Tracheomalacia Due to Esophageal Achalasia

To the Editor,

Primary esophageal achalasia, although rare, is a disorder of esophageal motility that is the result of a loss of ganglion cells responsible for motility and relaxation of the lower esophageal sphincter. As a result, patients present with worsening dysphagia to both liquids and solids and suffer from significant regurgitation of retained food, with weight loss and chronic cough. Achalasia symptoms may often mimic common diseases, and therefore, may delay the diagnosis.1

We report the case of a 10 years-old child who complained of barking cough from the age of 2 years and recurrent wheezing from the age of 3 years, with positive skin prick tests for alternaria. The child presented with frequent regurgitations in his first months of life, which subsequently disappeared.

Over the previous 6 months, the child had complained of episodic discomfort and sensation of heaviness while eating, nocturnal cough and occasionally nocturnal emesis. Based on the change of symptoms, the family’s pediatrician proposed treatment with long acting β2-adrenergic agonist (LABA)/ICS, without benefit. Admitted to a peripheral hospital, the child was transferred to our ward with the request to perform fiberoptic bronchoscopy and esophageal pHmetry for wheezing not responding to treatment and suspicion of gastroesophageal reflux disease. At admission he had body length of 141 cm (25–50th percentile), body weight of 31.5 kg (10th–25th percentile), transcutaneous oxygen saturation of 100%, respiratory rate 20 breaths per minute. Chest and abdomen examination were normal. Routine laboratory tests were normal, and allergy tests showed total IgE 193 KU/L with specific IgE for alternaria 23.2 KU/L. Spirometry revealed a flow/volume curve with reduction in PEF and FEV1 and a plateau consistent with tracheomalacia.

Fiberoptic bronchoscopy showed a tracheomalacia suggestive of aberrant innominate artery compression. The tracheal stenosis was fixed and pulsating, and the diameter remained constant with inspiration/expiration. The disease was localized where the vascular compression was present.

Spleunking the esophagus with the same instrument showed a severe dilatation. The child underwent esophagography, which showed a megaesophagus with “bird’s beak appearance” of the distal portion, typical of achalasia; esophageal manometry con-

References


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Tracheomalacia debida a acalasia esofágica

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Fig. 1. MRA showing displacement and compression of the trachea (T) against innominate artery (A) caused by esophageal dilatation (E).
Heller's esophagocardio-myotomy is the treatment of choice for children with achalasia because of its safety and long-term effective results after surgery. In our patient, chronic barking cough was caused by a secondary tracheomalacia resulting from a shift of the trachea against the innominate artery normally located.

References

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An Unusual Cause of Adult Respiratory Distress

Una causa no habitual de distrés respiratorio del adulto

To the Editor,

Acute fibrinous and organizing pneumonia (AFOP) is a rare histologic interstitial pneumonia pattern described in 2002 by Beasley.1 It is usually idiopathic, but among the possible causes associated with AFOP, infections, autoimmune diseases, drugs, environmental or occupational exposures, cancers and organ transplantation have all been previously reported.2 We report the case of AFOP in a previously healthy woman.

A 43-year-old woman presented to the emergency room (ER) with a 3-day history of worsening dyspnea along with intermittent fever since the previous week. She worked in a school as a teacher, and her personal history was unremarkable except for a smoking habit. She denied recent travels, environmental exposures and had no pets. Physical examination revealed bad general condition with fever, tachypnea, hypotension (95/69 mmHg) and desaturation (oxygen saturation of 89% on room air). Moreover, inspiratory crackles in the base of the right lung were detected. Thorax X-ray

![Fig. 1. (A) Thorax with contrast scan: Bibasilar alveolar consolidations, more prominent in the right lung. (B) Lung biopsy: multiple intra-alveolar fibrin “balls” and inflammatory thickening of alveolar septa along with areas of organizing pneumonia with patchy distribution.](Image)