ACQUIRED NONMALIGNANT TRACHEOESOPHAGEAL FISTULA

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INTRODUCTION

Tracheoesophageal fistula is defined as a pathological communication between the trachea and the esophagus. It is divided into congenital and acquired [1].

The most common fistulas are the congenital ones, with the diagnosis made early at a very young age [2]. The acquired ones are usually rare, divided into malignant and nonmalignant fistulas, occurring at any age. Acquired nonmalignant tracheoesophageal fistulas are secondary to infection, trauma, ruptured diverticula, or prolonged intubation [3].

Once diagnosed, the tracheoesophageal fistula is best managed with surgery [1].

In this paper, we present a case of an acquired nonmalignant tracheoesophageal fistula, a rare complication post prolonged intubation requiring a very challenging clinical diagnosis and surgical management.

CASE DESCRIPTION

A 59-year-old woman was admitted to our service with a history of dysphagia and cough for the past year.

The patient has a history of a bipolar disorder that is controlled with medications. One year ago, she was admitted to another hospital with an encephalopathy secondary to a change in her psychiatric medications. On admission, the patient was diagnosed having an aspiration pneumonia with mild hypoxemia that required intubation to protect her airways. She was admitted to the intensive care unit and remained intubated for a total of 10 days.

During those 10 days, the patient responded well to her medical treatment. The endotracheal tube was in good position and functioning well. She had no evidence of malnutrition and did not have any episodes of hypotension during that period.

The patient had an uneventful hospital course thereafter and was discharged home. She left the hospital with a buccal and lingual dyskinesia. Upon multiple clinic visits to her psychiatrist and neurologist, she continued to complain of this dyskinesia with a mild cough and dysphagia especially to liquids. All these symptoms were thought to be a side effect of her psychiatric medications.

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She continued to have symptoms for a total of one year with no improvement. Other causes of her symptoms were considered and a water soluble contrast swallow (Figure 1) and a neck-chest CT scan with oral contrast were obtained. These studies showed a tracheoesophageal fistula approximately 2 cm above the sternal notch.

Conservative treatment (enteral feeding at first then nothing per os) was tried for two weeks without any success.

Surgical intervention was then discussed with the patient who agreed to proceed.

After induction of anesthesia and during surgery, the tip of the tracheal tube was kept below the fistula to avoid air leakage and gastric distention through the tracheal fistula while ventilating the patient. The tracheal tube was positioned with the aid of bronchoscopy and did not interfere with the fistula.

The surgical approach was done through an horizontal neck incision above the sternal notch. Dissection of the tissues was done meticulously in order to avoid injury to the recurrent laryngeal nerves. The tracheoesophageal fistula was found along the left latero-posterior side of the trachea at the level of the 6th ring. Resection of the fistula was then done and the tracheal and esophageal defects were closed in two layers of interrupted 4.0 vicryl for the trachea and 4.0 vicryl and 3.0 silk for the esophagus. At the end a pedicled strap muscle flap was interposed between the trachea and the esophagus with a 4.0 vicryl running suture. The patient was extubated right after the surgery and kept in the ICU for 24 hours.

She had an uneventful postoperative course thereafter. On postop day 5 she had a normal upper esophageal swallow and an oral diet was started with no problems. She did not receive any enteral or parenteral nutrition because her preoperative nutritional status was adequate. Upon resumption of diet, she was receiving a caloric intake of 30 kcal/kg with 15% protein per day. She was discharged home on postoperative day 7.

At six months post surgery she had a normal neck-chest CT scan and at one year she remains free of symptoms.

DISCUSSION

Tracheoesophageal fistula is a congenital or acquired pathological communication between the trachea and the esophagus [1].

Congenital anomalies are much more common than acquired ones, and are usually diagnosed immediately after birth or during infancy [2].

Acquired tracheoesophageal fistulas are quite rare, divided into malignant and nonmalignant fistulas and may occur in individuals of any age.

Acquired nonmalignant tracheoesophageal fistulas are secondary to infection, trauma, ruptured diverticulae, or prolonged mechanical ventilation with an endotracheal or a tracheostomy tube [3].

Out of all these factors, the preventable causes are avoiding prolonged intubation and trauma on intubation. Several measures are routinely done in the intensive care unit to avoid these complications.

In the past, high-pressure endotracheal tube cuffs were used, but these cuffs highly contributed to tracheal injury and have been replaced by high-volume, low-pressure cuffs. The intracuff pressure in the endotracheal or tracheostomy tubes should be always measured [4], and kept below 20 cm H2O. The cuff volume should be limited to between 6 and 8 ml [5]. Also, any inflation with a volume exceeding 10 ml should raise concerns about tracheal injury.

When the cuff pressure exceeds the tracheal mucosal perfusion pressure, several injuries can occur as in mucosal erosions, ulceration, stenosis, tracheomalacia and tracheoesophageal fistula [6-8].

Our patient had a high-volume, low-pressure cuff and the cuff volume was 6 cc. The intra-cuff pressure was only measured intermittently. Thus the cuff pressure may have been high and caused the injury because our patient did not have a traumatic intubation. That’s why, tracheoesophageal fistula can occur even with the use of the newer endotracheal tubes with high-volume, low-pressure cuffs and even with close monitoring of the cuff pressure and volume.

Other factors that can cause decrease in mucosal blood flow to the tracheal tissue may contribute to the formation of the tracheoesophageal fistula. These include hypotension, hypoxemia, anemia, metabolic disorders, malnutrition, infection, sepsis, and steroid use.

Our patient had a pneumonia with mild hypoxemia but none of the other risk factors. She was also adequately sedated during mechanical ventilation.

The diagnosis of acquired nonmalignant tracheoesophageal fistula is usually made early, but in the case of our
patient, her symptoms were unrecognized for a year and mistaken for side effects of her psychiatric medications. Once the diagnosis of acquired nonmalignant fistula is confirmed, surgery is the treatment of choice. However, other options should be considered on a case per case basis.

Conservative treatment, keeping the patient with nothing per os is an option. In this case, conservative treatment was tried and failed. Stenting can be used for benign fistulas for a temporary period if the patient is critically ill and ventilated. This was not the case in our patient [9]. Endoscopic treatment has good results in the hands of experienced physicians in specialized centers. Through a bronchoscope, small fistulas can be closed using fibrin glue, small clips or argon plasma coagulation [10-12]. These options are definitely less invasive than surgery but couldn’t be offered to our patient due to lack of expertise in this field of endoscopic intervention in Lebanon.

The decision regarding the best surgical approach depends on the size of the defect, location of the fistula and any associated tracheal damage.

For small fistulas with a normal trachea, surgical treatment consists of simple division of the fistula with primary closure of the esophageal and the tracheal defects through a collar incision. A strap muscle pedicle flap is then interposed between the trachea and the esophagus to reinforce the two suture lines [13-14]. The muscles used are either the sternohyoid or the sternothyroid.

For fistulas around the carina, a right lateral thoracotomy is a more appropriate approach with the use of the pleura or the intercostal muscles for the flap.

For larger defects with tracheal damage, the surgical repair requires tracheal resection and reconstruction. The esophageal defect should be closed in two layers, and a viable muscle flap interposed between the trachea and the esophagus [14].

In rare situations, esophageal diversion may be needed [15].

CONCLUSION

Acquired nonmalignant tracheoesophageal fistula is a rare entity. A traumatic intubation should be avoided and the endotracheal tube cuff pressures should be routinely measured and kept below 20 cm H2O and the cuff volume limited from 6 to 8 ml [16].

Once it occurs, the treatment of choice of the tracheoesophageal fistula remains surgical resection with interposition of a muscle flap.

REFERENCES