

Title:

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Endoscopic observation of a rare duodenal tumor

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Dear Editor,

A 30-year-old young previously healthy man presented to our hospital with middle and upper abdominal discomfort. Abdominal computerized tomography (CT) showed no significant abnormalities. White light endoscopy showed the local mucosa in the descending part of the duodenum had granuloid uplift, some of which were fused into pieces with red color, and some other areas showed a fading tone (Fig. 1A). Magnifying endoscopy with indigo-carmin staining and narrow-band imaging showed a finger-like, loose villous structure with irregular microvessels on the surface (Fig. 1B-D). Pathological examination of biopsy specimens showed that lymphocytes were diffused and dispersed in the mucosa with relatively simple morphology. No lymphoid follicles were observed, and local compression was obvious (Fig. 1E). Immunohistochemical staining revealed a lymphoid population

highly positive for CD20 (Fig. 1F) and CD10 (Fig. 1G). These results were consistent with duodenal-type follicular lymphoma (D-FL).

Discussion

D-FL is a new type of lymphoma with low incidence and rare clinical features (1). It often occurs in the descending part of duodenum. Patients with D-FL often present with no obvious clinical symptoms or only non-specific gastrointestinal symptoms. The lesions are usually confined to the mucosal and submucosal layer of the intestinal wall, without muscular layer, other viscera and lymph nodes involved. Although it is an indolent lymphoma with a fairly good prognosis (2), D-FL in the descending duodenum is easily missed by endoscopists and sometimes easily mistaken for Brunner's gland. Therefore, it is necessary to improve the endoscopic understanding of this disease for early diagnosis and treatment.

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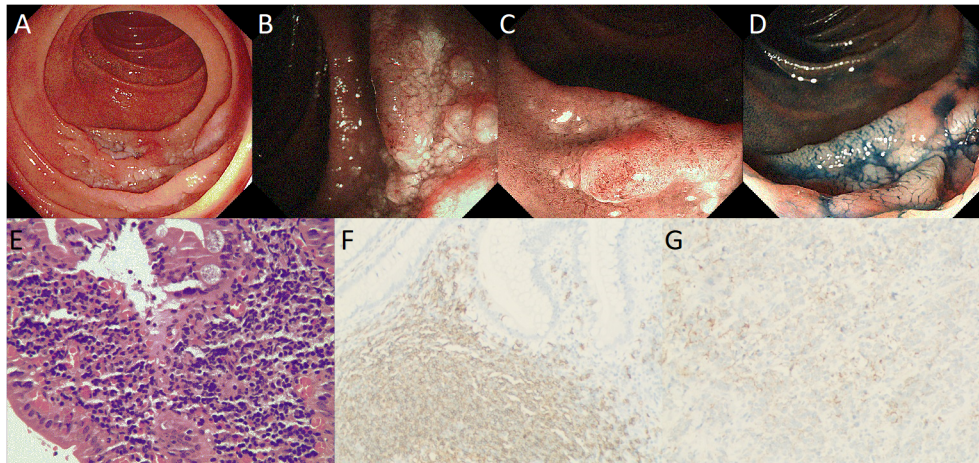


Fig. 1. A. The local mucosa in the descending part of the duodenum had granuloid uplift, some of which were fused into pieces with red color, and some other areas showed fading tone. B-D. Magnifying endoscopy with indigo-carmin staining and narrow-band imaging showed a finger-like, loose villous structure with irregular microvessels on the surface. E. Pathological examination showed that lymphocytes were diffused and dispersed in the mucosa with relatively simple morphology; no lymphoid follicles were observed, and local compression was obvious (40 x objective). F and G. Immunohistochemical staining revealed a lymphoid population highly positive for CD20 (F; 10 x objective) and CD10 (G; 10 x objective).