CASE REPORT

Rare cause of severe anemia due to pyogenic granuloma in the jejunum

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Abstract

Background: Pyogenic granuloma (PG) is a polypoid lobular capillary hemangioma rarely observed in the gastrointestinal tract. Only a few cases in the small bowel have been described in the literature.

Case presentation: A 58-year-old man had been suffering from general fatigue and severe anemia. Esophagogastroduodenoscopy and colonoscopy did not reveal any significant bleeding. Abdominal computer tomography revealed a hypervascular tumor in the small intestine. Oral double-balloon endoscopy (DBE) detected a polypoid lesion (2 cm in diameter) in the jejunum. We performed laparoscopic-assisted partial resection of the jejunum. The histological features of the tumor were consistent with PG. The patient’s anemia gradually improved without the need for oral iron after surgery.

Conclusion: In this case report, we present a case of pyogenic granuloma in in the jejunum that was detected by DBE.

Keywords: Pyogenic granuloma, Small intestine, Double-balloon endoscopy, Obscure gastrointestinal bleeding, Hypervascular tumor

Background

Obscure gastrointestinal bleeding (OGIB) is frequently observed in the small intestine. Double-balloon endoscopy (DBE) and capsule endoscopy (CE) are highly effective in diagnosing the origin of OGIB [1, 2]. DBE is particularly valuable for the detection and diagnosis of small bowel tumors, and biopsies and therapeutic procedures have also become possible [3].

Pyogenic granuloma (PG) is a capillary hemangioma that usually occurs on the skin or in the oral cavity; it is rarely observed in the gastrointestinal tract. We report a case of PG in the jejunum that was detected by DBE.

Case presentation

A 58-year-old man had been suffering from general fatigue and severe anemia for several months. His hemoglobin levels were 6.6 g/dl (normal range: 12–16 g/dl). He had no medical history and did not take any medicine. Esophagastroduodenoscopy and colonoscopy did not reveal any significant bleeding. Abdominal computer tomography revealed a 2-cm hypervascular tumor in the small intestine (Fig. 1). Oral DBE detected a 2-cm-diameter reddish, submucosal tumor-like lesion with surface ulceration in the jejunum, approximately 20 cm away from the Treitz ligament (Fig. 2). We did not perform biopsy because it can be difficult to stop bleeding in the case of hypervascular lesions. Under the diagnosis of a small bowel tumor, gastrointestinal stromal tumor (GIST), malignant lymphoma, or cancer, we performed laparoscopic-assisted segmental resection of the jejunum with the dissection of lymph nodes.

Examination of the resected tumor showed that it measured 19 × 16 mm in diameter (Fig. 3). Histology revealed the proliferation of blood capillaries and granulation tissue, which was consistent with PG (Fig. 4). The patient was discharged on postoperative day 9 without complication and his anemia improved gradually without the need for oral iron after surgery.

Discussion

PG, also called granuloma telangiectaticum or lobular capillary hemangioma, was first described in 1897.
Gastrointestinal PG is apparently a rare cause of hemorrhage in the digestive tract. PG is a benign lesion that is considered reactive, e.g., it occurs as a result of minor trauma. However, the precise etiology is unknown. Satellite lesions have been described on the skin, which argues against trauma as the sole cause. Lesions are also found more frequently during pregnancy on the skin and in the mouth, suggesting that hormonal influences may also play a role. All gastrointestinal PGs reported thus far have occurred in elderly patients, both male and female [4]. The distinctive histopathological characteristics of PGs are proliferation and lobular arrangement of capillary-sized vessels with inflamed and edematous stroma and endothelial cell swelling cells [5].

To our knowledge, only 16 cases of PG in the small intestine have been described in the literature, and most involved Japanese patients (Table 1). This may be due to the development of DBE for examination of the small bowel in Japan. Including our case, the average age of the 17 patients was 58.1 years (range: 26–86 years), with no significant difference between males and females.

The tumor size is usually less than 20 mm, with a clinical presentation of anemia. Of the 17 reported cases, 12 involved the ileum and 5 involved the jejunum. Most tumors had an irregular shape without surface ulceration.
Table 1  Summary of the reported cases of PG in the small intestine

| Author        | Year | Age/Sex | Symptoms | Hb (g/dl) | Location | Size (mm) | Treatment         |
|---------------|------|---------|----------|-----------|----------|-----------|-------------------|
| Payson [7]    | 1967 | 45/F    | Abdominal pain | 12.4     | Ileum    | 60        | Surgical resection |
| Meuwissen [8] | 1986 | 37/M    | None     | NA        | Ileum    | NA        | Laser ablation    |
| Iwakubo [9]   | 1989 | 30/F    | Melena   | 4.8       | Ileum    | 8 × 8     | Surgical resection |
| Hizawa [10]   | 1993 | 26/F    | None     | NA        | Ileum    | NA        | Surgical resection |
| Yao [11]      | 1995 | 71/F    | Anemia   | NA        | Ileum    | 24        | Surgical resection |
| Yao [11]      | 1995 | 56/M    | Melena   | NA        | Jejunum  | 20        | Surgical resection |
| Motohashi [12]| 1999 | 58/F    | Anemia   | 6.1       | Ileum    | 25 × 30   | Surgical resection |
| Van Eden [4]  | 2004 | 55/F    | Anemia   | NA        | Ileum    | 9         | Surgical resection |
| Shirakawa [13]| 2007 | 72/M    | Anemia   | NA        | Ileum    | NA        | EMR               |
| CHO [6]       | 2009 | 58/M    | Melena   | 5         | Ileum    | 10        | Surgical resection |
| Moffat [14]   | 2009 | 78/M    | Anemia   | 6.2       | Jejunum  | 20        | Surgical resection |
| Nagoya [5]    | 2009 | 63/F    | Anemia   | 4         | Ileum    | 7         | EMR               |
| Kuga [15]     | 2009 | 55/M    | Anemia   | 9.9       | Jejunum  | 4         | EMR               |
| Yamashita [16]| 2013 | 61/M    | Anemia   | NA        | Ileum    | 15        | Surgical resection |
| Katsurahara [17]| 2015 | 65/M    | Anemia   | NA        | Jejunum  | 33        | Surgical resection |
| Kikuchi [18]  | 2015 | 86/F    | Anemia   | 8.1       | Ileum    | 7         | Surgical resection |
| This case     | 2015 | 58/M    | Anemia   | 6.6       | Jejunum  | 19        | Surgical resection |

NA data not available

**Fig. 4** Microscopic appearance of the hematoxylin and eosin (H&E) stained resected specimen. 

- **a** Low-power view of the histological features. 
- **b** High-power view of the histological features (original magnification: x100). Proliferation and lobular arrangement of capillary-sized blood vessels can be observed along with inflamed and edematous stroma and endothelial cell swelling. Remarkable inflammatory cell invasion and granulation tissues are noted in the capillaries and granulation tissue. 
- **c** Positive immunostaining for CD31 in capillary endothelial cells.
and were reddish in color. Surgical resection is the mainstay for treating small bowel PGs, although endoscopic resection with an electrosurgical snare is an alternative technique if an endoscopic approach is possible. Endoscopic treatment is a relatively easy, safe, and low-cost procedure for small-sized PGs of the gastrointestinal tract [3]. However, when resecting PGs, it is important that arteriovenous anastomosis under the tumor is included endoscopically, because incomplete resection may cause recurrence [6].

In the present case, we performed laparoscopic-assisted partial resection because complete endoscopic resection of the 2-cm hypervascular submucosal tumor with ulceration seemed to be difficult.

Conclusion

PG in the small intestine is a rare cause of anemia and is difficult to detect. Awareness regarding this infrequent benign lesion will make it easier to diagnose and treat it properly.

Consent

Written informed consent was obtained from the patient for publication of this case report and any accompanying images.

Abbreviations

PG: Pyogenic granuloma; OGIB: Obscure gastrointestinal bleeding; DBE: Double-balloon endoscopy; CE: Capsule endoscopy; GIST: Gastrointestinal stromal tumor.

Competing interests

The authors declare that they have no competing interests.

Authors’ contribution

SN, TM, SY, MH, and OQ performed endoscopic procedures. SM, HS, AK, YK, and HT managed the patients and performed operation. HO diagnosed pathologically. SM wrote the manuscript and KY and HT reviewed it. All authors approved the final manuscript.

Authors’ information

Not applicable.

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