Plummer–Vinson syndrome was first described in 1912 [1], but seems to be extremely uncommon nowadays. We present the case of a 48-year-old woman referred for evaluation of dysphagia of 8 years' duration, which was worsening in frequency and intensity. The dysphagia initially involved only solids, but the patient now reported difficulty swallowing liquids as well. She also reported a weight loss of 10 kg over the past 2 years. The past medical history was essentially negative; medication intake included occasional iron supplements. Physical examination revealed no major abnormalities except for general pallor. A complete blood count showed a hemoglobin of 7.1 mg/dL and a mean corpuscular volume of 55.

A barium swallow revealed circumferential narrowing of the cervical esophagus just below the cricopharyngeal muscle, at the level of the C4 vertebra (Fig. 1). This was followed by an endoscopic gastroduodenoscopy (EGD), which showed a smooth, circular, whitish narrowing at 20 cm from the scope entry, suggestive of a web. The scope could not be passed further (Fig. 2).

Endoscopic dilation was performed under fluoroscopy, using dilators with serially increasing diameters. The web was easily disrupted without complications. The postendoscopic course was uneventful with resolution of the dysphagia after iron therapy [2]. Many other causes have also been proposed. Management of Plummer–Vinson syndrome is straightforward. Iron replacement should be continued until normalization of the hematocrit and ferritin levels. Although iron therapy may lead to considerable improvement of dysphagia [3], this is not the case in patients with longstanding disease; they usually require mechanical dilation [4]. One session is usually enough to give long-term relief like in our case, but, rarely, multiple sessions may be warranted.

References
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Fig. 1 Pretreatment barium swallow showing proximal narrowing in the cricopharyngeal area.

Fig. 2 Esophagogastrroduodenoscopy revealing a smooth, circular web in the proximal esophagus.