

A 25-Year Old Woman with Recurrent Anemia

Jennings J¹, Scott N² and Sandle GI^{3*}

¹Bexley Wing (Level 4), St James's University Hospital, Leeds LS9 7TF, UK

²Bexley Wing (Level 5), St James's University Hospital, Leeds LS9 7TF, UK

³Clinical Sciences Building (Level 8), St James's University Hospital, Leeds LS9 7TF, UK

*Corresponding author:

Sandle GI,
Clinical Sciences Building (Level 8),
St James's University Hospital, Leeds LS9 7TF.
Tel: 44-113-2068607,
E-mail: g.i.sandle@leeds.ac.uk

Received: 03 Oct 2021

Accepted: 19 Oct 2021

Published: 23 Oct 2021

Copyright:

©2021 Sandle GI, This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and build upon your work non-commercially.

Citation:

Sandle GI, A 25-Year Old Woman with Recurrent Anemia. J Clin Med Img. 2021; V5(17): 1-2

Case Report

A 25-year old Ghanaian woman presented with a 1-week history of fatigue and breathlessness. Menstrual and dietary histories were normal, there was no overt gastrointestinal blood loss, and she took no medication. Physical (including digital rectal) examination was normal, apart from pale conjunctivae. Investigations revealed haemoglobin 4.6g/dL, white cell count 7.09 x 10⁹/L, platelet count 442 x 10⁹/L, MCV 63fL, MCH 16.0pg, serum ferritin 2.0µg/L. Other routine laboratory investigations, including serum anti-tissue transglutaminase, hemoglobinopathy screen, serum haptoglobin and reticulocyte count, were normal. Chest X-ray was normal. After transfusion of 4 units of red cells, the hemoglobin rose to 10.0g/dL with resolution of symptoms, and she was discharged on oral iron.

Soon afterwards, latent tuberculosis was diagnosed after routine screening revealed a positive Quantiferon test, and she was treated for 3 months with rifampicin. Several further transfusions were required during the next five months, the hemoglobin falling at one point to 5.8g/dL. Screening for coagulopathy and Von Willebrand disease was negative, and gastroscopy and colonoscopy were normal, as was duodenal histology. After referral to the gastroenterology service, she underwent capsule endoscopy, which showed fresh blood in the mid-jejunum, with clot and altered blood more distally, but no obvious mucosal abnormality. Computed tomography of the abdomen was unremarkable. Double-balloon enteroscopy revealed an actively bleeding submucosal lesion 1.6m beyond the pylorus, the overlying mucosa appearing intact (Figure 1, upper panel). At laparotomy, a small intestinal polypoidal tumor was resected and histology revealed a lymphangioma (Figure 2, lower

panel). She made a good recovery and remains symptom-free with a normal blood count.

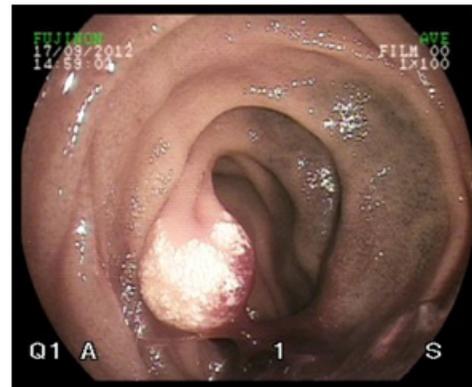


Figure 1: Upper Panel: Polypoidal lesion in jejunum during double-balloon enteroscopy

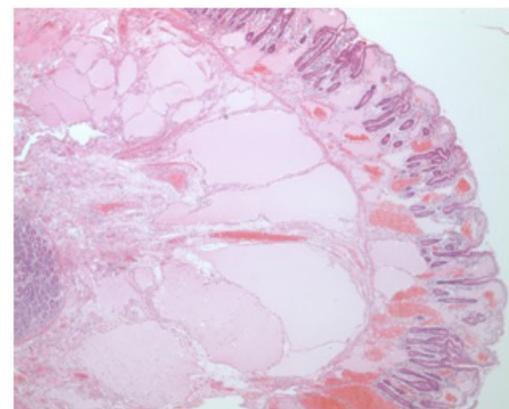


Figure 2: Lower Panel: Resected lymphangioma, showing dilated thin-walled channels covered by small intestinal mucosa (hematoxylin and eosin, x50 magnification).

Lymphangiomas are benign lesions of vascular origin with lymphatic differentiation and usually present in childhood, adult cases being rare. Only a minority (<5%) occur in the abdomen and are often incidental findings during endoscopy or imaging studies [1]. Mesenteric lymphangiomas may be large, causing abdominal distension, [2] or small intestinal obstruction secondary to volvulus, [3,4] but rarely cause significant intestinal blood loss. Our case is therefore unusual, and we are aware of only one other case of small intestinal lymphangioma diagnosed by enteroscopy, using the single-balloon technique [5]. Patients with recurrent iron deficiency anemia often present to specialties other than gastroenterology, and after a normal gastroscopy and colonoscopy, capsule endoscopy, followed if necessary by either single- or double-balloon enteroscopy, should be used undertaken to exclude more common pathologies such as Crohn's disease, tuberculosis, and radiation damage.

References

1. Mimura T, Kuramoto S, Hashimoto M, Yamasaki K, Kobayashi K, Kobayashi M, et al. Unroofing for lymphangioma of the large intestine: a new approach to endoscopic treatment. *Gastrointest Endosc*. 1997; 46: 259-63.
2. Khattala K, Boujraf S, Rami M, Elmadi A, Afifi A, Sbai H, et al. Giant cystic lymphangioma of the small bowel mesentery: case report. *Ann Pediatr Surg* 2008; 4: 107-9.
3. Campbell WJ, Irwin ST, Biggart JD. Benign lymphangioma of the jejunal mesentery: an unusual cause of small bowel obstruction. *Gut*. 1991; 32: 1568.
4. Day W, Kan DMY. A small bowel lymphangioma presenting as a volvulus. *Hong Kong Med J*. 2010; 16: 233-4.
5. Cai JT, Chen JM, Zhang XM, Chen Y, Wei SM, Du Q, et al. Small bowel lymphangioma diagnosed by single-balloon enteroscopy. *J Gastroenterol Hepatol*. 2012; 27: 1407.