Nine patients were diagnosed as stage IV and two as stages IA and IIB. Three patients were positive for EGFR mutations and three for EML4-ALK fusion gene. Two of them received targeted therapy with a TKI. All patients <40 years were found to have mutations. Regarding the follow-up of this sub-set of patients, 63.6% were not tracked and 27.4% died. Only one patient who received TKI with erlotinib, was followed-up for six months. We found a prevalence of 27% for EGFR mutations corresponding to the prevalence reported in Western and Asian populations. The prevalence of rearrangements in the EML4-ALK gene was 15.8%, higher than the reported worldwide.<sup>7</sup>

Patients with primary NSCLC are more likely to be non-smokers, women and have stage-IV adenocarcinoma, with a high rate of EGFR and EML4-ALK mutations. All patients < 40 years had genomic alterations, with EML4-ALK fusions being the most prevalent. We found an association between being a non-smoker and the presence of mutations, regardless of age or gender. A high prevalence of mutations in patients classified as never/light smokers has been reported in the literature before.<sup>8</sup> It is possible that the high rate of mutations discovered in this study is related to being a non-smoker and to the demographic characteristics of the study population in Colombia, specifically with the diverse ethnicity in this country. The probability of six-month survival was calculated from 40% of the initial sample, given a limited follow-up.

We recommend once the presence of EGFR and EML4-ALK mutations has been established,<sup>9</sup> follow-up should be undertaken at an institution that facilitates molecular diagnosis, treatment and management of the evolution and clinical response of the patients. This enables better decision-making and the creation of protocols for integral care. Issues in the healthcare system in Colombia may interfere with the implementation of these recommendations.

#### References

- 1. Travis WD, Brambilla E, Muller-Hermelink HK, Harris CC. Tumors of the lung International Agency for Research on Cancer (IARC); 2004. p. 9–124.
- Arrieta O, Cardona AF, Federico Bramuglia G, Gallo A, Campos-Parra AD, Serrano S, et al. Genotyping non-small cell lung cancer (NSCLC) in Latin America. J Thor Oncol. 2011;6:1955–9.
- Lindeman NI, Cagle PT, Beasley MB, Chitale DA, Dacic S, Giaccone G, et al. Molecular testing guideline for selection of lung cancer patients for EGFR and ALK tyrosine kinase inhibitors: guideline from the College of American Pathologists Interna-

tional Association for the Study of Lung Cancer, and Association for Molecular Pathology. J Thorac Oncol. 2013;8:823–59.

- Cardona AF, Ramos PL, Duarte R, Carranza H, Castro CJ, Lema M, et al. Screening for mutations in Colombian metastatic non-small cell lung cancer (NSCLC) patients (ONCOLGroup). J Clin Oncol. 2011;29 [suppl; abstr 7577].
- Fernandez L, Sua LF, Rodriguez LX, Munoz CA, Velasquez M, Restrepo JG. Epidermal growth factor receptor mutation in patients with non-small cell lung carcinoma in Fundacion Valle Del Lili A University Hospital of reference in Latin America [abstract]. Am J Respir Crit Care Med. 2016;193:A2589.
- Fernandez I, Sua LF, Munoz CA, Restrepo JG. Expression of protein kinase Eml4-Alk gene in non-small cell lung cancer in Fundacion Valle Del Lili A University Hospital of reference in Latin America [abstract]. Am J Respir Crit Care Med. 2016;193:A2588.
- 7. Chia PL, Mitchell P, Dobrovic A, Jhon T. Prevalence and natural history of ALK positive non-small-cell lung cancer and the clinical impact of targeted therapy with ALK inhibitors. Clin Epidemiol. 2014;6:423–32.
- Lindeman NI, Cagle PT, Beasley MB, Chitale DA, Dacic S, Giaccone G, et al. Molecular testing guideline for selection of lung cancer patients for EGFR and ALK tyrosine kinase inhibitors: guideline from the College of American Pathologists International Association for the Study of Lung Cancer, and Association for Molecular Pathology. J Thorac Oncol. 2013;8:823–59.
- Ministerio de Salud y Protección Social. Guía de Práctica Clínica para la detección temprana, diagnóstico, estadificación y tratamiento del cáncer de pulmón. Guía para profesionales de la salud. Colombia; 2014.

Liliana Fernández,<sup>a,\*</sup> Juan Fernando Henao,<sup>b</sup> Valeria Zuñiga,<sup>c</sup> Luz Fernanda Sua<sup>d</sup>

 <sup>a</sup> Internal Medicine, Pulmonology, Interventional Pulmonology, Biomedical Research in Thorax, Fundación Valle del Lili, Faculty of Health Sciences, Universidad ICESI, Cali, Colombia
<sup>b</sup> Internal Medicine, Biomedical Research in Thorax, Fundación Valle

del Lili, Cali, Colombia

<sup>c</sup> Medical Student, Universidad ICESI, Biomedical Research in Thorax, Fundación Valle del Lili, Cali, Colombia

<sup>d</sup> Anatomic Pathology and Clinical Pathology, Department of Pathology and Laboratory Medicine (Pulmonary Pathology), Biomedical Sciences, Biomedical Research in Thorax, Fundación Valle del Lili, Faculty of Health Sciences, Universidad ICESI, Cali, Colombia

\* Corresponding author.

*E-mail addresses*: lilianafernandeztrujillo@gmail.com, liliana.fernandez@fvl.org.co (L. Fernández).

1579-2129

© 2018 SEPAR. Published by Elsevier España, S.L.U. All rights reserved.

## 

# Evolución de los ingresos por bronquiectasias en un hospital de tercer nivel (2008-2017)

## To the Editor,

The incidence of bronchiectasis is increasing, especially among older patients.<sup>1</sup> Bronchiectasis occurs with exacerbations, which, when coupled with respiratory failure, hemodynamic instability, sepsis, or other complications, may require a hospital stay.<sup>2</sup> These hospitalizations generate a heavy economic burden,<sup>3,4</sup> which increases with severity and readmissions.<sup>4</sup> Our objective was to analyze trends in admissions for bronchiectasis exacerbations over a period of 10 years.

The study included all discharges coded 494.1 (bronchiectasis exacerbation) according to the International Code of Diseases version 9 (ICD-9) as primary diagnosis, between the years 2008 and 2017, in the Hospital Universitario Germans Trias i Pujol. The linear trend and trend in proportions were analyzed using the nptrend and ptrend commands in Stata<sup>®</sup> version 15, respectively. The study was approved by the Ethics Committee of the Hospital Universitari Germans Trias i Pujol.

During the study period, there were 304 admissions with bronchiectasis exacerbation as a primary diagnosis, in a total of 190 patients. Of these, 136 patients (71.6%) had 1 admission, 36 (18.9%) 2 admissions, 6 (3.2%) 3 admissions, and 6.3% 4 or more admissions. Of these, 75.1% were women with a mean age of 80.3 years (SD: 6.1). Mean stay was 9.5 (SD: 2.0). In-hospital mortality rate was 3.7%, with a 30-day readmission rate of 11.7%. Mortality was not significantly associated with age (P=.08) nor with the number of admissions (P=.11). The proportion of admissions with bronchiectasis as primary diagnosis compared to total admissions in our hospital was 2.29 per 1000. No significant changes occurred in readmission rates, mortality, and length of stay over the period

<sup>☆</sup> Please cite this article as: Garcia-Olivé I, Marin A, Rodríguez-Pons L, Abad J. Evolución de los ingresos por bronquiectasias en un hospital de tercer nivel (2008-2017). Arch Bronconeumol. 2019;55:217–218.



**Figure 1.** shows the progress of numbers of admissions for bronchiectasis and age over the study period. The left *y*-axis displays the number of admissions/1000 for bronchiectasis compared to total admissions during the time period analyzed. The right *y*-axis shows the average age of patients in years.

analyzed. On the other hand, a significant increase was observed in the age of patients (P<.01) and in the proportion of admissions for bronchiectasis compared to the total number of admissions in our hospital (P<.01) (Fig. 1).

Although some authors have described falling admission rates for acute exacerbations of bronchiectasis,<sup>5</sup> most of the existing studies coincide with ours, and report increasing rates of admissions for this reason.<sup>6,7</sup> Sanchez-Munoz et al.,<sup>8</sup> unlike us, described a decline in admissions when bronchiectasis exacerbation was the primary diagnosis, but when it was a secondary diagnosis, admissions increased. We did not include admissions for bronchiectasis exacerbation as a secondary diagnosis in our study.

With regard to the length of hospital stay of these patients, other authors in general describe a reduction in length of stay,<sup>8,9</sup> which is around 11 days.<sup>7,8</sup> Similarly to other reports,<sup>9</sup> when readmissions were analyzed, no changes were observed over the study period. As for mortality, we did not find significant variations over time, unlike other previously published studies that describe a decrease.<sup>9</sup> Again, Sanchez-Munoz et al. found differences depending on whether the diagnosis was the principal or secondary diagnosis.<sup>8</sup>

Our study has clear limitations. It is retrospective, so we were unable to analyze the criteria for admission of patients, associated co-morbidities, and antibiotic treatment (type, duration, etc.) received, factors that clearly have a role in patient prognosis. It is also possible that the increasing number of diagnoses is due not to a real higher incidence, but rather to a greater awareness of the disease and better coding in the hospital. In summary, we found an increase in the number of admissions for bronchiectasis, and in the age of patients over the time period analyzed. There were no changes in mortality, readmission rates, or length of stay. It would be interesting to perform a multicenter study to analyze in more detail the factors associated with the prognosis of patients admitted for bronchiectasis exacerbations.

### References

- 1. Quint JK, Millett ERC, Joshi M, Navaratnam V, Thomas SL, Hurst JR, et al. Changes in the incidence, prevalence and mortality of bronchiectasis in the UK from 2004–2013: a population based cohort study. Eur Respir J. 2016;47: 186–93.
- 2. Martinez-Garcia MA, Maiz L, Olveira C, Giron RM, de la Rosa D, Blanco M, et al. Normativa sobre el tratamiento de las bronquiectasias en el adulto. Arch Bronconeumol. 2018;54:88–98.
- Joish VN, Spilsbury-Cantalupo M, Operschall E, Luong B, Boklage S. Economic burden of non-cystic fibrosis bronchiectasis in the first year after diagnosis from a US health plan perspective. Appl Health Econ Health Policy. 2013;11: 299–304.
- **4.** De la Rosa D, Martinez-Garcia MA, Olveira C, Giron R, Maiz L, Prados C. Annual direct medical costs of bronchiectasis treatment: impact of severity, exacerbations, chronic bronchial colonization and chronic obstructive pulmonary disease coexistence. Chronic Res Dis. 2016;13:361–71.
- Säynäjäkangas O, Keistinen T, Tuuponen T, Kivelä SL. Bronchiectasis in Finland: trends in hospital treatment. Res Med. 1997;91:395–8.
- Seitz AE, Olivier KN, Steiner CA, Montes de Oca R, Holland SM, Prevots R. Trends and burden of bronchiectasis-associated hospitalizations in the United States, 1993–2006. Chest. 2010;138:944–9.
- Ringshausen FC, de Roux A, PLetz MW, Hämäläinen N, Welte T, Rademacher J. Bronchiectasis-associated hospitalizations in Germany, 2005–2011: a population-based study of disease burden and trends. PLoS One. 2013;8:e71109.
- Sanchez-Muñoz G, Lopez de Andres A, Jimenez-Garcia R, Carrasco-Garrido P, Hernandez-Barrera V, Pedraza-Serrano F, et al. Time trends in hospital admissions for bronchiectasis: analysis of the Spanish National Hospital Discharge Data (2004 to 2013). PLOS ONE. 2016;11:e0162282.
- Ford ES. Hospital discharges, readmissions, and ED visits for COPD or bronchiectasis among US adults: findings from the nationwide inpatient sample 2001–2012 and Nationwide Emergency Department Sample 2006–2011. Chest. 2015;147:989–98.

Ignasi Garcia-Olivé,<sup>a,b,c,\*</sup> Alicia Marin,<sup>a,b,c</sup> Laura Rodríguez-Pons,<sup>a,b,c</sup> Jorge Abad<sup>a,b,c</sup>

<sup>a</sup> Servicio de Neumología, Hospital Universitari Germans Trias i Pujol, Badalona, Barcelona, Spain

<sup>b</sup> Ciber de Enfermedades Respiratorias (CIBERES), Bunyola, Mallorca, Spain

<sup>c</sup> Fundació Institut d'Investigació en Ciències de la Salut Germans Trias i Pujol, Badalona, Barcelona, Spain

#### \* Corresponding author.

E-mail address: ignasi.g.olive@gmail.com (I. Garcia-Olivé).

#### 1579-2129/

© 2018 SEPAR. Published by Elsevier España, S.L.U. All rights reserved.