Mounier-Kuhn syndrome*

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dyspnea. He was diagnosed with pneumonia and was hospitalized, receiving intravenous cefazolin for seven days. He presented favorable evolution. The patient also reported having experienced three previous episodes of pneumonia, the first at the age of 22. The chest X-ray performed at the time of the evaluation revealed discrete areas of consolidation, of residual aspect, in the middle lobe, as well as a significant increase in the diameter of the trachea and of the main bronchi. The patient underwent empirical treatment with a bronchodilator for two weeks, with no improvement, and then returned for a reevaluation. The cough had worsened, becoming productive. The chest X-ray taken at that time revealed new areas of consolidation in both lower lobes and in the middle lobe. There was no alteration in the vital signs, and the patient presented good general status. A ten-day course of oral erythromycin was prescribed. Over the first three days, the cough and fatigue lessened. By the end of the ten-day period, the cough had almost completely disappeared.

A few days after the treatment with erythromycin, however, the patient again presented productive cough, accompanied by fever, and the X-ray revealed no improvement in the consolidation. The patient was again hospitalized and was submitted to sputum smear microscopy, which revealed innumerable polymorphonuclear cells and abundant gram-positive cocci. Sputum culture was unavailable. The patient started being treated with intravenous oxacillin, and a satisfactory clinical response was obtained in a few days, which was the response that had been expected after the administration of oral cephalaxin. After having received oral cephalaxin for 48 h, the patient again presented fever, which led us to return to the intravenous treatment and transfer him to a hospital possessing more advanced equipment in order to perform a more elaborate etiological investigation. The new sputum smear microscopy revealed polymicrobial flora, and the patient was started on ceftriaxone combined with amikacin. This treatment continued for four weeks, by which time the symptoms had been completely resolved. At that point, the patient was submitted to a computed tomography scan of the chest, which revealed that the areas of consolidation, although smaller in size, persisted in both lower lobes, and confirmed the abnormal widening of the trachea and main bronchi (Figures 1, 2 and 3). In the lower

Figure 1 - Anteroposterior chest X-ray revealing areas of consolidation in both lower lobes (predominantly on the right side) and widening of the trachea and bronchi
lobes, there was bilateral cylindrical bronchiectasis, which is consistent with a diagnosis of congenital tracheobronchomegaly, or Mounier-Kuhn syndrome. Pulmonary function tests revealed an increase in total lung capacity (123% of predicted) and residual volume (160% of predicted), with no other alterations. Currently, the patient remains asymptomatic, and has daily respiratory therapy sessions. Annual immunization against influenza has been recommended.

DISCUSSION

Congenital tracheobronchomegaly, or Mounier-Kuhn syndrome, is a rare clinical entity, described for the first time in 1932,[1] and characterized by marked tracheobronchial dilation and lower respiratory tract infections. It is predominantly found in male individuals in their 40s or 50s. Although the etiology of Mounier-Kuhn syndrome remains unknown, it is believed that it is related to a lack of smooth muscle and elastic connective tissue in the trachea and main bronchi, leading to herniation and even to the formation of diverticula between the cartilaginous rings. A finding of bronchiectasis, such as in the case in question, is uncommon.[2-3]

Mounier-Kuhn syndrome is occasionally found in asymptomatic individuals. When symptomatic, it is characterized by recurrent pneumonia, sometimes evolving to chronic productive cough, occasional hemoptysis, and progressive dyspnea as a result of the pulmonary involvement. More rarely, massive hemoptysis,[4] spontaneous pneumothorax, and digital clubbing can be observed. We believe that, in the case in question, the syndrome was responsible for the inappropriate response to the appropriate antibiotic therapy, which led us to suspect of a structural abnormality of the airways.

Diagnosis can usually be made by measuring the tracheal diameter, using only data from chest X-rays, in which the trachea can seen in profile and thus the diameter determined. Computed tomography of the chest, however, makes this measurement more precise. The limits are 3 cm for the transverse diameter of the trachea, and 2.4 cm and 2.3 cm for the transverse diameters of the main right and left bronchi, respectively. Pulmonary function tests typically reveal an increase in total lung capacity, to the detriment of residual volume, occasionally with signs of obstructive ventilatory disorder. Currently, there are no specific treatments for this condition, except for antibiotic therapy during the crises and respiratory therapy in order to eliminate secretions, if necessary. The use of permanent prostheses is reserved only for selected advanced cases since there are no precise indications for the practice.[5-6]

Other presentations of widening of the lower airways can be mistaken for Mounier-Kuhn syndrome.[7] Among such presentations are those...
caused by congenital disorders such as Ehlers-Danlos syndrome and Williams-Campbell syndrome (the latter, a rare form of cystic bronchiectasis resulting from cartilage deficiency from the fourth to the sixth bronchial order, with normal trachea and main bronchi). Disorders such as sarcoidosis, usual interstitial pneumonia and cystic fibrosis cause severe fibrosis of the upper lobes, leading to tracheal retraction. Inflammatory processes of the airways, such as allergic bronchopulmonary aspergillosis, caused by the colonization of the airway with Aspergillus sp., are also involved in the widening of the airways. In addition, it is known that smoking, chronic obstructive pulmonary disease, and repeated infections can worsen the profile.

A careful evaluation of the airway anatomy is very important in patients presenting recurrent pneumonia, productive chronic cough, or, as in the case in question, incomplete response to appropriate antibiotic therapy for pneumonia. Mounier-Kuhn syndrome, although rare, constitutes a diagnostic possibility in these patients and must be considered.

REFERENCES