Pharyngeal webs in a patient with dysphagia and Behcet’s disease

A 64-year-old woman presented with longstanding dysphagia to solids and a recent onset of nocturnal cough and change in her voice. The patient described restriction to the passage of solids at the level of the neck, but denied odynophagia or weight loss. She had a background of Behcet’s disease with previous colonic and oropharyngeal involvement. She had also been treated for many years with esophageal dilations for an upper esophageal web. Her symptoms of dysphagia improved temporarily after each dilation; however, typically there was recurrence within 2 weeks of treatment. Video fluoroscopy was performed, which demonstrated a small web in the region of the pharyngo-esophageal junction, above which there was pharyngeal dilation. At least one episode of aspiration occurred during this procedure. After informed consent had been obtained from the patient, repeat esophagogastro-duodenoscopy was undertaken using a 5-mm pediatric endoscope (GIF-180; Olympus, Ontario, Canada). Multiple fibrotic-looking rings were noted throughout the hypopharynx (Fig. 1a) and pharynx (Fig. 1b,c). There was no ulceration or erythema present. A small diverticulum was seen just above the larynx (Fig. 1d). The pediatric endoscope was passed into the esophagus, stomach, and duodenum with no other abnormalities being detected. The patient was referred to an ear, nose, and throat (ENT) surgeon for further management. ENT examination confirmed the presence of several fibrotic rings affecting the pharynx and hypopharynx with considerable narrowing toward the glottic opening. It was felt that the degree of restriction at the oropharynx was sufficient to account for the patient’s symptoms. The etiology of the rings remained unclear. Because of her persistent symptoms, it was recommended to the patient that she undergo excision of the rings during direct laryngoscopy using a laser. Histopathology of the specimens confirmed squamous mucosa with submucosal fibrosis and mild chronic inflammatory changes. At a follow-up visit after 2 months, the patient reported considerable improvement in her symptoms.

Behcet’s disease is a type of systemic vasculitis that may affect small, medium, and large vessels (arteries or veins). The diagnosis of Behcet’s disease is a clinical one and requires the presence of recurrent mouth aphthous ulcers (at least three times per year), and two of the following: recurrent genital aphthous ulcers, eye lesions, skin lesions, or a positive pathergy test [1]. In the oropharynx, pharyngeal stenosis may be associated with pharyngeal webs. In patients with persistent dysphagia once esophageal lesions and motility disorders have been excluded. Behcet’s disease may be associated with pharyngeal webs.

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M. Efthymiou, S. Raftopoulos, P. Korton
St Michael’s Hospital, Toronto, Ontario, Canada

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Corresponding author
M. Efthymiou, MD
Therapeutic endoscopy unit
St Michael’s Hospital
30 Bond St
Toronto
MSB 1W8
Ontario
Canada
Fax: +613-8678-1032
mariosefthymiou@hotmail.com

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