A 48-year-old man was diagnosed with an esophageal subepithelial tumor (SET) found incidentally. Esophagogastroduodenoscopy (EGD) revealed a SET in the lower third of the esophagus (36 cm away from the incisor) (Fig. 1). EUS confirmed a hypoechoic tumor, measuring 0.8 cm, in the submucosal layer (Fig. 2). Benign leiomyoma was the initial impression, and hence no aggressive therapy was performed. The tumor size and appearance remained apparently unchanged during the following 2 years. The last EGD examination showed some alteration. Another smaller SET was also found adjacent to the previous lesion (Fig. 3).

Endoscopic mucosal resection with the suction and cap method (EMR-c) was performed for tissue proof. A whitish tumor measuring 8 mm was discovered in the submucosal region of the specimen and residual tumor remained under macroscopic view (Fig. 4). The histological examination showed undifferentiated carcinoma. EMR-c was repeated 1 month later for a confirmatory diagnosis. Squamous cell carcinoma (SCC) was verified (Fig. 5). The patient therefore underwent esophagectomy and lymph node dissection. The final histological diagnosis was SCC invading the submucosal layer without lymph node metastasis or muscular layer invasion (Fig. 6). This was a rare case of esophageal SCC with an atypical presentation mimicking a SET. SETs of the esophagus are typically considered to be benign tumors [1]. In addition, histological diagnosis of esophageal SETs by biopsy is difficult. Therefore, SETs of the esophagus are frequently neglected unless symptoms occur [2].
EMR-c, initially designed for resecting mucosal lesions, could be used for removing SETs of the gastrointestinal tract [2–4]. Tissue specimens gathered by the EMR-c method are generally large enough for immunohistochemical diagnosis. The method is convenient and safe for the resection of the SETs within the submucosal layer. On the basis of our experience, an esophageal SET can be an atypical presentation of SCC that should not be overlooked. Early endoscopic resection to get tissue proof is pivotal to avoid missing diagnosis.

References

Bibliography
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