Loss of peristalsis of the esophagus due to diffuse esophageal leiomyomatosis

A 15-year-old girl was referred to our hospital with a tumor in the mediastinum that had been found on a chest radiograph during a medical check-up. An esophagram showed a large mediastinal mass (60 mm in diameter) (▶ Video 1). No peristalsis of the thoracic esophagus (smooth muscle area) was apparent, and dilatation of the esophagus and retention of the barium were observed on the proximal side of the tumor. High resolution manometry (HRM) findings showed complete loss of peristalsis of the smooth muscle area, whereas peristalsis of the striated muscle area was preserved (▶ Fig. 1). Although the esophageal lumen was narrowed by the tumor, the endoscope could be passed through it (▶ Fig. 2). Computed tomography (CT) scanning demonstrated a mildly enhancing tumor, which had circumferentially surrounded the lower esophagus (▶ Fig. 3). On T1-weighted magnetic resonance imaging (MRI), the tumor had a relatively high signal, which was the same as that of other muscular organs (▶ Fig. 4).

The patient underwent endoscopic ultrasound-guided fine needle aspiration (EUS-FNA), with the endoscopic ultrasound showing that the tumor had a mosaic echo, and that there was a clear boundary between the tumor and other organs (▶ Fig. 5). Biopsy specimens...
showed spindle cells without atypia on hematoxylin and eosin (H&E) staining; they were positive for smooth muscle actin and desmin, and negative for c-kit by immunohistochemistry (▶ Fig. 6). A diagnosis of diffuse esophageal leiomyomatosis (DEL) was made but, as she was asymptomatic, surgery was not indicated.

DEL is a rare, benign disease characterized by diffuse hypertrophy of the esophageal muscular layer; some cases of DEL are associated with Alport syndrome [1–4]. Although patients with DEL usually present with dysphagia [1, 2], our patient was asymptomatic with no signs of Alport syndrome. Esophageal motility in DEL investigated by HRM has rarely been reported [5]. This is the first report showing loss of peristalsis of the esophagus due to DEL, although the mechanism of this remains unknown.

References


Competing interests

None

The Authors

Kazuya Takahashi, Yui Ishii, Kazunao Hayashi, Satoshi Ikarashi, Hirokazu Kawai, Yuichi Sato, Shuji Terai
Division of Gastroenterology and Hepatology, Graduate School of Medical and Dental Science, Niigata University, Niigata, Japan

Corresponding author

Kazuya Takahashi, MD, PhD
Division of Gastroenterology and Hepatology, Graduate School of Medical and Dental Science, Niigata University, 1-757 Asahimachi-dori, Chuou-ku, Niigata City, Niigata, 951-8510, Japan
Fax: +81-25-2270776
kazuya911@med.niigata-u.ac.jp

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Fig. 4 The relatively high signal of the tumor was similar to that of other muscular organs on the T1-weighted magnetic resonance imaging (MRI) scan, indicating that the tumor tissue had similar properties to that of muscle.

Fig. 5 Endoscopic ultrasound image showing that the tumor had a mosaic echo and that there was a clear boundary between the tumor and other organs.

Fig. 6 Spindle cells without atypia are seen using hematoxylin and eosin (H&E) staining. Immunohistochemical staining shows the cells are positive for smooth muscle actin and desmin, but negative for c-kit, indicating the histopathological diagnosis of leiomyoma.