Case report

Internal laryngopyocele as a cause of acute airway obstruction: an extremely rare case and review of the literature

Il laringopiocele interno come causa di ostruzione acuta delle vie aeree: descrizione di un caso e revisione della letteratura

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SUMMARY

The laryngocele is an abnormal cystic dilatation of the saccule or appendix of the laryngeal ventricle, filled with air and communicating with the lumen of the larynx. When the neck of the laryngocele is obstructed, it becomes filled with mucus of the glandular secretion and is changed to a laryngomucocele. When this lesion becomes infected, a laryngopyocele is formed. The laryngocele is fairly rare and laryngopyoceles occur even more rarely. Overall, 39 cases of laryngopyocele have been reported in the world literature. Only in 4 cases was a laryngopyocele reported to have caused acute airway obstruction and only one case of internal laryngopyocele causing acute airway obstruction has been reported until now. This is the first case reported in the literature of an internal laryngopyocele in a female patient in a septic condition, which caused almost 100% obstruction of the airway. An emergency tracheotomy was performed in order to secure the airway. Computed tomography of neck was performed which revealed a cystic 29 mm hypodense mass extending from the right false vocal cord to the level of the epiglottis, narrowing the laryngeal cavity and causing an almost 100% airway obstruction. Laryngopyoceles may present with a rapid and alarming obstruction of the airway and, therefore, an urgent tracheotomy may be inevitable. It is an emergency case, in the field of otolaryngology, and should be included in the differential diagnosis of acute airway obstruction, especially when hoarseness, stridor and fever are present. Diagnosis requires a high index of suspicion for these lesions and scrupulous clinical and radiological evaluation. A computed tomography scan is critical in determining the nature and site of the lesion. The recommended treatment of laryngopyocele is immediate endoscopic drainage. Definitive management of laryngopyoceles is surgical excision which can be performed immediately after endoscopic drainage or some time thereafter.

KEY WORDS: Larynx • Laryngocele • Laryngopyocele • Airway obstruction

RIASSUNTO

Il laringocele è una dilatazione cistica del saccolo o appendice del ventricolo laringeo, piena di aria e comunicante con il lume laringeo. Quando il collo del laringocele si ostruisce, esso si riempie di muco e prende il nome di laryngomucocele, la cui infezione porta alla formazione di un laringopiocele. Il laringocele è piuttosto raro e ancora di più lo è il laringopiocele. Finora sono stati descritti 39 casi di laringopiocele, e soltanto 4 di questi hanno determinato un'ostruzione acuta delle vie aeree. Solo in un caso l'ostruzione acuta delle vie aeree è stata causata da un laringopiocele interno. In considerazione della rarità dell'ostruzione acuta delle vie aeree da laringopiocele, si riporta per la prima volta in letteratura un caso di laringopiocele interno che ha determinato un'ostruzione pressoché totale delle vie aeree in una paziente in stato settico, tanto da rendere necessaria una tracheotomia d'urgenza. La tomografia computerizzata ha evidenziato una massa cistica ipodensa di 29 millimetri di diametro, estesa dalla corda vocale vera fino al livello dell'epiglottide. Il laringopiocele rappresenta un'emergenza nel campo dell’otorinolaringoiatria e dovrebbe essere incluso nella diagnosi differenziale di ostruzione acuta delle vie respiratorie, soprattutto quando sono presenti disfonia, stridore e iperpiressia. È necessaria pertanto un'attenta valutazione clinica, che non può prescindere dall’esecuzione di una tomografia computerizzata, essenziale per la definizione della natura e della sede della lesione. Il trattamento raccomandato per il laringopiocele è l’immediato drenaggio per via endoscopica, seguito da asportazione chirurgica per una risoluzione definitiva della patologia.

PAROLE CHIAVE: Laringocele • Laringopiocele • Ostruzione delle vie aeree

Introduction

Laryngocele is an abnormal cystic dilatation of the saccule or appendix of the laryngeal ventricle, filled with air and communicating with the lumen of the larynx. Virchow introduced the term laryngocele, in 1867, to describe an abnormal dilatation of the saccule forming an air sac. Based on location, three types of laryngocele have been described. The internal, the external and the combined or mixed laryngocele. When the neck of the laryngocele is obstructed, it becomes filled with mucus of glandular secretion and is altered to a laryngomucocele. When this lesion becomes infected, a laryngopyocele is formed. A total of 39 cases of laryngopyocele have been reported in the world literature. Only one case of internal laryngopyoceles, causing acute airway obstruction, has been reported until now. In view of the rarity of the laryngopyoceles and the even rarer laryngopyocele that causes acute airway obstruction, it was decided that the report of another case would be worthwhile. Herewith, a rare case of an internal laryngopyocele is described in a 61-year-old female, in a septic condition which caused almost 100% airway obstruction.

Case report

A 61-year-old female presented to the emergency room, in respiratory distress, with a three-day history of sore throat, cough and odynophagia. She was a heavy smoker and had a past medical history of hypothyroidism and hypertension. Physical examination revealed marked respiratory distress with stridor, tachypnoea and hoarseness. Her temperature, on admission, was 38.3°C. Her white cell count was 23400 U/mm³ with 81.3% neutrophil leukocytes and C-Reactive Protein (CRP) was 17.8 mg/dl. Upon admission, the arterial blood gases on 21% O₂ were PO₂ 82 mm Hg, PCO₂ 52 mm Hg and pH 7.32. On palpation of the neck, no mass was noted. Indirect flexible laryngoscopy demonstrated a large mass which originated in the right false vocal cord and caused an almost total obstruction of the airway. The mucosal surface of the mass was smooth.

Waiting for the urgent computed tomography (CT) scan of the neck, she had a dramatic deterioration of dyspnoea and presented cyanosis. Since the large laryngeal mass prohibited intubation, an emergency tracheotomy was performed to secure a free airway. CT of the neck revealed a 29 mm low attenuation mass above the level of the true vocal folds which caused almost total obstruction of the airway (Figs. 1-2). Thickening of the walls was demonstrated and the lesion was confined within the larynx (Fig. 1). A diagnosis of internal laryngopyocele was made and confirmed with physical examination, indirect laryngoscopy and CT scan.

Direct laryngoscopy confirmed that the large laryngopyocele had caused an almost 100% obstruction and
it originated in the right ventricle. No visible neoplasm was noted. The laryngopyocele was drained and a large amount of purulent material was removed and cultures were collected. The definitive surgery was scheduled at a later time. The patient was treated with cortisone and intravenous antibiotics, including ampicillin, clindamycin and ceftriaxone. Culture revealed the presence of *Pseudomonas aeruginosa* and, therefore, ceftriaxone was substituted with gentamycin. The fifth post-operative day despite medical advice and the fact that she still had the cannula, the patient signed out in order to be transported to her homeland.

Discussion

A laryngocele is an air-filled herniation of the sacculus of the laryngeal ventricle which is in communication with the lumen of the larynx. The laryngeal ventricle is a fusiform fossa delimited by the true and the false vocal cord and extending from the thyroid notch to the arytenoid cartilage. The anterior part of the roof of the ventricle leads up into a blind pouch of the mucous membrane called the sacculus or appendix. Embryologically, the sacculus and ventricle of the larynx develop as a secondary outpouching from the laryngeal lumen towards the end of the second intrauterine month.

Several hundred laryngoceles have been reported in the English literature. A total of 39 cases of laryngopyocele have been reported in the world literature. Only in 4 of these cases had a laryngopyocele caused acute airway obstruction and only one case of internal laryngopyocele causing acute airway obstruction has been reported until now. The case described here is the first case reported in the English literature of a female patient in a septic condition, with an internal laryngopyocele causing acute airway obstruction (Table I).

In the case presented here, the patient was in a septic condition, at the time of admission, based on her clinical picture and the laboratory results. Indirect flexible laryngoscopy demonstrated a large smooth mass which originated in the right false vocal cord and caused an almost total obstruction of the airway. The differential diagnosis of acute upper airway obstruction includes several entities, except laryngopyocele (Table II). Diagnosis of an internal laryngopyocele was made by correlating the history, the clinical picture and the laryngoscopic and the radiographic findings (Table II).

Laryngoceles are rare entities, occurring in only one per 2.5 million population per year in the UK. The sex incidence is 5:1 in favour of male sex and the maximum age of incidence is in the sixth decade. Based on location, three types of laryngoceles have been described, the external, the internal and the combined type. The external laryngocele presents clinically as a swelling in the neck, at the level of hyoid bone, anterior to the sternocleidomastoid muscle. During Valsava’s manoeuvre, the swelling is increased and it becomes smaller on palpation. The internal and combined type, appear on laryngoscopy as a smooth swelling mass of the supraglottis.

The precise aetiology of laryngocele is unknown. Many authors have hypothesized that congenital or acquired factors may be responsible. It is believed that laryngoceles occur in subjects with congenitally dilated sacculles. A less important role is played by the narrowness of the periventricular connective tissue and the weakening of the thyroaryepiglottic muscles. A congenital weakness or defect predisposes to the formation of laryngoceles under the influence of acquired factors.

### Table I. Laryngopyoceles as a cause of airway obstruction.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age</th>
<th>Sex</th>
<th>Necessity of emergency tracheotomy</th>
<th>Laryngoscopic findings</th>
<th>Radiographic findings</th>
<th>Diagnosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>59</td>
<td>Male</td>
<td>Yes</td>
<td>Swelling of the left aryepiglottic fold and left false vocal cord</td>
<td>X-ray: large cavity with an air fluid level in the left neck (Thawley et al.)</td>
<td>Combined laryngopyocele</td>
</tr>
<tr>
<td>2</td>
<td>57</td>
<td>Female</td>
<td>No</td>
<td>A mass filling the right aryepiglottic fold and pyriform fossa</td>
<td>X-ray: right sided neck mass displacing trachea to the left (Weissler et al.)</td>
<td>Combined laryngopyocele</td>
</tr>
<tr>
<td>3</td>
<td>51</td>
<td>Male</td>
<td>Yes</td>
<td>Diffuse swelling over the right false cord and aryepiglottic fold</td>
<td>X-ray: right sided neck mass and an air-fluid level (Weissler et al.)</td>
<td>Combined laryngopyocele</td>
</tr>
<tr>
<td>4</td>
<td>34</td>
<td>Male</td>
<td>No</td>
<td>A mass originating in the left false cord caused a near total airway obstruction</td>
<td>CT: 18 mm low-attenuation mass within larynx that caused a significant airway obstruction (Fredrickson et al.)</td>
<td>Internal laryngopyocele</td>
</tr>
<tr>
<td>5</td>
<td>61</td>
<td>Female</td>
<td>Yes</td>
<td>A mass originating in the right false vocal cord, caused a near total airway obstruction</td>
<td>CT: 29 mm within larynx that caused an almost total airway obstruction in our case</td>
<td>Internal laryngopyocele</td>
</tr>
</tbody>
</table>
Internal laryngopyocele as a cause of acute airway obstruction

that increase intra-glottic pressure such as professional trumpet playing, glass blowing, singing, straining at passing of stools, weight lifting and carcinoma of the larynx are considered to promote the development of laryngoceles. The glandular serum-mucus secretion is evacuated through the ventricular opening. Some situations, such as chronic inflammations, laryngeal trauma or laryngeal carcinoma, lead to incomplete mechanical stenosis of the neck of the appendix. When the neck of the laryngocele is obstructed, the laryngocele becomes filled with mucus. If the mucus-filled laryngocele is infected, it is called a laryngopyocele. A review of the English literature showed that the most common type is the mixed laryngocele (44%), 30% were internal and 26% were external. Bilateral laryngoceles were found in 23%.

Laryngopyoceles may present with signs of rapidly progressive respiratory obstruction and/or an infected painful neck mass which may rapidly increase in size. The symptoms of laryngopyocele include hoarseness, dyspnoea, stridor, dysphagia, odynophagia, pain, sensation of a foreign body and fever. In the internal and combined forms, flexible nasolaryngoscopy can reveal a smooth mass of the vestiular fold, aryepiglottic fold and pyriform sinus which may displace the larynx to one side.

In the case presented here, after the emergency tracheotomy, computed tomography of the neck was performed which revealed a cystic 29 mm hypodense mass extending from the right false vocal cord to the level of the epiglotis, narrowing the laryngeal cavity and causing an almost 100% airway obstruction (Figs. 1-2). The lesion was confined within the larynx and it was diagnosed as an internal laryngopyocele. The cystic cavity was full of fluid and no air-fluid level was observed (Fig. 2).

Radiological evaluation includes neck ultrasound which may determine swelling dimensions and content and a CT scan that permits diagnosis. CT scan shows the characteristic intra-laryngeal and extra-laryngeal expansion and defines laryngopyocele content, the relationship with the laryngeal ventricle and thyroid membrane and the presence of a carcinoma. A contrast-enhanced CT scan can demonstrate signs of inflammation such as thickening of the walls or perimeter enhancement of the laryngocele and assist the differential diagnosis. In the differential diagnosis of laryngopyocele, it is necessary to take into consideration the saccular cyst, fluid filled laryngocele, branchial cysts, paraganglioma, schwannoma, and thyroglossal duct cysts which exist in the supraglottic area.

Laryngopyoceles are a rare complication of laryngoceles. They can present with rapid and alarming obstruction of the airway. Diagnosis requires a high index of suspicion, for these lesions, and careful clinical and radiologi-

**Table II. Causes of acute upper airway obstruction in adults.**

<table>
<thead>
<tr>
<th>Category</th>
<th>Causes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Congenital</td>
<td>Micrognathia, Macroglossia</td>
</tr>
<tr>
<td>Infections</td>
<td>Retopharyngeal abcess, Tuberculous laryngitis, Laryngopyocele, Epiglottis</td>
</tr>
<tr>
<td>Traumatic</td>
<td>Intubation trauma, Hematoma, Facial fracture, Laryngeal fracture, Subglottic stenosis</td>
</tr>
<tr>
<td>Allergic/autoimmune</td>
<td>Rhinitis, Sarcoidosis, Wegener’s disease, Angioedema, Asthma</td>
</tr>
<tr>
<td>Neoplastic</td>
<td>Nasopharyngeal carcinoma, Epiglottis carcinoma, Recurrent respiratory papillomatosis, Haemangiomia</td>
</tr>
<tr>
<td>Neurologic</td>
<td>Altered mental status, Vocal fold paralysis, Paralysis of respiratory muscles</td>
</tr>
</tbody>
</table>

Conclusions

Laryngopyoceles are a rare complication of laryngoceles. They can present with rapid and alarming obstruction of the airway. Diagnosis requires a high index of suspicion, for these lesions, and careful clinical and radiologi-
References


Received: August 26, 2010 - Accepted: January 6, 2011

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