



# Changing epidemiology of nontuberculous mycobacterial lung disease over the last two decades in a region of the Basque country

Sandra Pedrero<sup>1</sup>, Eva Taberbero<sup>1</sup>, Eunáte Arana-Arri<sup>2</sup>, Elena Urra<sup>3</sup>,  
Maialen Larrea<sup>3</sup> and Rafael Zalacain<sup>1</sup>

**Affiliations:** <sup>1</sup>Dept of Respiratory Medicine, Hospital Universitario Cruces, Barakaldo, Spain. <sup>2</sup>Research Unit, Hospital Universitario Cruces, Barakaldo, Spain. <sup>3</sup>Dept of Microbiology, Hospital Universitario Cruces, Barakaldo, Spain.

**Correspondence:** Sandra Pedrero, Dept of Respiratory Medicine, Hospital Universitario Cruces, Plaza de Cruces, S/N, 48903 Barakaldo, Bizkaia, Spain. E-mail: [sandra.pedrerotejada@osakidetza.eus](mailto:sandra.pedrerotejada@osakidetza.eus)

**ABSTRACT** Recent studies suggest an increasing prevalence of nontuberculous mycobacteria (NTM) lung disease. The aim of the present study was to describe incidence rates of NTM lung disease and trends therein in our area over a 20-year period.

This was a retrospective study of all cases of NTM lung disease between 1997 and 2016 that met the 2007 American Thoracic Society criteria. We analysed the annual incidence rates, species of mycobacteria isolated, trends over time and annual mortality in 327 patients.

*Mycobacterium kansasii* was the most common mycobacterium isolated (84%), followed by *Mycobacterium avium* complex (MAC) (13%). We compared two periods: 1997–2006 (257 cases, 79%) and 2007–2016 (70 cases, 21%). The incidence rates tended to decrease across these years, with a peak of incidence in 2000 with 10.6 cases per 100 000. There was a clearly decreasing trend in *M. kansasii* infection, not only in the first period (incident rate ratio (IRR) 0.915, 95% CI 0.88–0.90;  $p < 0.0001$ ) but also in the second (IRR 0.869, 95% CI 0.780–1.014;  $p = 0.080$ ), reaching 1.8 per 100 000 in 2016. In contrast, MAC infection tended to increase across the two periods (IRR 1.251, 95% CI 1.081–1.447;  $p = 0.003$ ).

In our region, the incidence of NTM lung disease has notably decreased in recent years. *M. kansasii* had high incidence rates in the first decade but clearly decreased in the second decade.



@ERSpublications

**This epidemiological study shows a clear decrease in the incidence of nontuberculous mycobacterial lung disease, unlike other studies that have described an overall increase** <http://bit.ly/2ZH46nv>

**Cite this article as:** Pedrero S, Taberbero E, Arana-Arri E, *et al.* Changing epidemiology of nontuberculous mycobacterial lung disease over the last two decades in a region of the Basque country. *ERJ Open Res* 2019; 5: 00110-2018 [<https://doi.org/10.1183/23120541.00110-2018>].



## Introduction

Lung disease due to nontuberculous mycobacteria (NTM), that is, environmental mycobacteria, is considered an emerging clinical entity [1]. Knowledge of this disease has improved thanks to advances in radiological and microbiological techniques, in particular, molecular identification methods, but there are still numerous uncertainties regarding both clinical management of the disease and the aetiopathogenic and other factors underlying the epidemiological differences between geographical areas.

Unlike tuberculosis, lung infection due to NTM is not a reportable disease, and hence, epidemiological data are not accurate, but most publications describe an increase in prevalence. On the other hand, diagnosis and epidemiological research are difficult as NTM are ubiquitous in the environment and their isolation in the laboratory does not necessarily imply disease [2, 3].

Environmental conditions and individual susceptibility play an important role in the prevalence of lung disease due to NTM, with areas of high disease risk and great variability in the causative species by geographical area. According to recent studies, the areas with the highest prevalence of the disease are regions with a high population density, a high level of education and a large proportion of the area covered by surface water with high evapotranspiration [4].

The importance of environmental mycobacteria lies in the fact that they require long antibiotic treatments with several drugs, generally for more than 1 year, with high associated costs and high toxicity [5]. Excess mortality has also been described in association with infection by NTM [6].

Thanks to the experience gained over 20 years in a tuberculosis control project at our hospital, communication and close cooperation between clinicians and the microbiology department have been strengthened in our daily work with scheduled information flow. Between 1997 and 2004, we detected a large number of cases of lung disease due to NTM, and since then, this condition has continued to be analysed.

The objective of this study was to describe the epidemiological pattern of lung disease due to NTM in our area, Bizkaia, and changes therein over the last 20 years. Bizkaia is a coastal and mountainous province of the Basque Country located in the north of Spain. It is an area with a high population density and is very industrialised, traditionally with chemical and metallurgical (especially steel) industries, though these have been progressively decreasing over the last 20 years. This area has not experienced significant population changes in the period studied.

## Methods

A retrospective study was conducted including all cases of lung disease due to NTM that met the current American Thoracic Society (ATS) criteria in people over 18 years old diagnosed between 1997 and 2016 in Bizkaia. The epidemiological data were provided by the Microbiology Laboratory of Cruces University Hospital, which has a catchment population of 420 000. We analysed the annual incidence rates (IRs), species of mycobacteria isolated, trends in each species over time and annual mortality.

The inclusion criteria used for defining a case as “lung disease” were those established by the ATS in 2007 [3]. That is, we included all patients with mycobacterium isolates who met clinical, radiological and bacteriological criteria as follows: 1) presence of compatible symptoms (cough, fever, weight loss, haemoptysis or dyspnoea) not attributable to other diseases; 2) characteristic findings on chest radiography (infiltrate, nodules and cavitation) or high-resolution computed tomography scan (multifocal bronchiectasis with multiple nodules); and 3) positive culture results from at least two separate expectorated sputum samples or one bronchial wash or lavage or lung biopsy sample. Extrapulmonary samples were not considered. We excluded patients with a single isolate or who had no symptoms or radiological signs. Radiological and clinical data were obtained from the unified electronic health record, which extends back to 1995.

### Microbiological analysis

The methods for identifying mycobacteria have changed substantially over the last two decades, except for culture techniques (BACTEC MGIT 960 and Lowenstein–Jensen medium) that have continued to be used throughout the study period. In the first 10 years, phenotypic methods were used, involving the assessment of growth rates and optimal temperatures, pigment production, colony morphology and biochemical features. In contrast, in recent years of the study period, other techniques have been adopted such as nucleic acid probes (AccuProbe, Gen-Probe); *hsp65* gene PCR amplification followed by restriction fragment length polymorphism analysis using *Bst*EII and *Hae*III enzymes; agarose gel electrophoresis (PCR restriction fragment length polymorphism assays); analysis of the sequence of the *hsp 65* fragment; line probe assays (GenoType, Hain Lifescience, used since 2015); and matrix-assisted laser desorption/ionisation time-of-flight mass spectrometry (MALDI-TOF, used since 2014).

### Statistical analysis

The inter- and intra-group comparisons were carried out using parametric or nonparametric tests depending on the characteristics of the data distribution. Qualitative variables expressed as proportions or percentages were compared with Chi-squared or Fisher's exact test. For perioperative and follow-up variables, means were compared with t-tests if data were normally distributed, and otherwise, with the Mann-Whitney U-test. The IR for each mycobacteria species was calculated as the number of new cases divided by the number of people at risk per 100 000 per year for the period 1997–2016. A Poisson regression model was used to analyse the incidence rate ratios (IRRs) by species and period of time included in the model, as well as to estimate temporal trends in the overall IRs. Results are reported as frequency rates (95% CI). A p-value  $\leq 0.05$  was considered statistically significant. Statistical analysis was carried out using IBM SPSS Statistics for Windows version 23.0 and STATA<sup>®</sup> 13.

### Ethical considerations

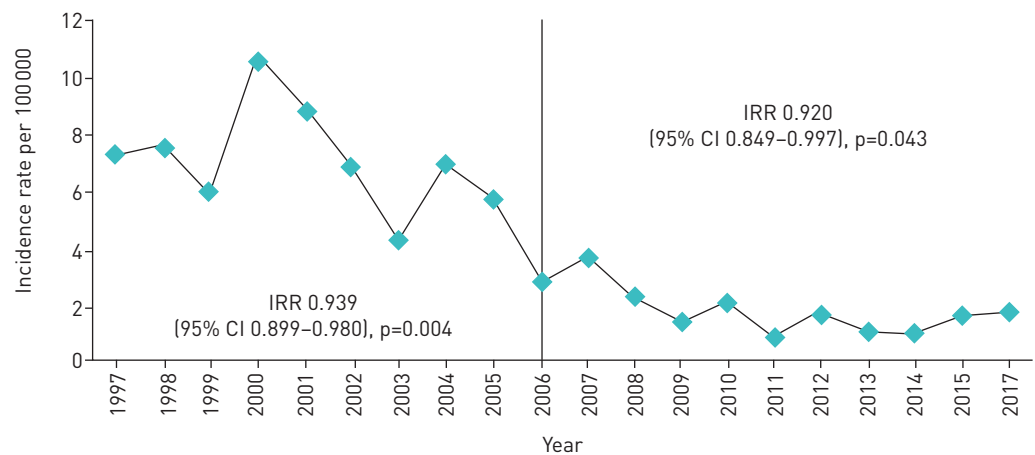
The study complied with the principles of the latest version of the Declaration of Helsinki and international regulations. Measures were taken to ensure the confidentiality and security of personal information concerning research participants.

### Results

A total of 327 patients were diagnosed with NTM lung disease during the 20 years studied: 274 cases were attributed to *M. kansasii* (83.8%), 43 to *M. avium* complex (13.1%), and 10 to other mycobacteria such as *M. xenopi*, *M. celatum*, and *M. abscessus* (3.1%). Any patients with a diagnosis of infection by *M. avium* complex, or specifically *M. intracellulare*, *M. avium* or *M. chimaera*, were classified as infected by MAC. All the patients met ATS diagnostic criteria for NTM lung disease and 85.6% of them were treated (89.4% of cases of *M. kansasii* and 58.1% of cases of *M. avium* complex).

The overall IRs of lung disease due to NTM over the 20 years of study are shown in figure 1. The highest IRs were recorded in 2000 and 2001, with 10.6 and 8.8 cases per 100 000, respectively. It can be observed that the IRs significantly decreased over the years to 1.8 cases per 100 000 in 2016. In this context, the high IRs during the first few years of the study are striking. Hence, we decided to compare two 10-year periods, after observing a very marked decrease in the first decade, compared with the decrease in the second decade. The first decade from 1997 to 2006, in which there were 257 cases (79%), and the second, from 2007 to 2016, in which there were 70 cases (21%). We confirmed a decreasing trend in both decades, with a yearly average decrease of 6.4% (IRR=0.939, 95% CI 0.899–0.980;  $p=0.004$ ) in the first decade, and of 6.2% (IRR=0.920, 95% CI 0.849–0.997;  $p=0.043$ ) in the second decade.

Regarding the species involved, there was a clearly decreasing trend in *M. kansasii* over the course of the period studied, with an overall IRR of 0.858 (95% CI=0.835–0.881;  $p<0.0001$ ) and an annual decrease of 6.5%. In contrast, there was a significant increasing trend in MAC in both decades, with an overall IRR of 1.133 (95% CI=1.058–1.213;  $p<0.0001$ ) and an annual rise of 3.3% (figure 2).



**FIGURE 1** Overall incidence rates per 100 000 population of lung disease due to nontuberculous mycobacteria between 1997 and 2016. IRR: incident rate ratio.

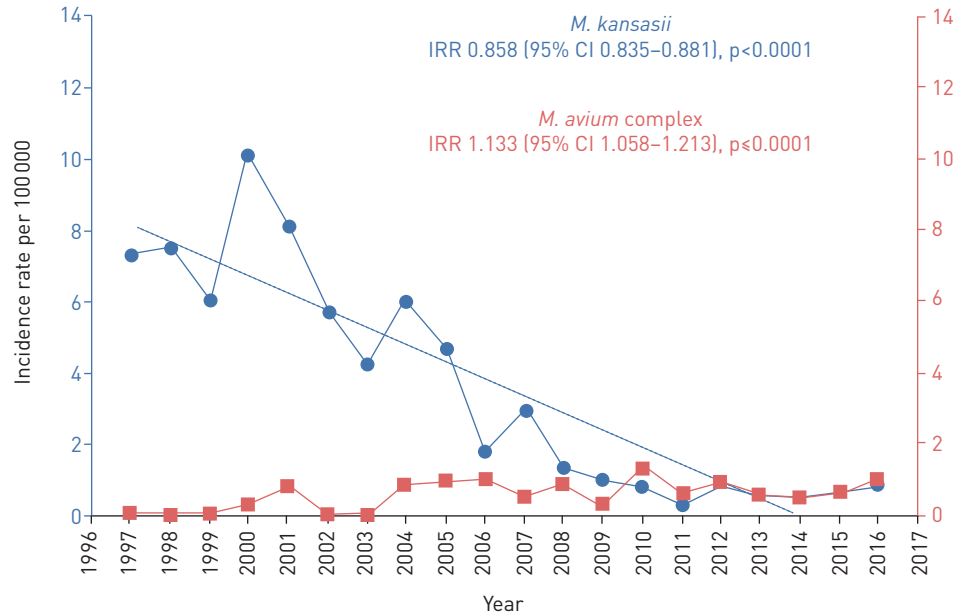


FIGURE 2 Incidence rates of lung disease due to nontuberculous mycobacteria between 1997 and 2016 by species involved: *Mycobacterium kansasii* and *Mycobacterium avium* complex. IRR: incident rate ratio.

Most patients with lung disease due to NTM were male (71.2%) and the mean age overall was 56 years old. Notably, in the first decade, patients were slightly younger (mean: 54 years *versus* 61 years in the second decade,  $p=0.01$ ) and more likely to be male (75.2% *versus* 57% in the second decade,  $p=0.03$ ) (table 1).

Concerning risk factors, no differences were found between the two decades in HIV, chronic obstructive pulmonary disease, history of lung disease, use of oral corticoids, alcohol use, liver disease, cancer or gastroesophageal reflux. On the other hand, the rate of active smoking was significantly higher in the first than the second decade (60.4% *versus* 44.8%;  $p=0.003$ ). Among radiological features, cavitation was the most common finding, especially in the first period, but no differences in radiological findings between the periods reached significance. Other findings included bronchiectasis, nodules and the tree-in-bud pattern. Notably, involvement was unilateral in almost half of the patients and overall upper lung lobes were the most common site of infection (table 2).

The overall annual mortality was 9.8% and the mortality attributed to mycobacteriosis was 1.8%, with no significant differences between study periods. Nevertheless, the microbiological cure rate was higher in the first decade (88.7% *versus* 70.1% in the second decade,  $p<0.001$ ) (table 3).

Analysing these data by species, we have noticed differences between *M. kansasii* and *M. avium* complex over the period studied. IRs per 100 000 population are shown in figure 2 by species. Concerning risk factors, no differences were found between NTM species except for the rates of active smoking and cancer,

TABLE 1 Characteristics of lung infections by nontuberculous mycobacteria (NTM), overall and in each decade

	Total	1st decade <sup>#</sup>	2 <sup>nd</sup> decade <sup>¶</sup>	p-value
<b>Patients with NTM lung disease</b>	327	257 (78.6%)	70 (21.4%)	<0.001
<b>NTM species</b>				
<i>Mycobacterium kansasii</i>	274 (83.8%)	236 (91.6%)	38 (54.3%)	<0.001
<i>Mycobacterium avium</i> complex	43 (13.1%)	15 (5.8%)	28 (40.0%)	
Others <sup>+</sup>	10 (3.1%)	6 (1.8%)	4 (1.2%)	
<b>Age years mean</b>	56.8±18	54±18.6	61±16.9	0.010
<b>Male</b>	71.2%	75.2%	57.1%	0.003

<sup>#</sup>: 1997–2006; <sup>¶</sup>: 2007–2016; <sup>+</sup>: *Mycobacterium celatum*, *Mycobacterium abscessus* and *Mycobacterium xenopi*.

TABLE 2 Risk factors, clinical and radiological characteristics of lung infections by nontuberculous mycobacteria, overall and in each decade

	Total	1st decade <sup>#</sup>	2nd decade <sup>¶</sup>	p-value
<b>Risk factors</b>				
HIV	12.2%	13.2%	8.6%	0.200
Smoking	53.6%	60.4%	44.8%	0.003
History of lung disease	56%	53%	60.6%	0.201
COPD	30.1%	34%	23.8%	0.113
Oral corticosteroid use	6.7%	8.2%	4.5%	0.283
Alcohol use	18.3%	20.2%	15.4%	0.285
Liver disease	10.1%	9.5%	11.1%	0.468
Cancer	22.6%	23.8%	20.6%	0.395
Gastroesophageal reflux	9.6%	11.5%	6.6%	0.233
<b>Signs and symptoms</b>				
Cough/expectoration	73.2%	75.3%	70.3%	0.308
Fever	40.4%	42.5%	37.3%	0.323
Asthenia	38%	36.1%	40.7%	0.354
Weight loss	24.1%	23.8%	24.6%	0.484
Haemoptysis	25.7%	25%	26.7%	0.484
<b>Radiological findings</b>				
Cavitation	37.9%	41.9%	32.8%	0.165
Bronchiectasis	29.3%	22.6%	38.1%	0.032
Nodules	28.4%	20.2%	39.1%	0.010
Tree-in-bud pattern	25.7%	15.7%	39.3%	0.001
<b>Lung involvement</b>				
Unilateral	45.7%	49.4%	40.9%	0.547
Upper lung lobes	52.4%	58.8%	43.3%	0.260
Middle lung lobes	15.2%	11.8%	20%	0.260
Lower lung lobes	15.2%	9.4%	6.7%	0.260
Multifocal	22.1%	18.8%	26.7%	0.260

<sup>#</sup>: 1997–2006; <sup>¶</sup>: 2007–2016.

which were significantly higher in the *M. kansasii* group (63.8% versus 31.0%;  $p < 0.0001$ , and 27.7% versus 9.5%;  $p = 0.011$ , respectively) (table 4).

Among radiological features, *M. kansasii* was associated with more cavitation, multifocal affectation and upper lobe predominance. On the other hand, *M. avium* complex was associated with bronchiectasis, nodules and tree-in-bud infiltrates with middle/lower lobe predominance and unilateral involvement. Microbiological cure rates were higher among patients with *M. kansasii* than MAC disease (89.0% versus 76.7%,  $p = 0.029$ ), with a lower overall 1-year mortality, including NTM-related and other-cause mortality.

## Discussion

The epidemiological data from our hospital show a clear decrease in the incidence of mycobacteriosis, especially since 2005, unlike other studies that have described a general increase in the incidence of this condition [7, 8]. In a study in Queensland, Australia, the incidence was observed to have increased from 2.2 cases per 100 000 in 1999 to 3.2 cases per 100 000 in 2005 [9]. Subsequently, in Oregon, USA, the

TABLE 3 Disease course and mortality from lung infections by nontuberculous mycobacteria (NTM), overall and in each decade

Disease course and mortality	Total	1st decade <sup>#</sup>	2nd decade <sup>¶</sup>	p-value
<b>Microbiological cure</b>	275 (84.1%)	228 (88.7%)	47 (70.1%)	<0.0001
<b>Overall mortality</b>	32 (9.8%)	22 (8.5%)	10 (14.3%)	0.153
<b>Deaths due to NTM lung disease</b>	6 (1.8%)	2 (0.8%)	4 (5.7%)	0.057
<b>Deaths due to other causes</b>	26 (79.5%)	20 (7.7%)	6 (8.6%)	0.339

<sup>#</sup>: 1997–2006; <sup>¶</sup>: 2007–2016.

TABLE 4 Comparison between *Mycobacterium kansasii* and *Mycobacterium avium* complex

	Total	<i>M. kansasii</i>	<i>M. avium</i> complex	p-value
<b>Patients with NTM lung disease</b>	317	274	43	
<b>Age years</b>				
Mean±SD	58.5±17.8	54.9±17.9	63.0±17.3	0.010
Median (IQR)	61 (44–73)	57 (42–69)	68 (47–78)	
<b>Male patients</b>	296 (70.8%)	204 (75.0%)	65 (60.2%)	0.003
<b>Risk factors</b>				
HIV	12.3%	11.3%	18.6%	0.136
Smoking	55.1%	63.8%	31.0%	<0.0001
History of lung disease	55.8%	53.9%	61.0%	0.276
COPD	32.0%	34.2%	26.2%	0.226
Oral corticosteroid use	6.5%	4.5%	11.9%	0.100
Alcohol use	18.7%	21.9%	9.8%	0.065
Liver disease	10.7%	9.3%	14.6%	0.252
Cancer	22.7%	27.7%	9.5%	0.011
Gastroesophageal reflux	10.2%	10.1%	10.5%	0.577
<b>Signs and symptoms</b>				
Cough/expectoration	73.6%	70.9%	80.5%	0.166
Fever	42.3%	39.6%	48.8%	0.209
Asthenia	37.6%	34.4%	45.0%	0.168
Weight loss	25.0%	22.8%	30.0%	0.253
Haemoptysis	26.7%	30.2%	17.9%	0.105
<b>Radiology</b>				
Cavitation	38.2%	48.1%	12.5%	<0.0001
Bronchiectasis	29.0%	19.2%	53.8%	<0.0001
Nodules	26.6%	22.2%	37.5%	0.053
Tree-in-bud pattern	25.0%	20.6%	35.9%	0.052
<b>Lung involvement<sup>#</sup></b>	49.7%	42.7%	67.5%	0.003
<b>Localisation</b>				
Upper lung lobes	53.4%	57.1%	42.9%	0.044
Middle lung lobes	15.0%	10.2%	28.6%	0.044
Lower lung lobes	8.3%	7.1%	11.4%	0.044
Multifocal	23.3%	25.5%	17.1%	0.044
<b>Disease course and mortality</b>				
Microbiological cure	87.3%	89.0%	76.7%	0.029
Overall mortality	9.8%	9.1%	14.0%	0.229
Deaths due to NTM lung disease	1.9%	1.5%	4.7%	0.189
Deaths due to other causes	7.9%	7.7%	9.3%	0.448

NTM: nontuberculous mycobacteria; IQR: interquartile range. <sup>#</sup>: all unilateral.

incidence of lung disease due to NTM was found to have increased from 4.8 cases per 100 000 in 2007 to 5.6 per 100 000 in 2012 [10], while over the same period, Shah *et al.* [11] observed an increase in pulmonary NTM isolates from 4.0 cases per 100 000 in 2007 to 6.1 cases per 100 000 in 2012 in England, Wales and Northern Ireland.

Our results also differ from those of a recent study in Asturias, Spain, which described a 35% increase in the number of NTM isolates over a period of 8 years, although there were no changes in the number of patients with clinically significant NTM disease [12]. The only study we have found in which, as in ours, no increase was observed in the incidence of NTM was conducted in Denmark [13].

We should emphasise that the incidence of lung disease due to NTM varies across studies due to differences in the study design, samples used, diagnostic criteria applied and classification of lung disease due to NTM *versus* colonisation, these differences making it difficult to compare studies [3]. Additionally, numerous publications report numbers of isolates and not cases of disease. In our series, both the number of isolates and the number of cases of lung disease due to NTM have tended to decrease over the years.

The highest rates of NTM infection were found in 2000 with 10.6 per 100 000 per year, decreasing to 1.8 per 100 000 per year in 2016. The incidence of lung disease due to *M. kansasii*, the leading causative agent of NTM, decreased over the study period, with 236 patients diagnosed with this disease (91.6%) in the first decade, and only 38 (54.3%) in the second period. This peak in incidence in 2005 was described

previously by a research group studying an overlapping geographical area [14], who attributed it to several factors yet to be demonstrated. In particular, the construction work for the underground train network (Metro Bilbao) might have led to there being more mycobacteria in the environment, while another nonexclusive explanation would be a change in water treatment, with more permissive temperatures and changes in water chlorination [15]. In this context, it has been stated that the main environmental niche for *M. intracellulare* appears to be in biofilms lining drinking water pipes and *M. abscessus* and *M. kansasii* have also been isolated from drinking water [9]. Another interesting possibility is that the high prevalence of *M. kansasii* coincided with a period of heavy industry, as has been also described in other studies [16, 18–].

Notably, in our study, despite the inclusion of individuals with structural lung damage who are more susceptible to infection, such as patients with cystic fibrosis or treated with immunosuppressive drugs (anti-tumour necrosis factor agents), and improvements in microbiological techniques in our hospital, we have not observed an increase in the prevalence of the disease. Indeed, NTM are considered emerging pathogens in patients with cystic fibrosis [19] and patients treated with anti-tumour necrosis factor agents [20], potentially causing significant morbidity and mortality.

Despite microbiological improvements, the increases observed in the detection of mycobacterial diseases do not coincide with the introduction of new microbiological techniques. In fact, newer techniques such as line probe assays (GenoType) and MALDI-TOF were introduced in 2014 and 2015 respectively, while the peaks of maximum incidence of NTM pulmonary disease were seen in 2000 and 2001.

On the other hand, this can be explained by the fact that in countries such as Spain, where the tuberculosis incidence is decreasing, the number of requested mycobacterial cultures also decreases and the awareness of mycobacterial disease as a differential diagnosis. In fact, in our study a decrease observed in the number of samples sent to the microbiology service was attributable to lower rates of suspicion of tuberculosis by clinicians. Specifically, the number of requests fell in the second decade to almost half (55%) that in the first decade of the study and this coincided with the decrease in cases of mycobacterial disease.

To our knowledge, this is the first study that indicates such a clear decrease in the incidence rate of mycobacterial infection, though it is important to emphasise that this is due to a decrease in *M. kansasii* specifically in our area. It is possible that the pattern observed between 1997 and 2004 in our setting was due to higher exposure to *M. kansasii*, the load subsequently decreasing; this underlines the potential importance of exposure to high loads of environmental mycobacteria in the aetiopathogenesis and development of lung disease due to NTM.

As noted in the Introduction, there is marked geographical variability in the species isolated, with differences even within the same country, considering coastal or inland areas [18]. In a recent study in 30 countries across six continents, the most commonly isolated pathogen was MAC [21], with *M. kansasii* being the sixth most isolated species, with higher rates in eastern Europe and the UK, as well as in France, in the centre of Paris, and South America. The leading causative agent of the disease in our region was *M. kansasii*, consistent with the findings of earlier research in Spain based on a small sample in which *M. kansasii* was the most commonly isolated species [22]. In contrast, in a more recent study in a region close to our study area, also coastal, the most common causative agent of NTM was MAC [12].

Similarly, a trend towards increasing MAC and decreasing *M. kansasii* was shown in a large multicentre study [23] and in a recent study in Spain [24], where the authors described a decline in *M. kansasii* infections progressively from the early 2000s onwards, MAC becoming the most frequently isolated mycobacteria. Nevertheless, in both studies, the total number of isolates increased.

As in other studies, lung disease due to NTM is common in patients with HIV, in whom these types of lung infections were first described [25]. Nevertheless, such infections are affecting a growing number of people without HIV (87.8% of patients in our series), this confirming the change in patient clinical profile noted previously [26, 27]. Patients who are immunocompetent commonly have underlying chronic conditions such as chronic obstructive pulmonary disease, a history of tuberculosis, a smoking habit [28] and/or other risk factors such as oral corticosteroid intake, alcohol use, liver disease, cancer and/or gastroesophageal reflux. In our series, it is notable that *M. kansasii* lung infections were significantly more likely to be in men and young people, which could be related to exposure to heavy industry.

Previous studies have demonstrated high rates of death due to NTM lung disease, but none have compared outcomes with matched population-based controls. Generally, research on mortality from this type of infection and associated factors is limited. Some studies have analysed overall 5-year mortality with very mixed findings, from 18% in a study from the USA [6] to 40% in Denmark [28]. More specifically, the aforementioned overall rate in Denmark was obtained in a comprehensive population-based study that

found death rates of 39.7% due to MAC lung disease and 51% for *M. xenopi* disease [28], while rates of 36.3% and 43.2% respectively were found in Ontario, Canada [29]. In our study, we found an overall 1-year mortality rate of 9.8%, and though the mortality rate was slightly higher in the second period of study (14.3% versus 8.5% in the first period), the difference was not significant. As in other studies, we found low rates of mortality associated specifically with mycobacteria, it being difficult to establish the cause of death due to the presence of comorbidities [28, 29].

It is plausible that, in our study, the differences in radiological and microbiological cure rates between the first and second period can be explained by the shift in the predominant NTM species, with *M. kansasii* being the most prevalent species in the first decade with higher cure rates.

### Limitations

This was a retrospective study with the disadvantages inherent to this type of study, in particular, concerning the quality of the data recorded for some clinical variables. Moreover, no information was gathered on the occupational history of the patients. Further, this study was carried out in the specific geographical area of Bizkaia, and hence, results cannot be extrapolated to other areas.

### Conclusions

In our region, the incidence of lung disease due to NTM has notably decreased in recent years with a shift in the predominant NTM species. The mycobacteria most commonly found in our region was *M. kansasii* with IRs that were high in the first decade of the study and tended to decrease in the second decade. Mycobacteriosis-associated mortality was low with high cure rates, especially in the first decade of study. Although our study is not able to confirm this, we believe that greater environmental exposure, as well as host susceptibility, may have contributed to the development of this condition.

Conflict of interest: None declared.

### References

- 1 Stout JE, Koh WK, Yew WW. Update on pulmonary disease due to non-tuberculous mycobacteria. *Int J of Infect Dis* 2016; 45: 123–134.
- 2 Larsson LO, Polverino E, Hoefsloot W, et al. Pulmonary disease by non-tuberculous mycobacteria-clinical management, unmet needs and future perspectives. *Expert Rev Respir Med* 2017; 11: 977–989.
- 3 Griffith DE, Aksamit T, Brown-Elliott BA, et al. An official ATS/IDSA statement: diagnosis, treatment, and prevention of nontuberculous mycobacterial diseases. *Am J Respir Crit Care Med* 2007; 175: 367–416.
- 4 Adjemian J, Olivier KN, Seitz AE, et al. Spatial clusters of nontuberculous mycobacterial lung disease in the United States. *Am J Respir Crit Care Med* 2012; 186: 553–558.
- 5 Ballarino GJ, Olivier KN, Claypool RJ, et al. Pulmonary nontuberculous mycobacterial infections: antibiotic treatment and associated costs. *Respir Med* 2009; 103: 1448–1455.
- 6 Fleshner M, Olivier KN, Shaw PA, et al. Mortality among patients with pulmonary non-tuberculous mycobacteria disease. *Int J Tuberc Lung Dis* 2016; 20: 582–587.
- 7 Winthrop KL, McNelley E, Kendall B, et al. Pulmonary nontuberculous mycobacterial disease prevalence and clinical features: an emerging public health disease. *Am J Respir Crit Care Med* 2010; 182: 977–982.
- 8 Kim RD, Greenberg DE, Ehrmantraut ME, et al. Pulmonary nontuberculous mycobacterial disease: prospective study of a distinct pre-existing syndrome. *Am J Respir Crit Care Med* 2008; 178: 1066–1074.
- 9 Thomson RM, on behalf of the NTM working group at the Queensland TB Control Centre, Queensland Mycobacterial Reference Laboratory. Changing epidemiology of pulmonary nontuberculous mycobacteria infections. *Emerg Infect Dis* 2010; 16: 1576–1583.
- 10 Henkle E, Hedberg K, Schafer S, et al. Population-based incidence of pulmonary nontuberculous mycobacterial disease in Oregon 2007 to 2012. *Ann Am Thorac Soc* 2015; 12: 642–647.
- 11 Shah NM, Davidson JA, Anderson LF, et al. Pulmonary *Mycobacterium avium-intracellulare* is the main driver of the rise in non-tuberculous mycobacteria incidence in England, Wales and Northern Ireland, 2007–2012. *BMC Infect Dis* 2016; 16: 195.
- 12 Martinez S, Cano A, Sota LA, et al. Micobacterias no tuberculosas ¿una amenaza emergente? *Arch Bronconeumol* 2017; 53: 554–560.
- 13 Hermansen TS, Ravn P, Svensson E, et al. Nontuberculous mycobacteria in Denmark, incidence and clinical importance during the last quarter-century. *Sci Rep* 2017; 7: 6696.
- 14 Leal Arranz MV, Gaafar A, Unzaga Barañano MJ, et al. Clinical and epidemiological study of disease caused by *Mycobacterium kansasii* in the metropolitan area of Bilbao, Spain. *Arch Bronconeumol* 2005; 41: 189–196.
- 15 Donohue MJ, Mistry JH, Donohue JM, et al. Increased frequency of nontuberculous mycobacteria detection at potable water taps within the United States. *Environ Sci Technol* 2015; 49: 6127–6133.
- 16 Hamada S, Ito Y, Hirai T, et al. Impact of industrial structure and soil exposure on the regional variations in pulmonary nontuberculous mycobacterial disease prevalence. *Int J Mycobacteriol Medknow Publications* 2016; 5: 170–176.
- 17 Isaac-Renton JL, Allen EA, Chao CW, et al. Isolation and geographic distribution of *Mycobacterium* other than *M. tuberculosis* in British Columbia, 1972–81. *CMAJ* 1985; 133: 573–576.
- 18 Corbett EL, Hay M, Churchyard GJ, et al. *Mycobacterium kansasii* and *M. scrofulaceum* isolates from HIV-negative South African gold miners: incidence, clinical significance and radiology. *Int J Tuberc Lung Dis* 1999; 3: 501–507.



- 19 Martiniano SL, Nick JA, Daley CL. Nontuberculous mycobacterial infections in cystic fibrosis. *Clin Chest Med* 2016; 37: 83–96.
- 20 Winthrop KL, Chang E, Yamashita S, et al. Nontuberculous mycobacteria infections and anti-tumor necrosis factor-alpha therapy. *Emerg Infect Dis* 2009; 15: 1556–1561.
- 21 Hoefsloot W, van Ingen J, Andrejak C, et al. The geographic diversity of nontuberculous mycobacteria isolated from pulmonary samples: an NTM-NET collaborative study. *Eur Respir J* 2013; 42: 1604–1613.
- 22 Martínez Moragón E, Menéndez R, Marco V, et al. Enfermedad pulmonar por micobacterias ambientales oportunistas en pacientes sin infección por el virus de la inmunodeficiencia humana. Factores de riesgo, clínica, diagnóstico y evolución. *Arch de Bronconeumol* 1996; 32: 170–175.
- 23 Martín-Casabona N, Bahrmand AR, Bennedsen J, et al. Non-tuberculous mycobacteria: patterns of isolation. A multi-country retrospective survey. *Int J Tuberc Lung Dis* 2004; 8: 1186–1193.
- 24 Santin M, Barrabeig I, Malchair P, et al. Pulmonary infections with nontuberculous mycobacteria, Catalonia, Spain, 1994–2014. *Emerg Infect Dis* 2018; 24: 1091–1094.
- 25 Marinelli DL, Albelda SM, Williams TM, et al. NTM infection in AIDS. *Radiology* 1986; 160: 77–82.
- 26 Maureen Cassidy P, Hedberg K, Saulson A, et al. Nontuberculous mycobacterial disease prevalence and risk factors: a changing epidemiology. *Clin Infect Dis* 2009; 49: 124–129.
- 27 Prevots DR, Marras TK. Epidemiology of human pulmonary infection with nontuberculous mycobacteria: a review. *Clin Chest Med* 2015; 36: 13–34.
- 28 Andrejak C, Thomsen VO, Johansen IS, et al. Nontuberculous pulmonary mycobacteriosis in Denmark: incidence and prognostic factors. *Am J Respir Crit Care Med* 2010; 181: 514–521.
- 29 Marras TK, Campitelli MA, Lu H, et al. Pulmonary nontuberculous mycobacteria-associated deaths, Ontario, Canada, 2001–2013. *Emerg Infect Dis* 2017; 23: 468–476.