Difficult To Treat Recurrent Esophageal Dysphagia Due To Secondary Esophageal Motility Disorders: A 24 Weeks Follow Up Case Report

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Abstract

Limited data are available on the prevalence of esophageal motility disorders (OMD), with a suggested range of 4-12% in cases with dysphagia. The proposed pathophysiology is either impairment of inhibitory innervation or overactivity of excitatory innervation. The optimal treatment is not defined until
now. This case represented one of the secondary OMD in a 70 years year diabetic patient with ischemic cardiomyopathy, who was complaining of recurrent daily intermittent dysphagia to solids and fluids, with a previous vague history of esophageal dilatation without available documentation. After endoscopy with multiple biopsies and barium imaging, the picture of the corkscrew esophagus was revealed. The patient, who was already on nitrates, failed to respond to twice-daily proton pump inhibitor for two months with a prokinetic drug, calcium channel blocker, and the first session of endoscopic esophageal dilatation. After the second dilatation session, we added sodium alginate to substitute nitrates with sildenafil. The patient reported a gradual improvement in dysphagia, especially fluids, decreasing daily attacks. We learn from this case that a different patient-to-patient response necessitates other treatment modalities, even switching between each patient's nitric oxide scavengers.

**Keywords:** Case Report; Dysphagia; esophageal Motility Disorders.
Introduction

Dysphagia is an alarming symptom that warrants prompt evaluation to define the exact cause and initiate appropriate therapy. It may be due to a structural or motility abnormality in the passage of solids or liquids. Dysphagia in older adults should not be attributed to aging. Aging alone causes mild esophageal motility abnormalities, rarely symptomatic (1).

Dysphagia to solids and liquids may be related to either an esophageal motility disorder (OMD) or a functional disorder that can be differentiated by investigations like upper endoscopy, barium image, and manometry studies. Symptoms of dysphagia may be intermittent or present after each meal (2).

Nonspecific esophageal dysmotility (including diffuse esophageal spasm, hypertensive peristalsis "nutcracker or corkscrew esophagus," hyper or hypotensive lower esophageal sphincter, and ineffective esophageal motility) is a type of OMD, which is considered if motility findings exceed two standard deviations from those found in a large group of healthy subjects (3).

OMD may occur as a primary or secondary to other diseases (like systemic sclerosis, Chagas’ disease, diabetes mellitus, and chronic gastroesophageal reflux disease), with limited data on the prevalence that is ranging, in some studies, from 4% up to 12% in other reviews, which were carried on individuals referred for esophageal manometry for evaluation of dysphagia or unexplained chest pain (4-6).

Although the underlying pathology is unknown, the supposed pathophysiology of OMD range from impairment of inhibitory innervation, leading to premature and rapidly propagated or simultaneous contractions and overactivity of
excitatory innervation or smooth muscle response to excitatory nerves. Patients may malfunction in endogenous nitric oxide synthesis and degradation (5, 7-10).

There is considerable controversy concerning the clinical implications of these abnormalities and whether they cause or explain the patient's symptoms, contrary to other clear ones like achalasia.

**Presenting Concerns**

Seventy-year-old male, a previous farmer, married and has four offspring with a medical history of diabetes mellitus for 20 years on insulin therapy, ischemic cardiomyopathy with moderate diastolic dysfunction, and preserved systolic function. For five years, the patient gave a vague history of endoscopic esophageal dilatation (without available documentation) for the same condition, recurrent intermittent dysphagia for solids and liquids with a gradual increase in daily attacks over the last three months from the presentation. There was no history of psychiatric illness. The patient was on the following treatment, Insulin Mixtard 30/70, Furosemide 250mg daily, Isosorbide Mononitrate 50mg daily, Losartan 50mg/Hydrochlorothiazide 12.5mg daily, Ivabradine 5mg daily, Rosuvastatin 10mg daily, Clopidogrel 75mg daily, and Omeprazole 40mg daily.

**Clinical Findings**

There is unremarkable general and local examination except for mild oral thrush, bilateral lower limb edema up to mid-leg, and body mass index of 36.7.
Timeline

**esophageal Dilatation for Dysphagia from 5 years**

<table>
<thead>
<tr>
<th>Date</th>
<th>Event Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>June 2018</td>
<td>Recurrent intermittent dysphagia for solids and liquids</td>
</tr>
<tr>
<td>September 2018</td>
<td>Upper endoscopy with multiple biopsies. HbA1c → 7.2%</td>
</tr>
<tr>
<td>September 2018</td>
<td>Barium swallow → Corkscrew - Biopsy → Mild non-specific inflammation</td>
</tr>
<tr>
<td>June-August 2018</td>
<td>Full dose Omeprazole 20mg/bid for 2 months. Itopride 50mg/tid for 15 days.</td>
</tr>
<tr>
<td>September 2018</td>
<td>1st session endoscopic dilatation. Lercanidipine 10mg/d.</td>
</tr>
<tr>
<td>October-November 2018</td>
<td>2nd session endoscopic dilatation. Change Isosorbide Mononitrate to Sildenafil 25mg/d. Add Sodium Alginate.</td>
</tr>
</tbody>
</table>

**Improvement of dysphagia especially to fluids with decreasing in daily attacks**
Diagnostic Focus and Assessment

Diabetes was controlled in the past three months as glycated hemoglobin (HbA1c) was 7.2%. Complete blood count, liver chemistry tests, and kidney functions were within the normal range. There are normal chest X-ray findings and abdominal ultrasonography except for fatty liver. Echocardiography revealed moderate diastolic dysfunction with preserved systolic function. Diagnostic upper endoscopy showed multiple circular narrowing in the mid and lower esophagus with the complex proceeding of the endoscope (fig 1).

*Fig 1: Endoscopic view of the multiple circular narrowing in the mid and lower esophagus.*

Multiple biopsies were taken, and the histopathological report revealed esophagitis, mild keratosis, and parakeratosis. Barium swallow imaging of the esophagus revealed the corkscrew appearance (Fig 2).
Therapeutic Focus and Assessment

We started with a twice-daily proton pump inhibitor Omeprazole 20mg before the meal for one hour for two months without changing the patient complaint. Then we tried a prokinetic agent, Itopride 50mg tid before the dinner by one hour for 15 days but no response. Then we proceed to the first session of dilatation with Savary-Gilliard® Dilator size 9-10mm (27-30Fr), and we add a calcium channel blocker Lercanidipine 10mg daily after the cardiologist consultation. The patient still had no response, so we performed the second dilatation session with a size 11-12mm (33-36Fr) plus changing the Isosorbide
Mononitrate to Sildenafil 25mg daily and adding Sodium Alginate 10ml tid before meals.

**Follow-up and Outcomes**

The patient, after one month, gave us positive feedback of a gradual improvement of dysphagia, especially to fluids with decreasing in daily attacks.

**Discussion**

One of the strengths of this report was the ability to manage this complex case associated with significant cardiac comorbidity that limits our choices in drugs and doses. Although, a considerable limitation was the absence of esophageal manometry study due to patient refusal and the unique treatment modality that needs validation on a large scale of similar conditions. Our approach is consistent with the American Gastroenterological Association, the American Society for Gastrointestinal Endoscopy, and the World Gastroenterology guidelines (11-14). The principal concern, in this case, was esophageal cancer due to the age of the patient. However, the intermittent nature of dysphagia and the absence of alarming signs like significant weight loss and anemia make it unlikely (15). Also, complications of gastroesophageal reflux disease, such as erosive esophagitis, peptic stricture, or adenocarcinoma of the esophagus, were precluded by the absence of chronic heartburn (16, 17). The presence of recurrent, intermittent dysphagia to solids and liquids with long-term diabetes mellitus makes it more reasonable for secondary motility disorders; however, functional conditions may cause. Still, it was excluded due to lacking Rome IV criteria. Also,
eosinophilic esophagitis or lymphocytic one should be excluded; we take multiple biopsies from the esophagus (15, 16). Lastly, we must not ignore the cardiovascular abnormalities in this age, like severe atherosclerosis or a giant aneurysm of the thoracic aorta can result in impingement on the esophagus (17). However, the endoscopic picture, chest X-ray, and echocardiography did not raise suspicion of this possibility.

**Conclusion**

We learn from this case that a different patient-to-patient response necessitates other treatment modalities, even switching between each patient's nitric oxide scavengers.

**Informed Consent**

The patient provided his informed consent for the publication of this case report.

**Conflicts of interest**

The author declares no conflicts of interest.

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