

Progress towards tuberculosis elimination: secular trend, immigration and transmission

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ABSTRACT

This study aimed to determine to what extent tuberculosis trends in the Netherlands depend on secular trend, immigration, and recent transmission. Data on patients in the Netherlands Tuberculosis Register in the period 1993-2007 were matched with restriction fragment length polymorphism (RFLP) patterns of *Mycobacterium tuberculosis* isolates. Index patients were defined as patients with pulmonary tuberculosis whose isolates had RFLP patterns not observed in another patient in the previous two years.

Among 8330 patients with pulmonary tuberculosis the isolates of 56% of native and 50% of foreign-born patients were clustered. Of these, 5185 were included in detailed analysis: 1376 native index patients, 2822 foreign-born index patients, and 987 secondary cases within two years after diagnosis of the index case. The incidence of native and foreign-born index patients declined with 6% and 2% per year, respectively. The number of secondary cases per index case was 0.24. The decline of native cases contributed most to the overall decline of tuberculosis rates and was largely explained by a declining prevalence of latent infection. TB among immigrants was associated with immigration figures. Progress towards elimination of TB would benefit from intensifying diagnosis and treatment of latent infection among immigrants and global TB control.

Key words: Infectious diseases epidemiology, molecular epidemiology, prevention & control, tuberculosis

INTRODUCTION

Tuberculosis (TB) is caused by *Mycobacterium tuberculosis* and may result from rapid progression, defined as progression to disease within one to five years after (re-) infection, or from endogenous reactivation of latent infection up to decades after infection [1-3].

Moreover, in particular among the foreign-born [4], cases occur after recent or remote infection outside the study area.

Genotyping of *M. tuberculosis* isolates using restriction fragment length polymorphism (RFLP) with IS6110 and the polymorphic GC rich sequence (PGRS) as markers in principle allows a separation of disease attributable to recent transmission and disease attributable to reactivation or importation [5]. The RFLP patterns of *M. tuberculosis* isolates of patients with disease due to recent transmission are likely to have been observed in at least one other patient in recent years, provided infection took place within the study area, the vast majority of patients was captured and their isolates subjected to genotyping [6]. If such a DNA fingerprint was not observed, disease is probably due to reactivation or importation.

Using molecular epidemiologic methods, declining tuberculosis rates in New York [4], San Francisco [7] and Madrid [8] in the 1990's were attributed to improved control which led to lower TB rates attributable to recent transmission. Declining TB rates in the Netherlands [9] and in Arkansas [10] around the beginning of this millennium were attributed to declining TB rates attributable to endogenous reactivation of latent infection, which in the Netherlands were attributed to a cohort effect [9].

In the Netherlands, annual tuberculosis case numbers declined from over 1500 in 1993 to less than 1000 in 2007 (Figure 1). The present study aimed to extend previous analysis [9] by covering a longer time period, exploring quantitative support for a cohort effect among natives, and assess the association of TB among the foreign-born with immigration figures. Specifically, we determine to what extent the decline of tuberculosis incidence in the period 1995-2007 was explained by the following three factors: (i) reduced TB rates attributable to endogenous reactivation in the native population, (ii) reduced importation, and (iii) reduced TB rates attributable to rapid progression after transmission within the Netherlands.

METHODS

TB is a notifiable disease in the Netherlands. Since 1993 information on risk factors and treatment outcome has been entered in the Netherlands Tuberculosis Register maintained by the KNCV Tuberculosis Foundation. Cross-matching with notification data suggested over 99% completeness up to 2003. Since 2004, notification and voluntary registration take place with a single internet-based system and discrepancies have been reduced to zero. For this study we used data on age, sex, year of diagnosis, pulmonary localisation of TB, direct sputum smear result for acid-fast bacilli, country of birth, urban residence, and risk group. We refer to those born in the Netherlands as the native population.

All *M. tuberculosis* complex isolates of patients diagnosed in the Netherlands are subjected to species identification, DNA fingerprinting and drug susceptibility testing. In the period 1993-2007 DNA fingerprinting was performed with standard restriction fragment length polymorphism (RFLP) typing using *IS6110* as a probe [11]. Strains with fewer than five *IS6110* copies in RFLP typing were subjected to subtyping with the PGRS probe [12].

Information in the two databases was matched using sex, date of birth, postal area code and year of diagnosis as identifiers.

In the period 1993-2007, 21,155 patients were diagnosed with tuberculosis, 14,818 (70%) were confirmed by culture. Matching patient data to DNA fingerprinting results yielded a total of 12,222 (82%) culture confirmed patients with information on the genotype of their *M. tuberculosis* isolate. Since tuberculosis is transmitted by patients with culture-positive pulmonary tuberculosis, the analysis of clustering was restricted to 8330 patients with pulmonary TB. Clusters were defined as groups of two or more patients who had isolates with identical DNA fingerprints at any time in the period 1993-2007. For each calendar year we determined the population strain diversity of *M. tuberculosis* as the number of different RFLP patterns divided by the total number of isolates [13].

Previously, we defined the transmission index as the number of secondary cases per potential source case using clustering over the complete study period [14,15]. As the study duration gets longer, this approach is unsatisfactory, since the duration of follow-up is shorter for potential source cases occurring late in the study period than for those occurring early. Moreover, the attribution of secondary cases to the potential source case becomes increasingly inaccurate with time, because of propagated transmission by second and further generation source cases within the cluster.

In order to choose a cut-off point for the definition of recent transmission we determined the probability that an isolate would be followed by one with an identical genotype with Kaplan-Meier survival analysis [7,9]. The Kaplan-Meier probability of finding another case with identical fingerprint within 5 years was 0.40 and within two years was 0.33, i.e., 83% of the 5

year probability. Therefore, we defined index cases as patients with strains not seen in other patients in the previous two years [9]. Most of these index patients are likely to have had reactivation or imported disease [16]. The transmission index was defined as the number of secondary cases within two years per index case. In a sensitivity analysis we explored the consequences of using a one or three year period to define recent transmission.

The trend of the incidence of native index cases was used as an indicator of secular trend, while among the foreign-born we determined the association with recent immigration figures. The number of secondary cases was used as an indicator of TB attributable to recent transmission. In this analysis we excluded index cases in 1993 and 1994 and their secondary cases (in total 1867), since we were unable to determine whether strains of these index cases had not been observed in the previous two years. We excluded 687 index cases and their secondary cases in 2006-2007, since we were unable to follow these index cases for a full two year period. Finally we excluded 591 patients occurring less than two years after a previous patient with that fingerprint, but more than two years after the start of a cluster in 1995-2005. Thus, 5185 patients remained: 3624 non-clustered index patients, 574 index patients who were the first patient of a cluster and 987 secondary cases within two years after the start of the cluster. Because in the sensitivity analysis we included the effect of a 3 year period to define recent transmission, we restricted this to index patients in the period 1996-2004 and their secondary cases.

Population denominators and immigration figures were obtained from the Central Bureau of Statistics (statline.cbs.nl/StatWeb/ accessed on 20 May 2009). Countries were classified by total TB incidence (<50, 50-199, \geq 200/100,000), estimated by the World Health Organization [17].

Person-years were estimated by the mid-year population. We determined the incidence of index cases and identified associated risk factors with Poisson regression for the native and foreign-born population [18]. We estimated to what extent the reduced incidence of native index cases may be attributed to a cohort effect, by relating the decline in index cases to the estimated decline in numbers of people with latent tuberculous infection (LTBI). The age-specific proportion of people with LTBI was estimated by Styblo, assuming that the risk of infection is independent of age and sex, and that the risk of infection since 1988 would have a negligible effect on the age specific prevalence of infection [19]. The expected number of native index cases in 2005 was estimated given the number of native index cases by age in 1995, and the decline in the number of people with LTBI. The expected number of index cases was then compared with the observed number. We used Poisson regression, with an offset of one for each index case, to identify characteristics of index cases associated with the number of secondary cases [18].

RESULTS

Clustering

Of the isolates, 56% (1756/3116) were clustered among native patients and 50% (2594/5214) among foreign-born patients (Table 1). Among natives the incidence of non-clustered pulmonary tuberculosis declined by 4.9% per year (95% confidence interval (CI) 3.7-6.1%) and of clustered tuberculosis by 3.7% (95% CI 2.7-4.8%). Among the foreign-born, the incidence of non-clustered tuberculosis declined by 1.9% per year (95% CI 1.0-2.7%) and that of clustered tuberculosis by 3.1% (95% CI 2.2-4.0%). Population strain diversity in each year was on average 0.81 and did not change over time (slope 0.0004, 95% CI -0.002 to 0.003).

We observed 4936 strains among 8330 patients. The probability that two randomly selected patients had the same strain was 0.0008.

Trends in incidence of native index cases

The incidence of native index cases was on average 0.9 per 100,000. The incidence was 1.7 times higher among men than women, and increased steeply with age (Table 2). The annual decline was 6% overall, and depended on the age of the index patient: in the age group 75+ it was 8 % (95% CI 5-10%), in age group 65-74 years 11% (95% CI 7-15%), 45-64 years 3% (95% CI 0-7%), and <45 years 5% (95% CI 2-8%).

From 1995 to 2005 the estimated size of the native population with LTBI declined from 1.70 to 0.98 million (Table 3). The expected and observed numbers of index cases in 2005 were 91 and 91, respectively (standardized ratio 1.00, 95% CI 0.81-1.23). The observed number of cases was higher than expected in those aged less than 65 years (standardized ratio 1.79, 95% CI 1.33-2.35) and lower than expected in those aged 65 years and more (standardized ratio 0.65, 95% CI 0.47-0.88).

Trends in incidence of foreign-born index cases

Among the foreign-born, the incidence of index cases was 17.6 per 100,000. The crude incidence ratio of foreign-born versus those born in the Netherlands was 20.4 (95% CI 19.1-21.7). The incidence was 1.7 times higher among men than women, and was highest in the age group 15-34 years (Table 2). Among the foreign-born, the incidence of index cases declined with 2% per year, irrespective of age. The number of immigrants in the period 1995-2005 from countries with estimated incidences of <50, 50-199, and \geq 200/100,000 was 318,000, 284,000 and 120,000 respectively. The number of index cases among the foreign-

born was significantly associated with the number of immigrants in the same year from countries with an estimated incidence of ≥ 200 per 100,000 ($r=0.69$, 95% CI 0.14-0.91) For countries with an estimated incidence of ≥ 50 per 100,000 the association was not significant ($r=0.55$, 95% CI -0.08-0.86).

Trends in disease attributed to recent transmission

On average, an index case had 0.24 (95% CI 0.21-0.26) secondary cases within two years. This number was similar for native and foreign-born index cases. Most cases attributable to recent transmission were aged less than 45 years and were seen in clusters with index cases aged less than 45 years (Figure 2).

Among native index cases, the number of secondary cases did not change over time, was higher for male than female index cases, declined with age, and was increased if the index case had smear-positive tuberculosis, lived in an urban setting, or abused alcohol (Table 4). In univariate analysis the number of secondary cases was increased if the index case was a drug user, but this was not significant after taking age and alcohol abuse into account.

Among the foreign-born, the number of secondary cases per index case declined with 3% (95% CI 0-5%, $p=0.045$) per year. The number of secondary cases declined with age of the index case, was increased if the index case was male, had smear-positive tuberculosis, lived in an urban area or was a drug user or homeless (Table 5). The number of secondary cases was much lower for index cases born in Asia than for those born outside Asia.

Sensitivity analysis

The incidence of index cases declined somewhat with an increasing time period to define recent transmission (Table 6). The transmission index tended to decline with an increasing time period among native index cases and to increase among the foreign-born. Time trends were not affected by the definition used (Table 6).

Summary results

From 1995 to 2005 the number of native index cases declined from 170 to 91 and their secondary cases within two years from 49 (32 native, 17 foreign-born) to 16 (10 native, 6 foreign-born). The number of foreign-born index cases declined from 250 to 222 and their secondary cases within two years from 53 (13 native, 40 foreign-born) to 37 (10 native, 27 foreign born). Thus, of the total reduction of 202 cases from 1995 to 2005, 42% was accounted for by fewer native index cases, 23% by fewer foreign-born index cases, 24% by fewer secondary cases from native index cases, and 11% by fewer secondary cases from foreign-born index cases. The proportion of native patients with TB attributable to recent transmission with a foreign-born index case increased from 13/45 (29%) in 1995 to 10/20 (50%) in 2005 (difference 21%, 95% CI 5 – 45%).

DISCUSSION

This study has shown that tuberculosis rates in the Netherlands declined in the period 1995-2005 as the result of a reduced incidence of index cases, while maintaining a low number of secondary patients per index case. The decline of index cases was stronger for natives than for the foreign-born. The average number of secondary cases per foreign-born index case declined slightly.

The reduced incidence of native index cases may be attributed mainly to a cohort effect as suggested previously [9]. This study provides quantitative support for this explanation, since the trend over time of the incidence of native index cases was fairly well predicted by the estimated time trend of the prevalence of LTBI. Discrepancies may have resulted from uncertainties in the age specific LTBI prevalence estimates, changes in the rate of progression to disease, and from the role of second generation immigrants, who have higher tuberculosis rates and presumably higher LTBI rates than other natives.

The incidence of index cases among the foreign-born was associated with the number of immigrants from high incidence countries in the same year. This was expected because incidence is highest during the first year after entry [20]. An important intervention to reduce tuberculosis incidence among immigrants is screening for and treatment of latent infection [4]. This strategy is not routinely implemented in the Netherlands although screening for active TB is in place.

Risk factors among index cases for a relatively high number of secondary cases such as young age, male sex, and smear-positivity were consistent with earlier findings [9,14,15,21,22]. Transmission among young adults is striking and may reflect they are socially active and tend to mix with those similar in age [23]. Among immigrants, those born in Indonesia and other Asian countries had a significantly lower transmission index. It will be of interest to determine whether the lower number of secondary cases among Asian index cases is attributable to lower infectiousness of the index patients, lower rates of rapid progression among contacts, host adaptation of the strains circulating in Asia [24], social mixing, or a combination of these factors. High rates of transmission among the homeless and drug users

have been attributed to low health care utilization and difficulties with contact investigation, necessitating active case finding in this group [25].

The contribution of foreign-born patients to transmission was not surprising in comparison with other settings. In San Francisco, using more detailed contact investigations, 2/19 (11%) of cases attributed to recent transmission were attributed to a foreign-born source [26]. Using comprehensive DNA-fingerprinting results, it was estimated that in San Francisco 61/(89+61) (41%) of US-born secondary cases were attributable to a foreign-born index case [27]. As in Norway [13], where 23/75 (31%) native secondary cases were attributed to recent transmission from a foreign-born index case, we observed no overall increase in cases attributable to recent transmission, despite the continuing influx of tuberculosis through immigration.

Clustering percentages and the proportion of patients attributed to recent transmission are underestimated if case finding is incomplete [6,28]. While case notification in the Netherlands is over 85% complete [29], failure to perfectly match the DNA fingerprint database to the NTR led to loss of 18% of registered patients. Moreover, in approximately 5% of patients with disease due to recent transmission, a transposition may have resulted in a slightly different RFLP pattern [30,31]. On the other hand, transmission may have been overestimated, since not all clustering based on RFLP typing may represent recent transmission. However, in a study in Amsterdam, with intensive efforts to identify epidemiological links, 86% of clustered patients were found to have such links [32]. Since the declining trend of all tuberculosis patients 1995-2005 was somewhat steeper than that of patients included in the study (Figure 1), our trend estimates may be slightly underestimated. The definition of TB attributable to rapid progression versus reactivation of latent infection

varies from one to five years between authors [1,2,7,33]. We have used a two-year period and found that time trends were not sensitive to this definition.

In conclusion, over the past 15 years, substantial progress has been made towards TB elimination in the Netherlands. However, given that an increasing proportion of TB cases is foreign-born, elimination of tuberculosis is unlikely with current tools and current levels of immigration. In order to accelerate progress towards elimination, the TB program in the Netherlands needs to explore strategies to expand the diagnosis and treatment of LTBI among the foreign-born. Furthermore, since global control of tuberculosis may lead to lower tuberculosis rates among immigrants [34], global tuberculosis control should be strongly supported by low incidence countries such as the Netherlands.

Major barriers for expanding the diagnosis and treatment of LTBI are the limited validity of diagnostics for predicting disease, side effects of current regimens, the risk of selecting for drug resistance, and the logistics and cost to ensure compliance. Therefore, there is a need to invest in research and development of better diagnostics to identify individuals with a high risk to progress to disease to allow focused preventive treatment, new drugs for preventive treatment and a post-exposure vaccine which would be of great benefit to eliminate tuberculosis in low incidence countries.

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References

1. Styblo K. Epidemiology of tuberculosis. Selected papers vol 24. The Hague: KNCV Tuberculosis Foundation, 1991.
2. Rieder H. Epidemiologic basis of tuberculosis control. Paris: International Union Against Tuberculosis and Lung Disease, 1999.
3. Lillebaek T, Dirksen A, Baess I, Strunge B, Thomsen VØ, Andersen AB. Molecular evidence of endogenous reactivation of *Mycobacterium tuberculosis* after 33 years of latent infection. *J Infect Dis* 2002;185:401-4.
4. Geng E, Kreiswirth B, Driver C, Li J, Burzynski J, DellaLatta P, LaPaz A, Schluger NW. Changes in the transmission of tuberculosis in New York City from 1990 to 1999. *N Engl J Med* 2002;346:1453-8.
5. Small PM, Hopewell PC, Singh SP, Paz A, Parsonnet J, Ruston DC, Schechter GF, Daley CL, Schoolnik GK. The epidemiology of tuberculosis in San Francisco. A population-based study using conventional and molecular methods. *N Engl J Med* 1994;330:1703-9.
6. Glynn JR, Vynnycky E, Fine PE. Influence of sampling on estimates of clustering and recent transmission of *Mycobacterium tuberculosis* derived from DNA fingerprinting techniques. *Am J Epidemiol* 1999;149:366-71.
7. Jasmer RM, Hahn JA, Small PM, Daley CL, Behr MA, Moss AR, Creasman JM, Schechter GF, Paz EA, Hopewell PC. A molecular epidemiologic analysis of tuberculosis trends in San Francisco, 1991-1997. *Ann Intern Med* 1999;130:971-8.
8. Iñigo J, Arce A, Palenque E, García de Viedma D, Chaves F. Decreased tuberculosis incidence and declining clustered case rates, Madrid. *Emerg Infect Dis* 2008;14:1641-3.
9. Borgdorff MW, van der Werf MJ, de Haas PE, Kremer K, van Soolingen D. Tuberculosis elimination in the Netherlands. *Emerg Infect Dis* 2005;11:597-602.

10. France AM, Cave MD, Bates JH, Foxman B, Chu T, Yang Z. What's driving the decline in tuberculosis in Arkansas? A molecular epidemiologic analysis of tuberculosis trends in a rural, low-incidence population, 1997-2003. *Am J Epidemiol* 2007;166:662-71.
11. Van Embden JD, Cave MD, Crawford JT, Dale JW, Eisenach KD, Gicquel B, Hermans P, Martin C, McAdam R, Shinnick TM. Strain identification of *Mycobacterium tuberculosis* by DNA fingerprinting: recommendations for a standardized methodology. *J Clin Microbiol*. 1993;31:406-9.
12. Kremer K, van Soolingen D, Frothingham R, Haas WH, Hermans PW, Martín C, Palittapongarnpim P, Plikaytis BB, Riley LW, Yakrus MA, Musser JM, van Embden JD. Comparison of methods based on different molecular epidemiological markers for typing of *Mycobacterium tuberculosis* complex strains: interlaboratory study of discriminatory power and reproducibility. *J Clin Microbiol* 1999;37:2607-18.
13. Dahle UR, Eldholm V, Winje BA, Mannsåker T, Heldal E. Impact of immigration on the molecular epidemiology of *Mycobacterium tuberculosis* in a low-incidence country. *Am J Respir Crit Care Med*. 2007;176:930-5.
14. Borgdorff MW, Nagelkerke N, van Soolingen D, de Haas PE, Veen J, van Embden JD. Analysis of tuberculosis transmission between nationalities in the Netherlands in the period 1993-1995 using DNA fingerprinting. *Am J Epidemiol* 1998;147:187-95.
15. Borgdorff MW, Nagelkerke NJ, De Haas PE, Van Soolingen D. Transmission of tuberculosis depending on the age and sex of source cases. *Am J Epidemiol* 2001;154:934-943.
16. Vynnycky E, Borgdorff MW, van Soolingen D, Fine PE. Annual *Mycobacterium tuberculosis* infection risk and interpretation of clustering statistics. *Emerg Infect Dis* 2003;9:176-83.

17. World Health Organization. Global Tuberculosis Control: surveillance, planning, financing. Geneva: World Health Organization, 2008.
18. Breslow NE, Day NE. Statistical methods in cancer research. Lyon: International Agency for Research on Cancer, 1987.
19. Styblo K. The elimination of tuberculosis in The Netherlands. *Bull Int Union Tuberc Lung Dis* 1990;65:49-55.
20. Vos AM, Meima A, Verver S, Looman CW, Bos V, Borgdorff MW, Habbema JD. High incidence of pulmonary tuberculosis persists a decade after immigration, The Netherlands. *Emerg Infect Dis* 2004;10:736-9.
21. van Soolingen D, Borgdorff MW, de Haas PE, Sebek MM, Veen J, Dessens M, Kremer K, van Embden JD. Molecular epidemiology of tuberculosis in the Netherlands: a nationwide study from 1993 through 1997. *J Infect Dis* 1999;180:726-36.
22. Behr MA, Warren SA, Salamon H, Hopewell PC, Ponce de Leon A, Daley CL, Small PM. Transmission of *Mycobacterium tuberculosis* from patients smear-negative for acid-fast bacilli. *Lancet* 1999;353:444-9.
23. Robins AB. The age relationship of cases of pulmonary tuberculosis and their associates. *Am J Public Health* 1953;43:718-23.
24. Gagneux S, DeRiemer K, Van T, Kato-Maeda M, de Jong BC, Narayanan S, Nicol M, Niemann S, Kremer K, Gutierrez MC, Hilty M, Hopewell PC, Small PM. Variable host-pathogen compatibility in *Mycobacterium tuberculosis*. *Proc Natl Acad Sci U S A* 2006;103:2869-73.
25. de Vries G, van Hest RA, Richardus JH. Impact of mobile radiographic screening on tuberculosis among drug users and homeless persons. *Am J Respir Crit Care Med* 2007;176:201-7.

26. Chin DP, DeRiemer K, Small PM, de Leon AP, Steinhart R, Schechter GF, Daley CL, Moss AR, Paz EA, Jasmer RM, Agasino CB, Hopewell PC. Differences in contributing factors to tuberculosis incidence in U.S. -born and foreign-born persons. *Am J Respir Crit Care Med* 1998;158:1797-803.
27. Borgdorff MW, Behr MA, Nagelkerke NJ, Hopewell PC, Small PM. Transmission of tuberculosis in San Francisco and its association with immigration and ethnicity. *Int J Tuberc Lung Dis* 2000;4:287-94.
28. Murray M. Sampling bias in the molecular epidemiology of tuberculosis. *Emerg Infect Dis* 2002;8:363-9.
29. van Hest NA, Smit F, Baars HW, De Vries G, De Haas PE, Westenend PJ, Nagelkerke NJ, Richardus JH. Completeness of notification of tuberculosis in The Netherlands: how reliable is record-linkage and capture-recapture analysis? *Epidemiol Infect* 2007;135:1021-9.
30. de Boer AS, Borgdorff MW, de Haas PE, Nagelkerke NJ, van Embden JD, van Soolingen D. Analysis of rate of change of IS6110 RFLP patterns of *Mycobacterium tuberculosis* based on serial patient isolates. *J Infect Dis* 1999;180:1238-44.
31. Eilers PHC, van Soolingen D, Lan NTN, Warren RM, Borgdorff MW. Transposition rates of *Mycobacterium tuberculosis* IS6110 restriction fragment length polymorphism patterns. *J Clin Microbiol* 2004; 42: 2461–2464.
32. van Deutekom H, Hoijng SP, de Haas PE, Langendam MW, Horsman A, van Soolingen D, Coutinho RA. Clustered tuberculosis cases: do they represent recent transmission and can they be detected earlier? *Am J Respir Crit Care Med* 2004;169:806-10.
33. Sanchez MA, Blower SM. Uncertainty and sensitivity analysis of the basic reproductive rate. Tuberculosis as an example. *Am J Epidemiol* 1997;145:1127-37.

34. Schwartzman K, Oxlade O, Barr RG, Grimard F, Acosta I, Baez J, Ferreira E, Melgen RE, Morose W, Salgado AC, Jacquet V, Maloney S, Laserson K, Mendez AP, Menzies D. Domestic returns from investment in the control of tuberculosis in other countries. *N Engl J Med.* 2005;353:1008-20.

Figure 1. Trends of tuberculosis (all cases), tuberculosis with genotyping results, and pulmonary tuberculosis (PTB) with genotyping results among native and foreign-born patients in the Netherlands, 1993-2007

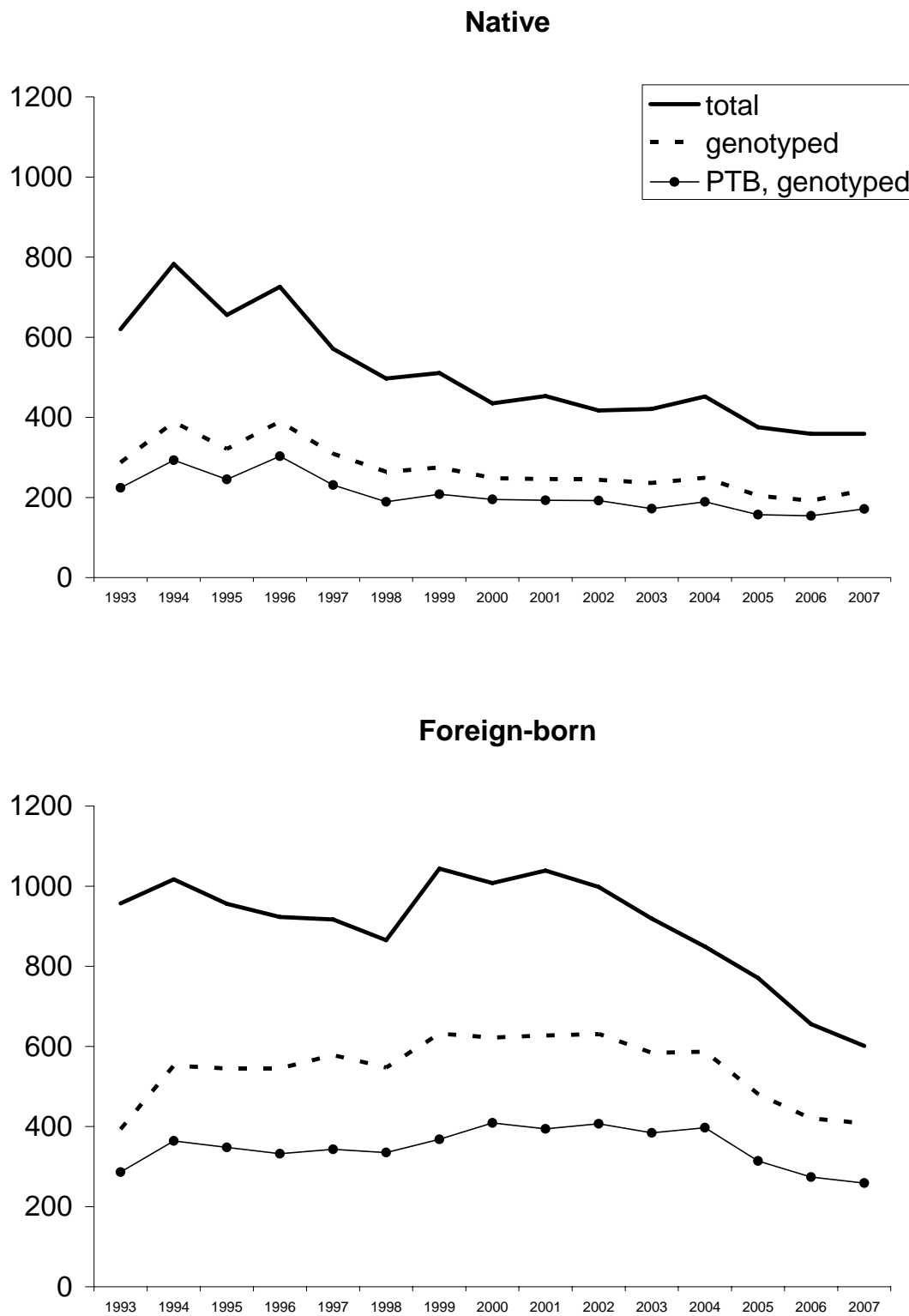


Figure 2. The age distribution of 987 secondary cases by age of their 4198 index cases in the Netherlands, 1995-2005.

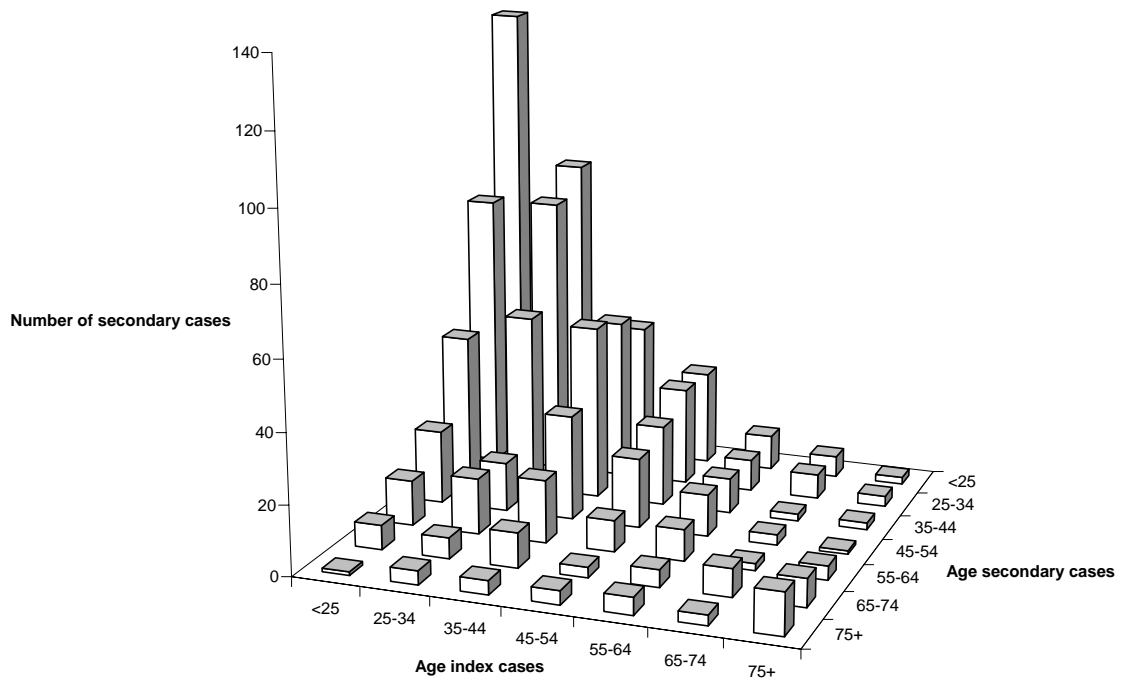


Table 1. Incidence of clustered and non-clustered pulmonary tuberculosis in the native and foreign-born population in the Netherlands, 1993-2007.

Native						
Year	Clustered	Non clustered	Person-years x 100,000	Rates per 100,000		% clustered
				Clustered	Non-clustered	
1993	127	97	141	0,9	0,7	57%
1994	163	130	141	1,2	0,9	56%
1995	116	129	142	0,8	0,9	47%
1996	178	125	142	1,3	0,9	59%
1997	111	120	143	0,8	0,8	48%
1998	102	87	143	0,7	0,6	54%
1999	132	76	144	0,9	0,5	63%
2000	115	80	145	0,8	0,6	59%
2001	119	74	145	0,8	0,5	62%
2002	120	72	146	0,8	0,5	63%
2003	88	84	146	0,6	0,6	51%
2004	110	79	147	0,7	0,5	58%
2005	98	59	147	0,7	0,4	62%
2006	83	71	147	0,6	0,5	54%
2007	94	77	148	0,6	0,5	55%
Total	1756	1360	3116	0,8	0,6	56%

Foreign born						
Year	Clustered	Non clustered	Person-years x 100,000	Rates		% clustered
				Clustered	Non-clustered	
1993	142	144	12	11,7	11,8	50%
1994	193	171	13	15,4	13,7	53%
1995	171	177	13	13,4	13,9	49%
1996	167	165	13	12,9	12,7	50%
1997	176	167	13	13,3	12,6	51%
1998	166	169	14	12,1	12,4	50%
1999	197	171	14	14,0	12,1	54%
2000	205	204	15	14,0	14,0	50%
2001	191	203	15	12,6	13,4	48%
2002	203	204	16	13,0	13,0	50%
2003	205	179	16	12,9	11,2	53%
2004	195	202	16	12,2	12,6	49%
2005	143	171	16	8,9	10,7	46%
2006	127	147	16	7,9	9,2	46%
2007	113	146	16	7,1	9,1	44%
Total	2594	2620	5214	12,0	12,1	50%

Table 2. Incidence of tuberculosis (TB) index cases in the native and foreign-born population in the Netherlands, 1995-2005

Native								
	Index cases	Person-years X 100.000	TB rate Per 100.000	Crude Rate ratio	95% CI	Adjusted Rate ratio	95% CI	
Year								
1995	170	142	1.2	0.94	0.93-0.96	0.94	0.92-0.95	
....	per year				
2005	91	147	0.6					
Sex								
male	865	787	1.1	1.73	1.55-1.93	2.12	1.90-2.37	
female	511	803	0.6	1		1		
Age group								
<15	22	312	0.1	0.13	0.08-0.19	0.13	0.08-0.20	
15-24	108	192	0.6	1		1		
25-34	137	230	0.6	1.06	0.82-1.37	1.05	0.81-1.35	
35-44	153	242	0.6	1.12	0.88-1.44	1.13	0.89-1.45	
45-54	136	222	0.6	1.09	0.85-1.40	1.10	0.85-1.42	
55-64	143	167	0.9	1.52	1.19-1.96	1.57	1.23-2.02	
65-74	209	125	1.7	2.98	2.37-3.77	3.11	2.47-3.93	
75+	468	100	4.7	8.33	6.79-10.32	9.52	7.75-11.80	
Total	1376	1590	0.9					
Foreign born								
	Index cases	Person-years X 100.000	TB rate per 100.000					
Year								
1995	250	13	19.6	0.98	0.97-0.99	0.98	0.97-1.00	
....	per year				
2005	222	16	13.8					
Sex								
male	1748	79	22.2	1.68	1.56-1.81	1.69	1.57-1.83	
female	1074	81	13.2	1		1		
Age group								
<15	72	12	6.2	0.19	0.15-0.24	0.19	0.14-0.24	
15-24	680	21	32.7	1		1		
25-34	957	38	24.9	0.76	0.69-0.84	0.76	0.69-0.84	
35-44	481	35	13.7	0.42	0.37-0.47	0.42	0.37-0.47	
45-54	250	25	10.0	0.31	0.26-0.35	0.31	0.26-0.35	
55-64	168	16	10.8	0.33	0.28-0.39	0.32	0.27-0.38	
65-74	132	8	16.3	0.50	0.41-0.60	0.51	0.42-0.62	
75+	82	6	14.4	0.44	0.35-0.55	0.50	0.39-0.62	
Total	2822	160	17.6					

Table 3. The prevalence of latent tuberculous infection (LTBI) and incidence of index cases in the native population in the Netherlands in 1995 and 2005.

	index cases observed 1995	Population with LTBI x 100,000		index cases expected 2005	index cases Observed 2005
	(i)	1995	2005	(i) x (iii) / (ii)	
		(ii)	(iii)		
<15	2	0.01	0.00	0.5	2
15-	11	0.05	0.02	3.7	6
25-	16	0.20	0.05	4.4	9
35-	17	0.53	0.20	6.3	8
45-	19	1.44	0.52	6.8	16
55-	14	3.27	1.36	5.8	9
65-	33	5.45	2.84	17.2	8
75+	58	6.08	4.83	46.1	33
TOTAL	170	17.02	9.82	91.0	91

The age specific proportion with LTBI was obtained from reference [18], the population denominator from the Central Bureau of Statistics.

Table 4. Risk factors among native index cases for number of secondary cases within two years.

	Index	Sec. cases <2 years	Crude		Adjusted		
			Transmission index (TI)*	relative TI	95% CI	relative TI	95% CI
Year							
1995	170	49	0.29	1.00	0.96-1.03		
....	per year			
2005	91	16	0.18				
Sex							
Male	865	218	0.25	1.34	1.06-1.71	1.44	1.13-1.84
Female	511	96	0.19	1		1	
Age group (yr)							
<15	22	13	0.59	1.33	0.70-2.33		
15-24	108	48	0.44	1.00	0.68-1.45		
25-34	137	61	0.45	1			
35-44	153	52	0.34	0.76	0.53-1.10	0.75	0.71-0.79
45-54	136	65	0.48	1.07	0.76-1.52	per age group	
55-64	143	26	0.18	0.41	0.25-0.64		
65-74	209	22	0.11	0.24	0.14-0.38		
75+	468	27	0.06	0.13	0.08-0.20		
Residence urban	305	118	0.39	2.11	1.68-2.65	1.43	1.13-1.81
Rural	1071	196	0.18	1		1	
Previous TB	289	36	0.12	0.49	0.34-0.68		
No	1087	278	0.26	1			
Smear positive	772	221	0.29	1.86	1.47-2.38	1.51	1.18-1.93
No	604	93	0.15	1		1	
HIV infection	36	12	0.33	1.48	0.78-2.51		
No	1340	302	0.23	1			
Alcohol abuse	37	19	0.51	2.33	1.42-3.60	1.94	1.17-3-01
No	1339	295	0.22	1		1	
Drugs	28	12	0.43	1.91	1.02-3.25		
No	1348	302	0.22	1			
Homeless	16	3	0.19	0.82	0.20-2.14		
No	1360	311	0.23	1			

* Transmission index is calculated as number of secondary cases divided by number of index cases.

Table 5. Risk factors among foreign-born index cases for number of secondary cases within two years in the Netherlands, 1995-2005

	Index	Sec. cases <2 years	Crude		Adjusted		
			Transmission index (TI)*	relative TI	95% CI	relative TI	95% CI
Year							
1995	250	53	0.21	0.97	0.95-0.99	0.97	0.95-1.00
....	per year			
2005	222	37	0.17				
Sex							
male	1748	475	0.27	1.47	1.25-1.74	1.29	1.09-1.53
female	1074	198	0.18	1		1	
Age group							
<15	72	19	0.26	1.28	0.77-2.00		
15-24	680	215	0.32	1.54	1.27-1.86		
25-34	957	197	0.21	1			
35-44	481	145	0.30	1.46	1.18-1.81	0.85	0.79-0.90
45-54	250	49	0.20	0.95	0.69-1.29		
55-64	168	34	0.20	0.98	0.67-1.39	per age group	
65-74	132	9	0.07	0.33	0.16-0.61		
75+	82	5	0.06	0.30	0.11-0.65		
Country of birth							
Somalia	308	111	0.36	4.10	2.95-5.79	3.72	2.66-5.27
Morocco	329	97	0.29	3.36	2.39-4.77	3.36	2.38-4.78
Turkey	239	78	0.33	3.72	2.61-5.34	3.52	2.47-5.07
Surinam	116	43	0.37	4.22	2.79-6.36	3.90	2.55-5.96
Indonesia	207	14	0.07	0.77	0.41-1.36	0.99	0.52-1.75
Former Soviet Union	91	19	0.21	2.38	1.37-3.97	2.50	1.44-4.18
Other Africa	572	158	0.28	3.15	2.30-4.38	2.72	1.99-3.80
Other Asia	558	49	0.09	1		1	
Other	331	92	0.28	3.17	2.25-4.51	2.92	2.07-4.17
Unknown	71	12	0.17	1.92	0.98-3.49	2.09	1.06-3.80
Residence urban	1022	302	0.30	1.43	1.23-1.67	1.25	1.06-1.47
rural	1800	371	0.21	1		1	
Previous TB	181	44	0.24	1.02	0.74-1.37		
no	2641	629	0.24	1			
Smear positive	1515	462	0.30	1.89	1.61-2.23	1.73	1.47-2.05
no	1307	211	0.16	1		1	
HIV infection	207	46	0.22	0.93	0.68-1.24		
no	2615	627	0.24	1			
Alcohol abuse	19	3	0.16	0.66	0.16-1.72		
no	2803	670	0.24	1			
Drug abuse and homeless	11	22	2.00	8.91	5.65-13.3	7.15	4.49-10.8
Homeless only	36	24	0.67	2.97	1.92-4.36	2.73	1.76-4.05
Drug abuse only	39	13	0.33	1.49	0.81-2.46	1.17	0.64-1.96
Neither	2736	614	0.22	1		1	

* Transmission index is calculated as number of secondary cases divided by number of index cases.

Table 6. Sensitivity analysis with varying definitions of recent transmission (index cases 1996-2004)

Recent transmission (within x years):	Native			Foreign-born		
	X=1	X=2	X=3	X=1	X=2	X=3
Index cases	1252	1115	1064	2563	2350	2237
Secondary cases	432	304	286	852	855	908
Incidence index cases per 100,000	0.96	0.86	0.82	19.49	17.87	17.01
Transmission index	0.35	0.27	0.27	0.33	0.36	0.41
Time trend index cases	0.94	0.95	0.95	0.99	0.99	0.99
Time trend transmission index	1.07	1.07	1.08	0.91	0.91	0.90