Ortner’s Syndrome Caused by Idiopathic Pulmonary Artery Aneurysm

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Abstract

The Ortner’s syndrome or cardiovocal syndrome is an uncommon entity characterized by hoarseness due to left recurrent laryngeal nerve paralysis caused by identifiable cardiovascular disease. The most common conditions which may lead to Ortner’s syndrome include mitral stenosis, aortic aneurysm, mitral valve prolapsed, cardiovascular surgery, aortic dissection etc. Herewith, we report the case of an atypical etiology of cardiovocal syndrome in a patient with a Pulmonary Artery Aneurysm (PAA).

Keywords

Dysphonic, Vocal cord palsy, Ortner, Pulmonary aneurism

Introduction

Hoarseness is a common condition underlying many different causes. The left recurrent laryngeal nerve paralysis caused by identifiable cardiovascular disease is known as Ortner’s syndrome or cardiovocal syndrome. Ortner’s syndrome was first described by Norbert Ortner in 1897, in a patient with mitral stenosis with dilated left atrium [1]. Many other etiologies have been identified as the cause of this entity. We present here the first Ortner’s syndrome due to idiopathic pulmonary aneurysm without primary pulmonary hypertension.

Case Report

A healthy 42-year-old woman was referred to the ear, nose and throat (ENT) department complaining about three-month history of hoarseness, with no other associated pathology. She was a non-smoker patient without any relevant past medical history.

The clinical examination with indirect laryngoscopy revealed a paralyzed left vocal cord in paramedian position while the rest of the otolaryngologic exam was normal. Tuberculosis, Lyme disease and syphilis infections were ruled out as well as collagen vascular disorders and sarcoidosis. No prior history of blunt or penetrating trauma neither surgery was identified. Contrast-enhanced computed tomography (CT) of the neck and chest was performed to rule out any cervico-thoracic process causing the symptoms. The CT examination showed a PAA that involves the trunk and the main left pulmonary artery, with a maximum diameter of 45mm. The echocardiogram study showed neither structural cardiopathy nor pulmonary hypertension, so it lead to the diagnosis of idiopathic PAA. A specialist for cardio-thoracic surgery was consulted but surgical intervention was dismissed due to the low risk of artery dissection according to the size of the aneurysm and the symptoms of the patient. Currently, the patient is followed by the speech therapist, the cardiothoracic surgeon and the ENT surgeon with improvement of the voice quality.

Discussion

The left vagus nerve supplies the innervation of the larynx with their two terminal branches, the laryngeal recurrent nerve and the superior laryngeal nerve. The vagus nerve emerges through the jugular foramen and runs within the carotid sheath together with the carotid artery and the internal jugular vein. The left recurrent laryngeal nerve is a branch of the left vagus nerve at the level of the aortic arch. This nerve curves below the aorta and ascends to the tracheoesophageal groove. The laryngeal recurrent nerve supplies all the muscles acting on the vocal cords, except the cricothyroid muscle, which is innervated by the superior laryngeal nerve. Due to this large trajectory, this nerve may be injured in many different locations. In unilateral vocal cord paralysis due to thoracic disease, left vocal cord paralysis was 1.75 times more frequent than the right side [2].

Neck and chest CT scan utility to identify the etiology of a vocal cord paralysis has been widely reported [2]. Sun and colleagues published a retrospective study to evaluate the profitability of this test to rule out the etiology of vocal paralysis, concluding that CT is a helpful tool for the early detection of malignant and non-malignant causes of vocal cord paralysis [2].

Many different causes have been attributed to Ortner’s syndrome. It was associated with mitral stenosis or regurgitation, atrial mixoma,
idiopathic PAA. Treatment can be either conservative or surgical, recommending surgical repair when they are symptomatic or larger than 6 cm.

This non previous published case should illustrates to the ENT surgeon the countless different causes which may lead to a vocal cord paralysis, revealing the importance role of CT scan to identify their etiology.

References