Esophageal natural killer (NK)/T cell lymphoma of true natural killer cell origin

A 53-year-old man with no prior medical history was transferred to our hospital because of a 2-month history of dysphagia with throat pain. A complete blood count showed hemoglobin 14.2 g/dL, and white blood cell count 11890/mL, and other laboratory findings were within normal limits. An enhanced computed tomography (CT) scan showed diffuse thickening of the esophageal wall without regional lymphadenopathy. An endoscopic examination revealed several well defined, variable-sized erosions and ulcers on the proximal esophagus (Fig. 1a) and distal esophagus (Fig. 1b). Endoscopic ultrasound (EUS) was performed to evaluate the esophageal wall. EUS showed a hypoechoic and homogeneous thickness along the entire length of the esophagus (Fig. 2). Endoscopic biopsy specimens revealed many neoplastic cells in the extensive necrotic tissue (Fig. 3) that showed positivity for CD3 (Fig. 4), CD56 (Fig. 5), and UCHL-1, and were negative for CD4, CD5, CD8, CD20, CD79a, and EBER-1. These findings were consistent with esophageal involvement by extranodal natural killer (NK)/T cell lymphoma of true natural killer cell origin.

The patient received two cycles of systemic chemotherapy with concurrent radiotherapy. However, his general condition gradually became worse and he died 4 months after admission.

The esophagus is the least commonly involved gastrointestinal organ, accounting for less than 1% of patients with primary gastrointestinal lymphoma [1]. Extranodal natural killer (NK)/T cell lymphoma of true natural killer cell origin is extremely rare, with only a few case reports in the literature [2,3]. In this case, the endoscopic picture showed mucosal erythema, a mass with ulceration, and wall thickness, similarly to previous reports of esophageal lymphoma. In a previous study, esophageal lymphoma showed lymphomatous involvement of the gastrointestinal wall to produce a typical hypoechoic transmural thickening [4], such as in this case. However, another report described imaging of a submucosal, heterogeneous, mainly hyperechoic mass [2].
Competing interests: None

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Fig. 4 Expression of CD3 antigen by lymphoid cells.

Fig. 5 Expression of CD56 antigen by lymphoid cells.