Esophageal lesions herald widely metastatic disease

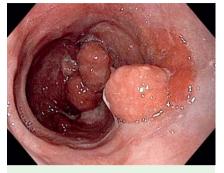


Fig. 1 Endoscopic view of the mid to distal esophagus showing multiple polypoid lesions.

A 59-year-old man with chronic hepatitis C and chronic obstructive pulmonary disease presented with an 8-month history of progressive solid food dysphagia, voice hoarseness, and weight loss. Prior ENT evaluation had revealed left vocal cord paralysis, with referral to gastroenterology. An upper endoscopy showed multiple polypoid masses in the mid and distal esophagus (> Fig. 1, > Video 1).

Biopsies demonstrated moderately differentiated adenocarcinoma of unknown primary (**> Fig. 2**).

Similar cells were found on fine-needle aspiration of the right superior chest wall (**>** Fig. 3).

A subsequent positron emission tomography (PET) scan revealed widely metastatic disease to the liver, bone, adrenal glands, and muscles (> Fig. 4).

Polypoid neoplasms, although common in the colon, are rare in the esophagus and when found are associated with Barrett's adenocarcinoma [1]. In addition, as a further distinction, our patient has multiple polypoid lesions and diffuse metastasis throughout the body. Apart from atypical adeno/squamous cell carcinomas, the differential diagnosis for multiple esophageal malignant lesions includes primary malignant melanoma, small and large cell carcinoma, and metastatic disease. Our case illustrates the difficulty in estimating the underlying pathology of Barrett's lesions solely based on the mucosal pit and microvessel pattern. Despite the use of

Video 1

Video demonstrating the endoscopic examination of the upper oesophagus.

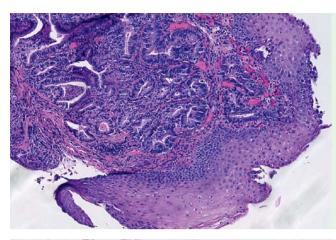


Fig. 2 Esophageal biopsy sample showing moderately differentiated adenocarcinoma.

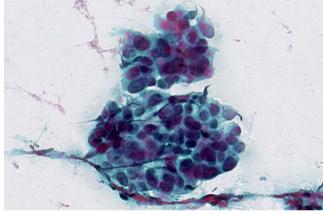
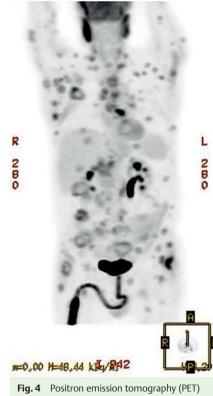


Fig. 3 Right superior chest wall fine-needle aspiration demonstrating similar cells to the esophageal biopsy specimen.

image-enhanced endoscopy, the mucosal patterns of the lesions were not significantly chaotic as has been reported [2,3]. The histology, showing high grade moderately differentiated adenocarcinoma, also did not seem to offer a satisfactory explanation for our patient's clinical findings, since differentiated lesions are less likely to metastasize. An explanation could be related to the unique mucosal penetration of esophageal lymphatics, which is very different from the rest of the gastrointestinal tract where it usually stops at the submucosa. The aggressive nature of superficial esophageal lesions has also been described as having an increased potential to metastasize to distant as opposed to local lymph nodes [4]. Our case highlights the continued importance for finding more effective screening modalities for Barrett's esophagus.

Competing interests: None

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