Laryngeal Manifestation of Forestier's Disease

Jasmina Stojanovic¹, Sandra Zivanovic², Suncica Sreckovic², Svetlana Jovanovic², Branislav Belic¹,², Sladjana Simovic³

¹Phoniatic Department of ENT Clinic, Clinical Centre Kragujevac, Kragujevac, Serbia; ²Faculty of Medical Science, University of Kragujevac, Kragujevac, Serbia; ³Department of Otorhinolaryngology Health Centre Kragujevac, Kragujevac, Serbia

Abstract

BACKGROUND: Forestier's disease is a rare disorder involving bony growths that can occur in various parts of the spinal column, mostly asymptomatic, but these osteophytes, very rarely have been associated with serious complications.

AIM: We report a 69-year-old man who was admitted at foniatric department for evaluation of presenting hoarseness, dysphagia and laborious breathing.

CASE PRESENTATION: Noninvasive endolaryngeal imaging and radiological examination revealed distortion of left side of the larynx pushing to the right due to bony mass of the anterior part of cervical spine which was prominent at the left side. The symptoms of the patient presented caused by Forestier's disease as found by the imaging.

CONCLUSIONS: In clinical practice it is advisable to take into consideration Forestier's disease as a possible cause of hoarseness and dysphagia in rare cases.

Introduction

Diffuse idiopathic skeletal hyperostosis (DISH), also known as Forestier's disease, is a noninflammatory ossification involving at least four contiguous vertebral bodies, with intensive formation of osteophytes affecting ligaments, tendons and fascia of the anterior part of the spinal column [1]. It mainly affects males over 50 years of age. The incidence has been estimated at 12% [2] and based on radiological surveys of the cervical spine 2.4-5.4% in older than 40 years of age [3]. Although Forestier's disease is asymptomatic in general, in the literature were reported dysphagia, dyspnea and dysphonia all together only in rare cases [4].

Case report

A 69-year-old man was referred to Foniatric department of ENT clinic, Clinical Centre Kragujevac, Serbia, for examination with the main complaints of hoarseness, dysphagia and exertional laborious breathing lasting for two months. We performed complete ENT examination and fiberoptic laryngoscopy revealed that the right side of the larynx was dislocated ventrally and left (Fig 1.). The airway was compromised and reduced to one-third but sufficient. Laryngeal videostroboscopy findings show passive vertical right vocal folds vibration, left vocal folds vibration was normal. In exertional inhalation there was stridor. Pulmonologist exclude any respiratory problems so there was no indications for further functional respiratory diagnostics or examinations The routine biochemical tests were normal.
Computed tomography of the neck (Fig. 2) revealed an elongated ossification of the frontal planes of the vertebral bodies, especially at the C 3-6 levels, with elongated and spiky spurs especially at the right side projecting into the soft tissues of the neck pushing hypopharynx, oesophagus and larynx ventrally and right. Computed tomography was also done with contrast to exclude possible malignant process. High density was not observed. Forestier's disease was also confirmed by X-ray examination of column performed in two directions.

Patient was examined by orthopedic surgeon and his assessment was that there were no indications for surgical treatment. Orthopedic surgeon advised just physical therapy for our patient. We decided to treat conservatively our patient with nonsteroidal anti-inflammatory drugs associated with intensive physical rehabilitation treatment. Physical therapy reduced pressure on larynx and nonsteroidal anti-inflammatory drugs probably decreased inflammation in that region. At the check up two months later, the clinical findings were better and the symptoms were reduced. At that time we performed multidimensional computerised voice analysis to quantify the voice characteristics of our patient. Acoustic voice-signal data were measured for fundamental frequency (F0): 268.08Hz elevation of F0 can be explained by increase of subglottic pressure, percent of jitter: 1.72% and percent of shimmer: 2.44%, SD: 8.16% and amplitude perturbation quotient: 1.94%. Multidimensional voice evaluation was done to make it possible to follow up improvement of voice in future. Vocal therapy was not conducted because there were no indications for vocal therapy.

**Figure 1**: The first fiberoptic laryngoscopy findings shown distortion of larynx to left

**Discussion**

The characteristic traits of DISH are ossification of anterior ligaments and the anterior osteophyte bridges that appear at both sides and symmetrically, but in literature we find them described very rarely asymetrically causing extraspinal symptoms such as laryngeal manifestation occurring with hoarseness, dysphagia and laborious breathing [5, 6].

The suggested pathogenesis of DISH indicates that ossification and new bone formation is the result of abnormal osteoblast cell growth/activity in the bony ligamentous region [7].

In our patient ossification and new bone formation were present at C3-6 pronounced on the left causing laryngeal manifestation. The possible mechanism is not only compression of the larynx and oesophagus but also paresis of terminal laryngeal nerve fibers, traumatic compromisation, and direct involvement of the cricoarytenoid joint [8]. Commonly appearing symptoms are dysphagia, which appears at about 3-20% of patients with DISH [8, 9] and airway symptoms where prevalence in patients with DISH was 11.2% [10].

Most patients with DISH can be managed conservatively, as we decided. Surgical excision is appropriate in patients with severe and progressive symptoms such as airway obstruction and progressive weight loss caused by dysphagia problems [5, 12, 13].

Surgical treatment may be followed by many complications. According to determination presence and impact of dysphonia and dysphagia following the anterior approach to the cervical spine in Winslow et al study, hoarseness was present in 51%, dysphagia was present in 60% while in 18% breathing difficulties were found postoperatively [13].
After appropriate history, clinical examination, diagnostic and radiological procedure we diagnosed Forestier's disease as a cause of the symptoms in our patient. We treated the patient conservatively and he responded well to the therapy. In clinical practice it is advisable to take into consideration Forestier's disease as a possible cause of hoarseness and dysphagia in rare cases.

Reference
1. Forestier J, Rotes-Querol J. senile ankylosing hyperostosis of the spine. Ann Rheum Dis. 1950;9: 321-330. http://dx.doi.org/10.1136/ard.9.4.321 PMid:14800245 PMcid:PMC1011670
2. Yochum TR, Rowe LJ. Essentials of Skeletal Radiology, bed. 3. Baltimore, Williams & Wilkins, 2005.
3. Weirfeld RM, Olson PN, Maki DD, Griffith HJ. The prevalence of diffuse idiopathic skeletal hyperostosis (DISH) in two large American Midwest Metropolitan hospital populations. Skeletal Radiol. 1997;26:222-5. http://dx.doi.org/10.1007/s002560050225
4. Ahn YJ, Hahn SH, Yang BK, Yi SR, Yoo JH, Yoon DJ, et al. Diffuse idiopathic skeletal hyperostosis associated with dysphonia and dysphagia: a case report. J Korean Soc Spine Surg. 2006:13:327–331. http://dx.doi.org/10.4184/jkss.2006.13.4.327
5. Rahimizadeh A, Taginedjadi O, Rahimizadeh S, Saghri M, Rahimizadeh A. Respiratory Distress and Dysphagia Secondary to Diffuse Idiopathic Skeletal Hyperostosis: Case Report and Review. World Spinal Column Journal, WScJ. 4:18-24, 2013.
6. Adnan V, Vikram Vel VR, Purushothaman PK, Rajesh Kumar MS. Crico Arytenoid Joint Fixation in Diffuse Idiopathic Skeletal Hyperostosis (DISH): A Case Report. Indian J Otolaryngol Head Neck Surg. 2011; 63(1): 55–57.
7. El Miedany YM, Wassif G, El Baddini M. Diffuse idiopathic skeletal hyperostosis (DISH): is it of vascular aetiology? Clin Exp Rheumatol. 2000;18(2):193–200. PMid:10812491
8. Giger R, Dulguerov P, Payer M. Anterior cervical osteophytes causing dysphagia and dyspnea: an uncommon entity revisited. Dysphagia. 2006;21:259-63. http://dx.doi.org/10.1007/s00455-006-9049-0 PMid:17216390
9. Aydin E, Akdogan V, Akkuzu B, Kibris I, Ozgingin ON: Six cases of Forestier syndrome: a rare cause of Dysphagia. Acta Otolaryngol.2006; 126:775-8. http://dx.doi.org/10.1080/00016480500504192 PMid:16803720
10. Resnick D, Niwayama G. Diagnosis of bone and joint disorders. Philadelphia, Pa:WB Saunders, 1982:1562-602.
11. Constantoyannis C, Papadas T, Konstantinou D. Diffuse idiopathic skeletal hyperostosis as a cause of progressive dysphagia: a case report. Cases J. 2008; 23;1(1):416.
12. Lin HW, Quesnel AM, Holman AS et al. Hypertrophic anterior cervical osteophytes causing dysphagia and airway obstruction. Annals of Otol, Rhinology & Laryngology. 2009; 118(10):703-7. http://dx.doi.org/10.1177/000348940911801004
13. Winslow CP, Winslow TJ, Wax MK. Dysphonia and Dysphagia Following the Anterior Approach to the Cervical Spine. Arch Otolaryngol Head Neck Surg. 2001;127(1):51-55. http://dx.doi.org/10.1001/archotol.127.1.51 PMid:11177014