A 29-year-old man presented with a 3-month history of dysphagia and chest pain. Esophagogastroduodenoscopy showed a narrowed mid-esophagus with smooth mucosa (Figure 1). Endoscopic ultrasound (EUS) was performed, which showed loss of echo layers, esophageal wall thickening (Figure 2), and enlarged mediastinal lymph nodes (Figure 3). EUS-guided fine-needle aspiration (FNA) of these lymph nodes was not performed at this stage because the patient was agitated. Endoscopic biopsies from the mid-esophagus showed eosinophilic esophagitis with >15 eosinophils per high-power field. He was treated with Advair (fluticasone/salmeterol) for 1 month by his physician and the dysphagia resolved.

Repeat esophagogastroduodenoscopy and EUS-FNA under monitored anesthetic care showed no esophageal luminal narrowing and significant reduction in the thickness of the esophageal wall (Figure 4). Enlarged subcarinal lymph nodes (15 mm and 27 mm) were seen once again. EUS-FNA of the larger lymph node (Figure 5) revealed polymorphic, small-lymphocytic proliferation, and numerous eosinophils, with a cytologic pattern that was most consistent with reactive lymph-node hyperplasia with eosinophilia (Figure 6). No granuloma, Reed-Sternberg cells, metastasis, or evidence of lymphoma was identified.

There have been few case reports describing the EUS findings in esophageal eosinophilia [1–3]. Fox et al. [1], performed high-resolution EUS on 11 children with esophageal eosinophilia and eight controls and found significant differences in wall thickness. Evrard et al. [2] described a 72-year-old man with dysphagia and weight loss, in whom EUS showed a localized infiltrating process between the muscular layers. He underwent esophagectomy because he was suspected to have a neoplasm, and was revealed to have esophageal eosinophilia. Stevoff et al. [3] described an 85-year-old man with an esophageal stricture, normal mucosa,
and circumferential thickening of the muscularis propria on EUS. Esophagectomy was performed because atypical cells had been identified in the fine-needle aspirate, and this revealed esophageal eosinophilic infiltration involving the muscularis propria on pathologic examination. Our patient’s biopsies were consistent with a diagnosis of esophageal eosinophilia. Esophageal thickening was noted, along with loss of the echo layer pattern and this subsequently resolved. The presence of eosinophils in the mediastinal lymph nodes on EUS-FNA in association with esophageal eosinophilia in our case is very interesting and to our knowledge has not been reported previously.

References


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