Cricopharyngeal peroral endoscopic myotomy for achalasia of the cricopharynx: “to do or not to do”

Cricopharyngeal achalasia is a rarely reported [1] entity traditionally treated by surgery [2]. The video shows two cases referred to our unit for cricopharyngeal peroral endoscopic myotomy (C-POEM).

The first case was a 40-year-old woman with a 2-year history of cervical dysphagia requiring enteral feeding. Previous gastroscopy, barium transit, and high resolution manometry (HRM) were compatible with cricopharyngeal achalasia. A neck ultrasound and computed tomography (CT) without intravenous contrast supported the diagnosis. During the endoscopy for C-POEM, an upper compression that flattened with the endoscope and presented a beat was observed. An urgent angio-CT diagnosed the patient with lusoria dysphagia. Therefore, no endoscopic treatment was performed and the patient was referred for vascular bypass surgery.

The second case was an 83-year-old man with upper dysphagia and laryngeal microaspirations. Barium transit showed an upper posterior imprint (Video 1) and HRM was compatible with cricopharyngeal achalasia. At gastroscopy, it was not possible to pass the upper esophageal sphincter. A CT scan with contrast ruled out extrinsic compressions. The patient was reluctant to undergo therapeutic maneuvers and accepted a treatment with botulinum toxin, which subsequently worsened the symptoms. Videoradiology and a new manometry reaffirmed the diagnosis of cricopharyngeal achalasia.

Fig. 1 Cricopharyngeal bar.

Fig. 2 Myotomy of the cricopharyngeal bar. a Exposed cricopharyngeal bar. b Complete cricopharyngeal bar myotomy. c Disappearance of the bar from the esophageal lumen after myotomy.
Finally, the patient agreed to undergo C-POEM. After initial tunneling without cap owing to the limited space, a myotomy of the cricopharyngeal bar (Fig. 1) was performed with subsequent closure of the mucosotomy with clips (Fig. 2). The patient experienced immediate symptomatic improvement that was confirmed by barium transit (Fig. 3), and remained asymptomatic after 5 months. Cricopharyngeal achalasia without Zenker’s diverticulum requires careful diagnosis to exclude other causes of upper dysphagia. Working without a cap at the beginning of the procedure led to successful completion of the myotomy.

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Competing interests

The authors declare that they have no conflict of interest.

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