An unusual case of jejunal choriocarcinoma detected by single-balloon endoscopy

A 41-year-old woman was admitted to our institution because of massive melena and symptomatic anemia (Hb 5.3 g/dL). The patient had previously undergone a cesarean section and tubal ligation 3 years ago while giving birth to her second child. Neither esophagogastroduodenoscopy nor colonoscopy identified the source of bleeding. Single-balloon endoscopy revealed a lobulated purplish tumor over the proximal jejunum (Fig. 1). Due to the tendency to bleed, only a limited biopsy specimen was obtained, and a poorly differentiated carcinoma was impressed. A computed tomography (CT) scan revealed several hypodense lesions in both lobes of the liver (1–2.5 cm in diameter), and a soft tissue mass over the right suprarenal region. No further lesion was detected by the systematic survey.

The patient underwent an emergency laparoscopy to control the continued heavy gastrointestinal bleeding. Laparoscopic reduction of jejunojejunal intussusception and segmental resection of the jejunum were performed (Fig. 2). The final histological diagnosis of the surgically resected specimen was choriocarcinoma of the jejunum (Fig. 3). The level of serum β human chorionic gonadotropin (β-HCG) determined after pathological diagnosis was 241 991 IU/L.

Choriocarcinomas of the small intestine, including both primary and metastatic tumors, are rarely encountered [1, 2]. They are seen most commonly in the jejunum, followed by the duodenum, and least commonly seen in the ileum [2]. The pure choriocarcinoma without other malignant elements identified by immunohistochemical analysis is not sufficient to exclude the possibility of primary choriocarcinoma of the jejunum [2, 3]. In the present case, a gestational origin for the jejunal choriocarcinoma was confirmed by analyzing genetic polymorphisms in the tumor [4] (Fig. 4). The majority of gestational choriocarcinomas develop shortly after the preceding gestation [5]. In the absence of uterine tumors, the diagnosis of the late onset of gestational choriocarcinoma at an unusual metastatic site is a challenge to clinicians. This is the first report of jejunal choriocarcinoma detected by single-balloon endoscopy.

**Endoscopy_UCTN_Code_CCL_1AD_2AB**

**Competing interests:** None

**Y.-J. Tai**, T.-C. Lee
L.-H. Tseng, L.-H. Wei

1 Department of Oncology, National Taiwan University Hospital and National Taiwan University College of Medicine, Taipei, Taiwan
2 Department of Internal Medicine, National Taiwan University Hospital and National Taiwan University College of Medicine, Taipei, Taiwan
3 Department of Medical Genetics, National Taiwan University Hospital and National Taiwan University College of Medicine, Taipei, Taiwan

**Acknowledgments**

The pathology review was made by Dr Kuan-Ting Kuo, Department of Pathology, National Taiwan University Hospital. A thoughtful review of this article was made by Dr Ruey-Jien Chen, Department of Obstetrics and Gynecology, National Taiwan University Hospital.

---

**Fig. 1** Single-balloon enteroscopy showed a 3-cm lobulated vascular tumor over the proximal jejunum 25 cm distal to the Treitz ligament.

**Fig. 2** Intraoperative findings included a 4-cm tumor at the proximal jejunum with serosal invasion causing intussusception.

**Fig. 3** Microscopic examination showed the tumor, consisting of two cell patterns, embedded in a hemorrhagic background with jejunal muscular and mucosal involvement. One group was composed of hyperchromatic pleomorphic tumor cells with vesicular nuclei and a medium amount of eosinophilic cytoplasm, the other was composed of multinucleate giant cells. (Hematoxylin and eosin stain.)
Fig. 4 Microsatellite analysis of Y chromosome loci of the jejunal tumor (left) showed identical polymorphic alleles to those of the paternal Y chromosome (right).

References

Bibliography
DOI http://dx.doi.org/10.1055/s-0032-1310018
Endoscopy 2012; 44: E360–E361
© Georg Thieme Verlag KG
Stuttgart · New York
ISSN 0013-726X

Corresponding author
L.-H. Wei, MD, PhD
Department of Oncology
National Taiwan University Hospital
7 Chung-Shan South Road
Taipei 100
Taiwan
Fax: +886-2-23711174
weilh1966@gmail.com