)
CXR grading		
Mean ± SD	–	33±29
Range	–	3–140
Sputum culture conversion, days		
Mean ± SD	–	242±108
Range	–	32–773
Outcome		
Alive	–	212 (87)
Death	–	33 (13)

**Abbreviations:** BMI, body mass index; CXR, chest X-ray; HIV, human immunodeficiency virus; SD, standard deviation; TB, tuberculosis.

grading ( $R^2=0.94$ ,  $P < 0.05$ ), whereas no correlation observed in PM<sub>10</sub> with the pulmonary cavity ( $R^2=0.73$ ,  $P=0.068$ ) or treatment success ( $R^2=0.14$ ,  $P=0.52$ ).

### PM<sub>10</sub> and treatment days

The association between different levels of PM<sub>10</sub> exposure and the time for sputum culture conversion from positive to

**Table 2** Levels of 1 year average of air pollutants among study subjects and the estimated ORs (95% CI) for the risk of TB associated with 1-unit increase in 1 year average

Variables	Concentration Mean ± SD	TB OR (95% CI) <sup>a</sup>
PM <sub>10</sub> , µg/m <sup>3</sup>	48.7±7.0	1.04 (1.01–1.08)*
SO <sub>2</sub> , ppb	3.2±0.8	1.24 (0.90–1.69)
NO <sub>2</sub> , ppb	21.6±3.2	1.03 (0.96–1.10)
CO, ppm	0.6±0.1	2.32 (0.19–27.9)
O <sub>3</sub> , ppb	25.7±1.8	1.02 (0.89–1.14)

Notes: <sup>a</sup>Adjusted for age, sex, and smoking. \*P<0.05.

Abbreviations: CI, confidence interval; CO, carbon monoxide; NO<sub>2</sub>, nitrogen dioxide; O<sub>3</sub>, ozone; OR, odds ratio; SD, standard deviation; SO<sub>2</sub>, sulfur dioxide; TB, tuberculosis.

negative was investigated, as shown in Table 3. In the present study, the WHO guideline for PM<sub>10</sub> of 50 µg/m<sup>3</sup> was applied to divide subjects with a positive sputum culture into two groups: <50 and ≥50 µg/m<sup>3</sup>. We observed an association between exposure to ≥50 µg/m<sup>3</sup> PM<sub>10</sub> and an increased risk for a longer time for conversion of the sputum culture from positive to negative (hazard ratio =1.28, 95% CI: 1.07–1.84, P<0.05).

**Table 3** HR for days of sputum cultures conversion in subjects with TB-positive cultures

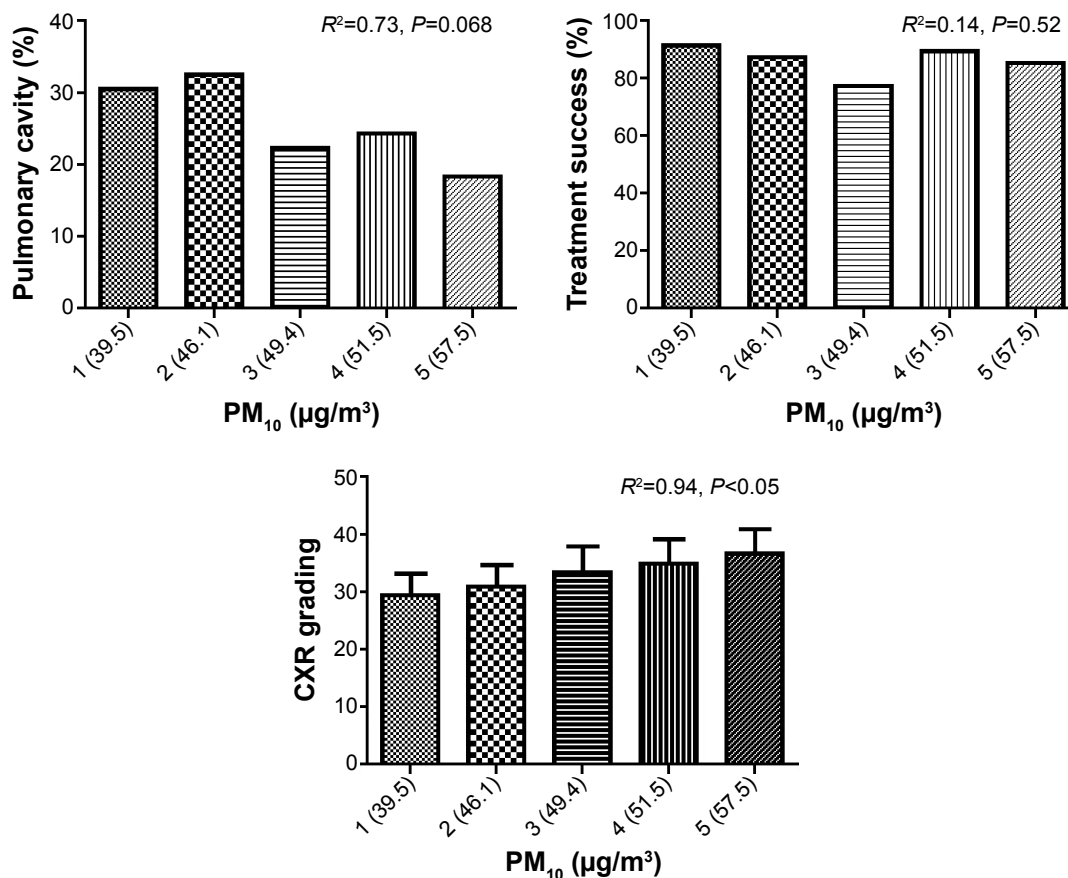
Smoking status	Sputum culture conversion HR (95% CI) <sup>a</sup>
<50 µg/m <sup>3</sup>	1
≥50 µg/m <sup>3</sup>	1.28 (1.07–1.84)*

Notes: <sup>a</sup>Adjusted for age, sex, smoking, and CXR grading. \*P<0.05.

Abbreviations: CI, confidence interval; CXR, chest X-ray; HR, hazard ratio; TB, tuberculosis.

## Discussion and conclusion

This analysis showed that exposure to PM<sub>10</sub>, but not other pollutants, was associated with TB that was sputum culture-positive. We found that the severity of pulmonary lesions, based on the score of the CXR grading, increased with increasing levels of PM<sub>10</sub>. Alteration of the immunity of TB patients by PM<sub>10</sub> exposure may be an explanation for our observations, suggesting the possibility that PM<sub>10</sub> is a risk factor for regulating sputum culture conversion in TB. We further showed that TB patients exposed to high PM<sub>10</sub> levels (≥50 µg/m<sup>3</sup>) had an elevated risk for prolonged sputum culture conversion from positive to negative.



**Figure 1** The relationship of pulmonary cavity, treatment success, and CXR grading to 1 year levels of PM<sub>10</sub>.

Notes: Study subjects were divided equally based on quintiles of PM<sub>10</sub>. The values in parentheses along the x-axis represent the mean PM<sub>10</sub> values for each quintile group. Linear regression was used to examine the correlation between quintiles of PM<sub>10</sub> and pulmonary cavity, treatment success and CXR grading. The CXR grading was correlated with 1 year PM<sub>10</sub> averages (R<sup>2</sup>=0.94, P<0.05).

Abbreviation: CXR, chest X-ray.

Examining mucus from the lungs of TB-suspected subjects provides the most accurate diagnosis for active TB. Thirty-six percent of all forms of new TB cases were diagnosed as positive sputum cultures in Taiwan in 2012,<sup>16</sup> suggesting that a significant portion of TB patients had sputum-positive cultures. To examine our hypothesis that exposure to air pollution delays sputum culture conversion, TB patients with positive sputum cultures were recruited for the study group. Compared with non-TB patients with negative sputum cultures, we found that a 1 µg/m<sup>3</sup> increase in PM<sub>10</sub> levels resulted in 4% higher odds of TB patients having positive sputum cultures. These results were consistent with other studies.<sup>7,9</sup> Clinically, higher bacterial loads in the sputum are associated with pulmonary lesions in TB patients.<sup>17</sup>

We investigated the effects of PM<sub>10</sub> levels on pulmonary cavity, treatment success rate, and CXR grading. There were no correlations of PM<sub>10</sub> levels with pulmonary cavity or the treatment success rate. It is noteworthy that CXR grading was positively associated with PM<sub>10</sub> levels. A previous report showed that the time between the appearance of TB symptoms and initiation of anti-TB treatment was related to smoking behavior.<sup>18</sup> Our results suggested that pulmonary lesions deteriorated along with exposure to PM<sub>10</sub>; consequently, the effects of PM<sub>10</sub> on the lungs of TB subjects may be associated with TB treatment.

If PM<sub>10</sub> causes a faster development of pulmonary lesions, then PM<sub>10</sub> may affect the treatment progress of TB patients. TB treatment may be extended for more than the standard 6 months in patients with risk factors for relapse. These risk factors include cavitation, extensive disease, immunosuppression, and a sputum culture that remains positive at 8 weeks.<sup>13</sup> The present study referenced the WHO PM<sub>10</sub> guideline of 50 µg/m<sup>3</sup> as the cutoff point to divide the sputum-positive culture cohort into two groups (<50 and ≥50 µg/m<sup>3</sup>). Importantly, we found an association between annual exposure to ≥50 µg/m<sup>3</sup> PM<sub>10</sub> and treatment days for sputum culture conversion. Results showed that clinical exposure to PM<sub>10</sub> higher than 50 µg/m<sup>3</sup> may prolong the treatment time for sputum culture to convert from positive to negative. This observation was also indicated in other pulmonary diseases. Lung cancer, for example, was linked to cigarette smoke, which may be associated with drug pharmacokinetics and treatment efficacy.<sup>19</sup> Based on our findings, we suspect that sputum culture-positive TB patients with PM<sub>10</sub> exposure ≥50 µg/m<sup>3</sup> may need longer standard treatment periods. However, further research is needed to explore the possible association.

The most crucial cause of the extended course of TB treatment is the dormant state of the *M. tuberculosis*.<sup>20</sup> When *M. tuberculosis* bacilli reach the alveolar space, dormant and

active *M. tuberculosis* may coexist. Clinically latent infection or active TB depends on the proportions of dormant and active *M. tuberculosis*. When dormant TB is resuscitated during treatment, the microenvironment changes, the proportion of active TB increases, and an extended period of treatment is necessary. Additionally, the respiratory tract is the system most frequently infected after PM<sub>10</sub> exposure, which can result in an increased risk of TB.<sup>21</sup> The possible mechanisms for PM<sub>10</sub>-related infections include an accumulation of chemicals, such as metals, by surface functional groups of PM<sub>10</sub>.<sup>21</sup> Polycyclic aromatic hydrocarbons were also shown to modulate drug-metabolizing enzymes.<sup>19</sup>

Our study has some limitations. First, factors besides outdoor air pollution could influence the treatment course of TB. Additionally, treatment side effects and multidrug-resistant TB may have affected study outcomes and were not investigated in this study. In addition, indoor air pollution and seasonal effects of air pollution were not determined in the present study, which should be addressed in future work. TB drug medications were not recorded; we could not assess possible interactions between the therapeutic regimens and PM<sub>10</sub>. In addition, other risk factors for TB, such as alcohol consumption, should be quantified in future work. In conclusion, PM<sub>10</sub> was significantly associated with the period required for sputum culture conversion and the severity of pulmonary lesions. Importantly, exposure to PM<sub>10</sub> at ≥50 µg/m<sup>3</sup> may prolong the TB treatment time. Our findings have important implications for TB treatment strategies, TB control policies, and public health.

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## Disclosure

The authors report no conflicts of interest in this work.

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