Non-cystic fibrosis bronchiectasis in an endemic area of pulmonary tuberculosis, Belém do Pará - Brazil - 2019

RESUMO

DESCRITORES: Bronquiectasia; pneumologia; tuberculose; tosse.

ABSTRACT
Objectives: Describe the clinical and epidemiological features of non-cystic fibrosis bronchiectasis patients treated at a specialized health-service in Belém-PA, an area with high rates of pulmonary tuberculosis. Method: The study evaluated 100 medical records, including 53, attended in 2019, with sociodemographic and clinical characteristics. Results: women from the capital, with onset of symptoms in adulthood, whose main etiology is post-tuberculosis. The most prevalent symptoms are cough, sputum and dyspnea. Radiologically diffuse and bilateral involvement, and the main morphological classifications: cylindrical and sacular. The most used medications: long-acting beta-2 agonist and acetylcysteine. According to the E-FACED severity score, most patients were classified as mild disease. Conclusion: women with a history of pulmonary tuberculosis, with persistent cough and sputum should be investigated for bronchiectasis.

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RESUMEN

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RECEIVED: 19/10/2021 APPROVED: 04/11/2021

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9115 saúdecoletiva • 2021; (11) N.71
INTRODUCTION

Bronchiectasis is a chronic, heterogeneous clinical condition of variable severity characterized by an abnormal and permanent dilation of the airways, whose etiologies can be congenital or acquired origin. The main classification divides: those of cystic fibrosis patients and those of other causes (Non-cystic fibrosis). The post-infectious and secondary cause of tuberculosis in Brazil is the main cause, resulting in chronic bronchial dilatation, with a peculiar clinical picture and compromised quality of life.

The state of Pará, in 2019, had the highest incidence of tuberculosis cases notified and monitored in Primary Care. Unlike what happens in the northern health region of Espírito Santo, which between 2018 and 2019 presented an incidence coefficient lower than the national average. In 2020, 4913 new cases of tuberculosis were confirmed in the state, of which 4221 were pulmonary.

Respiratory symptoms with persistent cough and large amounts of sputum, sometimes hemoptysis and recurrent infections, plus high-resolution computed tomography (HRCT) scans of the chest confirm the diagnosis, thus enabling early intervention and preservation of the pulmonary function.

Monitoring through spirometry with a bronchodilator test, in addition to evaluating the microbiota present in the airway of patients, is important for monitoring and scoring systems are proposed, thus estimating the severity of the patients. Among them, we have the so-called: E-FACED, which assesses the percentage (%) of predicted forced expiratory volume in one second (FEV-1), age, colonization by Pseudomonas aeruginosas, extent of impairment on HRCT, the dyspnea scale and presence of exacerbations.

Treatment must be individualized according to clinical and etiological particularities, and drug therapy includes: antimicrobials, bronchodilators and mucoviscous substances; and respiratory physiotherapy with maneuvers to clear this secretion is of paramount importance. Vaccination against influenza and pneumococcus, use of osmotic agents are also mainstays of treatment. From a surgical point of view, lung resections in localized and refractory disease, and even lung transplantation, may be considered.

The lack of studies on the subject in the state and the high burden of pulmonary infections (especially tuberculosis) in the region, calls attention to the need to describe the profile of patients with bronchiectasis in the region, their main etiologies, in addition to their clinical-epidemiological behavior. In this way, promoting academic knowledge on the subject, aiming to improve the quality of life of patients and reduce hospitalizations.

The objective of the research was to trace the clinical and epidemiological profile of patients with non-cystic fibrosis bronchiectasis treated at a specialized outpatient clinic at a university hospital in Belém-PA.

METHODS

Cross-sectional and retrospective study of appointments performed in 2019, which included patients over 18 years of age, with a diagnosis confirmed by high-resolution computed tomography (HRCT), and patients diagnosed with cystic fibrosis were excluded. Data collection started after approval by the Ethics Committee and was carried out based on electronic medical records of patients treated at the outpatient...
Clinic specializing in bronchiectasis. Data Use Commitment Terms (TCUD) have been completed.

Socio-demographic and clinical characteristics were evaluated, such as: period of onset of symptoms, probable etiology, signs and symptoms and tomographic alterations. The number of exacerbations, need for hospitalization, pulmonary colonization proven by sputum culture, degree of dyspnea by the modified dyspnea scale – Medical Research Council (mMRC) and severity assessment by the E-FACED score, in addition to the proposed treatment, were described.

The sample characterization information was calculated and entered into a spreadsheet created in the Microsoft® Office Excel® 2016 software. In the application of Descriptive Statistics, tables and graphs were built to present the results and position measurements were calculated as arithmetic mean and standard deviation.

Statistical analysis was performed using the G and Chi-Square Adhesion Tests for univariate tables. Descriptive and analytical statistics were performed using BioEstat® 5.4 software. For decision making, the significance level α = 0.05 or 5% was adopted, indicating the significant values with an asterisk (*).

RESULTS

One hundred medical records were analyzed, including 53 in the research, as the rest had insufficient data. Table 1 describes the epidemiological profile of patients.

The onset of symptoms predominantly started in adulthood (p<0.0001), and the etiology is probably more frequent after respiratory infection (p=0.004), especially after pulmonary tuberculosis. The other probable etiologies were described in Table 2, however it is important to emphasize that in some patients there was suspicion of two etiologies at the same time, due to history and clinical picture. Symptoms were described in Table 2, and cough was statistically significant (p=0.006).

Of the 53 patients studied, 40 (75.47%)...
were non-smokers, 10 (18.87%) were active smokers, with an average smoking load of 19.73 packs/year and standard deviation of about 24.31 packs/year (p < 0.00010) and the rest (3; 5.7%), passive smokers.

Regarding bacteria isolated in sputum culture, Pseudomonas aeruginosa was found in 8 patients (15.10%), followed by Klebsiella pneumoniae in 9.40% (5) and Staphylococcus aureus in 3.80% (2). As for the extent of pulmonary involvement on CT, most of it was diffuse, with more than two lobes affected in 49.1% (26) of the patients (p = 0.0004). When analyzing the tomographic morphological aspects and considering that there were concomitant findings, 47.2% (25) of the sample had cylindrical bronchiectasis, 43.4% (23) of the saccular type, followed by 24.5% (13) of the varicose type. Pulmonary function assessed by spirometry showed that 32 (60.4%) patients had FEV1 > 50%, and the rest (21; 39.6%) had FEV1 < 50%.

Regarding drug treatment, 62.3% (33) of the patients were using long-acting B2 agonist bronchodilators (LABA) (p = 0.0214), 49.1% (26) of mucoactive agents such as acetylcysteine, 34% (18) inhaled corticosteroids and 24.5% (13) used systemic antibiotics due to exacerbations. Among the non-pharmacological treatments, 15.1% (8) of the patients underwent respiratory physiotherapy. Surgical treatment, lobectomy, was observed in 5.7% (3) of the study population. Table 3 shows the number of exacerbations and hospital admissions for the disease.

The degree of dyspnea classified by the mMRC showed that 64.2% (34) was said to be degree zero (p<0.0001). The classification of patients according to the criteria of severity and prognosis provided by the E-FACED score showed a result of 81.1% (43) as classified as having mild bronchiectasis (Table 4).

### DISCUSSION

For some decades, bronchiectasis disease was considered an orphan disease due to the scarcity of studies and public-private policies on the subject. However, with the increase in the number of cases around the world, it is taking a robust statistical shape led by the European registry (EMBARC).

Women in the sample in question, as well as in a reference hospital in São Paulo, are the most affected. With the onset of symptoms in adulthood, and the imprecise assessment of the mean age, they are repeated in other cohorts around the world, such as in a tertiary hospital in Pakistan, 10 in which the majority of patients had the onset of symptoms over the age of sixty years, and another in Morocco,11 in which the age of onset of symptoms was 48 years of age.

A multicenter cohort study in Europe and India, 12 found that about 41.7% of 2195 patients had tuberculosis as the main probable cause of bronchiectasis, as well as Oliveira et al. 13 among those cases of post-infectious origin, tuberculosis was the main one involved. Clinically, cough is the most cited symptom, followed by expecto-
ration and dyspnea. It is noteworthy that the hodgepodge of symptoms is common similar to the registry data from the University Hospital of Baskent in Adana-Turkey, 14 where most patients had cough, spu-
tum and dyspnea, respectively, as the main symptoms.

The radiological pattern most observed in this study is the diffuse and bilateral one, with involvement of more than two lung lobes, that is, of severe extension on HRCT
scans. A Pakistani study shows a different result, showing that 41.8% of patients have focal lung injury, and only 20.4% have diffuse bilateral injury. 15

The predominant tomographic morphological type was the cylindrical shape, followed by the saccular pattern, which diverges from those found by Tejada et al., 16 in which the prevalent morphology was mainly saccular, followed by varicose and

This study showed that most patients, 75.47%, do not have a history of smoking. As in a study by Martinez-Garcia et al, 14 who analyzed 1912 patients with bron-
chectasis in which 58.4% were non-smokers.

There was a statistically significant proportion of patients undergoing drug therapy, especially LABA, and there was an expressive use of mucolytics such as
N-acetylcysteine by the sample. The use of oral antibiotics was used in cases of exacerbation by secondary infections. Since this drug therapy is similar to that found by
Dhar et al, 12 and the one proposed by the Brazilian Consensus on Non-Fibrocystic
Bronchiectasis. 2

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cenconsensus, 17 for patients with focal impairment, whose clinical treatment was
unsuccessful.

In a prospective cohort study of 608 patients, it showed that the majority (363) had at least one exacerbation over a year, 18 130 of these were hospitalized. Contrary to
what was seen in this research, 56.6% of the sample did not show exacerbation recorded in 2019.

Regarding dyspnea assessed by the mMRC scale, most patients in this study were classified as grade zero, that is, dyspnea after intense exercise, similar to the study by Clamers et al. 19 When analyzing the E-FACED score, most respondents were classified as mild, followed by moderate cases, similar to the result of a multi-
center observational study of a historical cohort, 20 in which 60.5% of the patients
were classified as having mild bronchiectasia by the same score, 30% and 8.6% as
having moderate and severe, respectively.

CONCLUSION

Thus, it is concluded that a history of pulmonary tuberculosis is an important predictor of bronchiectasis and in regions where the prevalence of tuberculosis is high, bronchiectasis should be investiga-
ted in patients with persistent symptoms such as coughing and pulmonary hyper-
secretion. Drug therapy shows benefit in terms of disease prognosis, as according to the E-FACED severity and prognosis score, most patients were classified as having mild bronchiectasis. It is expected that this cohort of patients will be evaluated in the future with a larger number of participants, over a longer period of time and with more adequate control of the variables involved.

REFERENCES

1 - Gomes ELFD, Gimenes AC, Lanza FC. Técnicas de fisioterapia e reabilita-
ção pulmonar na bronquiectasia não fibrocística. Pneumologia Paulista; 2016; 29(1).
2 - Pereira MC, Athanazio RA, Dalcin PTR, Figueiredo MRFS, Gomes M, Freitas CG, et al. Consenso brasileiro sobre bronquiectasias não fi-
brocísticas. J Bras Pneumol; 2019; 45(4). Disponível em: www.jornalde-
pneumologia.com.br/details/3030/en-US/consenso-brasileiro-so-
obre-bronquiectasias-nao-fibrocisticas.
3 - Brasil, Ministério da Saúde. I Lare da Tuberculose: evolução dos cenâ-
os epidemiológicos e operacionais da doença. 2019 (50).
4 - Soto, R. L. Gouveia, R. R., Cardoso, M. L., Gomes, A. M., Forza, D.
va. 2021; 69 (11).
5 - Brasil, Ministério da Saúde. Banco de dados do Sistema Único de Saú-
6 - Rocurek EG and Jagana R. Noncystic fibrosis bronchiectasis man-
com;2019, 25 (2).
7 - Pasteur MC, Bilton D, Hill AT. British Thoracic Society Bronchiectasis
non CFFGG. British Thoracic Society guideline for non-CF bronchiectasis.
nlm.nih.gov/20627931/
8 - Nucci MCN, Fernandes FLA; Salge JM, Steimach R, Cudler A, Athanazio R. Caracterização da gravidade da disneia em pacientes
com bronquiectasias: correlação com aspectos clínicos, funcionais e
scielo.br/j/pneu/v47n5/a5/pdf/pt
9 - Chalmers J, Alberti S, Polverino E, Vendrell M, Crichton M, Loebleing
M, et al. The EMBARC European Bronchiectasis Registry: protocol for an
international observational study, European Respiratory Society open
10 - Sharif N, Baig MS, Sharif S, Irfan M. Etiology, Clinical, Radiological,
and Microbiological Profile of Patients with Non-cystic Fibrosis Bronchi-
ectasis at a Tertiary Care Hospital of Pakistan. Cureus, 2020; 12(3). Available
11 - Bopaka AG, Khattabi WE, Janah H, Jabri H, Aff H. Bronchiectasis: a
bacteriological profile. The Pan African medical journal, 2015 [22]. Available
12 - Dhar R, Singh S, Talwar D; Mohan M, Tripathi SK, Swarnakar R, et al.
REFERENCES


